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Vibrant and Healthy Kids: Aligning Science, Practice, and Policy to Advance Health Equity

Committee on Applying Neurobiological and Socio-behavioral Sciences from Prenatal through Early Childhood Development: A Health Equity Approach

Jennifer E. DeVoe, Amy Geller, and Yamrot Negussie, *Editors*

Board on Population Health and Public Health Practice

Health and Medicine Division

A Consensus Study Report of

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COMMITTEE ON APPLYING NEUROBIOLOGICAL AND SOCIO-BEHAVIORAL SCIENCES FROM PRENATAL THROUGH EARLY CHILDHOOD DEVELOPMENT: A HEALTH EQUITY APPROACH

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Title: Chasing Sunshine

Artist: Stephanie Kohli (Weston, Wisconsin)

Artist statement:

This piece is based off of my daughter running in our garden. Feeding children the best quality food and letting them explore nature in community gardens is a beautiful way to help them shine.

This artwork was submitted as part of the National Academy of Medicine's Visualize Health Equity Community Art Project nationwide call for art. This call for art encouraged artists of all kinds to illustrate what health equity looks, sounds, and feels like to them. More information on this project can be found at nam.edu/VisualizeHealthEquity.

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REVIEWERS

This Consensus Study Report was reviewed in draft form by individuals chosen for their diverse perspectives and technical expertise. The purpose of this independent review is to provide candid and critical comments that will assist the National Academies of Sciences, Engineering, and Medicine in making each published report as sound as possible and to ensure that it meets the institutional standards for quality, objectivity, evidence, and responsiveness to the study charge. The review comments and draft manuscript remain confidential to protect the integrity of the deliberative process.

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Although the reviewers listed above provided many constructive comments and suggestions, they were not asked to endorse the conclusions or recommendations of this report, nor did they see the final draft before its release. The review of this report was overseen by **ROBERT M. KAPLAN,** Stanford University, and **BOBBIE BERKOWITZ,** University of Washington. They were responsible for making certain that an independent examination of this report was carried out in accordance with the standards of the National Academies and that all review comments were carefully considered. Responsibility for the final content rests entirely with the authoring committee and the National Academies.

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Preface

All children deserve the opportunity to meet their full health potential and lead a fulfilling life. Our nation's future depends on it. Yet, there are millions of children in the United States who are not afforded this opportunity today. While spending a record amount of money on health care services, the United States has the worst infant mortality rate among 19 similar wealthy nations, and the U.S. maternal mortality rate in 2018 was our highest since 2000. Although the United States is one of the richest nations in the world, in 2015 more than 9.6 million children lived in families with annual incomes below the poverty line (based on the Supplemental Poverty Measure), with approximately 2.1 million living in deep poverty. The highest rates of poverty were found among Hispanic, African American, and American Indian/Alaska Native families. This is deeply concerning because poverty during pregnancy and childhood is directly tied to poor health and developmental outcomes. Our nation's health disparities, of which there are many, are directly linked to what happens in early children and prenatally (and even earlier). For all children to lead fulfilling lives, we need to first achieve health equity as a nation, and to do so, we must focus on the youngest, and most vulnerable, in our nation. We also need to look beyond health care for solutions; while health care is necessary to improve health outcomes, fixing health care alone will not address health inequities.

A multitude of factors, from the macro to the micro levels, contribute to the divergent health trajectories that children experience. A child's health ecosystem is influenced by social, economic, cultural, and environmental factors that impact healthy development and well-being. These influences start before birth and have an impact throughout an individual's life and across generations. Exposure to positive influences consistently and longitudinally increases the likelihood of health production, while exposure to negative influences decreases opportunities to be healthy. The timing of these exposures in life also matters—the prenatal to early childhood period is one of the most sensitive times for children to get on the right track to meet their full health potential. Lifelong and multigenerational health disparities are a result of children in this critical age group lacking access to positive opportunities (such as high-quality early care and education, stable and safe housing, and healthy foods) that promote health combined with a preponderance of negative influences that harm health trajectories. Children's health is inextricably linked to family health and community health. For many communities, population health disparity gaps are widening. Persistent, additive disadvantages and early adversity are significant contributors to the widening gaps. Past historical injustices, such as segregated schooling laws, redlining, and assimilation policies, continue to impact children due to structural injustices put in place in the past that persevere today and continue to create barriers to health for those who live in contexts that undermine their opportunity to reach their health potential. This has led to persistent childhood (and lifelong) health disparities. Communities of color have much higher rates of preterm birth, infant mortality, chronic disease (e.g., diabetes), and exposure to adverse childhood experiences, to name just a few.

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In preparing this report, the committee took seriously its charge to review the ways in which early life stress affects health, the pathways by which health disparities develop and persist, and the roadmap needed to get all children on positive health trajectories. Scientific discoveries have built a solid base of evidence about what impacts children's health trajectories positively and negatively—now is the time to apply and advance science to chart a course of action to get all children back on track for health. During the committee's time reviewing the scientific evidence for how to translate the best science into action to positively impact health during early childhood, we strove to close the disconnect between evidence and practice in the nation today. While some scientific evidence has laid the groundwork for actionable practice, policy, and systems solutions, other emerging scientific findings are ripe for further research and inquiry. The committee also acknowledged that achieving and sustaining health equity is a long-term goal with many interrelated strategies and tactics. Thus, we included some recommendations that can be feasibly implemented more quickly by a focused group of actions, while other recommendations may take longer and will require broad support from many different actors at all levels of society.

This report details the latest scientific information about factors impacting health and how to achieve equitable promotion of health for all children. Multilevel and multipronged strategies focused on prevention, early detection and referral, and mitigation are needed to gain momentum toward achieving health equity. These strategies involve intervening at the policy, system, and program levels—this will ultimately require a concerted effort from the nation to distribute resources where they are needed and change policies to better align with the science and evidence. With this in mind, where possible, the committee sought to leverage existing resources or systems that serve children as platforms by which to improve and scale services for children. Further, intentional strategies to understand and reduce inequitable outcomes, access, and experiences across communities of different races, linguistic backgrounds, income groups, genders, and geography are needed. Taking action requires a life course lens, multisector collaboration, and ongoing measurement of outcomes that can be assessed longitudinally and across multiple generations. What science teaches us about sensitive periods and the plasticity of the brain and body provides a clear path for action—if we follow that path regarding prevention and mitigation of adversity during this crucial life period, we can turn the tide for our nation's children. This report provides a roadmap for doing so.

The committee is grateful to the Robert Wood Johnson Foundation for appreciating the need for this work and for supporting putting science into action. The committee welcomed this unique opportunity to shine a brighter spotlight on cutting-edge developmental science about how children develop and grow. Furthermore, we appreciated the opportunity to deepen our understanding about how the key principles and tenets of this critical scientific evidence base on optimal development can be made more accessible to prime the public, practitioners, and policy makers for action. It is the committee's hope that this report's bold recommendations will move our nation to practices and policies that center this science, hand in hand with equity, to advance health and well-being for all.

Jennifer E. DeVoe, *Chair*
Committee on Applying Neurobiological and Socio-Behavioral
Sciences from Prenatal Through Early Childhood Development:
A Health Equity Approach

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The committee wishes to thank and acknowledge the many individuals and organizations that contributed to the study process and development of this report. To begin, the committee would like to thank the Robert Wood Johnson Foundation—the study sponsor—for its support of this work.

The committee found the perspectives of multiple individuals and groups immensely helpful in informing its deliberations through presentations and discussions that took place at the committee's public meetings. Speakers provided presentations on the state of the science in several domains and offered promising models for action, which informed the committee's work; these include (in order of appearance) Dwayne Proctor, Paula Braveman, Fernando Martinez, Phil Fisher, Sarah Barclay Hoffman, Robert Kahn, Suzanne C. Brundage, Megan Smith, Lee Beers, Neal Halfon, Milton Kotelchuck, Ron Haskins, Greg Miller, Greg Duncan, Jessica Pizarek, Helena Sabala, Anne Mauricio, and Elisa Nicholas. The committee also heard policy perspectives from state Representative Ruth Kagi, state Senator Elizabeth Steiner Hayward, Bobby Cagle, and state Senator David Wilson—the committee greatly appreciates the perspectives they brought to the discussions.

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Importantly, the committee heard from a number of caregivers who shared their personal stories and experiences with the committee. These discussions helped ground the committee in the lived experiences of the complex issues that the committee needed to tackle in this report, and the committee is incredibly grateful for their bravery in sharing their experiences in a public forum. Thank you to Abraham Gomez, Shalice Gosey, Lori Hernandez, Ana De Jesus, Yesenia Manzo-Meda, Maria Rodgers, and discussants Alexa Bach, Jennifer Eich, Patricia McKenna, and Reggie Van Appelen.

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Acronyms and Abbreviations

ABC	Attachment and Biobehavioral Catch-Up Intervention
ACE	adverse childhood experience
ACH	Accountable Communities of Health
ASD	autism spectrum disorder
AI/AN	American Indian/Alaskan Native
BPH	bisphenol A
BRFSS	Behavioral Risk Factor Surveillance System
CDC	U.S. Centers for Disease Control and Prevention
CHIP	Children's Health Insurance Program
CPS	Child Protective Services
CVD	cardiovascular disease
DLL	dual-language learner
ECE	early care and education
ECHO	Environmental influences on Child Health Outcomes
ED	emergency department
EHB	essential health benefit
EITC	Earned Income Tax Credit
HAS	high-achieving schools
HHS	U.S. Department of Health and Human Services
HomVEE	Home Visiting Evidence of Effectiveness
HPA	hypothalamic-pituitary-adrenal axis
HUD	U.S. Department of Housing and Urban Development
IOM	Institute of Medicine
IPV	intimate partner violence
IUGR	intrauterine growth restriction
LBW	low birth weight
MBH	mental and behavioral health
MIECHV	Maternal, Infant, and Early Child Home Visiting Program
MLP	Medical-Legal Partnership
NFP	Nurse-Family Partnership
NHANES	National Health and Nutrition Examination Survey
NICU	neonatal intensive care unit
NRC	National Research Council
OECD	Organisation for Economic Co-operation and Development
RCT	randomized controlled trial
SDOH	social determinants of health
SES	socioeconomic status
SNAP	Supplemental Nutrition Assistance Program
SPM	Supplemental Poverty Measure

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SSA	U.S. Social Security Administration
SSI	Supplemental Security Income
TANF	Temporary Assistance for Needy Families
TIC	trauma-informed care
WIC	Special Supplemental Nutrition Program for Women, Infants, and Children

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Summary

ABSTRACT

Health inequities have persisted in the United States, and the factors that drive these inequities from preconception through early childhood are complex, interconnected, and systemic; they result from exposures and experiences that children and families encounter throughout their lives and across multiple generations. These exposures accumulate over the life course to exert a cumulative effect on health that is probabilistic, not deterministic. That is, the odds of good health are never fixed; individual exposures, experiences, and choices help set and adjust them over time. Specific subgroups of the population have varying rates of exposure to positive and negative experiences that shape choices and opportunities throughout the life course; therefore, from the very beginning, certain groups have different odds for good or poor health outcomes. Among the factors that may buffer negative outcomes in the early childhood period, supportive relationships between children and the adults in their lives are essential. Furthermore, reducing health disparities by addressing root causes, such as poverty and racism, is foundational to advance health equity.

Biologically, a number of critical systems develop in the prenatal through early childhood periods, and neurobiological development is extremely responsive to environmental influences during these stages. This report provides an overview of the core concepts of brain development and other body systems relevant to understanding the impact of early life adversity, including the mechanisms that link early life experiences to later outcomes. This information can be used by the public and policymakers to better inform effective actions for advancing health equity.

The committee provides both short- and long-term recommendations in several key areas that can be leveraged to improve health outcomes for children and families. Recommendations aimed at supporting caregivers include implementing paid parental leave and strengthening and expanding home visiting programs. Recommendations for creating supportive and stable early living conditions include improving economic security through increases in resources available to families to meet their basic needs; increasing the supply of high-quality affordable housing; and supporting and enforcing efforts to prevent and mitigate the impact of environmental toxicants. To maximize the potential of early care and education (ECE) to promote better health outcomes, the committee recommends developing a comprehensive approach to school readiness that explicitly incorporates health outcomes, developing and strengthening curricula that focus on key competencies of educators, and improving the quality of ECE programs and expanding access to comprehensive high-quality and affordable ECE programs. The committee recommends leveraging the health care system to make care in the preconception through early childhood periods more continuous, equitable, integrative, and comprehensive by transforming services to apply a life course perspective and address the social, economic, cultural, and environmental determinants of health. To mitigate the early-life drivers of health inequities, there is no one-sector solution—the complex and interconnected root causes call for

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coordination across multiple sectors. Therefore, the committee provides recommendations for sector alignment and collaboration, as well as the need for child and family serving sectors to enhance detection of early-life adversity, improve response systems, and develop trauma-informed approaches.

The committee identifies knowledge gaps and recommends multidisciplinary research efforts to bring new ideas and practical approaches to advance efforts to achieve health equity. However, substantial advances in knowledge in the past 20 years make it clear that policy makers, health providers, business leaders, and others in the public and private sectors do not need to wait any longer to take action.

Health inequities by race, ethnicity, socioeconomic status (SES), geography, and other important demographic characteristics have persisted in the United States despite increasing evidence about their contributions to poor health. Research shows that exposures to factors that shape health trajectories can start early and are multigenerational; thus, the preconception, prenatal, and early childhood periods are critical to setting the odds for lifelong health. Importantly, science can inform actions in policy and practice to advance health equity[1] and reduce health disparities. Neurobiological and socio-behavioral research indicate that early life experiences shape prenatal and early childhood development, and these experiences have a powerful impact on the developing brain and peripheral organ systems that impact health outcomes across the life course.

When different groups vary in their exposures to key experiences (both positive and negative), their odds for positive health diverge systematically over time, producing disparities in health outcomes across the life-span and across generations. While these exposures accumulate over the life course to exert a cumulative effect on health that is probabilistic, not deterministic. That is, the odds of positive or negative health are never fixed; individual exposures, experiences, resilience, and choices help set and adjust them over time. Individuals' distinct contexts also shape their choices and opportunities, and thus they have different odds of experiencing positive or negative health outcomes over time. Because the odds of these exposures are affected by policies and systems, advancing health equity will require more than individual-level interventions. It will necessitate systems-level changes, including changes to laws and policies and investment of resources, to improve the odds of positive experiences and reduce the odds of adverse exposures for all populations, especially those experiencing the most adversity.

Scientific evidence shows that prevention and early intervention are effective for children on at-risk developmental trajectories. Recent advances in science, technology, data sharing, and cross-disciplinary collaboration present opportunities to apply this emerging knowledge systematically to practice, policy, and systems changes. Given the burgeoning science available to advance health equity during early development, the Robert Wood Johnson Foundation (RWJF), as part of its Culture of Health Initiative, asked the Health and Medicine Division of the National Academies of Sciences, Engineering, and Medicine to:

1. Provide a brief overview of stressors that affect prenatal through early childhood development and health;
2. Identify promising models and opportunities for translation of the science to action;
3. Identify outcome measures to enable subgroup analyses;

[1] Health equity is the state in which everyone has the opportunity to attain full health potential and no one is disadvantaged from achieving this potential because of social position or any other socially defined circumstance.

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4. Develop a roadmap to apply the science to tailored interventions (i.e., policies, programs, or system changes) based on biological, social, environmental, economic, and cultural needs; and
5. Provide recommendations in these areas, including how systems can better align to advance health equity.

To respond to this charge, the Committee on Applying Neurobiological and Socio-Behavioral Sciences from Prenatal Through Early Childhood Development: A Health Equity Approach was formed. The committee applied a health equity frame and built on the concepts of the 2017 report *Communities in Action: Pathways to Health Equity*. As identified in the 2000 National Research Council and Institute of Medicine report *From Neurons to Neighborhoods: The Science of Early Childhood Development,* prenatal through early childhood are critical phases of development that have lifelong impacts on health and well-being. This report reviews the science that has emerged since that landmark report.

The health of both men and women before they have children is important for not only pregnancy outcomes, but also the lifelong health of their children; thus, the committee included the preconception period as an important focus of the report. The committee also adopted the life course approach to its work because an individual's health status and outcomes reflect the accumulation of experiences over the life-span. This approach takes into account an individual's larger social, economic, and cultural context and acknowledges that the life trajectory may be changed, negatively or positively, through interactions between the brain, body, and environment. Protective factors (such as stable, high-quality caregiver relationships and economic security) support positive, or flourishing, trajectories; risk factors (such as exposure to abuse, neglect, or racism) exacerbate the likelihood of poor trajectories.

Children in the United States may be perceived to be healthier now than in the past because they are much less likely to encounter the major infections and debilitating diseases of past generations and are typically able to recover fully from "acute" childhood illnesses. However, ailments of the past have been supplanted with chronic physical (e.g., diabetes, asthma, obesity) and socio-emotional (e.g., depression, anxiety) conditions, with large subsets of U.S. children facing barriers to positive mental and physical health and well-being as a result of poverty, food insecurity, unsafe or unstable housing, neighborhood segregation, and other substantial adversities (such as adverse childhood experiences [ACEs]) in the first few years of life.

Children who are born and raised in poverty are at particularly high risk for poor health outcomes, more problems in early development (e.g., lack of readiness for school at age 5, diagnoses of developmental delays and/or disorders), and higher rates of most childhood chronic conditions (such as mental illness, developmental disabilities, obesity, and asthma). Early adverse experiences may have intermediate effects on school readiness, weight, and physical and/or mental well-being and contribute to chronic disease and poor functioning in adulthood. In fact, these impacts are cumulative, and adults who experience adversity in childhood have substantially higher rates of heart disease, lung disease, metabolic syndrome, and other costly health conditions.

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CONCEPTUAL MODEL

The committee's conceptual model (See Figure S-1) served as a unifying framework for its approach to this report. It is important to note that risk and protective factors can be transferred intergenerationally, which makes parents and other family and community primary caregivers a central focus of interventions to improve child health. Within the context of the life course, the diagram's nested circles illustrate the complex sociocultural environment that shapes development at the individual level and the opportunities for interventions to improve individual health and developmental outcomes, as well as population health, well-being, and health equity. Individual social and biological mechanisms and culture operate and interact within and across the three levels.

Structural inequities operate at the outer level, the "socioeconomic and political drivers." Structural inequities are deeply embedded in policies, laws, governance, and culture; they organize the distribution of power and resources differentially across individual and group characteristics (i.e., race, ethnicity, sex, gender identity, class, sexual orientation, gender expression, and others). The next level represents social, economic, cultural, and environmental states (i.e., the social determinants of health [SDOH]). In the model, these interdependent factors are grouped into three domains—healthy living conditions, health care, and education. These domains were identified by the committee based on the available evidence and existing resources as important for targeting prenatal and early childhood interventions and are the primary foci of Chapters 5–7. The next level represents the factors that most directly and proximally shape children's daily experiences and routine patterns: family cohesion and social connections, which also affect access to critical resources for health, well-being, and development in early life (Chapter 4). The innermost circle and crosscutting arrows—biological, psychological, and socio-behavioral development—are the focus of Chapters 2 and 3.

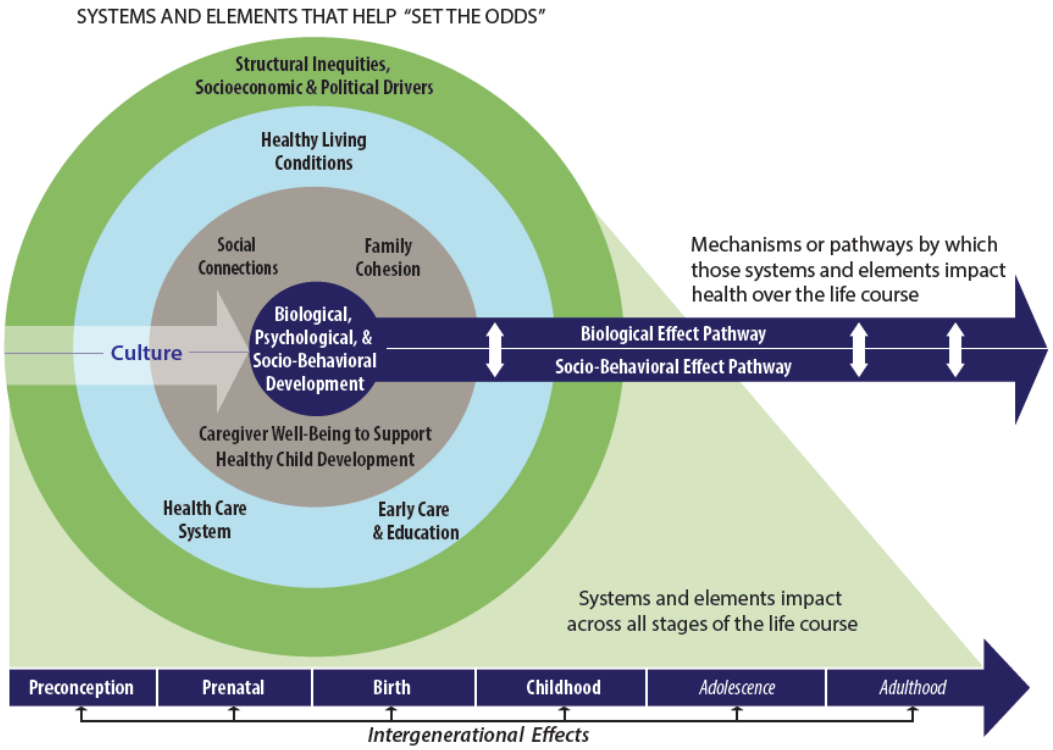

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FIGURE S-1 Leveraging early opportunities to advance health equity across the life course: A conceptual framework.

THE SCIENCE OF EARLY DEVELOPMENT: CORE CONCEPTS

Based on its review of the science, the committee updated and adapted the core concepts from the 2000 report *From Neurons to Neighborhoods* and identified 12 core concepts of early development, with a focus on health equity. The evidence underlying these concepts is described in Chapters 2–4, and this evidence guided the committee in its development of recommendations that apply the science of early development. In brief, these concepts include the following (see Chapter 1 for more detailed descriptions of each concept):

1. Biology–environment interaction impacts health and development.
2. Brain development proceeds in well-defined but continuous steps.
3. Major physiological systems develop rapidly during pregnancy and early childhood.
4. The early caregiving environment is crucial for long-term development.
5. The developing child plays an important role in interactions and development.
6. The development of executive functions is a key aspect of early childhood development.
7. Trajectories—positive or negative—are not immutable.
8. There is variability in individual and group development.
9. Experiences across environmental contexts play a significant role in early development.
10. Disparities in access to critical resources matter.
11. Health outcomes are the result of experiences across the life course.
12. Early interventions matter and are more cost effective than later ones.

A large body of recent research provides insights into the mechanisms by which early adversity in the lives of young children and their families can change the timing of sensitive periods of brain and other organ systems development and impact the "plasticity"[2] of developmental processes. In the past two decades, there has been a convergence of research that has led to many of the advances described in this report. First, a wave of neurobiological studies in model systems and humans found that responses to pre- and postnatal early life stress are rooted in genetic and environmental interactions that can result in altered molecular and cellular development that impacts the assembly of circuits during sensitive periods of development. The demonstration that certain systems involved in cognitive and emotional development are more sensitive to early disturbances that activate stress response networks, such as the frontal cortex, hippocampus, amygdala, and the hypothalamic-pituitary-adrenal axis (HPA), provided a basis for both short- and long-term functional consequences of early life stress.

Many of the new scientific advances in neuroscience are still in development, and more research is needed to apply these new findings in clinical and public health practice and to use them to inform policies. In particular, greater effort and support are needed to develop, implement, and evaluate programs based on scientific discoveries regarding the optimal timing for interventions. However, new research has clarified that altered nutrition, exposure to

[2] The process by which neurons within the brain change their gene expression, cellular architecture and connections with other neurons, and function in response to experiences and changes in the environment (i.e., change over time).

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environmental chemicals, and chronic stress during specific times of development can lead to functional biological changes that predispose individuals to manifest diseases and/or experience altered physical, socio-emotional, and cognitive functions later in life. The committee provides information about major biological responses to stressors and new discoveries that have contributed to advancing knowledge about how and when to intervene to improve health outcomes for children.

ROADMAP FOR APPLYING AND ADVANCING THE SCIENCE OF EARLY DEVELOPMENT

With the goal of decreasing health inequities, the broad question this report addresses is, "For those children who are placed at risk for negative outcomes, what can be done—guided by science-based evidence—to expediently and effectively move each of them toward positive developmental health trajectories?" In this report, the committee provides recommendations for practice, policy, and systems changes to achieve this goal. The roadmap the committee has put forth includes a suite of key strategies to advance health equity[3]:

- **Intervene early**—In most cases, early intervention programs are easier to implement, more effective, and less costly.
- **Support caregivers**—This includes both primary caregivers and caregivers in systems who frequently interact with children and their families.
- **Reform health care system services to promote healthy development**—Redesign the content of preconception , prenatal, postpartum, and pediatric care while assuring ongoing access, quality, and coordination.
- **Create supportive and stable early living conditions:**
    - **Reduce child poverty and address economic and food security,**
    - **Provide stable and safe housing, and**
    - **Eliminate exposure to environmental toxicants.**
- **Maximize the potential of early care and education to promote health outcomes.**
- **Implement initiatives across systems to support children, families, other caregivers, and communities**—Ensure trauma-informed systems, build a diverse and supported workforce, and align strategies that work across sectors.
- **Integrate and coordinate resources across the education, social services, criminal justice, and health care systems, and make them available to translate science to action.**

In this report, the committee provides a range of recommendations for practice, policy, and systems changes, including recommendations that will take time and sustained commitment to achieve and recommendations that could be implemented immediately or in the near term. Some of the committee's recommendations will be difficult to implement; however, the degree of difficulty in implementing any given recommendation does not determine the value of pursuing it. Where possible, the committee also recommends or highlights ways to leverage existing programs that either embrace the core scientific principles laid out above or have a

[3] Note that recommendations are not always presented in numerical order, as the summary has grouped them by topic in some cases; however, all report recommendations are presented.

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strong basic structure from which to build.

Supporting Family Cohesion and Social Connections

The construct of resilience[4] from developmental science is important, as it implies the ability to correct what otherwise might have been negative trajectories, given major life stressors. To set the foundation for the committee's considerations on the topic of supporting family systems, the committee discusses universal principles of human development pertaining to the broad domain encompassing children's psychological and behavioral adjustment (see Chapter 4). For example, for children, the single most important factor in promoting positive psychosocial, emotional, and behavioral well-being is having a strong, secure attachment to their primary caregivers—usually their mothers. Strong attachment presupposes effective parenting behaviors in everyday life, and "effective parenting" changes in complexity with development over time.

There is an urgent need to develop preventive interventions well suited for fathers and other male caregivers; existing approaches that are developed for and tested with women cannot be assumed to generalize to other caregivers with equal effectiveness (e.g., in the successful recruitment, retention, and support of men and fathers who take care of young children).

> **Recommendation 4-1: Federal, state, and local agencies, along with private foundations and philanthropies that invest in research, should include in their portfolios research on the development of preventive interventions that target fathers and other male caregivers. Special attention should be given to the recruitment, retention, and support of men and fathers parenting young children from underserved populations.**

Specific subgroups of children have unique needs and challenges when adjusting to adversity. Careful attention to potent subculture-specific processes need to be considered in working with subgroups well known to face serious inequities in relation to mental health—including families experiencing chronic poverty, immigrants, lesbian, gay, bisexual, transgender, and queer (LGBTQ) children, LGBTQ parents, children who are separated from parents due to incarceration, foster care, or other reasons, and children exposed to high achievement pressures, usually in relatively affluent communities.

> **Recommendation 4-2: Federal, state, local, tribal, and territorial agencies, along with private foundations and philanthropies that invest in research, should include in their portfolios research on the development of interventions that are culturally sensitive and tailored to meet the needs of subgroups of children known to be vulnerable, such as those living in chronic poverty, children from immigrant backgrounds, children in foster care, and children with incarcerated parents.**

In addition to addressing major goals relevant for children in general (e.g., fostering caregiver well-being and minimizing maltreatment), programs need to include components

[4] There are two essential conditions that make up resilience: (1) exposure to significant threat or severe adversity and (2) achievement of positive adaptation despite major assaults on the developmental process.

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that specifically address unique risk and protective processes within these subgroups of children.

A growing body of evidence suggests that home visiting by a nurse, a social worker, or an early educator during pregnancy and as needed in the first years of a child's life improves a wide range of child and family outcomes, including promotion of maternal and child health, prevention of child abuse and neglect, positive parenting, child development, and school readiness. These positive effects continue well into adolescence and early adulthood. Researchers, program leaders, and policy makers need to focus on expanding the concept of tailored home visiting to advance knowledge on which programs and activities are best for which family, in which communities, and for what outcomes.

Recommendation 4-3: To strengthen and expand the impact of evidence-based home visiting programs:

- **Federal policy makers should expand the Maternal, Infant, and Early Childhood Home Visiting Program.**
- **The Health Resources and Services Administration and the Administration for Children and Families should work with program developers to increase flexibility for states and communities, to tailor the program to the needs and/or assets of the community or population being served.**
- **Federal, state, local, tribal, and territorial agencies overseeing program implementation should continue to strengthen programmatic coordination and policy alignment between home visiting, other early care and education programs, and medical home.**

State policy makers should further expand support for evidence-based home visiting services through the use of general funds, Medicaid, and a combination of multiple funding streams. Health Resources and Services Administration (HRSA) and Administration for Children and Families (ACF) should support research to continue to assure program effectiveness and accountability of the expanded program. Expansion of home visiting programs should be done in conjunction with the expansion of other public investments and services.

Intervention trials have shown strong benefits of relational interventions, such as interventions to foster strong attachments, and group-based supports in communities for caregivers and their families. The core components of several effective interventions suggest that in addition to providing particular skills, improving the overall well-being of caregivers (especially mothers) is the most critical "engine" of change.

Recommendation 4-4: Policy makers at the federal, state, local, territorial, and tribal levels and philanthropic organizations should support the creation and implementation of programs that ensure families have access to high-quality, cost-effective, local community-based programs that support the psychosocial well-being of the primary adult caregivers and contribute to building resilience and reducing family stress.

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It is necessary to consider measures that should be included in evaluating results of large-scale preventive interventions targeting young children and their mothers. Given the need to identify individuals at risk for early adversity and the toxic stress response, regular brief assessments of the mothers' depressive symptoms, stress, feelings of rejection to the child, any involvement with child protective services, and the degree to which they have positive, buffering relationships in their lives, should occur routinely.

Recommendation 4-5: Health care providers who care for pregnant women and children should routinely track levels of individual health and social risk among mothers and children over time, using periodic assessments via a short set of scientifically validated measures.

Leveraging the Health Care System to Promote Health Equity

The health care system can serve as a platform, along with public health and other sectors, to address the social determinants that underlie many health inequities. However, the current health care system focuses mainly on clinical goals and addresses other determinants of health in fragmented and highly variable ways. U.S. health care provides only limited attention to integration of health care for the whole family, health care across the life course, or integration of mental and behavioral health into clinical care. Recognizing that the preconception through early childhood period are sensitive and important life periods to optimize health outcomes, care during these periods needs to become more continuous (access), equitable (quality), integrative (delivery), and comprehensive (content); therefore, the committee offers the following recommendations:

Improving Access to Health Care

Recommendation 5-1: The U. S. Department of Health and Human Services, state, tribal, and territorial Medicaid agencies, public and private payers, and state and federal policy makers should adopt policies and practices that ensure universal access to high-quality health care across the life course. This includes:

- **Increasing access to patient- and family-centered care,**
- **Ensuring access to preventive services and essential health benefits, and**
- **Increasing culturally and linguistically appropriate outreach and services.**

Achieving this recommendation will require actively promoting inclusion in coverage and care.

Improving Quality of Care

Recommendation 5-2: To expand accountability and improve the quality of preconception, prenatal, postpartum, and pediatric care:

- **Public and private payers should include new metrics of child and family health and well-being that assess quality using a holistic view of health and health equity. Federal, state, and other agencies, along with private foundations and philanthropies that invest in research, should support the**

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development and implementation of new measures of accountability, including key drivers of health, such as social determinants, along with measuring variations by key subgroups to determine disparities;

- **Public and private payers, including the Health Resources and Services Administration's (HRSA's) Bureau of Primary Care and Maternal and Child Health Bureau, Centers for Disease Control and Prevention, Centers for Medicare & Medicaid Services (CMS), and perinatal and pediatric quality collaboratives, should expand the use of continuous quality improvement, learning communities, payment for performance, and other strategies to enhance accountability; and**
- **Health care–related workforce development entities should expand efforts to increase diversity, inclusion, and equity in the health care workforce, including diversity-intensive outreach, mentoring, networking, and leadership development for underrepresented faculty and trainees.**

Needed metrics include social determinants and social risk measures; cross-sector developmental measures that move beyond common indicators of child development, including mental and behavioral health; and disparities as explicit measurement domains that hold providers accountable for not just delivering services but also improving outcomes. Workforce development (as noted in bullet 3) will need to be addressed by several entities, including the Accreditation Council for Graduate Medical Education (ACGME) and specialty boards, professional schools, training programs, teaching hospitals, including children's hospitals, and funders of graduate education in health professions (CMS, HRSA, and others).

Organization and Integration of Health Care Services

Recommendation 5-3: The U.S. Department of Health and Human Services, state, tribal, and territorial government Medicaid agencies, health systems leaders, and state and federal policy makers should adopt policies and practices that improve the organization and integration of care systems, including promoting multidisciplinary team-based care models that focus on integrating preconception, prenatal, and postpartum care with a whole-family focus, development of new practice and payment models that incentivize health creation and improve service delivery, and structures that more tangibly connect health care delivery systems to other partners outside of the health care sector.

Achieving this recommendation will require disseminating multidisciplinary team-based care models in community settings; developing integrated models for preconception, prenatal, postpartum, and pediatric care delivery modes; adopting and spreading integrated, whole-family and family-centered care models; developing and using new technologies that improve care and improve accessibility; aligning payment reform with health creation rather than service delivery; and developing systemic and cross-sector collaboration.

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Transforming the Content of Care

Recommendation 5-4: Transform preconception, prenatal, postpartum, and pediatric care to address the root causes of poor health and well-being—the social, economic, environmental, and cultural determinants of health and early adversity—and to align with the work of other sectors addressing health equity.

The U.S. Department of Health and Human Services should convene an expert panel to reconceptualize the content and delivery of care, identify the specific changes needed, develop a blueprint for this transformation, and implement a plan to monitor and revise the blueprint over time.

Implementation of this recommendation will require:

- **An update of clinical care guidelines and standards by the Women's Preventive Services Initiative, Bright Futures, American College of Obstetricians and Gynecologists, American Academy of Pediatrics, American Academy of Family Physicians, and others actively developing clinical care guidelines and standards to include this new content of care;**
- **Medical accreditation bodies, relevant programs, and agencies to develop performance monitoring and quality improvement based on this new content of care;**
- **Clinical care educational authorities, such as the Accreditation Council for Graduate Medical Education, to develop curricula, training, experiences, and competencies based on the updated guidelines; and**
- **Public and private payers to cover services reflecting this new content of care.**

This work should take place in a larger framework of social and reproductive justice and include more diverse voices, especially from communities most affected by adverse birth and child health outcomes. Such a shift will require that the health care system recognize the impact of both adverse and enriching experiences across the life course and cumulative effects on health and well-being by the health care system. It will also require integrating attention to social and environmental determinants as well as trauma assessment and response into clinical practice.

Although health care plays an integral role in advancing health equity, health care alone cannot meaningfully address health inequities, nor is it the primary actor or leader. Cross-sectoral and multidisciplinary collaboration is essential for decreasing health inequities.

Creating Healthy Living Conditions for Early Development

Reducing or managing caregiver stress is key to giving caregivers the capacity, supports, and resources to care for their children and serve as buffers against adversity. Addressing the primary needs of families and children is critical to achieving this goal. The committee identified four areas of fundamental needs that, if met, would have an impact on health inequities: (1) food security, (2) safe and stable housing, (3) economic stability and security, and (4) safe physical environments.

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Food Security

Given the importance of good nutrition for brain growth and development (during the preconception, prenatal, and early childhood periods), the committee concludes that providing resources to ensure families have access to sufficient and healthy foods can improve birth and child health outcomes. Because safety net programs such as WIC and SNAP have been shown to improve birth and child (and adult) health outcomes and to reduce food insecurity, the committee recommends:

Recommendation 6-2: Federal, state, local, territorial, and tribal agencies should reduce barriers to participation to the Special Supplemental Nutrition Program for Women, Infants, and Children (WIC) program and Supplemental Nutrition Assistance Program (SNAP) benefits. Receipt of WIC and SNAP benefits should not be tied to parent employment for families with young children or for pregnant women, as work requirements are likely to reduce participation rates.

Safe and Stable Housing

Evidence suggests that lack of affordable or quality housing, housing instability, and overcrowding have significantly detrimental effects to the health, well-being, and development of infants, children, and families. Housing affordability and quality is an acute problem that disproportionately affects people of color and contributes to health inequities among children. Over half of black and Hispanic renters live in unaffordable housing, and health issues related to poor-quality housing, such as elevated blood lead levels and asthma, are more prevalent among these renters. Current federal housing programs are not adequately funded, and there are not enough safe, affordable housing units in high-opportunity areas. Additional funding for programs can move families out of poverty and allow them to reallocate money for other basic needs that support child health and development. Incentives and/or regulations, along with enhanced programming, can increase the supply of affordable housing. Recognizing the centrality of housing to health and healthy child development, the committee recommends:

Recommendation 6-3: The U.S. Department of Housing and Urban Development, states, and local, territorial, and tribal public housing authorities should increase the supply of high-quality affordable housing that is available to families, especially those with young children.

Recommendation 6-4: The Secretary of the U.S. Department of Health and Human Services, in collaboration with the U.S. Department of Housing and Urban Development and other relevant agencies, should lead the development of a comprehensive plan to ensure access to stable, affordable, and safe housing in the prenatal through early childhood period. This strategy should particularly focus on priority populations who are disproportionately impacted by housing challenges and experience poor health outcomes.

Recommendation 6-5: The Center for Medicare & Medicaid Innovation should partner with states to test new Medicaid payment models that engage providers and other community organizations in addressing housing safety concerns,

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especially focused on young children. These demonstrations should evaluate impact on health, health disparities, and total cost of care.

Economic Stability and Security

Children's well-being and life course outcomes are strongly related to family income. Given the strong evidence that economic security matters, an important factor in reducing health disparities in early childhood is to ensure that families with young children have adequate resources. The committee concludes that public programs that provide resources to families in the form of cash, tax credits, or in-kind benefits improve childhood well-being and life course outcomes and that these effects are long lasting. Furthermore, while income-support programs that are contingent on employment status or based on earned income have positive benefits for families, they should avoid regulations and policies that might have unintended consequences for childhood outcomes through negative effects on family relationships and attachments, breastfeeding, and caregiver stress.

Additional income support for families with young children through paid parental leave would recognize the special needs of infants and their caregivers. Unpaid parental leave through the Family and Medical Leave Act does not cover all employees, and most families with low incomes cannot afford to take an unpaid leave.

Recommendation 6-1: Federal, state, local, tribal, and territorial policy makers should implement paid parental leave. In partnership with researchers, policy makers should model variations in the level of benefits, length of leave, and funding mechanisms to determine alternatives that will have the largest impacts on improving child health outcomes and reducing health disparities.

Recommendation 6-6: Federal, state, tribal, and territorial policy makers should address the critical gaps between family resources and family needs through a combination of benefits that have the best evidence of advancing health equity, such as increased Supplemental Nutrition Assistance Program benefits, increased housing assistance, and a basic income allowance for young children.

This recommendation focuses on strategies that are likely to have particularly important impacts on health outcomes for young children. A child allowance would fill in some of the gaps in the current safety net and particularly benefit the lowest-income children and those most at risk of poor health outcomes. The key advantage of a child allowance (over, for example, tax credits) is that funds are available to families on an ongoing, monthly basis rather than once per year. In addition, under the current structure of the child and working-family tax credits, the lowest-income families receive few benefits. Children whose parents are in unstable employment or not employed suffer the short- and long-term health consequences of living in poverty. Reducing health disparities requires reaching these children during their earliest years, regardless of parental employment. Increased SNAP benefits and housing allowances would address current inadequacies in both of these programs and provide targeted support for the critical food and housing needs of young children.

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Environmental Exposures and Exposure to Toxicants

There are numerous potential environmental toxicants that may be transmitted through the air, water, soil, and consumer products with which food and water come into contact. Many of these occur naturally in the environment (e.g., arsenic, radon, etc.), and many more are released through human-based processes (e.g., heavy metals, chemicals from plastic production and degradation, and particulates). The embryonic, fetal, and early childhood periods represent greater risk than adulthood for adverse mental and physical health outcomes from environmental exposures due to children's smaller size, proportionally large intake of food, air, and water to body weight, and rapid developmental processes that may be influenced and disrupted by chemicals and toxicants. As a result of toxicant exposures, children may suffer from a variety of developmental problems, chronic conditions, and even premature death. Poverty, substandard and/or unstable housing, minority racial/ethnic status, and proximity to known sources of pollutants heighten children's risk of exposure and poor health and developmental outcomes. The committee identified three areas where current efforts could be improved to prevent and mitigate the impact of environmental toxicants in the prenatal through early childhood periods:

Recommendation 6-7: The Administration for Children and Families, Maternal and Child Health Bureau, and federal and state regulators should strengthen environmental protection in early care and education settings through expanded workforce training, program monitoring, and regulations.

Recommendation 6-8: Professional societies, training programs, and accrediting bodies should support expanded or innovative models training of prenatal and childhood health care providers on screening, counseling, and interventions to prevent or mitigate toxic environmental exposures.

Recommendation 6-9: Federal, state, local, tribal, and territorial governments should support and enforce efforts to prevent and mitigate the impact of environmental toxicants during the preconception through early childhood period. This strategy should particularly focus on priority populations who are disproportionately impacted by harmful environmental exposures. This includes:

- **Environmental Protection Agency fully exercising the authorities provided by Congress to safeguard children's environmental health under the Toxic Substance and Control Act as amended by the Frank R. Lautenberg Chemical Safety for the 21st Century Act.**
- **Continued allocation of resources and technical assistance from the federal government through Center for Disease Control (CDC), EPA, Food and Drug Administration (FDA), and the U.S. Consumer Product Safety Commission to translate existing data and research findings into actionable policies and practices.**
- **Ongoing review and updating of environmental exposure levels by federal agencies to reflect health and safety standards specific to the unique vulnerability of children (from fetal development through early development).**

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In Chapter 6, the committee also discusses the role of civil rights strategies to promote healthy communities for developing children.

Promoting Health Equity Through Early Care and Education

While most of the attention on early care and education (ECE)[5] has focused on whether it improves children's cognitive and socio-emotional development and academic readiness, research shows that ECE affects various other child health outcomes, including children's physical, emotional, and mental health. ECE programs increase children's cognitive, social, and health outcomes through enhancing their motivation for school and readiness to learn and the early identification and intervention of problems that impede learning. This, in turn, helps children improve their cognitive ability and social and emotional competence, while increasing their access to and use of preventive health care. Access to ECE may lead to lower risk of dropping out of school, greater school engagement, and subsequently better educational attainment, which lead to increased income and decreased social and health risks, resulting in greater health equity.

Allocation of Adequate Resources to Support ECE Programs and Educators

Intentional policies and allocation of adequate resources to support these programs and educators are needed for ECE programs to contribute significantly to a health promotion and equity strategy.

Recommendation 7-1: The committee recommends that early care and education (ECE) systems and programs, including home visiting, adopt a comprehensive approach to school readiness. This approach should explicitly incorporate health promotion and health equity as core goals. Implementing this approach would require the following actions:

- **Federal, state, local, tribal, and territorial governments and other public agencies (e.g., school districts, city governments, public–private partnerships) that have decision-making power over ECE programs should establish program standards and accountability systems, such as a quality rating and improvement system, linked with better school readiness and health outcomes and provide adequate funding and resources to implement and sustain these standards effectively.**
- **The Office of Child Care and the Office of Head Start at the federal level, along with state, local, tribal, and territorial early care and other education agencies, should assess the full cost of implementing standards that promote health outcomes and equity as described above, including supporting educators' own health and well-being, and work with Congress to align funding levels of the major federal ECE programs—child care subsidy and Head Start—accordingly.**
- **Health and human service entities, the federal Early Learning Interagency Policy Board, state Early Childhood Advisory Councils, and federal, state,**

[5] ECE can be defined as nonparental care that occurs outside the child's home. ECE services may be delivered in center-, school-, or home-based settings.

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local, tribal, and territorial agencies that oversee home visiting and ECE programs should ensure greater programmatic coordination and policy alignment to ensure effective allocation of resources.

- **The Office of Planning & Research, and Evaluation in the Administration for Children and Families along with the U.S. Department of Education, should examine the feasibility and seek resources to conduct (a) an implementation study to examine the design and implementation of this comprehensive ECE approach that incorporates health standards and (b) an outcomes study that examines the impact on children's school readiness and achievement, and health outcomes, with particular attention to eliminating disparities and gaps prior to school entry.**

Health-Focused Competencies of the ECE Workforce

Policies and systems that prepare and support early childhood educators and program leaders, including those in public schools, need to incorporate the latest evidence about how to support children's school readiness and success by fostering their health and well-being. This would entail providing comprehensive supports and resources to degree granting institutions and preparation programs, including the development of curricula, textbooks, practicum experiences, toolkits, and fact sheets.

Recommendation 7-2: Building off the 2015 Institute of Medicine and National Research Council report *Transforming the Workforce for Children Birth Through Age 8*, the committee recommends that degree granting institutions, professional preparation programs, and providers of ongoing professional learning opportunities develop or strengthen coursework or practicums that focus on competencies of educators, principals, and early care and education program directors that are critical to children's health, school readiness, and life success.

Access and Affordability to ECE Programs

Maximizing the impact of ECE on positive early childhood development and health and well-being at the community or population level will require increasing public funds for ECE programs. Currently, eligibility for ECE programs is limited, and among eligible families, access is low due to lack of funding and availability of programs and services. Therefore, even if existing publicly funded programs have the resources to provide robust supports that improve young children's health and well-being, these will not reach most children, especially those who live in low-income households or confront adverse experiences and toxic stress.

Recommendation 7-3: Federal, state, local, tribal, and territorial policy makers should work with the U.S. Department of Health and Human Services (HHS), the Office of Head Start, and Office of Child Care develop and implement a plan to:

a. improve the quality of early care and education (ECE) programs by adopting the health-promoting standards discussed in Recommendation 7-

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1, such as building on the performance standards of Early Head Start and Head Start, and

b. within 10 years, expand access to such comprehensive, high-quality, and affordable ECE programs across multiple settings to all eligible children. Disproportionately underserved populations should be prioritized.

The Secretary of HHS should conduct a process evaluation to inform the expansion effort, and, once implemented, conduct rigorous and comparative outcomes studies to ensure that the expansion is having the intended impacts on children and families, with particular attention on what group(s) may be benefitting.

The strategic plan should be modeled after and build upon the relevant performance standards of Early Head Start and Head Start, which emphasize mixed settings, the whole child, family and community engagement, transition between home and school, and continuous quality improvement. It should also strengthen those program components discussed in Chapter 7 that lead to stronger school readiness and health outcomes, including mitigation of the impact of adverse experiences and toxic stress for children, families, teachers, and staff. Critical components include a comprehensive social-emotional strategy that encompasses both the classroom (curriculum, teacher training and support) and program/school (leadership, culture and climate) levels and educators who have competencies described in Recommendation 7-2.

Systems Approach

Advancing health equity in the preconception through early childhood periods cannot be achieved by any one sector alone—it will take action, collaboration, and alignment across all sectors that frequently interact with children, families, and the professionals who serve them. Systems are a collection of interacting, interdependent parts that function as a whole. For the purposes of this report, most of the systems are social constructs and are organized around a key functional area (e.g., education, health care, housing). Systems change is not an easy strategy, it seldom yields speedy returns, and it may not be sufficient without an investment of resources designed to take advantage of new and better aligned approaches. However, given that disparities are systematically generated, it is likely a necessary precursor to real and widespread advances in health equity. The committee identified eight crosscutting recommendation areas where multiple sectors need to take action, based upon review of the evidence in Chapters 1–7 and the committee's collective expertise. In brief (additional details available in Chapter 8), the committee recommends:

Policy makers and leaders in the health care, public health, social service, criminal justice, early care and education/education, and other sectors should:

- **Recommendation 8-1: Support and invest in cross-sector initiatives that align strategies and operate community programs and interventions that work across sectors to address the root causes of poor health outcomes. This includes addressing structural and policy barriers to data integration**

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and cross-sector financing and other challenges to cross-sector collaboration.

- **Recommendation 8-2: Adopt and implement screening for trauma and adversities early in life to increase the likelihood of early detection. This should include creating rapid response and referral systems that can quickly bring protective resources to bear when early-life adversities are detected, through the coordination of cross-sector expertise, as covered in Recommendation 8-1.**
- **Recommendation 8-3: Adopt best practices and implement training for trauma-informed care and service delivery. Sector leadership should implement trauma-informed systems that are structured to minimize implicit bias and stigma and prevent retraumatization. Standards for trauma-informed practice exist in a variety of service sectors, including health care and social services; those standards should be replicated and implemented across systems.**
- **Recommendation 8-4: Develop a transdisciplinary and diverse workforce to implement culturally competent service delivery models. The workforce should reflect the diversity of populations who will engage in sector services.**
- **Recommendation 8-5: Improve access to programs or policies that explicitly provide parental or caregiver supports and help build or promote family attachments and functioning by engaging with the families as a cohesive unit. For families with intensive support needs, develop programs or initiatives designed to provide comprehensive wraparound supports along a number of dimensions, such as health care, education, and social services, designed to address needs related to the social determinants of health that are integrated and community based.**
- **Recommendation 8-6: Integrate care and services across the health continuum, including the adoption of models that provide comprehensive support for the whole person in a contextually informed manner, leveraging and connecting existing community resources wherever possible, with a focus on prevention.**
- **Recommendation 8-7: Invest in programs that improve population health and in upstream programs that decrease long-term risk and poor health outcomes. These changes should be accompanied by accountability metrics to ensure that the spending is tangibly and demonstrably in service to the goals behind the original funding, but offer more flexibility in how those goals are achieved.**

Crosscutting Research Needs

A tremendous amount is known about what works to advance health equity in early development (and the lifelong benefits of doing so), and efforts to translate this science into action and to scale up effective interventions needs to be accelerated. Many interventions have shown promising results at small scale but have not been fully tested across multiple settings or in diverse communities and populations. Others have promising preliminary data but require more evidence. In addition, the evidence around systems and policy changes—the work needed

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to address inequities with a multisector and systems-based approach—remains less certain than programmatic evidence in many cases precisely because it is complex and set in shifting environments that make confident attribution of effects challenging. In Chapter 8, the committee recommends newly designed and adapted research strategies to help translate science to action across sectors, including needed data to inform subgroup analyses and elucidate the complex causality related to health inequities to better design interventions across sectors.

An important caution, however: although targeted research is needed to address population heterogeneity with more precision, enough is already known to act now to advance health equity in the prenatal and early childhood periods. The research recommended below is important to continually improve efforts and increase impact but should not impede action. Here the committee provides guidance on charting the course for future research to better meet the health and social needs of the nation's children in the future and, specifically, to advance health equity.

Recommendation 8-8: The National Institutes of Health, Agency for Healthcare Research and Quality, Centers for Disease Control and Prevention, Health Resources and Services Administration, Centers for Medicare & Medicaid Services, U.S. Department of Education, philanthropies, and other funders should support research that advances the state of the science in several critical ways to advance health equity. Specific actions and research to support include:

- Explore alternative methods to address complex causality
- Expand research into individual differences (heterogeneity) in response to adversity and treatment
- Promote scientific research that includes individuals and families from underrepresented communities
- Promote research that explicitly seeks to understand the interconnected mechanisms of health inequities
- Support research that addresses discrimination and structural racism
- Support research for trauma-informed care and implicit bias training
- Support systematic dissemination and implementation research
- The National Institutes of Health and other relevant research entities should support the development of public–private partnerships, or other innovative collaborations, to:
    - build multidisciplinary teams, including but not limited to researchers in neuroscience, endocrinology, immunology, physiology, and metabolism, behavior, psychology, and primary care to identify the most relevant factors in a child's complex environment that promote resilience and promote outcomes related to physical and mental health.
    - conduct research that measures the impact of chronic stress on all relevant organ systems and determines the specific molecular and biological pathways of interaction during the pre- and postnatal periods, which are directly relevant to potential interventions to address health disparities.[6]

[6] For reference, this is Recommendation 2-1 in this report.

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Many of the items in this recommendation will require recruiting diverse populations, with explicit attention to addressing racial/ethnic and socioeconomic inequities in developmental outcomes.

Measuring Success

The committee has identified a number of measures and indicators that can currently be measured and are important for tracking progress within each of the systems that act as key leverage points for early childhood development discussed in this report. For example, for caregivers, the committee proposes measuring maternal depression and stress, feelings of rejection or hostility to the child, available support for mothers, and any contact with child protective services. However, other measures will be needed. To further the ability for subgroup and other analyses and continuous data collection on both successes and failures, the following is needed (see Chapter 8 for more detail):

Understanding and measuring cumulative exposure. A number of factors impact early life development, ranging from influences in the microsocial or family environment, such as attachment, nurturing, and maternal well-being, to institutional levers, such as access to prenatal care or effective responses to trauma exposure, to macrosocial forces, such as racism and poverty. Effective tools already exist to measure exposure to some of these factors but there are few methods for empirically understanding how exposures to risks or protective factors accumulate and combine over time to establish a cumulative overall risk profile.

Understanding the interaction among developmental pathways. There are few frameworks for understanding the multidirectional relationship between the biological, social-behavioral, and psychological development of young children. In particular, understanding how these interactions may vary across the life course in response to changing plasticity of biological systems, different stages of personal and cognitive development, and different life conditions and accumulated experiences is critical to building a health equity strategy.

Measuring interactions between systems. Models that can estimate "integrated risk" by combining key data from across the sectors where people live their lives are needed. Similarly, measures that examine results from cross-sector collaboration can help in documentation and accountability.

Improving methods to assess complex causality. Perhaps the biggest challenge facing health equity research is complex causality. Many of the preferred tools of science, such as randomized controlled trials (RCTs), are designed to control for and isolate single causes rather than embrace complex, interrelated causality that may include multilevel, multidirectional, and nested effects—for which a larger toolbox of strategies is needed. For example, there needs to be greater exploration of effective community-based intervention approaches that use existing resources (e.g., as in "natural experiments").

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CONCLUSION

The advances in the science of early development are ready to be acted upon—there is no reason to wait for this additional science before taking action. Long-term psychological, behavioral, and physical health is shaped by biological and environmental factors, including their interactions, before conception and throughout the life course. This interplay necessitates action at the practice, policy, and systems levels that takes into account the full range of factors that shape health and well-being. These actions need to be taken before insults to early development occur. The science of plasticity shows that it is never too late to intervene but that early identification and intervention are generally more effective and cost-saving and require less effort. (See Box S-1 for a high-level overview of the report's findings and Table 9-1 for a summary of recommended actions in this report.) Further, these actions need to take a life course, multigenerational approach to decrease health inequity, as children's well-being depends on the well-being of the primary caregivers and the quality of their relationship. Progress toward health equity can be achieved through multipronged, cross-sector interventions that focus on prevention, early detection, and mitigation and that work at the practice, policy, and systems levels to address the SDOH. The committee hopes that the roadmap laid out in this report will catalyze the steps that need to be taken across systems to close the health equity gap and improve the lives of our nation's children.

BOX S-1 Chapter Key Messages

A. **Lessons from the science of early development are clear and actionable.** A tremendous amount is known about how development occurs in the prenatal and early childhood periods. When the science of early development is coupled with a health equity approach to inform decision making, it provides an opportunity to improve outcomes for children and families. (Chapters 1 and 2)
B. **Over time, biological and social-psychological development interact to shape the way health develops over the life course.** Neither is deterministic—health outcomes are never set in stone. Rather, they are probabilistic—together, they cumulatively "set the odds" for good health. (Chapter 2)
C. **Biology and environment work together to affect children's growth and development.** Intervening early—to both prevent and mitigate adverse outcomes—is crucial. During the prenatal and early life periods, critical biological systems that will help shape health across the life course are developed and affected by the early environment. Intervening early, when the plasticity of these systems is at its greatest, is the best way to improve chances of developing in ways that optimize health outcomes. (Chapters 2 and 3)
D. **Ensuring the well-being of caregivers by supporting and caring for them is critical for healthy child development.** Reducing children's exposure to maltreatment is a critical lever, as is promoting nurturing behaviors, fostering self-regulation, and developing coping skills for caregivers and children. (Chapter 4)
E. **Preconception, prenatal, postpartum, and pediatric care needs to be reconceptualized** to address the root causes of health inequities and to better meet the developing health and health care needs of children and their families. Content, quality, and access to care are critical components of change. (Chapter 5)
F. **Families need adequate resources available for meeting basic needs, especially when children are young.** Bolstering resources should not come at the expense of attachment or caregiver well-being, so programs such as paid parental leave, basic support, and housing stability are needed. (Chapter 6)

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G. **Early care and education (ECE) can be a platform for delivery or supporting services and interventions to advance health equity.** However, increasing the capacity and resources for the ECE professionals is needed. (Chapter 7)

H. **To advance health equity and meet the developmental needs of children, a systems approach, including collaboration and alignment across sectors, is needed,** such as workforce support and training, trauma-informed systems and care, enhanced detection of early life adversity and improved response systems, and integration of care and services across all dimensions of health. (Chapter 8)

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1

The Need to Intervene Early to Advance Health Equity for Children and Families

INTRODUCTION

Neurobiological and socio-behavioral research indicate that early life conditions, including social supports (e.g., supportive relationships) and adversity (e.g., chronic or severe stress), shape prenatal and early childhood health and development. These experiences have a powerful impact on developing biological systems that impact physical and mental health outcomes throughout life and are further influenced by the social determinants of health (SDOH) (e.g., education, housing, physical and social environment). Despite increasing evidence about what contributes to poor health, these health inequalities have persisted, and for some populations and outcomes, they are worsening. Scientific evidence can be used to better inform efforts to advance health equity; this report uses that evidence to support policy actions, program development, practice changes, systems reform, and research priorities. (See Box 1-1 for a high-level overview of this chapter.)

BOX 1-1
Chapter in Brief: The Need to Intervene Early

Health disparities during the prenatal through early childhood periods:

- Significant, long-standing disparities exist for many health outcomes by education, income, race, ethnicity, geography, gender, neighborhood, disability status, and citizenship status. These disparities put children on a course for poor health outcomes throughout the life-span.
- These health disparities include infant and maternal mortality, low birth rate, and chronic childhood diseases (such as diabetes, asthma, obesity, depression, and anxiety).
- Clinical care is necessary but not sufficient to address health inequities. To advance health equity, the root causes of poor health and chronic adversity—the social, economic, environmental, and cultural determinants of health—need to be addressed.

Scientific advances in the neurobiological and socio-behavioral sciences:

- The importance of the environment on biological processes (i.e., the complex interplay of biology and environment), including the impact of early adversity, poverty, and racism on lifelong health outcomes, is now well understood.

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- While more research is needed to develop a better understanding of tailoring interventions to address heterogeneity, research has advanced dramatically to apply new and more effective interventions.
- Evidence shows that prevention and early intervention for children on at-risk trajectories works, and it is generally more effective and less costly than intervening later in life.
- Negative and positive exposures accumulate over the life course to exert a cumulative effect on health that is probabilistic, not deterministic. That is, the odds of positive or negative health are never fixed; individual exposures, experiences, resilience, and choices help set and adjust these odds over time.

The committee's approach:

- This report takes a life course approach, which emphasizes that a temporal and social perspective—looking across an individual's life experiences or across generations—to gain a better understanding of health outcomes is needed. This approach takes into account an individual's larger social, economic, and cultural context and that the trajectory of an individual's life may be changed, negatively or positively, through interactions between the brain, body, and environment throughout the life-span.
- Achieving health equity for children will require attention and commitment from a range of sectors. Although there are many barriers, this report identifies many opportunities to make long-lasting reductions in long-standing and persistent health inequities.
- Building off 12 core concepts of early development, the committee used scientific evidence to guide its recommendations.
- This report provides a range of recommendations for practice, policy, and systems changes, including recommendations that will take time and sustained commitment to dismantle structural barriers and recommendations that could be implemented immediately or in the near term.

THE PROBLEM

The United States spends much more on health care than any other Organisation for Economic Co-operation and Development (OECD) nation, yet it ranks poorly on most measures of population health (NRC and IOM, 2013; OECD, 2017). Significant, long-standing disparities exist in most health outcomes by education, income, race, ethnicity, geography, gender, neighborhood, disability status, sexual orientation, and citizenship status (NASEM, 2017). Notably, the last few decades have marked a troubling rise in U.S. maternal mortality rates, including black-white disparities, while maternal mortality rates have declined globally (WHO, 2015). Similarly, disparities in infant mortality rates persist where non-Hispanic black, American Indian/Alaska Native (AI/AN), and Hispanic babies experience higher rates of mortality before their first birthdays compared to non-Hispanic white and Asian and Pacific Islander babies (CDC, 2019b). Children in the United States rank behind their peers in most OECD nations in health status and on key determinants of health, and they experience growing disparities on multiple measures of child well-being (OECD, 2009; Seith and Isakson, 2011) (see Box 1-2). For children living in both urban and rural communities, lack of access to critical resources is a major driver of increasing disparities, and this is compounded for economically disadvantaged groups. These disparities put children on a course for poor health outcomes later in life. For more information on maternal and child health disparities, see the sections on Early Childhood Health Disparities and Maternal Health below.

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BOX 1-2
United States Compared to Other OECD Countries on Key Health Indicators and Outcomes

Compared with other OECD countries, the United States has

- Higher infant mortality
- Higher child mortality
- Higher poverty
- Higher gun violence and death
- Higher rates of hunger
- Higher rates of obesity
- Much higher rates of incarceration of young adults
- Lower secondary school graduation.

SOURCES: Grinshteyn and Hemenway, 2016; OECD, 2014, 2017; Walmsley, n.d.

Many families in the United States do not receive health care when needed, receive it too late, and/or experience problems in quality; however, pathways to better health do not depend on health care alone. For example, children in relatively affluent communities, who ostensibly have easy access to the best mental health services, also show elevated distress compared to national norms—as do their low socioeconomic status (SES) counterparts, but due to a very different set of life stressors (Korous et al., 2018). The factors that ultimately contribute to good health (such as nutrition, stress, exposure to environmental toxicants) are on multiple interrelated causal pathways along the life-span (NASEM, 2017).

Over the past 100 years, there has been a strong trend toward the conflation of "health" and "health care," where the health of an individual is considered only through a biomedical lens, not taking into account the multiple social and developmental determinants that drive health (Lantz, 2018). Instead, health has been erroneously equated with health care and health disparities erroneously equated with health care disparities. The result is a narrow policy focus on health care interventions to improve health. More recently, research and practice has shifted away from this prevailing paradigm to one that targets upstream factors that shape health (Hahn, 2019; NASEM, 2017).

The 2017 National Academies of Sciences, Engineering and Medicine (The National Academies) report *Communities in Action: Pathways to Health Equity* reviewed the root causes of health disparities and concluded that health inequity arises from root causes that could be organized in two clusters:

1. Intrapersonal, interpersonal, institutional, and systemic mechanisms (also referred to as "structural inequities") that organize the distribution of power, and access to critical resources, differentially across lines of race, gender, social class, sexual orientation, gender expression, and other dimensions of individual and group identity, and
2. Unequal allocation of and access to power and resources—including goods, services, and societal attention—which manifests itself in unequal social, economic, and environmental conditions, also called the "determinants of health" (NASEM, 2017, p. 7).

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Therefore, health inequities are the result of more than individual choice or random occurrence. They are the result of the historical and ongoing interplay of inequitable structures, policies, and norms that shape lives. Interventions targeting the above factors hold the greatest promise for advancing health equity and promoting positive health outcomes at the population level.

Furthermore, the report concluded that

> **Health equity is crucial.** Health equity is fundamental to the idea of living a good life and building a vibrant society because of its practical, economic, and civic implications. Promoting health equity could afford considerable economic, national security, social, and other benefits. Yet, recent research demonstrates that worsening social, economic, and environmental factors are affecting the public's health in serious ways that compromise opportunity for all.
>
> **Health inequity is costly.** Beyond significant costs in direct health care expenditures, health inequity has consequences for the U.S. economy, national security, business viability, and public finances, considering the impact of poor health and disability on one's ability to participate in the workforce, military service, or society. Addressing health inequities is a critical need that requires this issue to be among our nation's foremost priorities (NASEM, 2017).

Given these findings, it is critical to address health disparities (differences in health outcomes) with a comprehensive approach—by treating all the factors that impact individual health, such as education, employment, health systems and services, family, community, housing, income and wealth, physical and social environments, public safety, and transportation (SDOH), in addition to racism, discrimination, segregation, and poverty. To achieve equitable health outcomes in the prenatal through early childhood periods and throughout the life course, all of these contexts need to be addressed.

Health inequities are systemic challenges, and chronic childhood adversities have biological implications and effect childhood development, with lifelong impacts on health and well-being. When exposures to key experiences (both positive and negative) differ for specific groups (e.g., black/African American, AI/AN, Hispanic), their odds for good health diverge systematically over time, producing disparities in outcomes. Because the odds of these exposures are impacted by systems, advancing health equity will require more than individual level interventions. It will require systems to change in ways that improve the odds of good experiences and reduce the odds of adverse exposures for specific populations.

The focus of this report is on how to best maximize well-being among all young children and families, especially those who are vulnerable at the outset—because life circumstances have rendered them statistically more likely to be on negative adjustment trajectories from early life onward. "Inequities" are operationally defined for the purpose of this report, in part, as the unequal likelihood of thriving or attaining positive adjustment outcomes over time because of differences in opportunity that lead to unfair and avoidable differences in health outcomes.

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OPPORTUNITIES

Failing to address the context in which children live, grow, and learn undermines the potential of so many children. Evidence shows that prevention and early intervention for children on at-risk trajectories works and is generally more effective and less costly than later intervention. Luckily, a great deal is known about the science of prenatal and childhood development and the biological mechanisms and effects of chronic adversity and adverse childhood experiences (ACES) (see Chapters 2 and 3). Recent advances in science—especially around epigenetics,[1] technology and data sharing, and cross-disciplinary collaboration—present an opportunity to systematically apply this knowledge to practice, policy, and systems changes. A large body of research now explicates the mechanisms by which early adversity can change the timing of sensitive periods of brain and other organ systems development, impacting the "plasticity"[2] of developmental processes that are driven by experiences in the life of the young child and their family. It is now known that what takes place in early development has lifelong impacts—both positive and negative—on health and well-being. While diseases may appear clinically throughout the life-span, it is known that many diseases originate during early development (Gluckman et al., 2007; Heindel and Vandenberg, 2015). For example, altered nutrition, exposure to environmental chemicals, or stress during specific times of development can lead to functional changes biologically, predisposing individuals to diseases that manifest later in life and affecting physical, mental, and cognitive functions.

This report employs the science of early development to inform multidisciplinary and developmentally appropriate systems to support optimal health and well-being for all children throughout their life-spans. The tremendous advances in the theoretical and empirical science in the past 30 years position practitioners and policy makers to take informed action to improve child health outcomes. There is now a firm understanding of the importance of the environment—the constant interplay of nurture–nature, biology–environment—on biological processes, including the impact of early adversity, poverty, and racism on lifelong health outcomes. While there is more to be discovered that will lead to a better understanding of best practices and address challenges of heterogeneity, research has advanced dramatically to apply new and more effective interventions now than ever before.

Given these advances and the understanding of how science can be used to advance health equity during early development, the Robert Wood Johnson Foundation (RWJF), as part of its Culture of Health Initiative, asked the Health and Medicine Division of the National Academies to (see Box 1-3 for the full Statement of Task)

1. Provide a brief overview of stressors that affect prenatal through early childhood development and health,
2. Identify promising models and opportunities for translation of the science to action,
3. Identify outcome measures to enable subgroup analyses,

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[1] The study of how genes are expressed due to changes in the environment and how these biological changes can be passed down from one generation to the next.

[2] The process by which neurons within the brain change their gene expression, cellular architecture, connections with other neurons, and function in response to experiences and changes in the environment (i.e., change over time).

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4. Develop a roadmap to apply the science to tailored interventions (i.e., policies, programs, or system changes) based on biological, social, environmental, economic, and cultural needs, and
5. Provide recommendations in these areas, including how systems can better align to advance health equity.

To respond to this charge, the Committee on Applying Neurobiological and Socio-Behavioral Sciences from Prenatal Through Early Childhood Development: A Health Equity Approach was formed. The committee applied a health equity frame and builds on the 2017 report *Communities in Action: Pathways to Health Equity*. As identified in the 2000 NRC and IOM report *From Neurons to Neighborhoods,* prenatal through early childhood are critical phases of development for the production of health. This report reviews the science that has been developed since *From Neurons to Neighborhoods.*

BOX 1-3
Committee on Applying Neurobiological and Socio-behavioral Sciences from Prenatal Through Early Childhood Development: A Health Equity Approach Statement of Task

Building on the science base described in the 2000 NRC and IOM report *From Neurons to Neighborhoods: The Science of Early Childhood Development* and the concepts in the 2017 National Academies report *Communities in Action: Pathways to Health Equity,* and drawing upon new insights from 21st-century science in the neurobiological and socio-behavioral fields in the prenatal to early childhood period, an ad hoc committee will

1. Provide a brief overview of
   - the key stressors that affect brain development and health outcomes during this period (e.g., structural inequities, income, housing, employment, access to health care, transportation, and others) and
   - the biological and environmental factors that lead to disparities in health and disease outcomes for subgroups of individuals and the pathways by which biological factors interact with and are influenced by sociocultural factors.
2. Identify promising models and opportunities for translation of the science to action and the intervention points during the prenatal and early childhood periods that will yield the greatest impact, with a focus on practice-based changes and the goal of facilitating broader systems change and alignment based on the science. The committee will draw from international examples as appropriate.
3. Identify the specific outcome measures needed to enable subgroup analyses based on the biological dynamics of the social determinants of health, and identify methods to continuously collect data on both successes and failures to enhance the knowledge base in the future.
4. Based on its review of the evidence and committee expertise, develop a roadmap to systematically apply the science to inform tailored interventions (i.e., policies, programs, or system changes) based on biological, social, environmental, economic, and cultural needs. The roadmap will identify pathways to implement the science in practice and policy.
5. Provide recommendations in the areas above as well as recommendations on how systems can better align to advance health equity and identify specific research needs, as deemed appropriate based on its review of the evidence and its collective expertise.

IOM = Institute of Medicine; NRC = National Research Council.

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The committee also adopted the life course approach to its work, which emphasizes a temporal and social perspective, while looking back across an individual's life experiences—or across generations—to gain a better understanding of health outcomes (Braveman and Barclay, 2009). This approach takes into account an individual's larger social, economic, and cultural context and that the trajectory of that individual's life may be changed, negatively or positively, through interactions between the brain, body, and environment throughout the life-span. An individual's health status and outcomes reflect the accumulation of experiences over the life course. Protective factors support positive, or flourishing, trajectories, while risk factors exacerbate the likelihood of poor trajectories (see Figure 1-1).

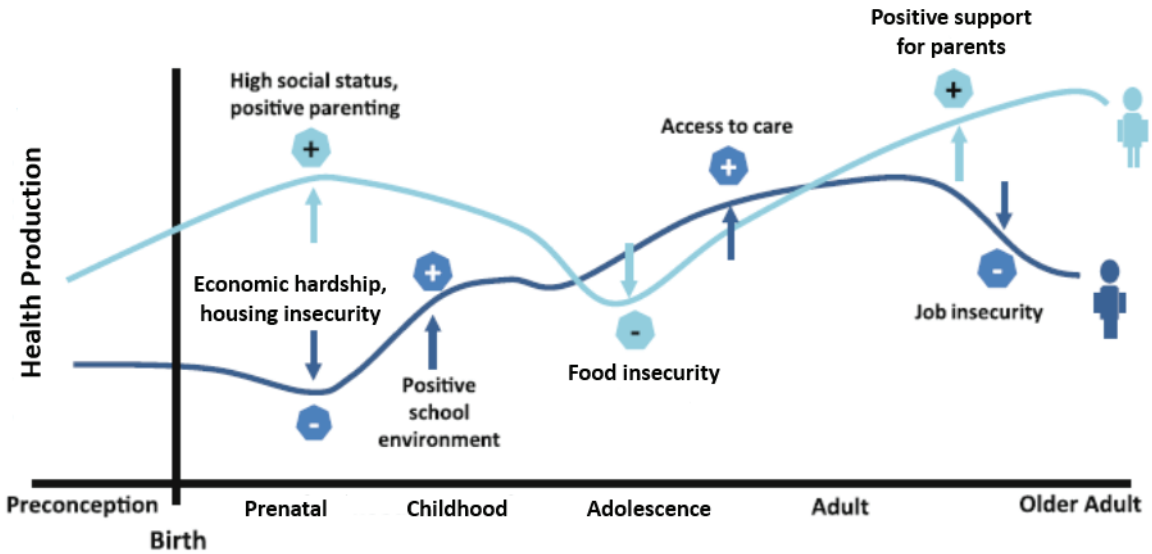

FIGURE 1-1 Variable health trajectories: Life course approach
NOTE: This figure includes several examples; however, there are many other variables that impact health trajectories (see Chapter 3).
SOURCE: Adapted from Halfon et al., 2014.

In line with the life course approach, the committee included the preconception period as an important focus of the report. The health of men and women before conception is important for pregnancy outcomes and the lifelong health of their children. During the preconception period, parents may be exposed to several types of stressors (e.g., chronic stress, environmental toxicants, poverty) that could have repercussions for the health of their future children even into adulthood.

For the purpose of this report, early childhood refers to birth through the onset of puberty and the beginning of adolescence (approximately 8 years of age). The committee uses the terms "preconception," "prenatal," "postpartum," and "early childhood" to refer to the various periods of development. The committee uses the term "early development" to refer to the preconception through early childhood periods. Another National Academies study that was under way during the course of this study—also looking at the neurobiological and closed-behavioral sciences—covered adolescence.[3]

[3] See Neurobiological and Socio-behavioral Science of Adolescent Development and Its Applications: http://sites.nationalacademies.org/dbasse/bcyf/adolescent-development/.

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The outcomes the committee seeks to improve across the life course fall into four categories, adapted from a 2016 report *Parenting Matters*: Supporting Parents of Children Ages 0-8:

1. Physical health and safety,
2. Emotional and behavioral competence,
3. Social competence, and
4. Cognitive competence.

See Box 1-4 for definitions of key terms used in this report.

BOX 1-4
Key Terms as Used in This Report

Early childhood: A time of tremendous physical, emotional, behavioral, social, and cognitive development. For the purpose of this report, early childhood encompasses from birth to approximately 8 years of age.

Early development: The period from preconception through early childhood.

Child health: The extent to which individual children or groups of children are able or enabled to (a) develop and realize their potential, (b) satisfy their needs, and (c) develop the capacities that allow them to interact successfully with their biological, physical, and social environments [a]

Health disparities: Differences that exist among specific population groups in the United States in the attainment of full health potential that can be measured by differences in incidence, prevalence, mortality, burden of disease, and other adverse health conditions.[b]**Health equity:** The state in which everyone has the opportunity to attain full health potential and no one is disadvantaged from achieving this potential because of social position or any other socially defined circumstance. [c]

Health inequities: Systematic differences in the opportunities that groups have to achieve optimal health, leading to unfair and avoidable differences in health outcomes. [b]

Social determinants of health: The conditions in the environments in which people live, learn, work, play, worship, and age that affect a wide range of health, functioning, and quality-of-life outcomes and risks. These include education, employment, health systems and services, housing, income and wealth, the physical environment, public safety, the social environment (including structures, institutions, and policies), and transportation. [c]

Structural inequities: Personal, interpersonal, institutional, and systemic drivers—such as racism, sexism, classism, ableism, xenophobia, and homophobia—that make those identities salient to the fair distribution of health opportunities and outcomes. For example, policies that foster inequities at all levels (from organization to community to county, state, and nation) are critical drivers of structural inequities. [b]

Toxic stress response: Prolonged activation of the stress response systems that can disrupt the development of brain architecture and other organ systems, and increase the risk for stress-related disease and cognitive impairment, well into the adult years. The toxic stress response can occur when a child experiences strong, frequent, and/or prolonged adversity—such as physical or emotional abuse, chronic neglect, caregiver substance abuse or mental illness, exposure to violence, and/or the accumulated burdens of family economic hardship—without adequate adult support.[c] Toxic stress is the maladaptive and chronically dysregulated stress response that occurs in relation to prolonged or severe early life adversity. For children, the result is the disruption of the development of brain architecture and other organ systems and an increase in lifelong risk for physical and mental disorders.

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[a]NRC and IOM, 2004.
[b]NASEM, 2017.
[c]Center on the Developing Child, n.d.

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WHY INVEST IN EARLY INTERVENTION AND PREVENTION?

There are many reasons to intervene in the prenatal through early childhood periods to prevent and mitigate adverse health outcomes both in early life and over the life course. First, preventing or reducing poor outcomes in the prenatal through early childhood periods generally leads to improved health outcomes later on and therefore can yield health care savings in the long term. For example, in 2007, the IOM reported that the cost associated with premature birth in the United States was $26.2 billion each year (IOM, 2007), and the CDC reports that the United States spends $147 billion in obesity-related health care costs each year (CDC, 2019a). Health care savings are also seen when preventing lead poisoning (Gould, 2009), and some childhood diseases. McLaughlin and Rank (2018) estimate that childhood poverty results in $192.1 billion in aggregate health costs (with another $96.9 billion due to child homelessness and $40.5 billion due to maltreatment). Further, inequity is costly. As noted in *Communities in Action* (NASEM, 2017), advancing progress toward health equity (across the life course) could produce economic, national security, and other benefits for the nation. The report made the case that beyond the dollar cost of health care services (which itself is significant at $3.5 trillion in 2017, accounting for 17.9 percent of the nation's GDP [CMS, 2017; Martin, et al., 2018a), health inequities contribute to overall poor health for the nation and therefore have consequences for the U.S. economy, including diminished productivity in the business sector. In 2009, the Urban Institute projected that from 2009 to 2018, racial disparities in health will cost U.S. health insurers approximately $337 billion in total (Waidmann and Urban, 2009). Further, the rising cost of health insurance and medical care for workers cuts into companies' ability to make a profit and stay competitive (IOM, 2015; Shak et al., 2013).

Investing in early development through prevention and early intervention yields cost savings because the investment costs are often less than the downstream costs of poor health and development. For example, investing in high-quality early care and education (ECE) is one way to improve outcomes related to child health. Not only do early childhood intervention programs yield benefits in academic achievement, behavior, educational progression and attainment, and labor market success, among other domains, but well-designed early childhood interventions have been found to generate a return to society (Karoly et al., 2005).[4] Garcia et al. (2017) found a 13.7 percent return on investment for comprehensive, high-quality, birth-to-5 early education.[5] From a public health perspective, a 2017 systematic review by Masters and colleagues found that national and public health interventions are highly cost-saving for interventions ranging from vaccination to larger determinants of health, though those focusing on the latter had a lower return on investment because they are more complex, resource intensive, and sustained.[6] Child maltreatment is costly to the nation as well. The total lifetime economic burden in the United States in 2008 resulting from new cases of fatal and nonfatal child maltreatment was approximately $124 billion (Fang et al., 2012). Further, Bellis and colleagues (2017) found that

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[4]Studies looking at cost-effectiveness often rely on different underlying assumptions, limiting comparisons across studies and programs. However, overall studies have shown that investments in early childhood appear to save money in the longer term.

[5] The study analyzed a wide variety of life outcomes, such as health, crime, income, IQ, schooling, and the increase in a mother's income after returning to work due to childcare.

[6] The review included studies from the UK, Western Europe, the United States, Canada, Japan, Australia, and New Zealand.

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disproportionate health expenditure in later life might be reduced through childhood interventions to prevent ACEs, showing the long-lasting costs of early adversity.
While the primary beneficiaries of prevention and early intervention efforts are children and their families, the nation as a whole also benefits through cost savings, a healthier and more productive workforce, and strengthened national security (NASEM, 2017).

CURRENT STATE OF CHILDREN'S HEALTH

Overall Well-Being of Children and Families in the United States

This section begins with a broad overview of the current state of child health in the United States, followed by a summary of key health disparities in child and maternal health in the country. Child health is the extent to which individual children or groups of children are able or enabled to (a) develop and realize their potential, (b) satisfy their needs, and (c) develop the capacities that allow them to interact successfully with their biological, physical, and social environments (NRC and IOM, 2004). Children in the United States are generally perceived to be healthier now than in the past because they are much less likely to encounter the major infections or debilitating diseases of past generations and are typically able to recover fully from "acute" childhood illnesses. However, ailments of the past have been replaced with chronic conditions (such as diabetes, asthma, obesity, depression, and anxiety), with large segments of U.S. children facing barriers to good health and well-being as a result of poverty, food insecurity, unsafe or unstable housing, and substantial adversity (such as ACEs) in the first few years of life. Children who are born and raised in poverty are at particularly high risk for poor health outcomes, more problems in early development (e.g., lack of readiness for school at age 5, diagnoses of developmental delays and/or disorders), and higher rates of most childhood chronic conditions. Early adverse experiences have intermediate effects on physical and/or mental well-being and contribute to chronic disease and poor functioning in adulthood (Hughes et al., 2017; Shonkoff et al., 2012). In fact, these impacts are cumulative, and adults who experience adversity in childhood have substantially higher rates of heart disease, lung disease, metabolic syndrome, and other costly health conditions (see Chapters 2 and 3 for more information). Many of these health disparities are rooted in historical practices and policies (such as segregation and redlining), and this historical legacy continues to shape the development of children today. (See Chapter 3 for a discussion on historical trauma and NASEM (2017) for an overview of historical injustices that impact health outcomes.)

Early Childhood and Maternal Health Disparities in the United States

This section provides a brief overview of child and maternal health disparities and indicators of health in the United States. It is not a comprehensive overview but rather highlights some of the key health disparities. The most recent data available are presented. Chapter 3 provides a detailed overview of health disparities and critical influences or factors that can either promote or hinder healthy development, with a focus on factors that shape inequities at the child/family level and the community and population levels.

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Infant Mortality Rates

In 2015, infant mortality rates per 1,000 live births by race and ethnicity were as follows: non-Hispanic black (11.3), AI/AN (8.3), Hispanic (5.0), non-Hispanic white (4.9), and Asian/Pacific Islander (4.2) (CDC, 2019b) (see Figure 1-2). In 2014, infant mortality in rural counties was 6.55 deaths per 1,000 births, 6 percent higher than in small and medium urban counties and 20 percent higher than in large urban counties (Ely et al., 2017). Neonatal mortality was 8 percent higher in both rural (4.11 per 1,000 births) and small and medium urban counties compared with large urban counties (Ely et al., 2017).

Mortality for infants of non-Hispanic white mothers in rural counties (5.95 per 1,000) was 41 percent higher than in large urban counties and 13 percent higher than in small and medium urban counties (Ely et al., 2017). For infants of non-Hispanic black mothers, mortality was 15 percent higher in small and medium urban counties and 16 percent higher in rural counties (12.08) compared with large urban counties (Ely et al., 2017).

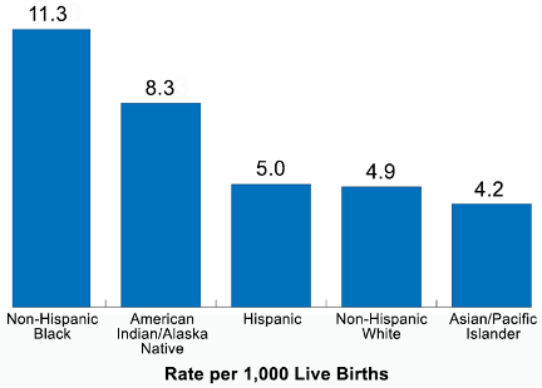

FIGURE 1-2 Infant mortality rates by race and ethnicity, 2015
SOURCE: CDC, 2019b.

Low Birth Weight

Low birth weight (LBW; less than 5.5 pounds at birth) babies are more at risk for many short- and long-term health problems, such as infections, delayed motor and social development, and learning disabilities (CDC, 2016). Causes of LBW include maternal smoking, use of alcohol, or lack of weight gain and social and economic factors, such as low income, low parental educational level, maternal stress, and domestic violence or other abuse (CDC, 2016).

LBW levels among race and Hispanic-origin groups in 2016 ranged from 6.97 percent for births to non-Hispanic white women to 13.68 percent for births to non-Hispanic black women. Rates among Hispanic subgroups ranged from 6.90 percent for births to Mexican women to 9.5 percent for births to Puerto Rican women (Martin et al., 2018b). In 2016, 14 percent of black infants were LBW, compared with 8 percent of Asian and Pacific Islander, 8 percent of AI/AN, 7.0 percent of white, and 7 percent of Hispanic infants. Among those of Hispanic origin in 2016, Puerto Rican infants were the most likely to be LBW (9 percent) (Child Trends, 2018b). In 2013, the most recent year that information for Asian and Pacific Islander subgroups was available, Asian Indian infants were the most likely to be LBW (11 percent), followed by Filipinos (9 percent). Black infants are also more than twice as likely as other infants to be very LBW—less than 3 pounds 5 ounces—at 2.9 percent in 2016, compared with between 1.1 and 1.2 percent for white and Hispanic infants, respectively (Child Trends, 2016b).

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Chronic Childhood Diseases

Conditions that rarely lead to death in children and youth are now more prevalent: obesity, asthma, mental health conditions (especially Attention-deficit/hyperactivity disorder, depression, and anxiety), and neurodevelopmental conditions (including autism spectrum disorders [ASDs]) (Perrin et al., 2007, 2014; Van Cleave et al., 2010). For example, in children less than 18 years of age, asthma was prevalent among 8.1 percent of white non-Hispanic children, 12.6 percent of black non-Hispanic children, 8.2 percent of other non-Hispanic children, and 7.7 percent of Hispanic children. Among Hispanic children, 11.3 percent of Puerto Rican children and 6.2 percent of Mexican/Mexican American children had asthma (CDC, 2019c). During 2016, asthma affected children living in families with incomes of less than 100 percent of the federal poverty level (FPL) (10.5 percent) more than those living in families with incomes of ≥250 percent of the FPL (250 to <450 percent FPL: 6.9 percent; ≥450 percent FPL: 6.7 percent) (Zahran et al., 2018).

Mental and Behavioral Conditions

Particularly noteworthy is the growth of mental and behavioral conditions among children and youth. Recent work has documented their a) high prevalence, b) major impact on youth well-being and functioning, c) common association with other chronic conditions, d) high costs, and e) complication of the course, treatment, and outcomes of most other conditions (Ghandour et al., 2012; Houtrow et al., 2014; Perrin et al., 2018). Importantly, most of these high-prevalence conditions occur at higher rates and usually higher severity among low-income children, even though the rate of growth has increased in all SES levels (Houtrow et al., 2014). Although all of these conditions have genetic components, they also often reflect the consequences of early childhood experiences and their influence through epigenetics and other physiologic mechanisms. Furthermore, less than optimal access to and use of health care in early years can negatively affect these conditions over time.

Early Life Adversity

The impacts of early life adversity and disparities are discussed in detail in Chapters 2 and 3; however, a highlight of a few disparities is provided here. ACEs (including physical, emotional and sexual abuse, physical and emotional neglect, and household stressors, including parental mental illness, substance use or incarceration, parental separation or divorce, and domestic violence) are highly prevalent in all racial and socioeconomic groups. Almost 50 percent of children and adolescents (age 0-17) have experienced at least one category of ACEs, according to national population-based studies (Bethell et al., 2014), and black, Hispanic, and poorer children are exposed to more ACEs relative to white or wealthier children (Slopen et al., 2016).[7] Foster care children are at between 1.5 and 7 times greater odds of having experienced any 1 of the 10 traditional ACEs compared to children not placed in foster care, even after controlling for race and ethnicity, parent education and employment, welfare services, and poverty status (Turney and Wildeman, 2017).

[7] ACEs are discussed in detail in Chapter 3.

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School Readiness

School readiness[8] is an important indicator for future child well-being, and there are deep disparities across race and ethnicity (see Chapter 7 for a detailed overview of the importance of ECE). Figure 1-3 provides examples of important measures of school readiness by race and Hispanic origin. Using 2016 National Survey of Children's Health data, Ghandour et al. (2018) found that only 41.8 percent of 3- to 5-year-olds in the United States were estimated to be on track in all four domains of school readiness. Overall, Hispanic children are less likely to show cognitive/literacy readiness skills than are white, black, or Asian/Pacific Islander children.

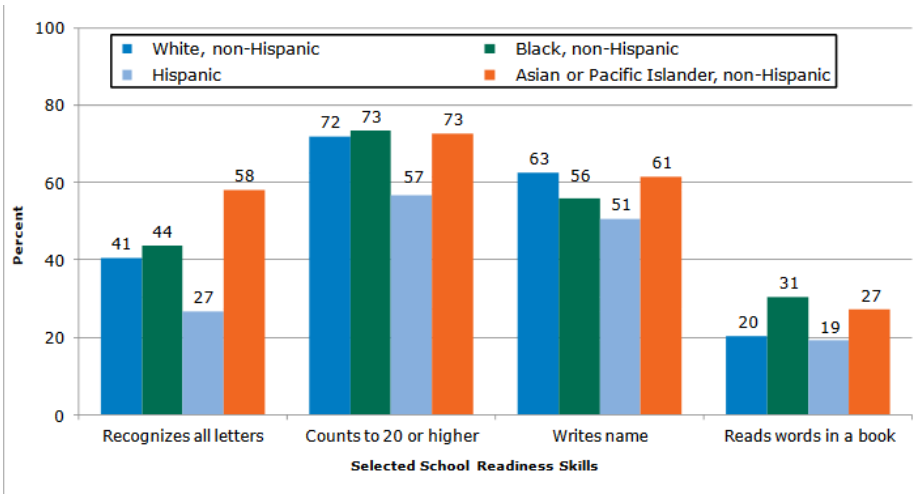

FIGURE 1-3 Percentage of children ages 3 to 6 with selected school readiness skills, by race and Hispanic origin, 2012
SOURCE: Child Trends, 2015.
NOTES: Data represent parent reports of specific cognitive and literacy skills. Data are sourced from Child Trends' analysis of the National Household Education Survey.

In 2012, 27 percent of Hispanic 3- to 6-year-olds could recognize all 26 letters of the alphabet, compared with 41 and 44 percent, respectively, of white and black children. Asian/Pacific Islander children had the highest rate of recognizing all the letters, at 58 percent. A similar pattern in the ability to count to 20 and write their name was seen by race, although Asian and Pacific Islander children were similar to their white and black counterparts (Child Trends, 2015). Young pre-K children living in poverty are much less likely to have cognitive and early literacy readiness skills than are children living above the poverty threshold. Disparities in all measures of early school readiness by income level were greatest in 1999, but these narrowed in 2007 (Child Trends, 2015).

[8] That is, children possessing the skills, knowledge, and attitudes necessary for success in school and for later learning and life. The concept of "school readiness" is broader than cognitive or pre-academic skills, such early literacy and math. However, comparable data across communities or states are limited mostly to such outcomes, which are what is presented here. All states have developed some form of school readiness assessments, and most of them focus on children's social-emotional development, approaches to learning, and physical health and development, as well as cognitive abilities (Daily et al., 2010).

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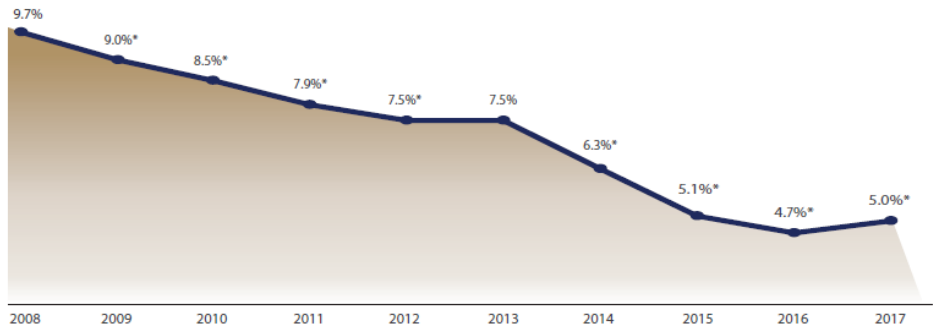
Poverty

Poverty affects large numbers of U.S. children. A recent National Academies report, *A Roadmap to Reducing Child Poverty*, estimated that in 2015, more than 9.6 million children under 18 years of age lived in poverty, as measured by the Supplemental Poverty Measure (SPM) (NASEM, 2019). Thus, 13 percent of the child population lived in households with annual incomes ranging from $22,000 to $26,000 for a family of four. Moreover, of those 9.6 million children, 2.1 million lived in "deep poverty," defined as having family resources below half the poverty line—$11,000 to $13,000 annual income. An additional 22 percent of U.S. children live in "near poverty" households, defined as between 100 and 150 percent of the SPM poverty line. These 16.7 million children live in households that frequently pay more in taxes than they receive in tax credits, reducing their net incomes (NASEM, 2019). Many U.S. families with children face persistent problems related to poverty, including inadequate housing, clothing, and food for their children, and health problems accompany those deficits.

Health Insurance Coverage

From 2008 to 2016, the rate of uninsured children steadily decreased from 9.7 to 4.7 percent. However, the percentage of uninsured children increased to 5.0 percent in 2017 (Alker and Pham, 2018). See Figure 1-4 for the rate of uninsured children from 2008 through 2017.



Source: Table HIC-5, Health Insurance Coverage Status and Type of Coverage by State - Children Under 19: 2008 to 2017, Health Insurance Historical Tables, U.S. Census Bureau American Community Survey (ACS).

*Change is significant at the 90% confidence level. Significance is relative to the prior year. 2013 was the only year that did not show a significant one-year increase or decrease in the national rate of uninsured children. The Census began collecting data for the health insurance series in 2008, therefore there is no significance available for 2008.

FIGURE 1-4 Rate of uninsured children, 2008–2017.
SOURCE: Alker and Pham, 2018.

For the percentage breakdown of the number of uninsured children by race and ethnicity in 2016 and 2017, see Figure 1-5. As the figure shows, for black, white, Asian/Native Hawaiian/Pacific Islander, and Hispanic children, the rate of children without insurance increased from 2016 to 2017. In the same 2-year period, the rate of children without insurance decreased for AI/AN children by 0.2 percent. However, the rate of children without insurance for this group is far higher than for other racial and ethnic groups at 12.8 percent in 2016 and 12.6 percent in 2017 (Alker and Pham, 2018). In 2017, 27 states and Washington DC, had significantly lower rates of children without insurance than the national rate of 5.0 percent, and 11 states had no statistically significant difference from the national rate. However, 12 states had significantly higher rates, with the highest rates of children without insurance in Texas (10.7 percent), Alaska (9.6 percent), and Wyoming (9.5 percent).

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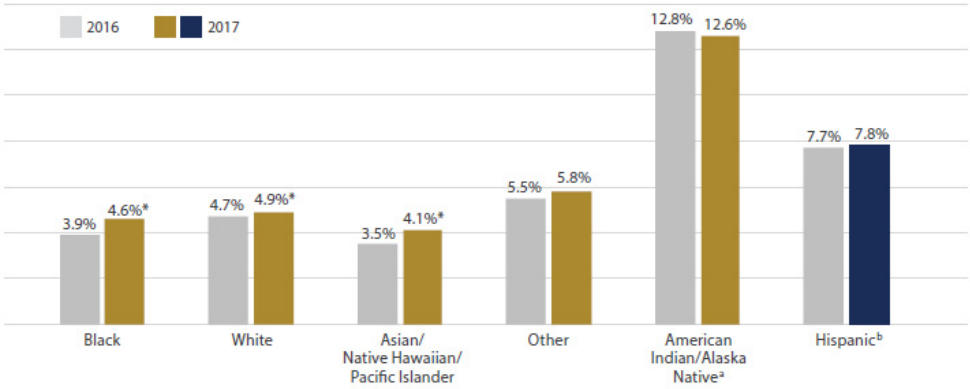

FIGURE 1-5 Children's uninsured rate by race and ethnicity, 2016–2017
SOURCE: Alker and Pham, 2018.

MATERNAL HEALTH IN THE UNITED STATES

Women in other high-income countries fare better in terms of access to health care and health status than women in the United States. U.S. women in have the highest rate of maternal mortality because of complications from pregnancy or childbirth and among the highest rates of cesarean sections, and maternal mortality rates are rising for them while declining in other countries (see Figure 1-6). For the past six decades, black women have died at a rate that ranges from three to four times that of white women, with 38.9 deaths per 100,000 live births among black women versus 12 deaths per 100,000 live births among white women as of 2010 (Creanga et al., 2015; MHTF, n.d.). AI/AN women also fare worse than white women, with approximately twice as many pregnancy-related deaths per 100,000 live births. Some researchers point to structural drivers of maternal mortality disparities in the United States, such as racism and discrimination (ACOG, n.d.) and adverse changes in chronic diseases and insufficient health care access (Nelson et al., 2018). See Chapter 5 for a detailed overview of preconception through postpartum care.

Prenatal Care

Access to prenatal care varies across race and ethnicity groups. In 2016, about 77 percent of women who gave birth initiated prenatal care in the first trimester. However, only 66.5 percent of black, 72 percent of Hispanic, 63.0 of AI/AN, and 55.9 percent of Native Hawaiian or other Pacific Islander women received it, whereas 82.3 percent of non-Hispanic white women did (Osterman and Martin, 2018). People of color are much more likely to be affected by late initiation of prenatal care, which is most common in Pacific Islanders at 18.4, with non-Hispanic white women at 3.3 percent. AI/AN women have a 9.2 percent chance of late prenatal care initiation, and black women have a 7 percent chance (Osterman and Martin, 2018). (See Chapter 5 for more statistics and information on prenatal care.)

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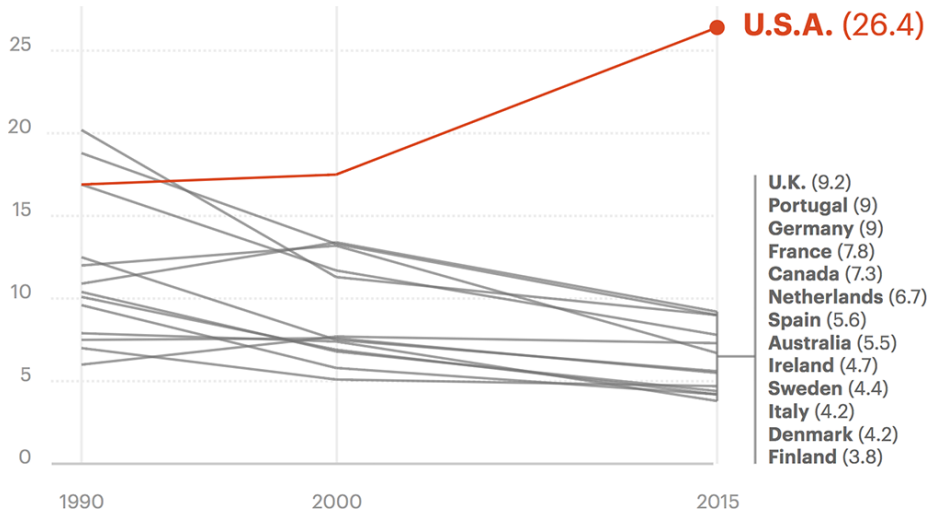

FIGURE 1-6 Global, regional, and national levels of maternal mortality, 1990–2015
NOTE: Data from GBD 2015 Maternal Mortality Collaborators, 2016.
SOURCE: Martin and Montagne, 2017 courtesy of Propublica.

THE CURRENT ENVIRONMENT FOR CHILDREN AND FAMILIES

In the past 20 years, much has changed in the landscape in which prenatal to childhood development takes place in the policy and funding environment and the social, economic, and cultural context. The following section outlines some of these important changes.

Changes in the Funding and Policy Environment

Federal spending (not including tax reductions) on children under 18 increased from $210 billion in 2000 to $375 billion in 2017, driven largely by increased health spending on Medicaid, which nearly tripled between 2000 and 2017, and the Children's Health Insurance Program (CHIP; Table 1-1) (Isaacs et al., 2018). Federal spending on nutrition, income security, ECE, and housing also increased substantially between 2000 and 2010, though many programs have seen a decline in federal support since 2010. Approximately 61 percent of federal expenditures in 2017 served children in low-income families through means-tested programs and tax provisions (Isaacs et al., 2018).

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TABLE 1-1: Federal Expenditures on Children by Program, Selected Years 1960–2017

	1960	1980	2000	2010	2017
1. Health	0.2	7.6	36.8	95.1	111.9
Medicaid	--	6.9	32.8	80.7	89.9
CHIP	--	--	1.7	8.5	15.4
Vaccines for children	--	--	0.7	4.0	4.4
Other health	0.2	0.8	1.6	1.9	2.1
2. Nutrition	1.5	22.5	30.9	60.8	58.0
SNAP (food stamps)	--	11.7	13.4	36.1	30.6
Child nutrition	1.5	9.1	12.7	18.3	22.3
Special Supplemental food (WIC)	--	1.6	4.8	6.4	5.0
3. Income Security	14.6	33.6	46.4	58.0	54.3
Social Security	7.0	17.7	18.6	22.3	20.8
Temporary Assistance for Needy Families	4.8	11.0	15.9	17.2	12.8
Supplemental Security Income	--	0.9	6.7	11.0	10.5
Veterans benefits	2.5	3.5	2.1	3.5	6.8
Child support enforcement	--	0.9	4.4	4.9	4.1
Other income security	0.3	-0.4	-1.3	-1.0	-0.6
4. Education	3.0	18.7	30.3	76.1	41.6
Education for the Disadvantaged (Title I, Part A)	--	8.3	11.8	21.9	16.2
Special education/IDEA	--	2.1	6.9	19.4	12.7
School improvement	--	2.0	3.5	6.0	4.4
Innovation and improvement	--	--	--	1.1	1.3
Impact Aid	1.7	1.8	1.2	1.4	1.5
Dependents' schools abroad	0.2	0.9	1.3	1.3	1.2
Other education	0.1	2.5	3.9	25.1	4.3
5. Early Education and Care	--	2.1	10.8	15.5	14.9
Head Start (including Early Head Start)	--	2.1	6.2	9.0	8.9
Child Care and Development Fund	--	--	4.6	6.6	5.7
Other early education and care	--	--	--	--	0.3
6. Social Services	--	4.6	10.7	11.2	10.1
Foster care	--	0.8	6.0	4.9	4.9
Adoption assistance	--	--	0.2	2.6	2.5
Other social services	--	3.9	4.5	3.6	2.7
7. Housing	--	2.8	8.3	10.7	9.5
Section 8 low-income housing assistance	--	1.4	6.5	8.0	7.7
Low-rent public housing	--	0.6	1.1	1.3	1.0
Other housing	--	0.8	0.8	1.4	0.7
8. Training	--	6.4	1.5	2.2	1.2
9. Refundable Portions of Tax Credits	--	3.1	34.5	81.8	74.0
Earned income tax credit	--	3.1	33.3	54.8	53.1
Child tax credit	--	--	1.1	25.4	19.4
Premium tax credit	--	--	--	--	0.6
Other refundable tax credits	--	--	--	1.6	0.8
10. Tax Reductions	41.2	50.1	93.1	105.1	106.2
Dependent exemption	40.6	42.3	39.7	36.0	37.8
Exclusion for employer-sponsored health insurance	NA	4.1	13.7	21.5	22.9
Child tax credit (nonrefundable portion)	--	--	26.8	33.4	29.9
Earned income tax credit (nonrefundable portion)	--	1.8	5.9	5.3	7.0
Dependent care credit	--	--	3.2	3.8	3.3
Other tax reductions	0.7	1.9	3.7	5.1	5.3
TOTAL EXPENDITURES ON CHILDREN	60.5	151.5	303.2	516.4	481.5
OUTLAYS SUBTOTAL (1–9)	19.3	101.4	210.1	411.3	375.3

NOTE: Numbers in billions of 2017 U.S. dollars.
SOURCE: Isaacs et al., 2018

Altogether, federal spending on children in 2017 accounted for 9.4 percent of total federal outlays (Figure 1-7), and it is projected to fall to 6.9 percent. In contrast, federal spending on adults through Social Security, Medicare, and Medicaid accounted for 45 percent of total federal outlays, which is projected to grow to 50 percent by 2028 (Isaacs et al., 2018). It should also be noted that federal spending on children (which accounted for 34 percent of total public spending on children) in 2017 represented only 2 percent of GDP in the United States, which is well below that of other developed nations. An international comparison of public spending on children from 1985 to 2000 ranked U.S. spending (2.4 percent of GDP) the second lowest among 20 OECD countries, much less than the 9.6 percent median across OECD countries (Lynch, 2006).

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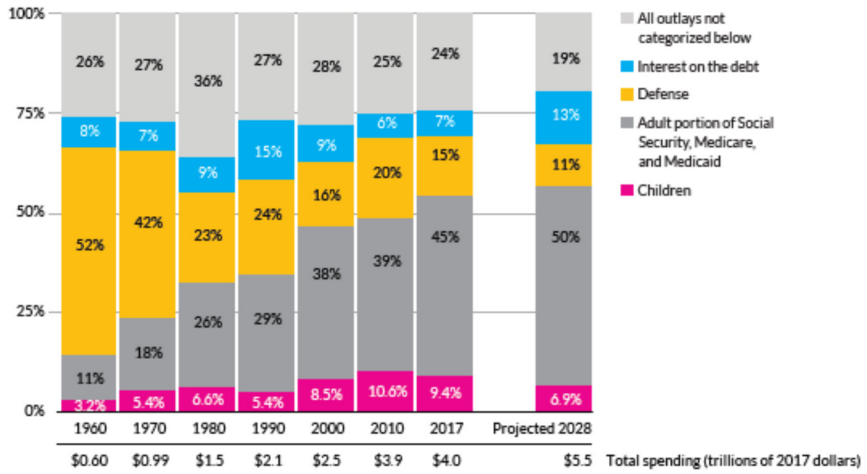

Source: Authors' estimates based primarily on Congressional Budget Office, *The Budget and Economic Outlook: 2018 to 2028* (Washington, DC: Congressional Budget Office, 2018), and Office of Management and Budget, *Budget of the United States Government, Fiscal Year 2019* (Washington, DC: US Government Printing Office, 2018) and past years. For more source information, see the appendix.

Note: Numbers may not sum to totals because of rounding.

FIGURE 1-7: Share of federal budget outlays on children and other items, selected years, 1960–2028
NOTES: Authors' estimates based primarily on Congressional Budget Office, The Budget and Economic Outlook: 2018 to 2028 and Office of Management and Budget, Budget of the United States Government, Fiscal Year 2019 and past years. (Republished with permission of Urban Institute, from Kid's Share 2018, Isaacs et al., 2018; permission conveyed through Copyright Clearance Center, Inc.).
SOURCE: Isaacs et. al., 2018.

One of the most important changes in the policy landscape over the past 20 years has been the Patient Protection and Affordable Care Act (ACA), which expanded health care access for children and families through a combination of Medicaid expansions, private insurance reforms, and premium tax credits (Kaiser Family Foundation, 2019). The uninsured rate among women ages 19-34 decreased from 25 percent in 2010 to 14 percent in 2016 (Gunja et al., 2017), which has significant implications for preconception and prenatal health. The ACA requires coverage for women's preventive services at no cost-sharing, including well-woman visits, and eliminates exclusions for preexisting conditions (for example, pregnancy and depression), which improves access for women with chronic conditions (Gunja et al., 2017). The ACA mandates coverage for essential health benefits (EHBs), which include maternity care and mental health services. For children, The ACA requires coverage for preventive services at no cost-sharing and EHBs, eliminated exclusions for preexisting conditions, and prohibits lifetime dollar limits, which improves access for children with special health care needs (National MCH Workforce Development Center, 2015). Even though the ACA did not substantially increase children's eligibility for Medicaid (or CHIP), the process of parents' seeking enrollment in Medicaid or exchange plans led to their learning of their children's eligibility, which increased child enrollment substantially. Presently, it is unclear how recent policy changes to deregulate consumer protections under the ACA and to restrict eligibility, enrollment, and benefits through Medicaid waiver will impact health care access and equity for children and families. (See Chapter 5 for more about the ACA.)

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Federal funding for ECE has steadily increased over the last 10 years; however, total spending on ECE in the United States remains limited. Only 0.5 percent of U.S. GDP is spent on ECE, whereas other peer nations spend 1 percent or more (OECD, 2019). Recent increases include lawmakers approving an $890 million increase for Early Head Start and Head Start, including $170 million for Early Head Start—Child Care Partnerships, from Fiscal Year 2016 levels to FY2019 levels (FFYF, 2018; see Chapter 7 for more on these programs). In addition, in response to decades of early childhood and brain development research, policy makers, advocates, program administrators, and other leaders in the child care community have advanced a number of policies and initiatives that recognize child care programs as opportunities to improve children's development, rather than solely as work-support programs. As a result, for example, the most recent reauthorization of Child Care Development Block Grant (CCDBG)[9] provides for more continuity of care by allowing parents to receive subsidies for a year, even if their income, work, or education status changes during that period. There has also been more attention paid to increased compensation for the workforce. For example, states and local communities have developed a number of strategies to enhance compensation for early childhood educators, including tax credits, wage supplements (often tied to attaining higher education or credentials), targeted increases for child care subsidy rates, salary scales, provision of benefits, and parity with K–12 teachers. More recently, there has been greater recognition from policy makers, advocates, and ECE practitioners of the importance of ECE program leaders (including elementary school principals).

CHANGES IN THE ECONOMIC, SOCIAL AND CULTURAL ENVIRONMENT

Income inequality has been growing—the income gap between higher- and lower-income individuals has increased substantially over the past 30 years (NASEM, 2017).[10] Those with incomes in the top 10 percent average 9 times the income of those in the bottom 90 percent, and those with incomes in the top 0.1 percent have more than 188 times the income of the bottom 90 percent (Inequality.org, n.d.). Lasting effects of the 2008 recession include displacement of vulnerable populations, which exacerbated the impact on both their health and their economic well-being and resulted in greater income inequality and wealth inequality (Smeeding, 2012). People of color continue to face structural barriers when it comes to securing quality housing, health care, employment, and education (NASEM, 2017; Pager and Shepherd, 2008). For example, data suggest that schools are becoming increasingly segregated by poverty and race, which has implications for which communities have access to high-quality, well-funded education (Boschma and Brownstein, 2016; Darling-Hammond, 1998). Racism also continues to be a pressing problem, and it is built into systems, as seen by racial profiling by law enforcement officers, disproportionate suspension and expulsion rates of young boys of color (see Chapter 7), and the difficulty that some subgroups have in breaking the cycle of poverty (see Chapter 6).

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[9] See, https://www.acf.hhs.gov/occ/ccdf-reauthorization.

[10] There are several different ways to measure income inequality, including the use of different data sets, that lead to different results (with no single source that illustrates all of the major trends in inequality) (CBPP, 2018). However, regardless of the method or data used, results consistently show an increase in income inequality in the last three decades. See a draft paper by Auten and Splinter (2018) for additional analysis that argues that income inequality has not grown as much as others have estimated.

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Chapter 3 expands on these issues and the impact they have on the health of children and families.

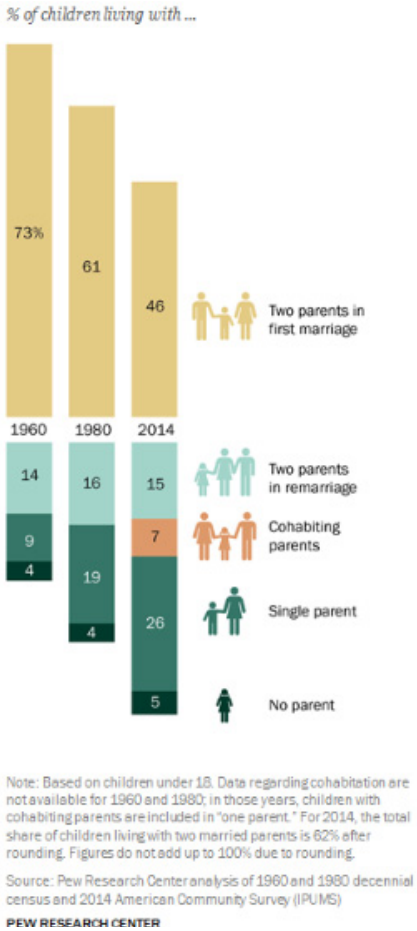

FIGURE 1-8: For children, growing diversity in family living arrangements.
SOURCE: Pew Research Center, 2015b

In terms of family structure, American families have changed, with a PEW Research Center (2015b) survey finding that there is no longer one dominant family type in the United States today (as compared to 1980, where 61 percent of children were living with married parents in their first marriage—today, that number is 46 percent) (see Figure 1-8). PEW also found, for example, that two-parent households are no longer the norm, as rates of divorce, remarriage, and cohabitation have increased. This decline has been offset by an almost threefold increase in those living with just one parent—typically the mother (10.5 percent of children age 18 or younger lived with only a mother in 2018, compared to 2.4 percent living with the father only [U.S. Census, 2018]). Women who are single or living with a nonmarital partner account for 4 in 10 births in the United States. Single-parent families are more than 4 times more likely to be poor than are two-parent families (Pascoe et al., 2016), and this can affect their ability to invest time and resources in their children, which would have longer-term implications for the persistence of poverty across generations. More mothers (of children 18 or younger) have also entered the workforce—70 percent, with 40 percent of them being the primary earner (Pew Research Center, 2015a).

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The makeup of children in the United States has changed as well. In 2000, 61 percent of all U.S. children were non-Hispanic white; in 2016 that number was 51 percent, with the proportion of children with Hispanic origins growing from 17 to 25 percent between 2000 and 2016. The percentage of non-Hispanic black children has stayed relatively constant since 1980 (14–15 percent) (Child Trends, 2018c). From 1994 to 2017, the population of children of immigrants[11] grew by 51 percent, from 18 to 27 percent (Child Trends, 2018a). These children are mainly second-generation immigrants (16.7 million), with first-generation immigrants making up a much smaller number (2.9 million, which is 3–5 percent of all children). In 2016, 33 percent of U.S. children lived in a household where more than one language is spoken (dual-language learners [DLLs]), with Spanish being the most common other language for these children (Child Trends, 2016a).

Evidence shows that populations from immigrant backgrounds and ethnic and racial enclaves might bring or develop particular child-rearing practices that lead to strengths and weaknesses in their children's adaptations and success in other environments. For example, Galindo and Fuller (2010) have shown that Latino children have higher social-emotional skills in preschool but some of the academic skills of low-income Latino children lag beyond other groups. School success as measured by grades is associated with different family constellations and practices in immigrant populations, such as Cambodian, Dominican, and Portuguese populations (Garcia Coll and Kerivan Marks, 2009). When examining cultural variation, no particular parenting or household composition is associated with success across all domains of child outcomes, aside from preventing extreme neglect, abuse, or lack of stimulation.

A more general cultural shift has been the increase in personal-use technology (e.g., computers, tablets, mobile phones), which has advanced at a rapid pace in the last 20 years. The divide in access to digital devices has decreased significantly, with more families having access to smartphones across socioeconomic lines. Access to other devices, however, continues to create a digital divide that children in non-mainstream populations have to surpass. School districts across the country are integrating technology into the classroom. The health care field has advanced in the area of telehealth, providing greater remote access to health care providers. This access can have positive and negative effects, and it can even increase inequities when not used properly.

ABOUT THIS REPORT

Report Conceptual Model

The committee's conceptual model (Figure 1-9) served as a unifying framework for the committee's approach to the report, aiming to improve prenatal through early childhood development with a health equity approach. The model adapts elements and concepts from the World Health Organization (WHO) Commission on Social Determinants of Health conceptual framework (WHO, 2010), the closed-ecological model (Bronfenbrenner, 1979; McLeroy et al., 1988; NASEM, 2017; Velez-Agosto et al., 2017), a model of children's health and its influences (NRC and IOM, 2004), and the life course health development model (Halfon et al., 2014). Building on the life course model, this figure shows early development within the context of the life course stages, beginning with preconception and ending with adulthood along the bottom. It

[11] Immigrant children are defined here as those who have at least one foreign-born parent.

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is important to note that risk and protective factors can be transferred intergenerationally, which makes the parent or primary caregiver and/or the parent–child dyad a central focus of intervention. With the necessary supports and conditions, healthy biological, socio-emotional, cognitive, and socio-behavioral development increase across the life course.

Within the context of the life course, the diagram's nested circles illustrate the complex sociocultural environment that shapes development at the individual level and the opportunities for interventions to improve individual health and developmental outcomes and population health, well-being, and health equity. The context and conditions here continue to play an important role in health and well-being throughout the life course. Individual social and biological mechanisms operate and interact within and across the three levels.

The outer level, "socioeconomic and political drivers," is adapted from the WHO social determinants framework and represents the level at which structural inequities operate. These structural inequities are deeply embedded into policies, laws, governance, and culture; they organize the distribution of power and resources differentially across characteristics of identity (i.e., race, ethnicity, gender, class, sexual orientation, and others) (NASEM, 2017; WHO, 2010).

The next level represents social, economic, and environmental conditions (i.e., the SDOH): education, employment, health systems and services, housing, income and wealth, physical environment, transportation, public safety, and social environment. In the model, these nine interdependent factors are grouped into three domains that the committee has identified, based on the available evidence and existing resources, as important for targeting prenatal and early childhood interventions: healthy living conditions, ECE, and health systems and services. These domains in the gray and light blue circles are the primary foci of Chapters 5–7, and each is discussed in terms of evidence-based solutions, opportunities for intervention, barriers, promising models, and research needs.

The next level represents the factors that most directly and proximally shape children's daily experiences and routine patterns; these include caregiver well-being and support and attachment and family cohesion, which affect social connections in early life.

Culture, according to Velez-Agosto and colleagues (2017), operates at various levels, including all the levels in the committee's conceptual model. Systems, such as ECE and health care, have cultures and also enact in their daily operations ways of conceptualizing the child's development, how to promote it, and what resources have to be accessed in order to correct illness and developmental problems. Cultures can also be sources of strength, providing support for coping with life demands and toxic stress (Garcia Coll et al., 1996). Families and communities also have cultures that are similarly enacted in daily routines and in developmental goals and expectations. Inequities might arise from lack of understanding of normative cultural frameworks between any of these levels, lack of respect and acceptance for different ways of being that are considered normative in other settings, and lack of access to critical resources, such as high-quality ECE and health care, due to racism and discrimination (Garcia Coll et al., 1996).

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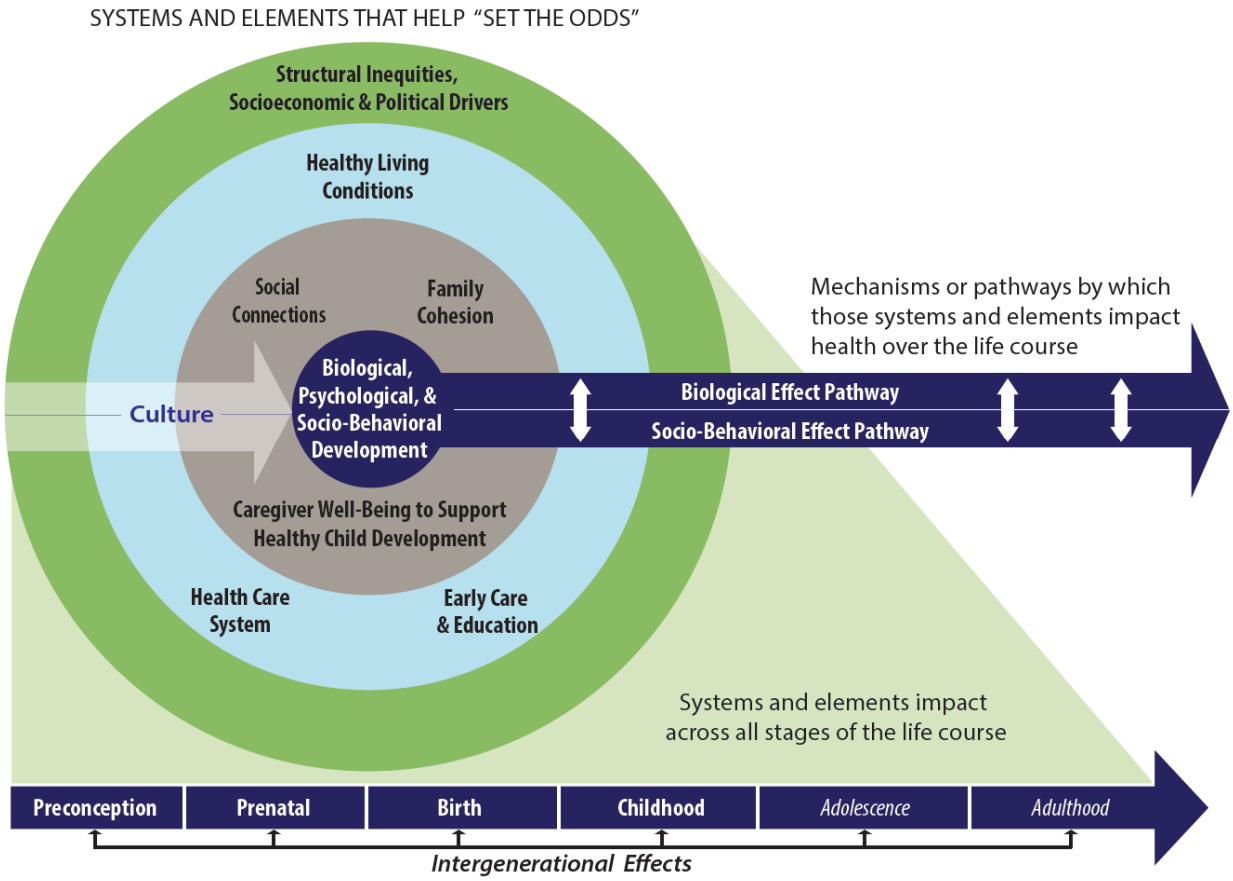

FIGURE 1-9: Leveraging early opportunities to achieve health equity across the life course: A conceptual framework

NOTE: The elements and systems included in the nested circles impact every stage of the life course.

Types of Interventions

The committee deployed lessons from the latest insights in neurobiological and closed-behavioral sciences that define how early childhood experiences translate into health outcomes and inequities across the life course in order to offer promising practices or transformational levers that might help move the needle in a positive direction. In its Statement of Task (see Box 1-3), the committee was asked to look at interventions defined as policies, programs, or system changes. What is meant by each of these is briefly described below (see Box 1-5 for examples).

Systems

Systems are a collection of interacting, interdependent parts that function as a whole. For the purposes of this report, most of the systems are social constructs and are organized around a key functional area (education, health care, criminal justice). Systems have existing patterns and structures that define how people tend to move through them. A few essential features that define a system include

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- **Structures:** The essential organizational components of a system, including how it is governed.
- **Flows:** How information, money, or people move through and are configured in relation to each other.
- **Goals:** The actions the system is trying to accomplish—its purpose or function.
- **Rules and Norms:** How a system is organized and what can and cannot happen within it.
- **Paradigms:** Assumptions that system makes that inform how things are organized and flow within it.

These features could be thought about in the context of a single system or considered through a social ecosystem lens and applied to how systems interact with each other.

Programs

A program is a targeted or specialized pathway by which a specific group of people move through a system or between a set of systems—a change in the system's structures or flows for certain kinds of clients. It is often driven by a recognition that the standard approaches are not achieving the system's goals for some people. Programs usually do not change the goals or rules or the system itself; rather, they generally offer a new way to achieve improved outcomes within the current system's structures (such as a support group for new mothers offered in a primary care setting).

Policy

Policy is a shift in the overarching legal or regulatory structure that governs how systems operate or interact with one another. Policies usually set the rules, goals, norms, or paradigms within which systems establish structures and flows, and they may also set the limits on the types of exceptions or changes that systems can put in place. Some policies can impact large swathes of the population, such as laws, regulations, court rulings, administrative rules, or executive orders, and often have consequences when not followed. Some policies have a smaller scope, such as guidelines issued by professional organizations, recommendations of expert panels, or local programs.

In general, policies, systems, and programs can be thought of as nested within each other. The impact profile of efforts at any given level may be constrained by what is happening in the others—promising programs that do not consider the potential limitations that the policy environment places on potential adoption and spread will not be as useful as those that do.

BOX 1-5
Example of Systems, Policy, and Programmatic Changes

Conceptual Example: How do we build more integrated care that helps support people in both the clinical and social drivers of poor health outcomes?

Systems Change: Altering the basic or fundamental structures or patterns by which information, money, people, or other resources flow through or between systems.

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- Example: A health care system implements standardized social determinants screening (for example, asking questions at medical appointments about housing and food security) for all patients and builds out a standard referral system to community agencies.

Policy Change: Altering the overarching legal or regulatory structure that governs how systems operate or interact with each other.

- Example: Data and privacy regulations are changed to allow partners in other sectors to see and share important information on their mutual clientele.

Programmatic Change: Creating specific alternative structures or patterns that alter the default experience for targeted subsets of a population.

- Example: A system builds an enhanced model of intensive case management for its clients with the most intensive needs. These clients receive extra supports, but the experiences of others remain mostly unchanged.

Committee's Methodology

As discussed above, an intervention could consist of policies, programs, or system changes or a combination of those approaches. The literature on the effectiveness and applicability of interventions provides important information for assessing which interventions are most effective and suitable for a general or more specific population. However, many interventions have not been adequately evaluated for their effectiveness—in general, for a specific outcome, or when brought to scale. In addition, studies vary in dimensions such as appropriateness of design and setting, quality of execution, interactions with other interventions, and consideration of economic consequences.

With the above in mind, the committee examined the available literature, conducting a comprehensive review from the peer-reviewed and gray literature. [12] This entailed a systematic search of academic and governmental databases and websites for studies that evaluated or assessed the effects of interventions. Where possible, the committee relied on existing systematic reviews and meta-analyses with strong methodologies and existing comprehensive reviews.

Comparisons across studies are needed to better assess intervention opportunities; however, such comparisons are often not available or difficult to interpret due to methodological strengths and limitations relating to the nature of the study design and the data collected. For example, what is considered the gold standard research design to show that a program does in fact lead to the results it is trying to achieve—the randomized controlled trial (RCT)—may not always be feasible. This is because RCTs are not appropriate in all settings, particularly not to study community-level interventions, and for some interventions, randomization can be considered ethically objectionable if it denies a service or treatment known to be beneficial (Center on the Developing Child, 2016). While large-scale, multiyear RCTs will continue to be important to demonstrate program impacts, there is also an important role for short-term, flexible study methodologies to test program components and subgroup variability and allow for mid-course corrections and enhancements (such corrections are not possible in traditional RCTs) (Center on the Developing Child, 2016). While there is nothing fundamental in the design of an

[12] The date range for the search was 2013–2019. Search terms included the following categories: prenatal and early childhood, neurobiological, socio-behavioral, biology/biomarkers, health indicators/outcomes, structural inequities, social determinants of health, and model/intervention.

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RCT that precludes investigating heterogeneous effects in different populations, to do so requires careful consideration of the sample size and identification of subgroups of interest prior to the randomization. As a result, many RCTs have not been able to answer important questions about variability in effects across subpopulations or which components of an intervention are driving the results. While RCTs can be used to study the effect of separate program components, it is difficult to use RCTs to examine complex programs with multiple dimensions (or for community-level interventions). RCTs remain a critical tool for studying the effectiveness of programs, but it is important to note their limitations as well. (See Chapter 8 for a research recommendation on this topic.)

Because of the limitations of RCTs in some contexts, and limitations of observational studies that are correlational in nature, researchers have increasingly used "natural experiments" to estimate causal effects. Such studies harness changes in state and local policies that generate plausibly random or quasi-random variation in exposure to a given service or treatment to estimate its causal effect on outcomes of interest (see Angrist and Pischke, 2009) for an overview of these methods). Despite the limitations of estimation based on observational data, careful use of observational data has many advantages. First, it is very useful for identifying associations that can be more rigorously studied using other approaches. Second, in some cases, careful use of natural experiments or other research designs can minimize the bias from confounding. Finally, some questions are, by their nature, not amenable to randomized trials and so can only be studied using observational data. The way data is gathered across study designs also varies, as data collection may occur for different outcomes at different points in time, so the data may be incomplete and therefore hard to compare.

When reviewing the science on a certain intervention/policy, there may be cases with strong evidence for one or more important outcomes and weaker (or mixed, or null) evidence for one or more other outcomes. This does not mean that the intervention/policy should not be considered or implemented. If there is a strong theoretical basis for making the change (for example, from a biological pathway or development standpoint) and strong evidence for one or more salient health outcomes that is being targeted and has a large population health impact, the intervention/policy could be deemed appropriate despite its limitations. This is true for several interventions and policies recommended in this report.

About the Report Recommendations

The committee provides a range of recommendations for practice, policy, and systems change, including recommendations that will take time and sustained commitment to achieve and recommendations that could be implemented immediately or in the near term. Some of the committee's recommendations will be difficult to implement; however, the degree of difficulty in implementing any given recommendation does not mean it is not worth pursuing. Achieving health equity for children will require attention and commitment from a range of sectors. For example, in Chapter 8, the committee discusses the need for better alignment among the many children- and family-serving sectors in the United States. Although there are many barriers to achieving alignment, there is also great opportunity to make long-lasting reductions in long-standing and persistent inequities. Furthermore, inequities that originate at the system and institutional levels will require solutions that target policy and structures. Given this understanding, the committee provides the recommendations with the long-term outcomes in mind and the goal of both improving the current state and making strides today to establish systems where inequities will no longer be the status quo. Where possible, the committee also

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recommends or highlights ways to leverage existing programs that either embrace the core scientific principles laid out in this report (see the Core Principles below and Chapter 2) or have a strong basic structure from which to build (for example, Home Visiting). Therefore, the committee makes some recommendations for improving programs that are not optimally or extensively implemented but have the potential to be updated, scaled, or better used to promote related services that advance health equity.

Given the fragmented nature of many systems in the United States (for example, the health care system has both public and private payers and multiple entities that set guidelines for care and accountability), many of the committee's recommendations are directed to multiple actors with varying roles and responsibilities, each of whom is important to advancing child and family outcomes. Further, as a matter of jurisdiction, some recommendations are better targeted to varying levels of government (e.g., local, state, tribal, and territorial). To address deeply rooted inequities, which play out across multiple sectors, some recommendations entail a comprehensive approach that requires partnerships across sectors and levels of government—this is consistent with *Communities in Action* (NASEM, 2017), which identifies cross-sector collaboration as a key element for promoting health equity in communities.

Resources Resources will be needed to implement many of the recommendations in this report. The root causes of structural inequities are found in differential access by virtue of race or ethnicity to resources for some groups, due in part to certain laws and policies (NASEM, 2017; see also Chapter 3). Calling for change of any kind to advance health equity will require changes to laws, policies, and other sources of inequity so as not to perpetuate structural racism and discrimination. Decision makers and leaders can coordinate, integrate services, and educate on all of these issues, but if the resources are not there to back the needed changes, the change likely will not occur. The committee was asked by RWJF to identify *what* practice, policy, and systems changes need to be implemented, based on what is known from developmental science; however, the committee was not tasked to identify the sources or mechanisms for funding those changes. In a few instances, the committee was able to provide estimates of the potential costs for implementing its recommendations using existing estimates. Doing so, however, was complicated when those estimates did not include cost savings from improved health outcomes resulting from the intervention, producing a one-sided picture.

Although the committee was not asked to identify the mechanisms and sources for implementing the report recommendations, there are many extant mechanisms and proposals for funding various public or population health activities. For example, taxes on tobacco, alcohol, and other products (such as sugary beverages) have been used to channel resources and shape economic incentives (NASEM, 2018b). An IOM (2012) committee proposed establishing a tax on medical care transactions to provide a long-term financing structure to cover public health services. Other mechanisms—both within and outside the health sector—include federal or state Wellness Trusts (Prevention Institute, 2015), hospital community benefit expenditures (Rosenbaum et al., 2015), community wealth-building (including the role of anchor institutions, see Porter et al., 2019), and Children's Services Councils[13]. (A 2018b National Academies proceedings provides an overview of several of these options and others.) Each of these financing mechanisms has advantages and disadvantages that would need to be weighed for the specific intervention or program for which it is considered.

[13] See, for example, http://flchildrenscouncil.org/about-cscs/overview (accessed June 17, 2019).

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Promising Models

In its statement of task, the committee was asked to identify promising examples of models that apply the science of early development to health equity. The committee adapted the selection criteria used in the *Communities in Action: Pathways to Health Equity* (NASEM, 2017) report to guide this process.[14] The committee used the three sets of criteria from the National Academies 2017 report (see Appendix A for a list of all of the selection criteria), which were informed by research and practice-based evidence and the expertise of the committee members. The committee made a few additions based on the focus of this report (the prenatal through early childhood period) and removed those not relevant to this study. The first set consists of six core criteria, which need to be met by all the promising models. These core criteria assure that the examples chosen are substantively significant. The committee used these promising models as examples throughout the report to highlight bright spots that have been able to use what is known from the science to advance health equity in the preconception through early childhood periods.

Furthermore, "promising" does not imply that the model is new but rather that it is a program or intervention that met the committee's core criteria, and each promising model has a unique approach and is at a different phase of development: some have been around for more than 30 years and have changed based on evaluations or input from users, while others have emerged in the past few years. In Chapters 4–7, three promising models were identified for each chapter (each model is summarized in a box). These examples are not blueprints, and exact replicas might not work with all populations or locations; however, the lessons learned and approaches used prove valuable to those working to create positive change toward health equity during the preconception through early childhood periods. Note that throughout the report, the committee cites many other examples of current practices and programs that illustrate the topic being discussed; however, those examples may not meet all of the committee's criteria (and are not labeled as promising models).

The first core criterion requires that the model's main focus be during the prenatal and/or early childhood period, as that is the focus of this report; this inevitably involves the mother (or other primary caregiver). The second notes that the intervention is informed by findings from the neurobiological, socio-behavioral, and/or biological sciences—also a focus of this report. The third core criterion is that the model addresses at least one (preferably more) of the nine SDOH identified by the NASEM 2017 report: education, employment, health systems and services, housing, income and wealth, the physical environment, public safety, the social environment, and transportation. "This criterion was informed by the wealth of literature suggesting the importance of targeting the social and economic conditions that affect health, especially at the community level" (Bradley et al., 2016; Galea et al., 2011; Heiman and Artiga, 2015; Hood et al., 2016; NASEM, 2017; Wenger, 2012). Furthermore, this criterion is basic to the committee's charge, which posits that the SDOH need to be addressed to reduce health inequities. The fourth criterion is that the model is designed to have or has evidence of having an impact on a group or population that experiences health inequities.

The fifth core criterion "states that the solution needs to be multisectoral, meaning that it engages one or more sectors" (ideally, at least one "nontraditional" sector, meaning other than public health or health care). Multisector collaboration is a powerful lever for addressing health

[14] The original criteria from the National Academies (2017) report are available at https://www.nap.edu/read/24624/chapter/7#323.

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inequities and building a culture of health (APHA, 2015; Danaher, 2011; Davis et al., 2016; Kottke et al., 2016; Mattessich and Rausch, 2014). Engaging stakeholders across multiple sectors provides the opportunity for innovative and cost-effective methods to sustain solutions at the community level (NASEM, 2017).

The sixth core criterion requires the solution to be evidence informed. "This entails an assessment of evidence or the best available information to identify a problem and develop a solution that has a measurable outcome. Here, there is considerable flexibility in terms of the type of evidence that will qualify. This flexibility is based on the understanding that low-resource communities that suffer from health inequities often do not have the infrastructure, personnel, or financial resources to provide the highest standard of evidence" (NASEM, 2017, p. 326). The sixth core criterion is that there needs to be an evaluation plan with identified outcome measures to track the impact of the intervention.

The second set of criteria—aspirational criteria—reflects the elements, processes, and outcomes of interventions that the committee identified as valuable for promoting health equity. This set of criteria highlights important features of interventions, such as nontraditional partners or nonhealth domains (e.g., community organizers, public libraries, Parent Teacher Association groups) and an intervention being interdisciplinary and/or multilevel (the intervention has multiple levels of influence, such as individual, family, organizational/institutional, or governmental).

THE SCIENCE OF EARLY DEVELOPMENT: CORE CONCEPTS

Based on its review of the science, the committee updated the core concepts from the 2000 report *From Neurons to Neighborhoods* and identified 12 core concepts of early development, with a focus on equity. The evidence behind these concepts is described in Chapters 2–4, and this evidence guided the committee in developing recommendations that are responsive to the science of early development.

1. **Biology–environment interaction impacts health and development:** Human long-term social, emotional, behavioral, cognitive, and physical health is shaped by genetic, epigenetic, and environmental factors that integrate biological information at the level of molecular, cellular, and organ systems with the family, neighborhood, and culture in which the individual is embedded. These developmental processes start before conception through a dynamic and continuous interaction between biology and aspects of the environment and extend throughout the life-span.
2. **Brain development proceeds in well-defined but continuous steps:** Human developmental processes are now conceptualized as continuous, rather than occurring in discrete stages as was originally thought. A sequence of temporally well-defined sensitive periods (sometimes referred to as "*critical periods"*) of brain maturation—in tandem with environmental influences—lead to the acquisition of social, emotional, and cognitive skills. During these sensitive periods, early adverse experiences influence brain development and can alter the trajectories of development in each of these functional domains, impacting long-term well-being.
3. **Major physiological systems develop rapidly in pregnancy and early childhood:** It is now known that physiological systems other than the brain, such as the immune system, microbiome, and endocrine system, can also be influenced preconceptionally. Like the

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brain, these systems begin to develop prenatally, with early sensitive pre- and postnatal periods that can be disrupted by early adversity. These early influences can have long-term consequences for mental and physical health.

4. **The early caregiving environment is crucial for long-term development:** The family and community caregiving professionals have a central role in early childhood development, indicating the need for a multigenerational approach to assuring optimal growth and development. The child thrives in healthy relationships throughout childhood, but relationships with parents are the building blocks for subsequent relationships and child outcomes. For example, maternal–child interactions impact the well-being of both mother and child, where a positive mother–child relationship rests on good maternal well-being (whereas prolonged maternal distress impairs parenting and the quality of attachment).
5. **The developing child plays an important role in interactions and development:** Children are active participants in their own development, reflecting the intrinsic human drive to explore and master one's environment. Early experiences create biological structures, and shape psychological and behavioral adjustment, in ways that influence how the child will react and act in response to later environmental demands.
6. **The development of executive functions[15] is a key aspect of early childhood development:** The emergence and maturation of executive functions (inhibitory control, mental flexibility, and working memory) are a cornerstone of early childhood development. Executive function maturation continues through early adulthood and contributes to the health, well-being, and productivity of adults.
7. **Trajectories—positive or negative—are not immutable:** The developing child remains vulnerable to risks and open to protective influences through adolescence and young adulthood, although early life represents the stage at which interventions are most effective in positively influencing a child's development.
8. **There is variability of individual and group development:** Individual children demonstrate significant variability in these sensitive periods of brain development and in response to interventions. This variability often makes it difficult to distinguish among typical development, maturational delays, transient disorders, and persistent impairments or to predict response to treatment. Understanding this heterogeneity of development and individual differences in access and responsiveness to interventions is a major goal.
9. **Experiences across environmental contexts play a significant role in early development:** Children can be more or less sensitive to experiences due in part to their biological makeup, but this is impacted powerfully by multiple family and community factors. These include family interactions and adversity, accessibility to community programs and interventions, and environmental factors, such as quality and stability of housing, toxic environmental exposures, food accessibility, early childhood education, family support, and culture. Effective interventions can alter the course of development

[15] Executive function (and self-regulation) skills "are the mental processes that enable us to plan, focus attention, remember instructions, and juggle multiple tasks successfully. Just as an air traffic control system at a busy airport safely manages the arrivals and departures of many aircraft on multiple runways, the brain needs this skill set to filter distractions, prioritize tasks, set and achieve goals, and control impulses" (Center for the Developing Child at Harvard University, see https://developingchild.harvard.edu/science/key-concepts/executive-function/, accessed June 17, 2019).

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throughout childhood by changing the balance between risk and protective processes, leading to more adaptive outcomes.

10. **Disparities in access to critical resources matter:** Disparities in health and developmental outcomes result from not only health care disparities but also disparities and inequities in family and community factors. Achieving equity in health and developmental outcomes in young children requires addressing health care disparities and also assuring those family and neighborhood conditions in which all children and families can be healthy and thriving.
11. **Health outcomes are the result of experiences across the life course:** Birth and early developmental outcomes are influenced by the health of the mother and father not only during pregnancy but also before and between pregnancies and across the life course, starting with their own prenatal and early life experiences. Disparities in the child's birth and early developmental outcomes are therefore the consequences of not only differential exposures during pregnancy and early childhood but of differential parental life course trajectories set forth by early programming mechanisms and influenced by cumulative life adversity.
12. **Early interventions matter and are generally more cost effective than later ones:** Family-based interventions that consider both risk and resilience and are informed by or sensitive to the mores and values of the target population can potentially positively change developmental pathways. The integration of various levels of services that are community based and have supportive policies in place tends to be more effective.

OVERVIEW OF THE STUDY PROCESS AND REPORT

To address its charge, the committee gathered information through a variety of means. It held two information-gathering meetings that were webcast live, in August and October 2018 (the meeting agendas are listed in Appendix B). The committee also held deliberative meetings throughout the study process. The committee received public submissions of materials for its consideration throughout the course of the study.[16] The committee's online activity page provided information to the public about its work and facilitated communication with the public.[17]

Throughout this report, the committee provides conclusions and recommendations. Chapter 2 provides an overview of healthy development from conception through early childhood, including what is new in the science of development since the publication of *From Neurons to Neighborhoods* (NRC and IOM, 2000). It aims to demystify the "black box" of development for the public, practitioners that work with children, and policy makers and to convey that we know enough about early development to act now to advance health equity for children and families. Chapter 3 provides an overview of the critical influences or factors that can either promote or hinder healthy development, with a focus on factors that shape inequities at the child/family level and the community and population levels across the SDOH. Chapter 4 focuses on how to best foster children's healthy psychosocial development, emotional adjustment, and physical health using what science has shown about risk and resilience and the importance of healthy relationships among children and families in high-risk contexts. Chapter 5

[16] Public access materials can be requested from paro@nas.edu.
[17] See nationalacademies.org/earlydevelopment.

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explores the role of the health care system in advancing health equity and what the system would need to look like from preconception through early childhood to meet the developmental needs of children. Chapter 6 provides recommendations on how to better meet the fundamental needs of families and children through economic security, stable and safe housing, and protection from environmental toxicants. Chapter 7 discusses the critical role of ECE and how it can serve as a platform for advancing health equity for children and families. Chapter 8 addresses needed systems changes and summarizes the opportunities to overcome barriers to strengthen a systems approach, the key stakeholders who need to be involved, and necessary alignment, measures, and research based on the committee's assessment of the literature in Chapters 2–7. Finally, Chapter 9 highlights the main findings and concepts discussed throughout this report and summarizes the report recommendations, laying out a roadmap for applying and advancing the science of early development.

CONCLUDING OBSERVATIONS

The science of early development is clear. Long-term physical health and emotional, behavioral, social, and cognitive competence is shaped by genetic, epigenetic, and environmental factors, including their interactions, before conception and through the life-span. There are risk factors that necessitate action at the practice, policy, and systems levels that take into account the full range of factors that impact health and well-being. These actions need to be taken before insults to early development occur; however, the science of plasticity indicates that it is never too late to intervene. These actions need to take a life course, multigenerational approach to make progress on health inequity, because the well-being of a child depends on the well-being of a parent/caregiver. Multipronged, cross-sector interventions, focused on prevention, early detection, and mitigation and working at the policy, system, and program levels, are needed to move toward health equity.

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2

Healthy Development from Conception Through Early Childhood

INTRODUCTION TO THE SCIENCE FROM PRENATAL THROUGH EARLY CHILDHOOD

Fetal and early child development are often conceptualized as a "black box," in which children simply learn by soaking up information, much like a sponge and water. Developmental science has demonstrated that development is, in fact, an active process and that it starts early. Thus, understanding early development, when biological systems and functions are coming "online," is essential for ensuring that prevention, interventions, policies, and systems are responsive to the needs of children. This will provide children the best opportunity for healthy outcomes. When the science of early development is coupled with a health equity approach to inform decision making, it provides an opportunity to improve outcomes for children and families in at-risk contexts. The goal of this chapter is to reveal core concepts of brain development and other systems relevant to understanding the impact of disparities on cognitive, social-emotional, and physical health and to describe new scientific advances in this area over the last 20 years. This information can be used by the public and policy makers to better inform effective actions for promoting healthy outcomes. (See Box 2-1 for an overview of this chapter.)

BOX 2-1
Chapter in Brief: Healthy Development

There are theoretical, technical, and scientific advances over the last 20 years that have been instrumental in rapidly advancing the science of child development. This chapter highlights those advances and provides an overview of the core concepts of brain development and other body systems relevant to understanding the impact of early life adversity on cognitive, social-emotional, and physical health, including the mechanisms that link early life experiences to later outcomes. This information can be used by the public and policy makers to better inform effective actions for advancing health equity.

Scientific advances in the last 20–30 years include

- Early experiences are essential for building brain connections that underlie biobehavioral health, and current understanding of whole-child development relies on an interplay of organ systems with each other and the environment.
- Responses to pre- and postnatal early adversity, such as poverty or maltreatment, are rooted in gene–environment interactions that can result in altered molecular and cellular development during sensitive periods of development.

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- Chronic early adversity has negative impacts on the assembly of certain brain circuits and emotional control, cognitive growth, and stress responsiveness.
- Certain body systems involved in cognitive and emotional development are more sensitive to early disturbances that activate stress response networks and provide a basis for both short- and long-term functional consequences of early life stress.
- Early adversity can change the timing of critical periods of brain development, impacting the "plasticity" of developmental processes that are driven by experiences in the life of the young child and the family.
- Deciphering the mechanisms through which adversity disrupts foundational developmental processes of the whole child and the long-term impact on health, wellness, personal and community relationships, and life-span productivity is being achieved through multidisciplinary, longitudinally designed research that has incorporated principles of human development, including individual differences and differential sensitivity to context.

Chapter conclusions in brief:

- Healthy development of the child begins in the preconception period and is dependent upon a strong foundation built prenatally.
- Among all the factors that may serve to buffer negative outcomes produced by toxic stress, supportive relationships between the child and the adults in life are essential.
- When considering the entire life course, it is early experiences, pre- and postnatally, that are the most powerful in working together with individual genetic makeup to influence the physical, mental, and cognitive development of the child.
- Science shows that maternal health and well-being is a major contributing factor to the development of the fetus and establishes a foundation for positive or negative child health outcomes.
- Based on the abundant science, the influence of access to basic resources prenatally, particularly nutritional, psychosocial, and health care components, is powerful. Resources to help families to limit chronic stress may reduce risk for disrupted development and help close disparities based on race, ethnicity, and socioeconomic status (SES).
- New data, combined with previous studies, provides a compelling demonstration that toxic stress substantially increases later-life risk for lower educational achievement and physical illnesses, including obesity and type 2 diabetes, cardiovascular disease (CVD), substance abuse, mental illness, cancer, and infectious disease. In addition to the adverse effects of toxic stress on the brain and nervous system, it also affects every other organ system in the child's body, impacting short and long-term health.

Early understanding of children's development, as postulated by Piaget and many others in the 20th century, consisted of age-specific stages, ruled primarily by maturation and biology (Flavell, 1963). This characterization has undergone profound revisions. The 2000 publication by the National Research Council (NRC) and the Institute of Medicine (IOM), *From Neurons to Neighborhoods: The Science of Early Childhood Development*, provided a transformational synthesis of existing research to reveal a complexity of factors that influence development beyond previous simplified models. This landmark report integrated a wealth of scientific knowledge on early childhood development to emphasize the continuous, dynamic interaction between the environment and the biological systems of each individual across the life course. The report offered clear recommendations for policy and the future of developmental science that were organized around core principles of child development that remain accurate to this day (NRC and IOM, 2000). The report inspired new science and novel ways to communicate the science to the public, service providers, and policy makers. *From Neurons to Neighborhoods* spurred research to refine the fields' understanding of how environmental influences shape child

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development, even at a molecular resolution. The report motivated research to understand how health-promoting environments and responsive relationships enable a strong foundation for healthy child development (i.e., learning, adaptive behaviors, and optimal health), while chronic adversity in the absence of supporting caregivers can lead to increased risk for negative learning and health outcomes, both in childhood and well into adulthood.

In the two decades since publication of the report, there has been a convergence of activities that led to many of the advances that will be described in this chapter. First, a wave of neurobiological studies in model systems and in humans underscored that responses to pre- and postnatal early life stress are rooted in genetic and environmental interactions that can result in altered molecular and cellular development that impacts the assembly of circuits during sensitive periods of development (Cameron et al., 2017; McEwen and Morrison, 2013; Shonkoff and Levitt, 2010). The demonstration that certain systems involved in cognitive and emotional development are more sensitive to early disturbances that activate stress response networks, such as the frontal cortex, hippocampus, amygdala, and hypothalamic-pituitary-adrenal (HPA) axis, provided a basis for both short- and long-term functional consequences of early life stress (Chen and Baram, 2016; Shonkoff et al., 2012). Second, life course theory (Ben-Shlomo et al., 2014; Kuh et al., 2003) and the developmental origins of an adult health and disease framework (Gluckman et al., 2007; Halfon et al., 2014) have become more widely adopted. Third, a key policy statement from the American Academy of Pediatrics in 2012 emphasized the important life-span health implications of the impact of early adversity (Shonkoff et al., 2012), declaring a "need for the entire medical community to focus more attention on the roots of adult diseases that originate during the prenatal and early childhood periods" (p. e233). Numerous research teams have clearly documented the effects and some mechanisms through which early childhood adversity affects development. Early adversity, such as maltreatment (Cicchetti, 1996; Teicher et al., 2016) and poverty (Cicchetti and Curtis, 2006; Council on Community Pediatrics, 2016; Gabrieli and Bunge, 2017; Johnson et al., 2016; McEwen and McEwen, 2017), has negative impacts on selective brain circuits and emotional control, cognitive growth, and stress responsiveness. A very new discovery, from multiple animal studies, shows that early adversity can change the timing of critical periods of brain development (Bath et al., 2016; Cameron et al., 2017; Hensch, 2016a; Heun-Johnson and Levitt, 2018), impacting the "plasticity" of developmental processes that are driven by experiences in the life of the young child and their family (Cicchetti, 2015a,b; Cicchetti and Blender, 2006; Pollak et al., 1997) (see Box 2-2). The extensions of the original retrospective Adverse Childhood Experiences (ACEs) Study by CDC and Kaiser Permanente fostered sustained scientific interest in the long-term implications of early childhood adversity (see Chapter 3 for more information). In addition, the prospective Dunedin Multidisciplinary Health and Development (Poulton et al., 2015), the Perry Preschool longitudinal study, and the Abecedarian studies (see Chapter 7 for more information on the Perry Preschool and Abecedarian projects) added to a growing body of scientific evidence that early experiences, positive or negative, have profound long-term effects on physical, mental, and cognitive functions. The U.S. Department of Health and Human Services introduced Healthy People 2020—and, subsequently, Healthy People 2030—which elevated the importance of research on strategies to intervene on health disparities, including among pregnant women and children, by declaring "the elimination of health disparities" as a national goal.[1]

Taken together, these research endeavors have generated many new questions about *when* and *how* early childhood adversity is incorporated biologically to influence long-term outcomes

[1] For more information, see https://www.cdc.gov/dhdsp/hp2020.htm (accessed June 17, 2019).

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and about optimal approaches to address early adversity and racial/ethnic and socioeconomic health disparities. These activities, and others, have pioneered and proliferated a new field of research on the importance of prenatal and early childhood experiences for health across generations.

Beyond scientific advances, *From Neurons to Neighborhoods* can also be credited as the impetus for the establishment of the National Scientific Council on the Developing Child in 2003, which uses science to catalyze public and private activities to promote the healthy development of children (Center on the Developing Child at Harvard University, 2014), to "close the gaps" between what the science says and which actions will promote child well-being. Composed of subsets of members of the *From Neurons to Neighborhoods* committee and the MacArthur Research Network on Early Brain and Child Development, the Council has a central objective of communicating the science to the public and policy makers. Through an ongoing collaboration with FrameWorks Institute, applying qualitative and quantitative research on communicating the science of brain and child development, a suite of effective "explanatory metaphors" have been developed, including a taxonomy of the stress response: positive, tolerable, and toxic stress (National Scientific Council on the Developing Child, 2014; Shonkoff and Bales, 2011). The term "toxic stress" was proposed to describe "excessive or prolonged activation of the stress response systems" in the absence of buffering protection from adult caregivers (National Scientific Council on the Developing Child, 2014). Unlike positive or tolerable stress, "toxic stress" refers to biological changes in the child that can result in disruption of developing brain architecture and other maturing organs, dysregulation of metabolic processes, and excessive stress system activation. These responses can lead to increased risk for chronic diseases later in life (Shonkoff et al., 2009). These explanatory metaphors have been adopted by the American Academy of Pediatrics (Garner et al., 2012; Shonkoff and Garner, 2012) and helped the science of child development to be incorporated into federal and state legislation that generated evidence-based policies (Thompson, 2016; Center on the Developing Child at Harvard University, 2014).

This chapter highlights a selection of important scientific advances since *From Neurons to Neighborhoods* (NRC and IOM, 2000). Next, key processes of healthy development from conception through early childhood are described, and some of the major biological responses to stressors are summarized. The chapter concludes with a discussion of individual differences in responsiveness and susceptibility. Throughout, the chapter identifies key points in development where disparities in health by race/ethnicity or SES can emerge, which could inform strategies for prevention-oriented interventions.

The goal of this chapter is to provide essential information about the state of the science on human development from preconception through early childhood and some of the exciting findings in the past two decades within the fields of neurobiology, social psychology, epidemiology, and others that have contributed to advancing knowledge about how and when to intervene to improve outcomes for children. All of this information needs to be read through the lens of life course theory (Ben-Shlomo and Kuh, 2002), which recognizes that early experiences influence health outcomes within and across generations. It is also important to keep in mind the interconnectedness of brain and body health, as brain activity controls peripheral systems, which in turn affect brain structure and function. The committee provides this evidence to help "bridge the gap" between what science tells us and what actions need to be taken—by policy makers, health care providers, educators, religious leaders, parents, and others—to close disparities and improve outcomes for all children in the United States. At present, many of the new scientific

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advances in neuroscience are still in development, and more study is needed to apply these new findings in clinical and public health practice and to use them to inform policies. In particular, greater effort and more support is needed to develop, implement, and evaluate programs based on scientific discoveries regarding the optimal timing for interventions. The committee could not be comprehensive with regard to coverage of all scientific advances, biological processes of healthy development, biological responses to stressors, and factors associated with individual differences, as each is a complex and active area of research.

BOX 2-2
Key Chapter 2 Terms

- **Adaptive immune system**: A portion of the immune system that uses highly specialized cells to eliminate foreign pathogens in the body and that, through exposure to certain pathogens, is able to eliminate those pathogens again in future infections.
- **Allostasis**: The process by which the body responds to internal or external stressors to return to homeostasis, or balance.
- **Amygdala:** A brain structure that is responsible for detecting threats in the environment and preparing the individual for emergency events.
- **Autonomic nervous system**: A network of nerves and ganglia that uses specific neurotransmitters (see below) to regulate internal organ function. The autonomic nervous system has two complementary components:
    - **Parasympathetic nervous system**: Often referred to as the “rest and digest” system, the parasympathetic nervous system activates the digestive system, slows breathing, and decreases heart rate.
    - **Sympathetic nervous system**: Often referred to as the “fight or flight” system, the sympathetic nervous system prepares the organism for action by propelling blood to the brain and limbs to provide increased oxygen and glucose for energy, slowing digestion, and increasing attentiveness.
- **Axon**: A main component of neurons; relatively long, extended structures that conduct a signal down the length of the neuron and play a key role in how neurons communicate with each other.
- **Brain plasticity**: The process by which neurons within the brain change their gene expression, function, cellular architecture, and connections with other neurons in response to experiences and changes in the environment.
- **Cellular homeostasis**: The notion that, when perturbed by some internal or external stimulus, cells react via feedback loops to achieve physiological balance with their immediate environment.
- **Chemical signature**: A general term used to highlight how epigenetics results in changes to DNA by placing chemical marks to modify how genes are expressed.
- **Connectome**: A general term for the amalgamation of connections between different parts of the brain.
- **Cortex**: The outermost region of the brain that exhibits the greatest expansion in evolution, where billions of neurons reside, trillions of connections are made, and much of the brain’s most complex computational activity takes place.
- **Cortisol:** A key hormone involved in the physiologic stress response. It elevates cell metabolism by binding to receptors in all organs to influence how they function during a “flight or fight” response. In the human brain, specific brain circuits express high levels of the glucocorticoid receptor. When cortisol activates the receptor, this impacts mood, motivation, and threat detection. Cortisol also helps to restore physiological balance after stress and regulates how the body uses carbohydrates, fats, and proteins; inflammation; blood pressure and blood sugar; and the sleep/wake cycle. Excessive exposure to cortisol prenatally also impacts development.
- **Endogenous versus exogenous**: Endogenous materials are produced within an organism or cell (e.g., hormone or neurotransmitter), whereas exogenous materials are introduced from the outside environment (e.g., medication or environmental toxicant).
- **Epigenetics**: The study of how genes are expressed or suppressed based on chemical changes to cells’ DNA due to biological factors that come from changes in the environment, including how these biological factors can be passed down from one generation to the next.

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- **Gene expression**: The mechanism by which DNA (composed of individual genes) is read and then turned into proteins that perform various functions throughout the body.
- **Genome**: An organism's complete set of DNA, including all of its genes. In humans, all cells that have a nucleus contain a copy of the genome, which consists of a highly organized arrangement of more than 3 million DNA base pairs.
- **Genome sequencing**: The scientific technique of identifying the genes that are present in a genome. This can be a sequence of either an organism's entire genome or certain parts of it.
- **Genomics**: The study of the structure, expression, and function of specific genes and of the entire collection of genes (genome).
- **Germ Cells**: Embryonic cells that contain half the genetic material (chromosomes) necessary to form a complete organism. In men, they produce sperm, and in women, they produce eggs/ova.
- **Glia**: A category of cells within the brain that do not make connections (synapses) like neurons but do perform many important functions: supporting neuronal metabolic health, contributing to the integrity of the blood-brain barrier to mediate access by blood-borne substances, viruses, and bacteria, controlling access by neurons to chemical neurotransmitters that are released by other neurons, aiding in the conduction of electrical signals, and removing cellular debris.
- **Innate immune system**: A portion of the immune system that removes foreign substances in tissues throughout the body and acts as a physical or chemical inflammatory barrier to infections but is not highly adaptable to a wide variety of infections.
- **Intergenerational transmission of risk**: The concept that epigenetic changes from certain behaviors in a parent may be transmitted from one generation to the next, separate and apart from the genes that are passed down through reproduction.
- **Lymphatic system**: The network of vessels through which waste products and toxins from tissues are drained for removal from the body.
- **Microbiome**: The collection of bacteria that comprise the microbial communities that exist on and within human beings.
- **Myelination**: The process by which oligodendrocytes (a type of glial cell in the brain) and Schwann cells (a type of glial cell outside of the brain) wrap axons with myelin, a protein- and fat-rich insulation substance that increases the speed of information conduction between neurons.
- **Neuroimaging**: A technique that provides structural, functional, or metabolic information and images from living human beings within a brain scanner.
- **Neurons**: Cells within the brain that communicate with each other via chemical signaling and underlie consciousness, sociability, emotion, motivation and reward, and cognition.
- **Neurotransmitters**: Substances synthesized and released by neurons to excite or inhibit other neurons at their points of connection—synapses.
- **Oligodendrocyte**: A glial cell responsible for wrapping the axons of neurons in myelin sheaths in order to aid the speed of electrical conduction between neurons.
- **Sensitive period**: A time in development in which the developing organism is especially likely to undergo change (either positive or negative) in response to some experience or environmental change (e.g., an environmental toxicant, intervention training, or certain nutrients). Sensitive periods for certain functions can extend into adulthood.
- **Synapse**: The connection between neurons, separated by a small gap, in which neurotransmitters and other chemicals are released by one neuron to influence the activity of other neurons with which they connect. This forms the basic unit of communication, resulting in the activation of circuits that process information and leading to a specific functional readout.
- **Transcriptomics**: The scientific field aimed at understanding how and when certain genes are expressed or suppressed.

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SCIENTIFIC AND TECHNOLOGICAL ADVANCEMENTS SINCE *FROM NEURONS TO NEIGHBORHOODS*

The big questions driving research on child development and interventions to improve well-being are the same today as they have been for decades. "What are the influences of genes and experience on child health and development?" "To what degree does the brain exhibit **plasticity** (i.e., change over time)?" "What are the brain circuits that control specific kinds of behaviors, and how can their functioning be influenced through experience-based interventions and/or medications?" "Why does early experience have lasting disease risk and functional effects into adulthood?" "How does a child's environment impact development of the brain, immune system, endocrine, metabolism, and cardiovascular systems, as well as physical and mental health?" (See Box 2-3 for highlights of applications of neurobiological research in practice.)

Much more is known today than two decades ago about the process of child development, and the explosion of research reports on child development since the publication of *From Neurons to Neighborhoods* (NRC an IOM, 2000) is remarkable. A PubMed search using just the term "early adversity" generated a list of 2047 research articles,[2] 96 percent of which were published since 2001. Advances in several related fields of study have developed simultaneously to refine the understanding of child health and development and the responses that each child may have to early adverse experiences. The following section outlines a selection of major developments, including genetics, transcriptomics, epigenetics, neuroimaging, computational modeling, maternal/fetal interactions, advances in longitudinal research, and the study of social interactions and physical environments. Refer to Box 2-2 for definitions of the key terms in **bold** below.

Genetics

Technological advances have led to cost-effective methods for capturing whole **genome sequences**. This has been completed for tens of thousands of human genomes, highlighting remarkable genetic variation among all individuals. **Genome sequencing** also has revealed a small but consistent set of genes, including *HSP90* and *FKPB51*, both of which are involved in regulating responses to stress activation through the HPA axis. Genome sequencing led to the discovery of specific sequence variations, some of which alter gene expression and are correlated with specific psychiatric disorders and stress lability. The variations in these genes contribute to individual differences in vulnerability and heterogeneity in response to early adverse experiences (Criado-Marrero et al., 2018). For complex outcomes, such as psychopathology, researchers are increasingly exploring polygenic risk (i.e., the combined contribution of many genes that influence brain development and function), rather than single gene variants that have very small contributions to risk. One idea being tested is that polygenic risk "scores" may better predict an individual's likelihood of a given outcome (Brikell et al., 2018). **Genomics** has developed in tandem with experimental animal models and clinical discoveries in humans to provide opportunities to study i) genes that control the formation and physiological maturation of neural connections, ii) the molecules responsible for the onset and termination of **sensitive periods**, iii) the molecular basis for experience-driven **synapse** formation, elimination (pruning), and stabilization, iv) regulation of inflammatory responses, and v) regulation of hormonal activation

[2] Accessed November 12, 2018.

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and feedback of the stress response (Bae et al., 2015; Geschwind and Rakic, 2013; Hensch, 2016b; Sudhof, 2018; Sudhof and Malenka, 2008).

Epigenetics[3]

Genomics discoveries were followed rapidly by advanced methods to identify the consequences of differences in how, where, and when specific genes are expressed—that is, the epigenetic ("above the genome") profile of specific cells (Waddington, 1942, 1957). Initial discoveries recognized that epigenetic changes are **chemical signatures** (National Scientific Council on the Developing Child, 2010) placed on an individual's inherited DNA due to environmental experiences and related contributing factors (e.g., nutrition, mental and physical stressors, immune activation, environmental toxicants, enriched environments) (Goldberg et al., 2007). Once thought to be limited in scope and immutable, there are millions of sites on DNA that can be chemically modified epigenetically, and the modifications at these sites can be gained and lost rapidly, or, in many instances, remain stable over the lifetime (Goll and Bestor, 2005). Recent studies have shown that environmental experience influences the genome with specific molecular changes, such as DNA methylation and modifications to chromatin (Goldberg et al., 2007; Heim and Binder, 2012; Kalish et al., 2014; Kumsta, 2019; Miguel et al., 2019). While evidence of these molecular changes has been observed in many studies, the process by which these changes are controlled is complex and not completely understood. Exposures and experiences can result in profound epigenetic modifications that alter the timing and location of expressed or suppressed genes (McEwen, 2017; Miguel et al., 2019). For instance, epigenetics is one mechanism that results in identical twins exhibiting distinct physical and mental traits. Preliminary data suggest that epigenetic changes may explain how early experiences get "under the skin" of individuals for whom physical and brain-related disruptions and disease risk can last a lifetime (McEwen, 2017; Miguel et al., 2019). Research shows that epigenetic changes that alter the DNA chemistry in **germ cells** of adult humans may be one way that experiences are passed on to the next generation through their children (Donkin et al., 2016). However, the best evidence for such intergenerational transmission comes from animal studies. Epigenetic changes occurring during preconception or during pregnancy can impact the germ cells of the offspring and result in intergenerational transmission of the parental experiences (Weber-Stadlbauer, 2017). Moreover, it has been hypothesized that germ cells of the fetus may be altered during pregnancy, meaning that the mother, and even the child when that child is grown and has children of their own, can pass on influences that occurred several generations previously (Bale, 2015; Rodgers et al., 2015; Rowold et al., 2017).

Transcriptomics

Keeping in mind that one's entire genome is in every cell of the body, the study of **transcriptomics** uses methods to identify how particular genes are expressed over time, in specific types of cells, and under specific circumstances (Li et al., 2018). New methods facilitated the organization of a research consortium that led to a remarkably detailed, open-access atlas[4] of gene expression in 16 different human brain areas during pre- and postnatal

[3] For a more in-depth discussion on epigenetics and how the environment gets under the skin, see Chapter 3 of the recent report, *The Promise of Adolescence: Realizing Opportunity for All Youth*, by the Committee on the Neurobiological and Socio-behavioral Science of Adolescent Development and Its Applications (NASEM, 2019).
[4] For more information, see http://www.brainspan.org (accessed June 16, 2019).

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development and into adulthood, building a unique bridge between human and model system brain research (Carlyle et al., 2017; Enoch et al., 2014; Li et al., 2018; Vied et al., 2014). For example, the expression of genes responsible for producing the thousands of neuron types in the brain is enriched during the first trimester of pregnancy. As development continues, many of those genes are turned off because neuron production ceases, and other genes are newly expressed that control the initial wiring and building of the unique architecture of the brain. After birth, genes are turned on to cause the rapid formation of synapses, pruning, and the signaling between neurons to process more and more complex information. The ability to relate specific events in development to the changing transcriptome has been accomplished by creating enormous databases of gene expression in the pre- and postnatal human brain and in experimental animal models. These new data offer a detailed directory of **gene expression** in space and time in the complex and rapidly changing developing brain, thereby providing the framework for ongoing research to determine how the interplay of genetic and environmental factors and experience alters the gene expression that contributes to typical or atypical developmental maturation.

Neuroimaging

Human and animal **neuroimaging** have provided the basis for a new understanding of the brain **connectome**, which defines the breadth and complexity of circuits that relay information between ensembles of neurons. New technologies provide unprecedented resolution of the organization of the connectome in humans, painting a far more complex picture that begins prenatally as early as the third trimester of pregnancy (Kostović et al., 2014; Krsnik et al., 2017) and continues in infancy and toddlerhood as remarkable periods of growth for the brain. Functional maturation of circuits continues throughout childhood, but connectome research, combined with genomics, has identified adolescence as a second period of dynamic changes in brain wiring.[5] In humans, it is now understood that while classic diagrams of brain development present a well-delineated blueprint, there is far more variability in infants and toddlers than expected in the timing and extent of development of connectivity within the standard connectome blueprint. This basic neuroimaging research in humans has provided a foundation for determining the impact of disparities resulting from poverty and other social factors on the trajectory of brain growth and development of circuitry (Gabrieli and Bunge, 2017; Hanson et al., 2013; Johnson et al., 2016; Luby et al., 2012). Given the vast heterogeneity between individuals in terms of brain development, drawing generalizations about the overall connectome and its influence on social, emotional, and cognitive maturation remains a key research challenge.

Maternal/Fetal and Infant Interactions

Prenatal development is marked by a unique relationship between the mother, the fetus, and a transient organ, the placenta. Once thought of as a filtration system of cells contributed mostly by the fetus and blood vessels from both mother and fetus to control the passage of nutrients and other substances, the placenta has been shown by current genomic, cell biological, and physiologic evidence to be an active partner in fetal development and influenced by maternal health. For example, maternal immune activation or persistent stress during pregnancy can alter

[5] For a comprehensive discussion of the developing adolescent brain, see NASEM, 2019—*forthcoming, May 2019*

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placental cell gene expression and function (Hoffman, 2016). Depending upon stage, the placenta expresses 40-60 percent of the human genome and produces cytokines, hormones, **neurotransmitters**, and growth factors that are necessary for healthy fetal development (Gonzalez et al., 2018; Jones, 2019; Paquette et al., 2018; Zhao et al., 2019). These genes are subject to environmentally regulated epigenetic changes; exposure to toxicants, nutrient status, high stress, or infection can result in altered gene expression through epigenetic changes (Bale, 2015). Thus, while the organ clearly is involved in regulating transport of maternally produced substances, the placenta also serves a critical resource for organ development and maturation of functions to ready the fetus for birth (Bonnin et al., 2011; Bonnin and Levitt, 2011; Goeden et al., 2016; Nugent and Bale, 2015). Developmentally, the maternal-placental-fetal relationship begins between the third and fourth week postconception, through a complex process of extraembryonic cell differentiation and vascularization that results in the implantation in the maternal uterus. Given that the main functional cell of the placenta, the trophoblast, is embryonic in origin and that there are thousands of genes expressed in the placenta, any gene mutation that could functionally disrupt the embryo also can impact placental function (McKay, 2011).

Intergenerational Transmission of Risk

Evidence from both animals and humans indicates that adverse social/psychological and physical exposures in one generation can alter risks for psychopathology, maladaptive behavior, and chronic disease in the next generation (i.e., the children of individuals who endured severe threat have elevated rates for psychiatric disorders and dysregulated physiology, despite an absence of direct exposure) (Boyce and Kobor, 2015; Burton and Metcalfe, 2014; Franklin et al., 2010; Matthews and Phillips, 2012; Santavirta et al., 2018; Yehuda and Bierer, 2007; Yehuda et al., 2014). For example, research has documented that 1) adolescents born to women who experienced severe child abuse are at higher risk of smoking and overweight/obesity(Roberts et al., 2014) and 2) there is altered **cortisol** expression among children who were in utero at the time of the 2001 World Trade Center attack in New York City (Yehuda et al., 2005), and 3) there are differences in stress reactivity in children of women affected by the Dutch Hunger Winter (Painter et al., 2006). These excess risks may emerge through a variety of mechanisms that are not mutually exclusive, such as altered parenting behaviors, lower socioeconomic position, epigenetic pathways (described above), or an altered fetal environment.

Adverse experiences before and during pregnancy can influence maternal stress-related physiology, including immune activation and endocrine disruptions (Entringer, 2013; Entringer et al., 2010, 2015; Hantsoo et al., 2019; Howerton and Bale, 2012). Increasing evidence suggests that maternal immune and endocrine activity during pregnancy (including the under- or overproduction of cortisol and levels of pro- or anti-inflammatory cytokines) are associated with adverse birth outcomes (Bastek et al., 2011; Wadhwa et al., 2011), the child's brain (Buss et al., 2017) and cognitive development (Gilman et al., 2017; Rudolph et al., 2018), and future risk for chronic disease (Entringer et al., 2013, 2015), providing support for embryonic and fetal life as sensitive periods for the intergenerational transmission of adverse maternal experiences (Buss et al., 2017; Entringer et al., 2010; Meaney, 2001). Much of this work is preliminary, and further research needs to be conducted before drawing concrete conclusions about **intergenerational transmission of risk**. At present, there is a limited understanding of how plastic or modifiable these intergenerational risk factors are and why some individuals are resilient to intergenerational risks whereas others are not. This is an area of active research that has elevated the importance of applying the life course approach to health and development, and by further understanding the

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pathways of both risk and resilience, scientists may uncover social, behavioral, or pharmacologic interventions that can interrupt the intergenerational transmission of the effects of stress on the developing embryo and fetus. Later in the chapter, the section on Biological Mechanisms of Healthy Development presents a more extensive discussion on the role of the placenta and how it can be influenced by social experiences.

Computational Methods

There has been a rapid increase in the capacity to collect and analyze enormous datasets with social and biological information. These datasets may be developed through research consortia that can involve dozens of investigators from around the world or from the linkage of administrative files, de-identified electronic health records (McGregor et al., 2013; Roden et al., 2008), and other sources (McCarty et al., 2011). In parallel, engineers and computer scientists have advanced sophisticated computational methods, including signal processing and machine learning, to extract predictable patterns that relate to outcomes of interest (Krishnan et al., 2017; Tseng et al., 2013; Van Essen et al., 2013). For example, connectomics and genomics have depended heavily on these methods, and the rapid advances in collecting physiological data from children (EEG, eye tracking, wearable devices to monitor physical activity, sleep habits, stress, and metabolic activity) also have benefited from the advanced analytical methods that help identify patterns of maturation of brain architecture and function (Bagot et al., 2018; Frazier et al., 2018; Hosseini et al., 2016; Medland et al., 2014; O'Driscoll et al., 2013; Wee et al., 2017). These measures have been used to predict risk for behavioral and cognitive disturbances later in development. Relatedly, there is growing interest in using machine learning and predictive risk modeling methods to identify children in adverse environments and predict risk or resilience (Amrit et al., 2017; Gillingham, 2016; Schwartz et al., 2017). For example, advanced analytical methods have been applied to produce highly sensitive and specific metrics for the prospective diagnosis of fetal alcohol spectrum disorder (Zhang et al., 2019). However, there is still much more research needed to determine how well these methods may be applied to determine underlying causal mechanisms. Finally, meta-analytic approaches are using new analytical methods to synthesize the results of multiple independent studies and are now widely employed to inform evidence-based recommendations, policies, and programs for children. For example, meta-analyses have been used to summarize the effects of center-based early education (Grindal et al., 2016; Schindler et al., 2015) and parenting programs (Casillas et al., 2016; Chen and Chan, 2016).

BOX 2-3
To What End—Examples of Application of Neurobiological Research in Practice

1. The discovery of molecules responsible for putting the “brakes” on critical period timing have led to clinical studies that manipulate the function of certain brakes in order to demonstrate that critical periods can be reopened far later in life than previously believed.
2. Eye tracking, coupled with machine learning and neural network algorithms, has been used to discover patterns of scene viewing that serve as a sensitive and specific signature to identify children who have normal facial morphology but were exposed to alcohol during fetal development and exhibit cognitive difficulties.
3. Neuroimaging and behavioral data were combined to demonstrate continued maturation of executive functions. These studies clarified the component skills and have led to a flurry of

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intervention strategies to train working memory, mental flexibility, and inhibitory control in toddlers, children, adolescents, and young adults.

4. The neurobiology of early adversity led to discoveries of genetic variants that increase vulnerability in the response to ACEs. Hsp-90 and FKBP5 were discovered as partner regulators of the hormonal response to stress by controlling glucocorticoid receptor signaling, and sequence single nucleotide polymorphisms (SNPs) predict response to early environmental stressors, which can increase the risk for adult depression and anxiety disorders. To support preventative efforts, current research studies in clinical settings are underway to determine the effectiveness of genetic screening, combined with history of ACEs, to identify those individuals at greatest risk for recurring and treatment-resistant depression.
5. The advances in the neurobiology of sociability and the burgeoning research area of infant–mother and infant–father synchrony have led to foundational discoveries about the importance of a child's environment of relationships. There are many more studies and direct community applications of targeting both child and caregiver in interventions—for example, multidimensional therapeutic foster care or coaching to promote healthy caregiver–infant bidirectional, responsive interactions.

Advances in Longitudinal Research

Investments by National Institutes of Health, the Department of Education, numerous private foundations, and other agencies have resulted in a substantial growth in child development research using longitudinal data from prospective cohort studies with large subgroups of racial/ethnic minority children and stratified-probability samples. These studies have documented more extensively the pervasiveness of health disparities in these populations and the mechanisms by which these disparities arise. For example, the Eunice Kennedy Shriver National Institute of Child Health (NICHD) supports the Fragile Families and Child Well-being Study, a birth cohort of nearly 5,000 children in 20 large U.S. cities that used a stratified random sample, with an oversample of nonmarital births (McLanahan et al., 2003). Baseline interviews for this study occurred between 1998–2000, and follow-up is still ongoing. The Institute of Education Sciences, an independent evaluation arm of the Department of Education, conducts the Early Childhood Longitudinal Study (ECLS) program, which has generated large representative cohorts for studies of child development for the United States, from birth onward (Tourangeau et al., 2009).

Longitudinal studies are critical to understand both normative developmental processes and those involving deviations from the norm. These studies facilitate examination of changes across development within individuals and subgroup differences in within-person changes over time (Lerner et al., 2009). Prospective studies are critical for documenting what changes—in brain, behavior, or health—are due to normal developmental progression and what changes are caused by chronic family-level stressors, structural inequalities, and/or positive environmental stimuli. Longitudinal data, combined with advances in statistical methods for identifying underlying pathways, have made it possible to show the lasting impact of early adverse experiences and to ask questions about mediating processes that underlie the association between early childhood adversity and poor health or educational outcomes (Danese et al., 2009; Iacono et al., 2008; Poulton et al., 2015). In parallel, longitudinal data have shown that for children who have been exposed to severe adversity, such as maltreatment, providing appropriate care can reverse early damage, even to biological systems, such as brain functioning (Cicchetti and Curtis, 2006; Cicchetti and Handley, 2017).

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In spite of these advances, it is important to note that U.S. investment in large birth cohort studies lags severely behind European countries. One new study, launched in 2016, is the Environmental influences on Child Health Outcomes (ECHO) Program. It is a 7-year research initiative focused on understanding the effects of environmental exposures on child health and development, and it draws on existing U.S. cohorts. The cohort studies that compose ECHO collectively include approximately 50,000 children and will address environmental exposures in relation to four pediatric outcomes with high relevance to public health: pre-, peri- and postnatal outcomes, neurodevelopment (i.e., cognition, emotion, and behavior), upper- and lower-airway function, and obesity.

Of note, many longitudinal studies of child development have been the product of interdisciplinary collaborations, which have become increasingly common. An example is the long-term follow-up of the Perry Preschool[6] project, which started in the 1960s (Gramlich, 1986; Schweinhart et al., 1985). This longitudinal study of the effects of a randomized control trial (RCT) is the result of a long-term collaboration between educational researchers and practitioners, developmental psychologists, and economists. The latest follow-up of the intervention administered at 3 and 4 years of age still yields positive results on the life trajectory of its participants at age 40, including more positive health outcomes, improved employment trajectory, and less crime involvement (Belfield et al., 2006; Social Programs That Work, 2017). Consistent with these findings, the Carolina Abecedarian Project—a randomized study of an early childhood intervention for economically disadvantaged children with long-term follow-up—found that children randomly assigned to the early intervention displayed lower cardiovascular and metabolic risk factors in their mid-30s, with effects most pronounced for men (Campbell et al., 2014). Baby's First Years,[7] launched in 2018 and funded by both private and public grants, is a new longitudinal study that seeks to establish causal links between parental income level and brain development in very young children.

Advances in the Study of Social Interactions

Since *From Neurons to Neighborhoods* was published, researchers have introduced a number of novel topics and measurement approaches that have advanced the understanding of early development and consequences for later physical and mental health outcomes. In particular, there has been a shift of focus to the well-being of salient caregivers as "a dependent variable" in research and associated interventions. In other words, rather than continuing to report (as psychologists had over decades) that a good relationship with the primary caregiver is the single most important protective process in promoting resilience among children whose life circumstances render them at risk, there is now explicit focus on the question of what it is that helps parents maintain good parenting when they are struggling with high ongoing stress. How can we promote resilience among the adults primarily responsible for raising children in stressful life circumstances (for reviews, see Luthar et al., 2015 and Luthar and Eisenberg, 2017)? Relatedly, intervention approaches are going beyond simply teaching parents what they should and should not do in didactic parenting classes or even imparting approaches to regulate their own affect (via mindfulness or trying to cope effectively with their own ongoing life stressors). Increasingly, programs are also seeking to provide the support that parents need to successfully negotiate the many challenges in sustaining good parenting behaviors with multiple children and

[6] For more information, see https://highscope.org/perry-preschool-project (accessed March 29, 2019).
[7] For more information, see https://www.babysfirstyears.com (accessed April 18, 2019).

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across multiple decades (for examples of such programs, see Corso et al., 2015; Kaminski et al., 2013; Luthar et al., 2007, 2017).

An emerging field of research suggests that there is a biological basis to healthy caregiver–child relationships and that caregiver–child synchrony in biological processes such as EEG or heart rate variability can help children to develop control of their behavior and emotions. The science of caregiver–child synchrony advances previous views that a child's central nervous system is primarily responsible for developing self-regulation (Welch, 2016). This research suggests that children display dysregulated behavior due to deficient coregulation from nonoptimal caregiver relationships and that coregulation can be improved via interventions, such as the Family Nurture Intervention, developed by Martha Welch and colleagues (2014, 2015). Although there are a number of neuroendocrine changes that occur during the process of parenting, a substantial amount of this research has focused on oxytocin measures in the infant and caregiver. In addition to oxytocin's roles in the birthing process, lactation, and maternal care in mammals, studies have demonstrated that oxytocin (and its close neuropeptide relative vasopressin) facilitates affiliative social behavior through modulation of the activity of specific forebrain circuits (Feldman and Bakermans-Kranenburg, 2017; Hammock et al., 2005; Insel, 2010b; King et al., 2016; Numan and Young, 2016). Both observational and experimental research studies suggest that oxytocin influences the synchrony between caregivers and children, and it is also associated with sensitivity of caregiving and amount of contact (Feldman and Bakermans-Kranenburg, 2017). Researchers are exploring how specific caregiver health conditions (e.g., postpartum depression) or social experiences (e.g., traumatic experiences in early childhood) influence oxytocin, other neuroendocrine hormones related to nurturing behaviors, and caregiver–infant synchrony and how interventions may target these processes to improve caregiver–child relationships.

Another major advancement is that there has been a growing appreciation for the role of culture for the development of children (Coll et al., 1996). In the study of children and families in the United States outside of the white middle class (i.e., minority families, based on race/ethnicity, immigrant status, sexual orientation, and others at both socioeconomic [SES] extremes) there has been a growing emphasis on potent culturally specific risk and protective factors and processes (see Velez-Agosto et al., 2017), especially those that can be harnessed in beneficial interventions. As an example, the *American Psychologist* devoted a special issue to marginalization, defined "as a multidimensional, dynamic, context-dependent, and diverse web of processes, rooted in power imbalance and systematically directed toward specific groups and individuals, with probabilistic implications for development" (Causadias and Umaña-Taylor, 2018). The special section contains a rich collection of articles addressing issues related to various marginalized groups, including recent immigrants and youth from different ethnic and racial minority groups. In another recent special section in *Research in Human Development,* Cunningham (2019) compiled papers addressing "myths and realities associated with human development research and theorizing" encompassing "diverse perspectives on sexual minority youth, resilience and risk for youth in high-achieving schools (HASs), a reconceptualization of hostility in African American parenting styles, a critical examination of diversity and contact for students attending racial/ethnically diverse schools, and a thoughtful consideration of contextual factors associated with aggressive attitudes and prosocial behaviors in African American males" (p. 1). Many of these studies move the conceptualization of "children or family at risk" to "children and family living in at-risk conditions." The "problem" to be "fixed" is not the child and/or family but the at-risk conditions within which they live, an important paradigm shift for

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prevention, early identification, and prevention. (On a related topic, there have also been advances in connecting racism to child and adult health outcomes; see Box 2-4.)

BOX 2-4
Racism and Child Health

Another major scientific advancement in the science of health disparities since *From Neurons to Neighborhoods* is the research linking racism to child and adult health outcomes (Pachter and Coll, 2009; Paradies et al., 2015; Priest et al., 2013; Williams and Mohammed, 2013). Much of the impetus for the work on the effects of racism on children's development stemmed from the critical analyses of the prevalent use of the deficits model to understand health disparities in minority children (García Coll, 1990; McLoyd, 1990; Ogbu, 1981; Spencer and Markstrom-Adams, 1990).

Beginning in the 1990s, several theoretical models and empirical work started to place the burden on the social and environmental conditions where the majority of the minority populations in the United States lived (Coll and Szalacha, 2004; García Coll et al., 1996; Stein et al., 2016). McLoyd's work showed that the stress associated with poverty explained harsh parenting among African American families. These segregated communities had a majority of families living below the poverty line, with schools, health care, recreation, housing, and transportation facilities below standards or with fewer resources than where predominantly white, middle-class populations lived. Health disparities were now considered a function of these environmental conditions and not the parents and children themselves. Most of these social and environmental conditions are now seen as a function of pervasive institutional and interpersonal racism (Pager and Shepherd, 2008). Similarly, scientists have denounced the use of race and ethnicity as variables in biological research (Collins, 2004; Foster and Sharp, 2004), noting that "historical racial categories that are treated as natural and infused with notions of superiority and inferiority have no place in biology" (Yudell et al., 2016, p. 565).

A large body of scientific evidence has documented that both institutional racism and interpersonal experiences of discrimination can influence the health and well-being of both children and adults in multiple ways, including reducing access to material resources and services that promote long-term health and development and acting as a psychosocial stressor that can lead to worse outcomes over time (Priest et al., 2013; Williams and Mohammed, 2009). Studies of children and adolescents indicate that experiences of discrimination are positively associated with depression and anxiety and negatively associated with positive mental health outcomes, such as self-esteem and resilience (Priest et al., 2013). Research also suggests that parental experiences of discrimination are associated with child and adolescent mental health and socio-emotional outcomes (Heard-Garris et al., 2018; Tran, 2014), as well as some indicators of physical health (Priest et al., 2012; Slopen et al., 2019). There are also a diverse set of health outcomes in adults resulting from racism (Williams et al., 2019).

Measures for experiences of racism and discrimination in children have relied mostly on parent- or self-reported answers to survey questions (Fisher et al., 2000; Pachter et al., 2010; Williams et al., 1997). These instruments are subjected to social desirability, recall biases, and other potential issues associated with self-reports. Some research in adults has begun to combine traditional measures of racial discrimination with implicit measures, in order to advance understanding of how conscious and unconscious experiences of racism influence outcomes (Krieger, 2012; Krieger et al., 2010). Future research is needed to examine whether similar approaches are useful for children and adolescents.

Advances in the Study of Physical Environments

Although the family remains a major determinant of the child's outcomes, studies following more comprehensive theoretical models (e.g., Bronfenbrenner, 1979; Bronfenbrenner and Evans, 2000; Ferguson et al., 2013) have documented the importance of neighborhood contexts on family interactions and child development. Neighborhood context can be characterized as positive and negative, and it can reflect the physical or built environment and psychosocial characteristics. It has become increasingly common for researchers to link

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measures of neighborhood social and structural environments to individual-level data on children. As long as families are willing to share an address, researchers are able to draw on information from the Census and other free and proprietary data sources in order to characterize a child's social and physical environment. When this field of research began, most studies focused on neighborhood SES (e.g., the proportion of the population in the Census tract that is below the federal poverty line), as this measure is associated with the physical infrastructure of the community, noise and pollution, amenities such as parks and groceries, and availability of health centers and early childhood education (Evans, 2004; Evans et al., 2010). However, use of neighborhood SES as a proxy for neighborhood quality and conditions does not provide sufficient information about the mechanisms of action—that is, the specific pathway through which neighborhoods influence child health. The majority of research on neighborhood SES and health is observational, but there are a few exceptions, including the Moving to Opportunity study (Chetty et al., 2016; Ludwig et al., 2011).

The Project on Human Development in Chicago Neighborhoods (PHDCN)—an interdisciplinary study that began in 1995 and aimed to explore how families, schools, and neighborhoods influence child and adolescent development—was among the first to provide an intensive investigation of neighborhood conditions, related to economic, social, organizational, political, and cultural structures and how communities change over time. The investigators conducted systematic social observations by videotaping one side of each block within the selected 80 Chicago neighborhood clusters, and they maintained observer logs about the activities of residents and the presence of detracting elements (e.g., garbage, abandoned cars). This study identified neighborhood collective efficacy and social capital as important aspects of neighborhood social organization for child and adolescent outcomes (Sampson et al., 1999; Sampson and Raudenbush, 1997, 1999).

Since PHDCN, researchers have begun to characterize neighborhood social and physical information for health research using Google Street View (Rundle et al., 2011; Rzotkiewicz et al., 2018), which has the benefits of being lower cost and more efficient, although it can be limited by resolution and availability for a given temporal period. There has also been a substantial amount of research to examine child health in relation to neighborhood violence or crime, which can be derived from police crime reports (Beck et al., 2016; Goldman-Mellor et al., 2016; McCoy et al., 2015). Some studies use surveys to assess parental perceptions of neighborhood safety (Christian et al., 2015). Other research has focused on the physical neighborhood environment, such as opportunities for physical activity, access to healthy food, density of fast food, air quality, and presence of natural environments (e.g., green space) (Gascon et al., 2016; Gorski Findling et al., 2018; McCracken et al., 2016; Sallis et al., 2018). While some studies use single measures of neighborhood context, other studies rely on more complex indexes, such as the Childhood Opportunity Index (Acevedo-Garcia et al., 2014, 2016). This measure—developed by diversitydatakids.org and the Kirwan Institute for the Study of Race and Ethnicity—combines 19 component indicators related to education, health and environment, and social and economic context to rank neighborhoods (defined by Census tract) to other neighborhoods within the metropolitan area (Acevedo-Garcia et al., 2014; Beck et al., 2017a,b).

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BIOLOGICAL MECHANISMS OF HEALTHY DEVELOPMENT

Introduction

Development is an interplay between biology and the profound impact that experiences have on biological systems. There are basic biologically driven mechanics that involve the production and specialization of cells that serve as the ingredients to build different organ systems, such as the brain, heart, digestive system, and kidneys. These systems are ultimately responsible for performing specific functions and become specialized over different periods of developmental time. This developmental process begins prenatally, and for the most complex organ—the brain—it extends into young adulthood. The developmental process is designed to promote maximal functioning and survival of the individual. The challenges an individual faces from infancy to adulthood are numerous and varied, and through the powerful biology-experience relations, development establishes both functional and adaptive capacities in order to adjust to the environment and to future experiences. Optimal functioning and robust adaptability to the environment are signatures of healthy development. No matter which organ system or cell type are considered, a core principle of development involves the regulation of the expression of the 23,000 genes in each person's genome at specific times, and in specific combinations, in order to achieve optimal functioning of each organ system.

It is important to re-emphasize here that the genome is composed mostly of regions of DNA that do not code for proteins but rather are responsible for using information from experiences and the environment to regulate gene expression. Gene expression is the first step toward producing proteins, which are the primary functional units of specialized cells that characterize each organ system. Thus, while all cells contain the identical genetic material (DNA code), gene expression and production of proteins varies greatly between types of cells, which specialize during development to perform specific functions. Because the brain is massively complex, with billions of **neurons** and trillions of connections, at least 85 percent of the genome is expressed at various times and locations during development (Negi and Guda, 2017), whereas, other organ systems may express far fewer genes in order to perform their functions. This orchestrated temporal and spatial expression of genes depends upon maintaining healthy **cellular homeostasis** and accessing nutrients that serve as the building blocks for proteins and eventually organ systems. Research has shown that regulation of fundamental metabolic processes, including mitochondrial function in the brain and periphery, is essential for cellular adaptations to developmental psychosocial stress that can impact mental and physical health (Eisner et al., 2018; Picard and McEwen, 2018). The most productive means to facilitate early healthy development include providing access to healthy nutrition, avoiding factors that can be detrimental to nutritional status for mother and fetus over the course of pregnancy, limiting chronic stress, and offering access to quality health care. Proper nutrition is needed postnatally through childhood as well. Moreover, disrupting the fetal-placental-maternal biological relationship through infection, toxicants, alcohol, nicotine, or poor diet increases risk for altering a typical, healthy developmental trajectory. Thus, improving physical environments to limit exposure risk and providing access to high-quality maternal health care during pregnancy is an additional essential ingredient for healthy development. These well-researched factors make up a strategy for viewing early development through a "prevention lens" that ultimately leads to reduced risk for later physical, mental, and cognitive disorders.

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Recent studies, mostly in animal models, indicate that the developmental process starts even prior to conception (Chan et al., 2018). Certain stressors, including psychosocial factors, toxicants, or drug exposure, can impact reproductive cells through an epigenetic process that ultimately may alter the expression of the genes that are inherited by the embryo. This suggests that parental behavior may impact how a child's genes are ultimately expressed and therefore what the child's biology and behavior will be. This transgenerational impact on development requires studies in humans before firm conclusions are made, but the research indicates that maternal and paternal experiences prior to conception may be an important factor to consider for programs that promote family health and welfare.

The following sections outline how several biological systems, those most relevant to this committee's report, develop between gestation and puberty and serve as a baseline for understanding how early environmental stressors and adversity affect child development and the emergence of health disparities: maternal physiologic adaptations during pregnancy and the nervous, immune, endocrine, and reproductive systems. Note that in addition to psychosocial stressors in the environment of a developing child, adversity may include limited or no physical activity, exposure to toxicants, including alcohol and nicotine, and chronic disruption of bodily physiological functions, such as sleep (Feldman et al., 2014; Jones et al., 2019; McEwen and Getz, 2013; Thompson et al., 2009). Each of the following sections describe typical processes from the prenatal period through early childhood and how the system may be impacted by experiences that disrupt typical development.

Maternal Physiologic Adaptations During Pregnancy

Remarkable changes take place in women during pregnancy to accommodate the needs of the developing fetus. A woman's heart pumps out 30 to 50 percent more blood every minute, while her lungs take more air with each deep breath, in order to deliver oxygen and nutrients to the baby (Cheung, 2013; Costantine, 2014). Her kidney functions also improve by 50 percent to filter out more waste products and toxins from the blood (Cheung and Lafayette, 2013). Her immune system is dampened to avoid rejection of the fetus, while she becomes progressively insulin resistant, which allows an increasing amount of blood glucose to stay in the bloodstream to nourish the growing fetus.

A comprehensive review of maternal adaptations is beyond the scope of this report, but it is important to note that how well a woman's body can adapt to these physiologic changes during pregnancy is closely related to her overall health going into the pregnancy. For example, a healthy heart can readily adjust to the increased workload, but a diseased heart will have a much more difficult time; heart disease is a leading cause of maternal death in the United States. Pregnancy predisposes blood to clotting, but the risk of severe complications from thromboembolism and pulmonary embolus is substantially increased in women with pre-existing conditions, such as hypertension, diabetes, or obesity. Pregnancy causes a physiologic anemia, which can be exacerbated by ongoing iron deficiency, thereby increasing the risk of having a low birth weight (LBW) baby. The insulin resistance in pregnancy is made worse if a woman enters pregnancy overweight, obese, or with preexisting diabetes mellitus; fetal exposure to excess blood glucose has been associated with adverse birth outcomes and obesity and diabetes later in life (Garcia-Vargas et al., 2012). As we will discuss in Chapter 5, advancing health equity in birth and child health outcomes first requires improving preconception health for all women.

One of the most important adaptations during pregnancy is the formation of the placenta, an organ that is responsible for controlling access of maternal-derived factors to the fetus and

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also serves as a source of growth factors and other molecules that support healthy fetal development. The placenta begins to form very early in pregnancy, and disruption of this process has been associated with pregnancy complications, such as intrauterine growth restriction (IUGR) and preeclampsia (Sharma and Sharma, 2016). At term, approximately 600–700 mL of blood flows through the placenta, delivering oxygen and nutrients to the baby (Wang and Zhao, 2010). The placenta also acts as a critical regulator of maternal–fetal interactions and resource allocation. For example, if a pregnant women experiences acute stress, the placenta shields the fetus from overexposure to maternal cortisol by turning up the activity and expression of a placental enzyme called 11β-hydroxysteroid dehydrogenase type-2 (HSD11B2), which inactivates cortisol as it passes through the placenta. However, chronic stress has been found to be associated with reduced activity and expression of placental HSD11B2, suggesting that the placenta's built-in ability to limit fetal exposure to maternal cortisol may be diminished in the face of chronic stress (Cuffe et al., 2012; O'Donnell et al., 2012; Welberg et al., 2005).

Nervous System Development

The neurobiological processes that build the brain share some common elements with other organ systems, such as producing specialized cells (e.g., neurons and **glia**) with functions specific to that organ system. However, understanding of neurodevelopment lags behind the understanding of other organ systems, in large part because the brain is extraordinarily complex. Experimental studies in model systems have identified basic mechanisms and the genes that are involved in specializing areas of the developing brain into what will make up function regions, the production of a diversity of neuron types, and the initial wiring of circuits that occurs prenatally (Kast and Levitt, 2019; Kolodkin and Tessier-Lavigne, 2011). Identification of mutations in some of these same genes in humans have validated the highly conserved nature of the ingredients that are responsible for the initial brain blueprint (Doan et al., 2018; Geschwind and Rakic, 2013; Jayaraman et al., 2018; Rubenstein and Rakic, 2013). Genes and their protein products involved in later events, including the extended period of synaptogenesis (the formation of synapses) have been identified and studied experimentally (Akins and Biederer, 2006; Favuzzi and Rico, 2018; Sudhof, 2018). For all the advances in the basic understanding of neurodevelopment, there is a knowledge gap in determining the many ways in which the environment and experience are woven into the developmental process. Figure 2-1 is a diagram showing development from the first trimester to puberty, with respect to both neurodevelopment and the development of other biological systems.

Within a few weeks of conception in the first trimester, neurons begin to be produced. By 10–12 weeks gestation, most of the neurons that make up the brain are generated (Bystron at al., 2008; Stiles, 2008). Specific types of neurons in structures like the cerebellum, hippocampus, and olfactory bulb continue to be produced prenatally. Unlike many other types of the cells in the human body, once formed, neurons lose the capacity to renew themselves, even if injured (a small number of neurons in the olfactory bulb and hippocampus represent limited exceptions). Because the neurons are "born" at sites in the developing brain that are different from their final position, all neurons migrate using a combination of mechanical guides, the extended processes of radial glial cells, and molecules that serve as guidance cues. This migration process begins as soon as neurons are produced, and as the brain grows through the second trimester, the later-produced neurons take longer to reach their final position. This developmental process is further complicated because newly formed neurons not only migrate but simultaneously extend a long process—the **axon**—that will connect each neuron to its appropriate target. At the same time, the

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receiving end of the neuron—the dendrite—develops but takes much longer to reach maturity, in sync with the formation of individual neural connections—the synapses. Throughout each of these developmental stages, additional sets of genes are expressed that drive a specific process to completion. Many of the earliest born neurons will regulate peripheral organ function (e.g., heart, lungs, gastrointestinal tract). Connections are formed between these neurons and their targets in the first trimester as the targets continue to develop. By the beginning of the second trimester, long connections from the hypothalamus to the autonomic brain stem are formed, providing the capacity for top-down regulation of what are called the **sympathetic** and **parasympathetic nervous systems**—in essence, the systems that control the "fight or flight" and "rest and digest" responses.

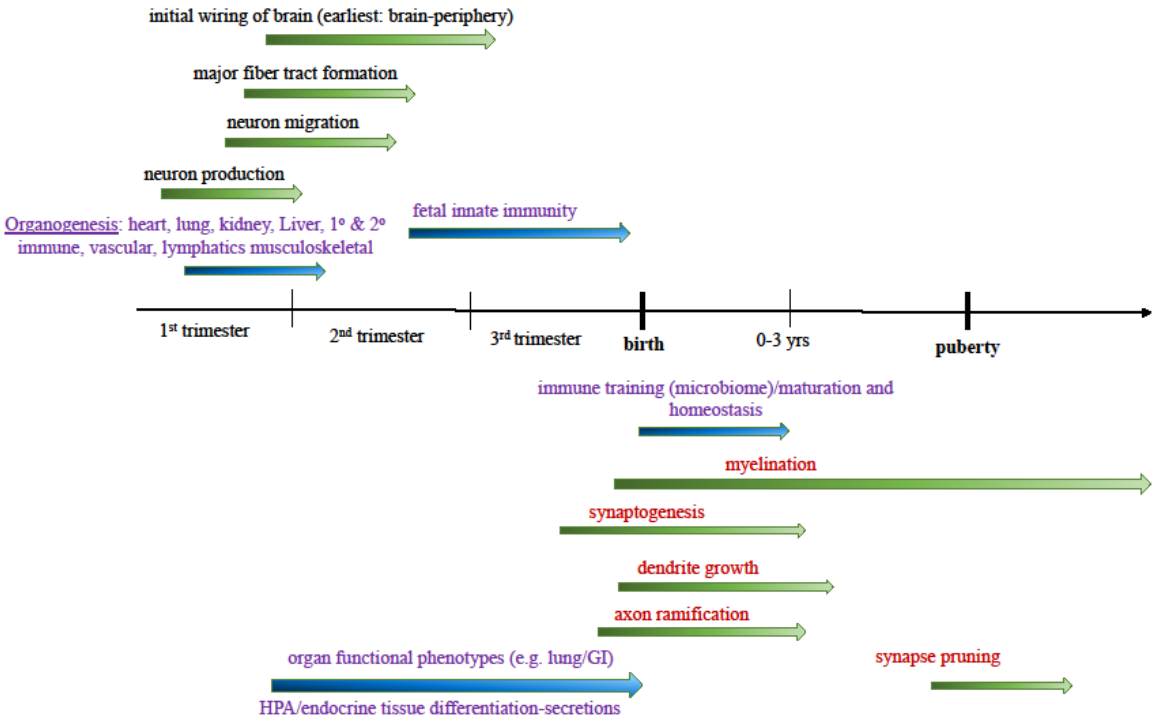

FIGURE 2-1: Human development of the brain and other organ systems from prenatal to pubertal periods
NOTE: Arrows represent time frames of approximate onset (darker shading) and completion (lighter shading) of specific developmental events for neural (green) and nonneural (blue) systems. For the neural components, above the line represents events that occur within the prenatal period, and below the line represents events that extend into postnatal periods.

By the second and third trimesters, sensory systems are connected from the periphery to the brain's cerebral **cortex**—the most complex part of the brain with the greatest neuronal diversity, which processes complex information from all of the senses. Toward the end of fetal development in the third trimester, the basic blueprint of neuronal connections is initiated, during a temporally extended process of synapse formation that continues through 2–3 years postnatally (Levitt and Eagleson, 2018; Silbereis, 2016). The human neocortex contains approximately 20 billion neurons, each with an average of several thousand synapses and connected with thousands of kilometers of dendrites and axons (Tang et al., 2001).

By birth, sensory systems are sufficiently wired to be able to take in information from the infant's environment and begin to process the most salient stimuli (for example, visual, tactile,

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and auditory stimuli of the primary caregiver that are experienced in temporal lockstep with satisfying the hunger, thirst, and warmth needs). The nervous system continues to build by adding more synapses, ultimately reaching its greatest synapse density during toddlerhood. Experimental studies in animals shows that experiences can influence the initial organization of and sheer number of synapses (Berbari et al., 2005; Turner and Greenough, 1985). In early human infant development, differences in growth parameters of gray and white matter have been shown to be influenced by experience in the human cerebral cortex (Brito and Noble, 2014; Hair et al., 2015), which assumes synaptogenesis changes due to experience. In humans, more than 1 million synapses are added per second for this 2–3 year period (Center on the Developing Child at Harvard University, n.d.-a). During early childhood, and as an individual approaches the onset of puberty, the number of synapses remains relatively stable, but this does not imply that no developmental activity is taking place. Instead, experience-driven processes impact maturation by influencing which specific genes are expressed and therefore how synapses process information. Over time, synapses that become activated more frequently and in concert with other synapses processing the same information become more stable and better at processing; those that are used less, on the other hand, are eliminated (Katz and Shatz, 1996). Remarkably, both early spontaneous and patterned activity (from sensory stimuli in the environment) influence synapse formation and stabilization (Leighton and Lohmann, 2016). In all mammalian species examined, including humans, synapse pruning (i.e., the controlled elimination of certain synapses) in the cerebral cortex begins before puberty, with a reduction of approximately 40 percent by the end of adolescence (Bourgeois, 1997). A core concept that has emerged from decades of neurobiological studies of the development and reorganization of connectivity is that environmental factors, which include fetal, postnatal, and child experiences, regulate the expression of genetic predispositions, and these experiences can dramatically change outcomes. This underlies the mechanism through which early exposure to chronic stressors and trauma so powerfully influences developmental processes in the brain and periphery. The research also makes clear that brain development and maturation depends upon experience—what Greenough labeled *"experience-expectant"* development (Greenough et al., 1987).

Myelination is an often-overlooked developmental process. Myelination entails forming the insulation around axons and is essential for increasing the capacity for rapid and efficient transmission of information in neural circuits (Bercury and Macklin, 2015). During the entire postnatal period discussed above (Gogtay et al., 2004), the glial cells responsible for myelination (Miller et al., 2012)—the **oligodendrocytes** in the brain and Schwann cells in the periphery—becomes its most active, forming the sheath around axons that promotes more rapid signaling between neurons. Myelin is composed mostly of protein and lipids (i.e., fats) to form the ensheathing membranes; myelination is highly dependent upon nutritional status, experience, and other environmental factors (McLaughlin et al., 2017). Neurons are among the most metabolically active cells in the body, so they too are highly dependent upon nutritional status. Myelination begins and ends in different regions of the nervous system at different times postnatally (Yeung et al., 2014), with the axon tracts beneath the frontal lobes taking the longest; myelination in this region can extend into the third decade of life (Miller et al., 2012). The timing of myelination completion corresponds to the reduction or ending of sensory critical periods, suggesting that myelin not only facilitates improved information processing but also participates in the maturation of synapse networks. This correlation has been tested experimentally by reversing myelination, resulting in reopening the critical period (McGee et al., 2005). This suggests that for human brain regions that have long periods of circuit myelination (such as

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frontal-lobe-associated circuits, in which executive functions continue to develop into young adulthood), extended periods of plasticity continue (Diamond, 2013).

Plasticity

The circuits of the brain undergo initial construction prenatally and remain relatively immature until postnatal periods, when experience drives remarkable growth that is reflected in formation of extensive connections (synapses) that form at a rate of at least 1 million per second during the first 2 years (Center on the Developing Child at Harvard University, n.d.-b,c). The role of experience in the development of functional circuits was established half a century ago in landmark studies of the visual system (reviewed by Hensch, 2016b; Hensch and Bilimoria, 2012). Binocular vision requires the use of both eyes by the infant to form an accurate representation of the visual world. This occurs during a sensitive or critical period, when there is heightened plasticity for changes in the fine details of circuits that will establish functioning for a lifetime. Since that time, neuroscience research has shown that all sensory and motor circuits, as well as those involved in social and emotional behaviors, cognition, motivation and reward, executive functioning, and even stress responsiveness, develop through experience-dependent mechanisms that involve heightened plasticity. Moreover, circuits remain open to change for different periods of time—sensory functions that are essential for the infant and toddler to perceive and respond to the environment are built first, followed by gross and fine motor and basic cognitive functions that provide the baby with opportunities to respond in more and more complex ways to experiences that provide input to the brain. This concept reflects the basic developmental rule that simple skills beget more complex skills, the paramount exemplar being language: from phoneme discrimination, to motor imitation, to making sounds that become part of a repertoire in which sounds become associated with objects, to objects becoming associated with words as the child begins the process of integrating language with more complex cognitive skills. It is important to emphasize that certain experiences, including physical exercise, are not necessarily limited in impact within certain periods and can be drivers of brain plasticity across the life-span. The evidence in basic and clinical research is compelling, showing effects on both physical well-being and cognitive and social-emotional capacities (Khan and Hillman, 2014; McEwen and Getz, 2013).

For all of these steps, there are heightened sensitive periods when experiences that occur over and over are the most potent. Periods have a beginning, a peak of varying length, and then a stretch of reduced plasticity, when change can occur but in a far more limited fashion and with much greater effort. Think about learning a second language for the first time as an adolescent or adult. For all circuits that will be responsive to either positive or negative experiences, critical period shifts can occur that result in more rapid maturation or longer times of remaining open. Neither outcome of changes in plastic periods is beneficial. Note that genetic factors also can change circuit maturation (Heun-Johnson and Levitt, 2018), which then causes a different response to early experience, as if events were occurring in an older individual. In a model of early adversity (Bath et al., 2016), precocious maturation of molecular and behavioral measures were induced by early adversity. Disrupting neuronal metabolism in certain types of neurons prolongs the visual system's critical period plasticity (Morishita et al., 2015). All of these experimental studies mean that the normal periods of heightened plasticity are out of synch with the timing of when normal experiences are supposed to have optimal impact on circuit development. In humans, early adversity affects circuits that underlie specific functions in different ways (Nelson et al., 2014), with the timing of interventions (e.g., the Bucharest Early

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Intervention Project, which has done extensive research on the timing of placement in quality foster care after experiencing neglect in an orphanage) (Almas et al., 2018; Nelson et al., 2007; Wade et al., 2018; Zeanah et al., 2003) having a lasting effect on the quality of executive functioning, stress responsiveness, and attachment-related behaviors. Here, the impact on critical periods can only be hypothesized, but based on measures years later, it is likely that the timing of optimal sensitivity to experiences is changed in relation to specific functional domains.

The concept that altered neural development, through genetic and experience-dependent mechanisms, establishes different sensitivity to later-life adversity has been suggested for postadolescent onset psychiatric disorders (Keshavan et al., 2014) and more recently for ACEs (Danese et al., 2009; Loria et al., 2014; Morrison et al., 2017; Shalev and Belsky, 2016). The second adversity may be related to normal challenges or other life experiences that all of us endure and to which those without early adverse experiences are able to adapt. The mechanisms through which this vulnerability is read out has recently been addressed in a new set of animal studies (Peña et al., 2017). These demonstrate that there are early molecular changes in specific brain reward circuits due to early life stress that are responsible for the negative impact of later-endured juvenile social defeat (bullying) to produce depressive-like symptoms. For both human and animal studies, it is important to note that while early adverse experiences can have both acute and long-term impacts on mental and physical health, men and women may express these changes in different ways (for example, externalizing or internalizing behaviors, respectively), which need to be accounted for in any intervention programs.

Immune System Development

The newborn immune system is shaped largely by the gut **microbiome**, which is initially primed by the vaginal or skin maternal microbiome during childbirth (depending upon birth process) and early life, such as through breastfeeding (Miller, 2017). The gut microbiome refers to the vast collection of microbes that populate the gastrointestinal system and contribute to whole-body physiology. Molecules produced by the immune system impact various aspects of brain development, with strong evidence for mediating the formation and molecular adaptation of neural connections. The immune system also plays a role in the response of the brain and body to early life stress. In addition to the development of neurons and their connections with the periphery, the fetus' immune system undergoes rapid development in the first years of life and is yet another system that may be impacted by early adversity and stress (Simon et al., 2015). The **innate immune system**—which includes specialized cells, such as resident neutrophils, macrophages, and monocytes—develops in the fetal stage, though it is not robust enough to ward off most external pathogens. Of course, this lack of strong innate immunity may also provide the substrate for healthy microbiota to develop symbiotically with the newborn infant; the bacteria colonizing the gut, skin, and mucosa, for instance, assist in digestion and protection from other pathogens. Children's **adaptive immune systems**—governed by the **lymphatic system**—are relatively undeveloped at the time of birth, rendering them prone to bacterial, viral, and fungal infections soon after birth. At this time, the fetal adaptive immune system is prepared for "training" by the maternal microbiome, which occurs rapidly after birth via vaginal, skin, or mammary exposure (Lynch and Pedersen, 2016). Throughout childhood, children are exposed to countless microbes that are either summarily dismissed by the immune system without notice or produce a robust immune response that primes the immune system to protect the body against similar pathogens in the future.

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The immune system can be trained to cope with certain challenges. For example, the "hygiene hypothesis" suggests that the prevalence of asthma has increased dramatically in certain parts of the world where children are no longer exposed to microbes that can train the immune system (Harding, 2006). In the face of other challenges, the nervous system activates the HPA axis, which can lead to suppressed immune functioning and thus increased susceptibility to infections (Cohen et al., 2007; Thompson, 2014). When children experience sustained exposure to challenges (e.g., chronic stressors, such as poverty or maltreatment), HPA axis activity may become blunted, which can reduce inhibition of inflammation and lead to elevated levels of chronic low-grade inflammation (Koss and Gunnar, 2018; Miller et al., 2011). In the past two decades, researchers have attempted to clarify (1) the relationship between social stressors and inflammation and other immune markers in children (Slopen et al., 2012) and (2) the role of immune system in the development of neurodevelopmental (Entringer et al., 2015; Gilman et al., 2017) and psychiatric disorders (Danese and Baldwin, 2017; O'Connor et al., 2014) and the cardiovascular and atherosclerotic disease process (Hansson and Hermansson, 2011; Pearson et al., 2003). There is now evidence from longitudinal research that inflammation tracks from childhood to adulthood (Juonala et al., 2006) and that chronic inflammation is a risk factor for a wide range of diseases, including cardiovascular diseases and depression (Dantzer et al., 2008; Libby et al., 2002).

Endocrine System Development

The endocrine system is composed of several glands that secrete hormones—signaling molecules—that reach their target organs through the circulatory system to modulate every physiological function, such as growth and development, reproduction, stress responses, and metabolism. A key characteristic of the endocrine system is that it operates on feedback loops to allow the hormones to perform a function (e.g., the "fight or flight" stress response) and then resets to be ready for the next event that may trigger an endocrine response. In all mammals, including humans, the hypothalamus and pituitary are the center of control for the endocrine system. Because they impact many stages of development, the organs of the endocrine system themselves develop early on, in the first trimester of gestation. Moreover, early in the first trimester, the placenta produces the same releasing factor proteins that are synthesized in the hypothalamus of nonpregnant women. These proteins control the production and release of hormones from the pituitary and peripheral tissues in the mother and the fetus.

Neuroendocrine hormone markers are evident by the second trimester. The endocrine systems work in concert with the **autonomic nervous system** to regulate organ function. Autonomic regulation can be measured in the late second trimester and throughout the third trimester and is intact and operational at the time of birth (DiPietro, 2015). The **autonomic neurotransmitters** norepinephrine, epinephrine, and acetylcholine and the endocrine hormones reach their targets via body circulation. Thus, they affect metabolic processes through their binding to membrane protein receptors that are expressed by target cells in the brain and other organs as early as the second trimester. Related to the stress response via cortisol production, activation of the glucocorticoid receptors in the hippocampus and hypothalamus serve as classic feedback loops to reduce hormone production in order to reset the stress response capacity.

Research in humans and animals shows that overexposure to stress hormones and autonomic neurotransmitters during prenatal or early postnatal development can result in a fetal programming response, which includes epigenetic changes in the genes that encode the stress hormone receptors. This process produces long-term changes in receptor expression and less

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robust negative feedback to limit stress hormones' chronic effects (for examples, see Maternal and Fetal/Infant Interactions, above) (Meaney and Ferguson-Smith, 2010). In contrast, high levels of maternal care documented in animal studies shows that the stress response system is capable of better management of later-life stress (Meaney, 2001; Plotsky et al., 2005; Plotsky and Meaney, 1993).

Reproductive System Development

The development of the reproductive system also begins early in gestation, in the first trimester. In the first 2 months of pregnancy, differentiation between male and female gonadal development occurs, which continues into the second and third trimesters. As noted above, animal studies show that intergenerational transmission of the impact of adversity during pregnancy can influence the mother, the fetus, and, if the developing gonads are affected, the offspring of the fetus when he or she matures into a reproductive adult. Sexual differentiation is controlled primarily by levels of testosterone, estrogen, and androgen, a process that can be influenced by **exogenous** chemicals similar in structure to these hormones. Among women, the ovarian follicle pool develops during the prenatal period; therefore, in-utero exposures may impact the size and quality of the follicle pool and influence the timing of ovarian loss and menopause (Bleil et al., 2018). For example, studies show that in-utero cigarette smoking exposure (Strohsnitter et al., 2008), famine (Yarde et al., 2013), and extremes of birth weight (Tom et al., 2010) are associated with earlier menopause. There is also evidence to suggest that maternal metabolic factors during pregnancy, including obesity and pregnancy hyperglycemia, are also associated with the timing of puberty (Kubo et al., 2018).

There has been a trend toward an earlier age of puberty for boys and girls in the United States that is not yet fully understood, and currently 9 years of age is within the normal range for onset of puberty (Herman-Giddens et al., 1997, 2012). Adverse social experiences in early and middle childhood—including socioeconomic adversity (Hiatt et al., 2017; Kelly et al., 2017; Sun et al., 2017) and child maltreatment (Mendle et al., 2016; Noll et al., 2017)—as well as exposure to endocrine-disrupting chemicals (Buttke et al., 2012) are associated with earlier pubertal development (Ellis and Giudice, 2019). In the United States and elsewhere, the age of puberty is earlier among racial/ethnic minority children (Herman-Giddens et al., 1997, 2012; Kelly et al., 2017; Reagan et al., 2012). These patterns by early childhood adversity and race/ethnicity may have implications for health disparities into adulthood (Bleil et al., 2017; Golub et al., 2008), as earlier puberty is associated with increased risk for depression (Wang et al., 2016) and substance use (Cance et al., 2013) during adolescence and numerous chronic disease outcomes in adulthood, including diabetes, cardiovascular diseases, and cancer (Canoy et al., 2015; Day et al., 2015; Elks et al., 2013).

> *Conclusion 2-1: Scientific research demonstrates healthy development of the child begins preconception and is dependent upon a strong foundation built prenatally. Therefore, access for the family to high-quality resources to limit chronic stress reduces risk for disrupted development and has the potential to close disparities based on race/ethnicity and socioeconomic status. Research findings show that specific types of resources are key to best outcomes, including healthy food, standard of care with a woman's health professional, maternal stress-reducing strategies, parenting education, and coaching. Research also has revealed that supportive relationships after birth are major contributors to healthy child development and building resilience.*

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BIOLOGICAL MECHANISMS OF STRESS

When the brain perceives a stressor, it induces a response that calls on multiple biological systems that, in the short term, are critical for adaptation and survival—sometimes defined as the "fight or flight" response. This stress response is a normal and healthy part of human biology. The spectrum of the stress response includes positive, tolerable, and toxic stress (Center on the Developing Child at Harvard University, 2005) and is governed by a complex interplay between the severity and duration of the stressor, individual genetic differences, gene–environment interaction, family environmental factors, developmental experiences, and the availability of buffering and coping strategies and resources. Positive and tolerable stress responses are characterized by a return to homeostasis, while the toxic stress response may induce lasting changes in brain architecture and function and in organ system development. Specific endocrine, immune, and brain cell populations produce bioactive chemicals that signal through their receptor proteins, which are present on cells throughout the body. These signals control cellular metabolism and physiological activity, which, in the short term, is a positive adaptation to the stressor.

Chronic activation of the stress response system can lead to risk for short- and long-term poor health outcomes beyond the early childhood period, including dysregulation of neuroendocrine and immune system development, altered cardiovascular functioning, metabolic dysregulation related to obesity and type 2 diabetes, changes to the gut microbiome, and epigenetic modifications that alter gene expression (Black, 2003; Campbell et al., 2014; Dinan and Cryan, 2012; Taylor et al., 2011; Vaiserman, 2015). There are individual differences in risk caused by chronically activated stress response systems, due in part to variation in the intrinsic sensitivity of the child, and physiological adaptive capacities that can be impacted by supportive environments (as discussed in greater detail below).

Activation of the stress response involves circuits in the brain that are essential for threat detection (**amygdala**), emotional regulation (frontal cortex), neuroendocrine regulation to produce cortisol (hypothalamus, pituitary, and adrenal—HPA axis), feedback to shut down the stress response (hippocampus), and the vagal complex in the brain stem that sends information to peripheral organs, such as the heart, lungs, and gastrointestinal systems (Cameron, 2009; Dedovic et al., 2009; Forsythe et al., 2014; Goodman et al., 2013). The Sympatho-Adreno-Medullary (SAM) responds to stress by producing and secreting adrenaline and noradrenaline through the vascular system. Adaptive responses of the SAM axis include increased arousal, alertness, and vigilance, improved cognition, focused attention, enhanced analgesia, and inhibition of appetite, feeding, digestion, growth, reproduction, and immunity (Chrousos, 2009). Adaptive responses of the HPA axis include regulation of cognitive, behavioral, affective, cardiovascular, and immune system functioning (Kudielka and Kirschbaum, 2005; McEwen et al., 2015). See Figure 2-2 for a diagram illustrating the stress response pathway.

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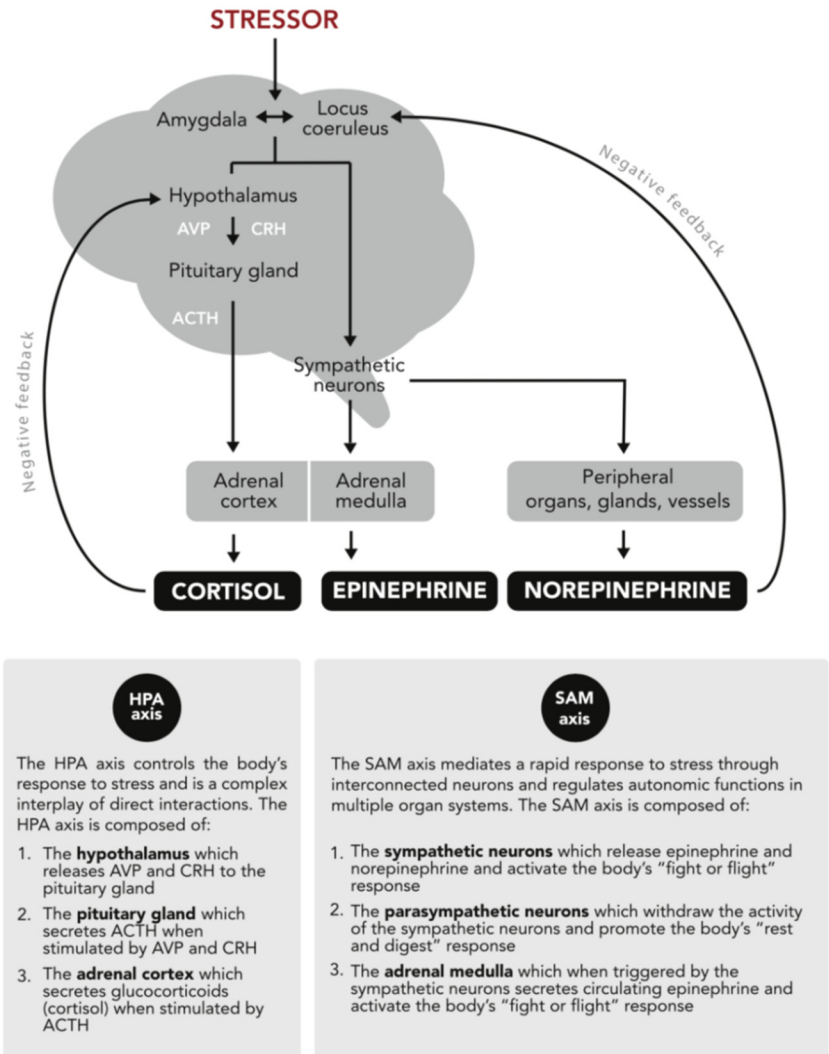

FIGURE 2-2: Stress response pathway
NOTES: HPA axis = hypothalamic pituitary adrenal axis; SAM axis = sympathoadrenomedullary axis; AVP = arginine vasopressin; CRH = corticotropin-releasing hormone; ACTH = adrenocorticotropin hormone. Reprinted from Advances in Pediatrics, 63/1, Bucci et al. Toxic Stress in Children and Adolescents, 403-428, Copyright (2016), with permission from Elsevier.
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Dysregulation of the Stress Response

Severe or chronic activation of the stress response, in the absence of adequate caregivers who serve as buffers to the stress activation, can lead to disruption of homeostatic mechanisms and long-term changes to brain architecture and organ systems (the toxic stress response) (Shonkoff et al., 2012). Early life stress can also result in epigenetic changes that can sensitize the individual to stress and adversity, alter the development of many organ systems, and impact the response to subsequent stressors later in the life course (see Epigenetics, above) (National Scientific Council on the Developing Child, 2014). Exposure to high levels of adversity in early childhood is associated with the following multisystemic disruptions, with some of these changes

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occurring as early as infancy (e.g., changes to brain structure), while others are not evident until later in life (e.g., hypertension):

- Neurologic: There are structural and functional changes to stress-sensitive regions of the brain (McEwen and Morrison, 2013; National Scientific Council on the Developing Child, 2014).
- Endocrine: There is excessive activation of both the SAM and HPA axes, associated with loss of feedback inhibition of the HPA axis, increased levels of CRH, and disruption of the daily cortisol pattern, leading to abnormally lower morning cortisol levels, elevated afternoon cortisol levels, and an overall increase in cortisol exposure. Over time, excess HPA activation may recede, leading to low or deficient HPA axis response. HPA axis overactivity is associated with suppression of thyroid function. High doses of early adversity are associated with changes in reproductive function through the HPA axis' effect on gonadotropins and/or cytokine suppression of reproductive function (Dedovic et al., 2009; Kudielka and Kirschbaum, 2005; Slopen et al., 2014).
- Immune function: There is dysregulation of neuro-endocrine-immune relations, leading to increased proinflammatory cytokines and inhibition of anti-inflammatory pathways. In addition, humoral (antibody production) and cell-mediated acquired immunity can be impaired. High levels of early adversity in childhood are associated with increased risk of autoimmune disease in adulthood (Nusslock and Miller, 2016; Padgett and Glaser, 2003).
- Cardiometabolic: There is insulin resistance, obesity, glucose intolerance, reduced control over blood lipid levels, and hypertension (Non et al., 2014; Woo Baidal et al., 2016).

Buffering the Stress Response

From a population health perspective, ameliorating the toxic stress response and understanding the biological mechanisms by which toxic stress can be prevented or mitigated is of high priority. Importantly, research suggests several mechanisms to buffer against toxic stress. One mechanism—studied in depth in animal models—is social relationships (Gunnar and Fisher, 2006). Research on resilience has clearly demonstrated that the single most important protective factor for children facing adversity is a strong, secure relationship with at least one parent; this helps foster positive outcomes across domains ranging from psychological adjustment to positive peer relationships (Cicchetti, 2013; Luthar et al., 2015; Masten, 2018; Sroufe, 2005). Moreover, a child who is securely attached to a primary caregiver is more likely to have high self-esteem, emotion regulation skills, and a positive outlook later in life (Sroufe, 2005). It also has been demonstrated that supportive relationships early in life play an important role in buffering stress responses, thereby allowing children to more easily confront stressful situations (Hostinar et al., 2014). (See Chapter 4 for an in-depth discussion on caregiver and social connections.) Some studies have shown an effect of social buffering on regulating the HPA axis, autonomic nervous system, and immune functioning, though more research is needed for more in-depth assessment of neuro-endocrine-immune, metabolic, and genetic regulatory functions (Hennessy et al., 2009; Hostinar et al., 2014; Kikusui et al., 2006).

While the mechanisms through which nurturing relationships buffer against toxic stress are likely to be complex, some studies have shown that the neuropeptide hormones oxytocin and

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vasopressin play important roles in affiliative social behavior (Gunnar and Hostinar, 2015) (Bartz et al., 2011; Ross et al., 2009a,b; Ross and Young, 2009), though experimental work in social neuroscience is challenging (Insel, 2010a). Studies in animals and humans have shown that oxytocin and/or vasopressin play a role social memory, pair bonding, empathy, altruism, emotion regulation, and trust (De Dreu and Kret, 2016; Donaldson and Young, 2008; Hostinar et al., 2014; Israel et al., 2012; Quirin et al., 2011; Rodrigues et al., 2009; Snowdon et al., 2010). It is thought that increased oxytocin counteracts the biological underpinnings of toxic stress, including increased inflammation, cardiovascular dysfunction, oxidative stress, and alterations in brain structure and function (Detillion et al., 2004; Donaldson and Young, 2008; Heinrichs et al., 2003; Hostinar et al., 2014; Mantella et al., 2004). Social buffering to toxic stress, and its action through oxytocin and vasopressin (Gobrogge and Wang, 2015), is still an ongoing area of research, as is intergenerational transmission of trauma and trauma-related stress (Kim and Strathearn, 2017). For example, how such mechanisms change with development still remains to be determined (Gunnar and Hostinar, 2015). Still, this evidence suggests that interventions to create strong social relationships may ward against the long-term impacts of ACEs and other chronic adversity over the course of child development and into adulthood. More research is needed to understand other buffers against toxic stress (e.g., sleep, exercise, nutrition) and the biological mechanisms through which they operate; prefrontal cortex plasticity is one such mechanism (McEwen and Morrison, 2013).

> *Conclusion 2-2: Among all the factors that may serve to buffer negative outcomes produced by toxic stress, supportive relationships between the child and the adults in their life are essential.*

Biological Measures of Stress in Children

In the decades since *From Neurons to Neighborhoods*, there have been substantial efforts to understand the biological changes that occur in response to early adversity (Danese and McEwen, 2012; NRC and IOM, 2000), at least in part inspired by the goal of developing valid, minimally invasive, low-burden measures of biological stress and neurobiological functioning in children that can be used to identify individuals at risk for poor long-term outcomes and, importantly, the impact of early life intervention (Shonkoff, 2010; Shonkoff et al., 2009). An increasing number of developmental studies with children now collect biological measures of stress, aiming to study how social experiences relate to changes in the biology of the child that may not cause immediate, gross disturbances (i.e., biological alterations that increase risk for subsequent chronic diseases) but rather set up the brain and peripheral systems for later emergence of physical and mental health problems over time. This research shows that traumatic experiences (such as child maltreatment) or prolonged exposure to chronic stress (such as poverty) can result in a chronically activated physiological stress response, which has a host of downstream consequences across biological systems. Some of this research has applied the concept of "**allostasis"** or "allostatic load," which provides a method to evaluate the cumulative biological impact, or "wear and tear" on the body, as the result of a given social exposure (Danese and McEwen, 2012; McEwen, 1998). As noted in previous sections of this chapter, both animal and human research has demonstrated that early life stress contributes to altered brain development (i.e., structure and function) and therefore to allostatic load.

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Synthesis

Advances in science technologies, big data collection and information technology, and new strategies for analyzing complex data have been instrumental in rapidly advancing the science of brain and child development over the nearly 20 years since *From Neurons to Neighborhoods* was released to the public (NRC and IOM, 2000). Research findings have revealed the details of how early experiences are essential for building brain connections that underlie biobehavioral health, the role of genetics in development, and a new understanding of whole-child development that relies on organ systems interacting with each other and the environment to establish health and biobehavioral fitness or risk due to early influences of adversity.

Conclusion 2-3: Research has changed the discourse to include scientific facts in child development that have determined that when considering the entire life course, it is early experiences, pre- and postnatally, that are the most powerful in working together with an individual's genetic makeup to influence the physical, mental, and cognitive development of the child. Science also shows that maternal health and well-being is a major contributing factor to the development of the fetus and establishes a foundation for positive or negative child health outcomes.

Conclusion 2-4: Based on the abundant science, the influence of access to basic resources prenatally, particularly nutritional, psychosocial, and health care components, is powerful. The new understanding defines three interrelated parts:

1) Chronic activation of the stress response system of primary caregivers, and in the infant and toddler, can occur due to a large number of factors in the environment of the family, including lack of access to quality health care, child care, economic security, community support programs, transportation, stable housing, and healthy nutritional sources; institutional and individual racism and sexism; and community violence;

2) Sustained stress activation can produce a response in the child known as toxic stress, which affects the development of the structural organization and functioning of brain circuits that impact the quality of cognition and social and emotional regulation, including the child's own stress response, reward, and motivation. The toxic stress response therefore directly impacts the behavioral and psychological well-being of the child; and

3) New data from longitudinal studies, combined with previous work from retrospective analyses of the early life experiences of adults, provides a compelling demonstration that toxic stress response substantially increases later-life risk for physical illnesses, including obesity and type 2 diabetes, cardiovascular disease, substance abuse, mental illness, lower educational achievement, cancer, and infectious disease. In addition to the adverse effects of toxic stress on the brain and nervous system, it also affects every other organ system in the child's body, impacting short and long-term health.

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Individual Differences in Responsiveness and Susceptibility

Research over the past two decades clearly shows that what were previously considered to be universal developmental processes (see Chapter 4) do not occur uniformly. Relevant to this focus of this chapter, in part, this can be due to variation in the development trajectory of stress-sensitive brain circuits in infancy, as well as the interplay between genes and environments that mediate the neurobiological and physical impact of positive or negative early life experiences. Thus, heterogeneity with substantial variations is often observed, such as across groups that differ by SES or ethnicity. Diversity brought about by individual and subgroup differences, based on factors such as personal experiences, individual temperament, culture, and the family and community context, can alter the pace and trajectories of developmental biological and psychological processes. For example, humans are predisposed to become attached to our main caregivers, but the quality of interactions between caregiver and child and other environmental conditions can contribute to the development of various types of infant attachment. To consider another example, a rich literature supports the notion that children differ in their temperament: behavioral predispositions can permeate how children react to and process environmental input. From conception, children bring differential responsiveness to—and processing of—their physical and relational context. Infants are described as difficult, inhibited or uninhibited, shy or slow to warm up (Conture et al., 2013). These dispositions carry consequences for adaptability to new environments, relational difficulties, and even long-term mental health problems (Clauss et al., 2015). One research notion from the temperament literature proposes that there are behavioral predispositions that are normative variations across individuals (Durbin and Hicks, 2014). Another theory sees the origin of these individual differences as consequences of exposure to early stress, whereby stress experienced by the mother and fetus before birth conditions the nervous system to respond, perhaps in an exaggerated way, to neutral or mild environmental input (Miller et al., 2011).

Recently, there has been an emphasis on elucidating the mechanisms underlying the differential susceptibility to the environment hypothesis. Work by both Belsky and Boyce has advanced understanding of how individual differences in how children process and react to the environment contribute to individual risk and resilience (Boyce, 2019; Bush and Boyce, 2016). Behavioral genetics research defined the contributions of both heritability and environment to the variance in the symptoms and incidence of behavioral disorder diagnoses (Plomin et al., 2001). This core principle is evident in the assessment of typical behavioral characteristics, such as affiliative social behavior. Studies of twins highlight the remarkable variability in specific traits that make up affiliative social behavior (Ebstein et al., 2010), with some elements exhibiting high (~0.7) heritability (prosocial behavior; social responsiveness) and others low (~.3) heritability (empathy; secure attachment). In animal studies, heritability of the behaviors that make up affiliative social behavior exhibits very similar genetic contributions (Knoll et al., 2018).

These studies also showed that environmental context plays a particularly important role in modifying behavior—even those with moderate to high heritability. As Bush and Boyce (2016) note, the accumulating biological data from studies of humans and animals has changed the concept from the long-standing diathesis stress model of sensitivity to environment to one that focuses on determining factors that confer individual differences that affect susceptibility and resilience. Stress reactivity, which had been presumed to influence behavioral outcomes in a single direction—negatively—is an important individual difference that can influence responses to different environmental contexts in either a positive or negative direction (Bush and Boyce,

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2014; Ellis et al., 2011; Obradovic et al., 2010). Again, context appears to be a major moderator of biological predispositions, and an ongoing topic of study is why some children adhere to healthy developmental trajectories despite profoundly adverse circumstances while others do not (NASEM, 2015; Cicchetti, 2013; Masten, 2014; Rutter, 2012; Shonkoff, 2016).

The importance of individual differences in development points out lessons for practice and policies in efforts to prevent developmental problems and promote positive trajectories (Shonkoff, 2017). A key fact is that universal interventions will be more or less effective depending upon the individual(s) being targeted. Thus, well-designed interventions are those that are sufficiently flexible to take individual differences and heterogeneity into account. The challenge continues to be predicting what specific changes are most likely to produce the best outcomes. For example, the New Hope Project, a welfare reform demonstration based in Milwaukee, WI, was designed to raise families out of poverty, using a work-support intervention that was tailored to the specific needs of each family (Huston et al., 2011). The intervention had multiple components that families would choose from over 3 years, and it was expected that the combination of benefits and services would increase parents' employment, income, and use of child care and health insurance. In a report to summarize effects over 8 years (i.e., 5 years after the intervention ended), researchers concluded that work supports can positively impact low-income parents and children, even though the economic effects of employment and income were not sustained beyond the 3 years of the intervention. Importantly, positive effects for children appeared during the intervention period and extended to the 8-year follow-up.[8]

Another example is *Filming Interactions to Nurture Development* (FIND), a 10-week video-feedback program to promote acquisition of developmentally supportive parenting skills (Fisher et al., 2016; Nese et al., 2016). This behavioral training program is tailored to address individual differences in the types of challenging parenting styles via weekly structured coaching sessions, and findings suggest that it is effective for improving caregiver skills (Fisher et al., 2016).

In some cases, individuals have different responses to social or environmental stimuli as the result of age of exposure. Specific time points in development represent "sensitive periods" when risk or protective factors have a maximal influence on cognitive capacities, biology, emotions, and behavior (Ben-Shlomo and Kuh, 2002; Hertzman and Boyce, 2010; Knudsen, 2004; Zeanah et al., 2011). The strongest evidence for the heightened importance of specific periods comes from experiments using laboratory animals (Hertzman and Boyce, 2010) because it is challenging to isolate sensitive periods within observational studies in humans (given that risk factors are often continuous and may in turn elicit subsequent risks). As noted in detail above, the prenatal, postnatal, and early childhood phases are recognized as having sensitive periods that result in brain and behavioral changes. Research is beginning to elucidate how risk factors at different time points may have varying impacts. Although it is generally accepted that developmental timing can influence the effect of a social or environmental risk factor on child health, the majority of studies do not consider how risk or protective factors vary by age.

[8] Although the positive effect of this personalized intervention on children's academic performance and test scores at the 2- and 5-year follow-ups faded, at the 8-year follow-up, children in families assigned to the intervention condition were more likely to be engaged in school and to show positive social behaviors relative to those in families assigned to the control condition, and they were less likely to have to repeat a grade, to be placed in special education, to receive poor grades, or to have cynical attitudes about work (Miller et al., 2008).

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CONCLUSION

With advances in science emerging from different disciplines, now more than ever, there is a major need to facilitate new public–private-nonprofit sector relations to build substantial opportunities for the fields of neuroscience, behavioral sciences, and psychology to collaborate. Therefore, the committee recommends:

Recommendation 2-1: The National Institutes of Health and other relevant research entities should support the development of public–private partnerships, or other innovative collaborations, to

- **Build multidisciplinary teams, including but not limited to researchers in neuroscience, endocrinology, immunology, physiology, and metabolism, behavior, psychology, and primary care, to identify the most relevant factors in a child's complex environment that promote resilience and promote outcomes related to physical and mental health.**
- **Conduct research that measures the impact of chronic stress on all relevant organ systems and determines the specific molecular and biological pathways of interaction during the pre- and postnatal periods, which are directly relevant to potential interventions to address health disparities.**

This will require recruiting diverse populations, with explicit attention to addressing racial/ethnic and socioeconomic inequities in developmental outcomes. Research with these populations will require researchers and practitioners as co-investigators that are knowledgeable of theoretical models, measures, and the realities of these families and communities. For all studies that typically use expensive biological assessments (e.g., biomarkers as mediators, moderators, or outcomes) for use in interventions, researchers need to strongly consider the ability to apply measures within the context of scaled, community-based interventions to have the broadest possible impact on outcomes for children and their caregivers. For example, there is a major gap in research support for projects that combine modern, noninvasive technologies to measure key domains of brain, behavioral, and psychological development with clinical interventions that promote best outcomes for children and their caregivers.

In summary, research advances in neuroscience, immunology, endocrinology, behavior, and psychology in the last two decades provide a new depth of understanding regarding the impact of adverse experiences on developmental processes throughout the life course. Because of these advances, the fields of developmental neuroscience, behavior, and psychology are recognizing the trajectories of development in the context of racial, ethnic, and socioeconomic health disparities. Deciphering the mechanisms through which adversity disrupts foundational developmental processes of the whole child, and the long-term impact on health, wellness, personal and community relationships, and life-span productivity across the life course, is being achieved through multidisciplinary, longitudinally designed research that has incorporated principles of human development, including individual differences and differential sensitivity to context. With this foundation, the chapters that follow provide advances in the research that addresses the role of macro- and micro-level variables in influencing development and outcomes.

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3

Development Happens in Contexts: Overview of Early Life Critical Influences

INTRODUCTION

To ensure healthy and optimal development for all children, there is a need to understand how the context that a child grows in and the cumulative risk associated with that specific context shape the odds for thriving. Sir Michael Rutter, the first professor of child psychiatry in the United Kingdom, is credited as one of the first to bring the concept of cumulative risk to the study of child development. In his now classic work studying 10-year-old children, he documented that children who had two or more risk factors had a fourfold risk for a psychiatric disorder (Rutter, 1979). Approximately a decade later, Sameroff and colleagues extended this work to young children's social and emotional competences (Sameroff et al., 1987a) and cognitive outcomes (Sameroff et al., 1987b). Since these studies, and the subsequent publication of *From Neurons to Neighborhoods* (NRC and IOM, 2000), there has been a rapid increase of cross-disciplinary research using an accumulation of risk models to assess early childhood social risk factors in relation to outcomes across the life course. Understanding the origins and mechanisms of the contextual factors and cumulative risk that produce inequities for children and families is a prerequisite to advance health equity.

While Chapter 2 discusses the mechanisms of healthy development within the growing child, this chapter provides an overview of the key early life protective and risk factors associated with development, as indicated in the committee's statement of task. This chapter discusses how each of these factors and conditions shape health and safety, mental and emotional well-being, and cognitive health during the prenatal and early childhood phases by mapping each factor to both the conceptual model and the forthcoming solution-driven Chapters 4–8 (see Figure 3-1 and Table 3-1 for how the content in this chapter maps to the conceptual model and the following chapters, and Box 3-1 for a high level chapter overview). The committee identified domains by which to group these critical influences, with a focus on factors that shape inequities at the child, family, community, and population levels. These domains include family cohesion and healthy social connections (see Chapter 4); health care (see Chapter 5); healthy living conditions (i.e., economic security, nutrition and food security, housing, and environmental safety) (see Chapter 6); and early care and education (ECE) (see Chapter 7).

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BOX 3-1
Chapter in Brief

This chapter provides an overview of the key external influences that affect brain development in early life to either promote or hinder a child's opportunity to achieve optimal health and wellbeing. The chapter summarizes the evidence for the way multiple domains (family cohesion and healthy social connections; health care; healthy living conditions including economic security, nutrition and food security; neighborhood conditions, housing, and environmental safety; early care and education) converge to create an accumulation of risk. This composite risk is heavily influenced by racism and discrimination and affects outcomes across a child's entire life course. A review of the epidemiology of risk and protective factors, prenatal and childhood outcomes, and evidence-based mechanisms (when known) are included.

Key findings on early life critical influences:

- There are specific risk and protective factors that affect health and development at multiple levels (e.g., individual, family, neighborhood, and systems/policies).
- Racism and discrimination are crosscutting factors that perpetuate structural inequities and thwart healthy development for specific groups of children.
- Exposure to multiple risk factors discussed in this chapter can lead to an accumulation of risk over the life course and ultimately result in poor health outcomes in adulthood. Conversely, exposure to positive exposures and buffering experiences can promote health and resilience for children and adults.

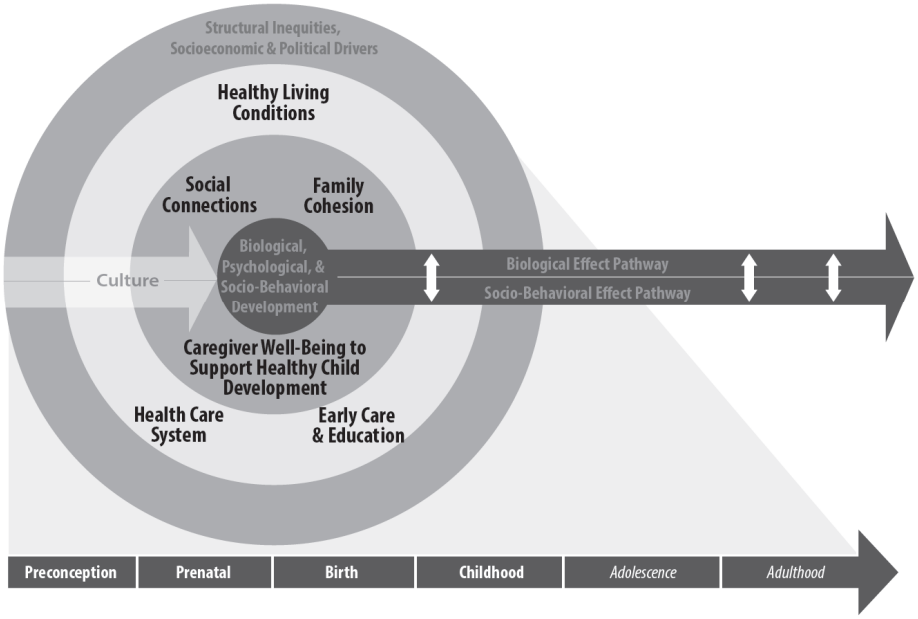

FIGURE 3-1 Mapping Chapter 3 early life protective and risk factors for child development to conceptual model and solution-driven chapters.
NOTES: The domains of focus in this chapter are reflected in the second and third rings in the model. These elements are bolded for emphasis.

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TABLE 3-1 Mapping Chapter 3 Content to Chapters 4–7

Critical Influences in Prenatal and/or Early Childhood Stages Discussed in This Chapter	Corresponding "Action" Chapter
Family cohesion and social connections	Chapter 4
Health care	Chapter 5
Economic security	Chapter 6
Neighborhood conditions (e.g., concentrated disadvantage, physical and social environments, violence and crime, housing, environmental exposures)	Chapter 6
Early care and education	Chapter 7

Committee's Approach to Early Life Critical Influences

While there are many critical factors that shape development, the committee limited the scope of this chapter to include those with strong evidence for shaping affecting outcomes from the prenatal through early childhood periods. Where possible, the committee relied on high-quality systematic reviews and meta-analyses to provide a brief overview. To the extent possible, this chapter includes data and outcomes specific to the prenatal through early childhood periods; however, when those data are not available, data for caregivers and families are presented. The committee takes a life course approach in this report (see Figure 1-1), so, as such, it is important to note that many of the influences discussed in this chapter manifest in adolescent and adult outcomes. In each domain, critical factors are addressed in terms of definition, overall prevalence and disparities, and prenatal, birth, and early childhood outcomes, including information on potential mechanisms when the evidence points to these.

Crosscutting Elements: Discrimination and Racism

Across all the critical factors in this chapter, there are two crosscutting elements that the committee has identified as being pervasive and rooted in health inequities—discrimination and racism—that can be thought of as the mechanisms by which structural inequities operate. (See Box 3-2 for a description of the root causes of health inequities and key definitions from a related report.) The crosscutting elements operate at multiple levels (i.e., intrapersonal, interpersonal, institutional, structural) and shape the experiences of children and families across the domains discussed in the rest of this chapter (and this report). While race is considered a social rather than biological construct (i.e., created from prevailing social perceptions, historical policies, and practices), the consequences of racism and the experiences of racial and ethnic minorities have psychological, biological, and social consequences (NASEM, 2017a). For example, historical policies and practices, such as residential segregation, redlining of districts, and discriminatory banking practices, are structural forms of racism that have long-lasting ramifications for the health of communities of color (NASEM, 2017a). This type of structural racism unfolds across many of the domains discussed in this chapter, including housing, economic stability, physical environment, and community violence. Furthermore, structural racism has been linked to historical trauma, which manifests from past treatment of specific

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racial or ethnic groups (NASEM, 2017a). This type of trauma is an important yet often "invisible" context that sets the stage for risk and resilience. Another example of a crosscutting element that is salient for this chapter is immigration-related experiences and challenges. For immigrant children or children of immigrants, there are unique barriers to achieving optimal health that play out at multiple levels and have serious implications for food security, housing stability, safety, and access to quality health care services. These important crosscutting elements will be highlighted throughout the chapter as they relate to the various factors that influence early development. The chapter will also discuss specific subgroups with unique needs or circumstances as they relate to these crosscutting elements, where appropriate.

BOX 3-2
Key Terms and Concepts from *Communities in Action: Pathways to Health Equity*

The 2017 report *Communities in Action: Pathways to Health Equity* examined the root causes of health inequities in the United States. The authoring committee identified two clusters of root causes:

1. The intrapersonal, interpersonal, institutional, and systemic mechanisms that organize the distribution of power and resources differentially across lines of race, gender, class, sexual orientation, gender expression, and other dimensions of individual and group identity and

2. The unequal allocation of power and resources—including goods, services, and societal attention—which manifests in unequal social, economic, and environmental conditions.

This report defines structural inequities as the systemic disadvantage of one social group compared to other groups with whom it coexists, encompassing policy, law, governance, and culture.

SOURCE: NASEM, 2017a

FAMILY COHESION AND SOCIAL CONNECTIONS

Relationships in early childhood form the foundation for how children interact with their environment and other individuals across the life-span. This section discusses key influences that exist within the most proximal microsystem for children, consisting of family and peers. The following discussions include the scientific findings on experiences or factors that either enhance or undermine family cohesion and healthy social connections. Chapter 4 describes the interventions and mechanisms needed to support the family and promote healthy relationships. See Figure 3-2 for a visual of how this section ties to the report conceptual model.

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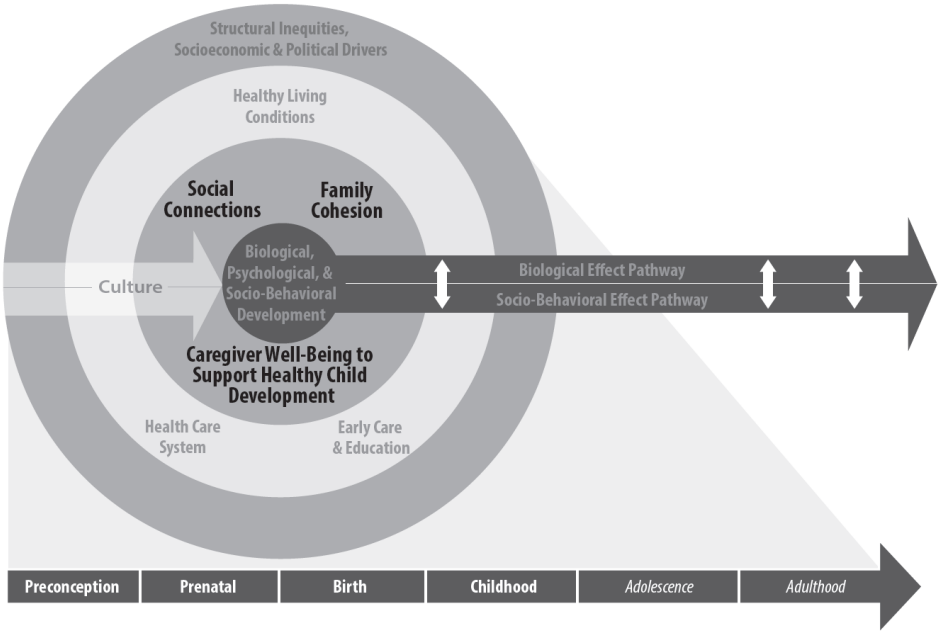

FIGURE 3-2 Leveraging early opportunities to achieve health equity across the life course: A conceptual framework.
NOTES: The elements of focus in this section are reflected in the second innermost ring in the model: caregiver well-being and support, family cohesion, and social connections. These elements are bolded for emphasis.

Parental Factors

Parental well-being is a critical determinant of child health and developmental outcomes; therefore, what happens to the parent before, during, and after pregnancy has serious implications for the child. Parental mental health is inextricably linked to child well-being and there is consistent evidence that maternal depression compromises healthy child development (Center on the Developing Child at Harvard University, 2009; Goodman and Garber, 2017). Estimates show that 10–20 percent of mothers have lifetime prevalence of depression and the rates are much higher for mothers in low-income households. Because of the socioeconomic disparities in maternal depression and the implications for children's development, maternal depression has been thought of as a contributing factor to the cycle of intergenerational poverty (Reeves and Krause, 2019). For children, the consequences of maternal depression include neurodevelopmental and other biological disruptions and psychological and behavioral difficulties. Postpartum maternal depression specifically has been linked to the neurobiological pathways that shape emotional regulation, cognitive and executive function, and physiologic stress response systems—all critical functions and systems for ensuring optimal development (Drury et al., 2016). The Center on the Developing Child at Harvard University (2009) published a paper on the effects of maternal depression that indicated the following:

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- Chronic depression can manifest in two types of problematic parenting patterns that disrupt the "serve and return"[1] interaction that is essential for healthy brain development: hostile or intrusive, and disengaged or withdrawn.
- Children who experience maternal depression early in life may have lasting effects on their brain architecture and persistent disruptions of their stress response systems.
- Maternal depression may begin to affect brain development in the fetus before birth.

While evidence exists that maternal depression is significantly related to children's behavioral and emotional functioning, more research is needed to better understand moderating effects for various subgroups of children (Goodman et al., 2011). In addition to maternal depression, paternal depression has also been shown to negatively affect parenting behaviors and child developmental outcomes (Gutierrez-Galve et al., 2015; Ramchandani et al., 2011; Wilson and Durbin, 2010). Sweeney and MacBeth (2016) identify the following mediators of the effect of paternal depression on children: paternal negative expressiveness, hostility, and involvement and marital conflict. There are other aspects of parental well-being for which evidence exists of an association with child health and well-being (e.g., parental alcohol or substance abuse or incarceration) (Conners et al., 2004; Lieb et al., 2002; Luthar et al., 2007; Nichols and Loper, 2012; VanDeMark et al., 2005). Additional select parent-specific factors that have been shown to be associated with child health and well-being outcomes include incarceration (see Chapter 4), interaction with the welfare system (see Chapter 4), adverse childhood experiences (ACES) (see the Adverse Childhood Experiences section at the end of this chapter), and intimate partner violence (IPV).

IPV has adverse effects on pregnancy and birth outcomes (Boy and Salihu, 2004; Iiiopoulou et al., 2012). A systematic review found that pregnant women who have experienced IPV are more likely to suffer from low birth weight (LBW), maternal mortality, and infant mortality when compared to women who have not experienced IPV (Boy and Salihu, 2004). Research also suggests that children can suffer from harmful consequences associated with exposure to IPV, even if they have not directly observed the violence (Wathen and Macmillan, 2013). These consequences include social, emotional, and behavioral problems, such as mood and anxiety disorders, posttraumatic stress disorders, substance abuse, and school-related problems in childhood and adolescence. Rates of comorbidity between exposure to IPV and child maltreatment are high—some data show that 60–75 percent of families that have experiences of IPV also include children exposed to maltreatment (Osofsky, 2003).

Child Maltreatment

In the absence of safe and nurturing relationships, children are vulnerable to the effects of maltreatment, or child abuse and neglect. The Child Abuse Prevention and Treatment Act defines child abuse and neglect as

[1] "Serve and return" interactions occur when young children innately reach out for connection by babbling, using facial expressions, and gestures (i.e., serve), and adults respond with similar vocalization or gestures (i.e., return) (National Scientific Council on the Developing Child, 2004).

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> any recent act or set of acts or failure to act on the part of a parent or caretaker, which results in death, serious physical or emotional harm, sexual abuse or exploitation, or an act or failure to act, which presents an imminent risk of serious harm.[2]

Generally, child abuse and neglect are grouped into four types: neglect and physical, sexual, and emotional abuse. See Box 3-3 for descriptions of the four types of maltreatment. Despite this definition, there remain challenges related to varying state legal definitions, data collection, and calculating accurate incidence and prevalence rates in population-based studies (NRC and IOM, 2014). Therefore, the magnitude of child abuse and neglect is more than likely underestimated in the United States (Fortson et al., 2016). Not only is child maltreatment detrimental to children's health and well-being, it is also costly for the United States. Data from 2008 indicate that the total lifetime economic cost associated with child abuse and neglect amounted to approximately $124 billion (Fang et al., 2012).

BOX 3-3
Defining Common Types of Child Maltreatment

- *Physical abuse* is the intentional use of physical force that can result in physical harm. Examples include hitting, kicking, shaking, burning, or other shows of force against a child.
- *Sexual abuse* involves pressuring or forcing a child to engage in sexual acts. It includes behaviors such as fondling, penetration, and exposing a child to other sexual activities.
- *Emotional abuse* refers to behaviors that harm a child's self-worth or emotional well-being. Examples include name-calling, shaming, rejection, withholding love, and threatening.
- *Neglect* is the failure to meet a child's basic physical and emotional needs. These needs include housing, food, clothing, education, and access to medical care.

SOURCE: Excerpted from CDC, 2019.

Trends and Disparities

Despite the well-known consequences, child maltreatment is still too common today. At least one in seven children have experienced child abuse or neglect in the past year, which is likely an underestimate (Finkelhor et al., 2015). State-level data on reports of child abuse and neglect show a 2.7 percent increase in the national rate of victims of child maltreatment from 2013 to 2017 (HHS, 2019).

Disparities in child maltreatment exist across groups by race and ethnicity, age, and SES. According to 2017 data from the National Child Abuse and Neglect Data System (NCANDS), rates of child abuse or neglect were reported to be highest for American Indian/Alaska Native (AI/AN) children (14.3 per 1,000 children). Following those are rates for African American (13.9 per 1,000), multiracial (11.3 per 1,000), Pacific Islander (8.7 per 1,000), white (8.1 per 1,000), Hispanic (8.0 per 1,000), and Asian (1.6 per 1,000) children. Likewise, the rate of fatalities due to maltreatment for African American children (4.86 per 100,000 children) is 2.6 times greater than that of white children and 3.1 times higher than that of Hispanic children (HHS, 2019).

Young children are the most vulnerable to child maltreatment (HHS, 2019), and more than one fourth (28.5 percent) of child maltreatment victims are younger than 3 years. Overall,

[2] 42 U.S.C. § 5101 note.

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children under the age of 1 year are most likely to die from child abuse or neglect (see Figure 3-3). Very young children with disabilities have increased risk of child maltreatment when compared to children without disabilities—and rates of disability are higher among low-income populations that experience higher rates of preterm birth (Hibbard et al., 2007; NASEM, 2015a). Children living in poverty are disproportionately exposed to child abuse and neglect and experience five times the risk of child abuse or neglect as children from higher SES households (Sedlak et al., 2010).[3] It is important to note that disparities in child maltreatment have been attributed to other social, environmental, or economic factors, pointing to the need for a multilevel approach to preventing maltreatment (Fortson et al., 2016; Gilbert et al., 2009).

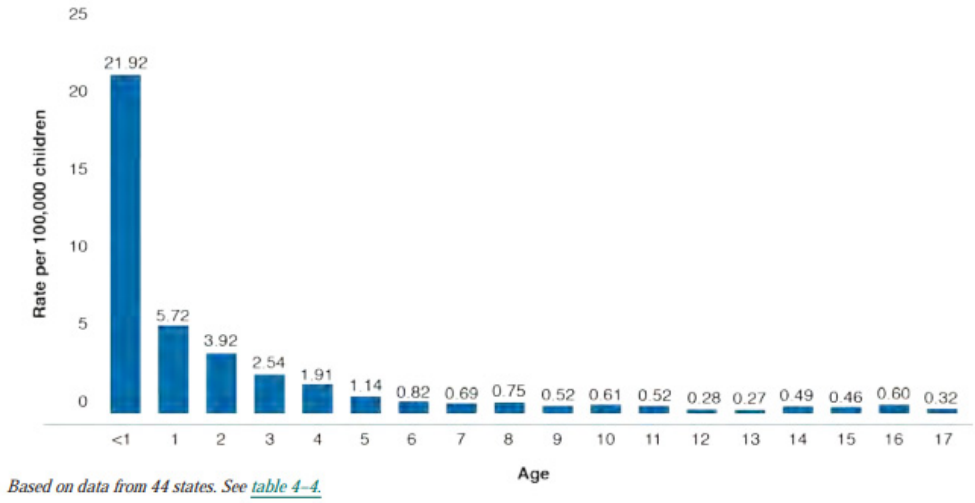

FIGURE 3-3 Child fatalities due to maltreatment by age, 2017
SOURCE: HHS, 2019

Outcomes Related to Child Maltreatment

Child maltreatment has a pervasive and harmful influence on many aspects of development and children who are exposed to a certain type of maltreatment are frequently exposed to others (Gilbert et al., 2009). Exposure to child abuse and neglect has serious ramifications for a child's biological, behavioral, cognitive, and emotional development (Cicchetti and Handley, 2017).The relationship between child abuse and neglect and physiological, psychological, and behavioral outcomes has been well documented throughout the child development literature, and a comprehensive review was published in the 2014 report IOM and NRC *New Directions in Child Abuse and Neglect Research.* Therefore, the literature review is not repeated here; key findings related to the effects of maltreatment from the NRC and IOM report are provided in Box 3-4. See Chapter 2 for more on biological mechanisms of development that can be hindered or enhanced through early experiences.

[3] In the *Fourth National Incidence Study of Child Abuse and Neglect (NIS-4): Report to Congress*, a low-socioeconomic status household was defined as having household income below $15,000 per year; parent education level less than high school; or any member of the household receiving poverty program benefits (e.g., Temporary Assistance for Needy Families, food stamps, public housing, energy assistance, or subsidized school meals) (Sedlak et al., 2010).

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BOX 3-4
Key Findings from the 2014 IOM and NRC Report *New Directions in Child Abuse and Neglect Research*

- Across human and nonhuman primate studies, perturbations to the hypothalamic–pituitary–adrenal axis (HPA) system often are seen to be associated with child abuse and neglect. The findings are complex, moderated by a number of factors, and seen at some ages and not others.
- Abused and neglected children show behavioral and emotional difficulties that are consistent with effects on the amygdala, such as internalizing problems, heightened anxiety and emotional reactivity, and deficits in emotional processing.
- Despite mixed evidence regarding structural changes in the prefrontal cortex, a number of studies suggest that abuse and neglect are associated with functional changes in the prefrontal cortex and associated brain regions, often affecting inhibitory control.
- Abuse and neglect have profound effects on selected aspects of children's cognitive development. Although many attempts have been made to disentangle the effects of abuse and neglect, the balance of findings suggests that severe neglect may interfere with the development of executive functioning, and both neglect and abuse increase the risk for attention regulation problems and attention deficit hyperactivity disorder (ADHD), lower IQ, and poorer school performance.
- Children who experience abuse or neglect have been found to be at higher risk for the development of externalizing behavior problems, including oppositional defiant disorder, conduct disorder, and aggressive behaviors. Abused and neglected children also have been found to be at increased risk for internalizing problems, particularly depression, in childhood, adolescence, and adulthood.
- A number of studies have found elevated rates of Posttraumatic Stress Disorder (PTSD) among individuals with a history of abuse and neglect. PTSD has been associated with physical, cognitive, psychological, social, and behavioral problems among youth who were abused or neglected in childhood.
- Experiences of child abuse and neglect have effects on many health outcomes, including risks for long-term chronic and debilitating diseases and, in extreme cases, stunted growth.
- Experiences of abuse and neglect in childhood have a large effect on delinquency, violence, and suicide attempts in adolescence and adulthood.
- Adolescents and adults with a history of child abuse and neglect have higher rates of alcohol abuse and alcoholism than those without a history of abuse and neglect, although this relationship has been found most frequently in women.
- Not all children who experience abuse or neglect show problematic outcomes. Factors that influence resilience among abused and neglected children have been identified at the level of the individual child, the family, and the child's broader social context. These factors, along with risks and stressors at each level, interact with one another to predict resilient outcomes.
- The timing, chronicity, and severity of child abuse and neglect, as well as the context in which they occur, have been shown to impact the associated outcomes.

SOURCE: IOM and NRC, 2014

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Peer Social Connections

For children, an important source of social connections is the peer group that they interact with. After experiences with primary caregivers and family members, peers are the most proximal interpersonal relationships for young children. Through peer relationships and interactions, young children learn and reinforce notions of reciprocal interactions. For example, young children learn how to share, take turns, and give and receive from interactions with other young children. From these experiences, they can also learn to interpret and take into account the needs and desires of others and to manage their own impulses (National Scientific Council on the Developing Child, 2004). From early on, peer interactions can have positive and/or negative effects on children's mental, behavioral, and emotional health. (See Box 3-5 for findings and conclusions from a report on preventing bullying.)

BOX 3-5
The Effects of Bullying in Early Childhood

The 2016 report *Preventing Bullying Through Science, Policy, and Practice* makes the case that bullying is not a normal aspect of childhood, as it is commonly thought to be. Bullying——unwanted aggressive behavior repeated (or likely to be repeated) by another youth or group of youths that involves a power imbalance—can be thought of as a public health issue. In fact, the report draws the following findings and conclusions about the serious development effects and implications of being the victim or perpetrator of bullying behavior at a young age:

- Although the effects of being bullied on the brain are not yet fully understood, there are changes in the stress response systems and in the brain that are associated with increased risk for mental health problems, cognitive function, self-regulation, and other physical health problems.
- The long-term consequences of being bullied extend into adulthood, and the effects can be more severe than other forms of being maltreated as a child.
- Bullying has significant short- and long-term internalizing and externalizing psychological consequences for the children who are involved in bullying behavior.
- Existing evidence suggests that both social-cognitive and emotion regulation processes may mediate the relation between being bullied and adverse mental health outcomes.
- Although genes appear to modulate humans' response to being either a target or a perpetrator of bullying behavior, it is still unclear what aspects of these experiences are interacting with genes and which genes are implicated to produce the variability in outcomes. Examining the role of genes in bullying in the context of the environment is essential to providing meaningful information on the genetic component of individual differences in outcomes from being a target or a perpetrator of bullying behavior.

See Chapter 7 in this report on Promoting Health Equity Through Early Care and Education for more on the role of educators in preventing bullying.

SOURCE: NASEM, 2016b

Safe, Stable, Nurturing Relationships and Environments

While child abuse and neglect increase the risk for a host of mental and physical ailments, lifelong mental and/or physical anguish is not a foregone conclusion. There is clear scientific evidence that the presence of safe, stable, and nurturing relationships is critical to healthy development and can buffer the mechanisms of adversity and support positive trajectories (Bornstein and Leventhal, 2015; Bronfenbrenner and Morris, 2006; CDC, 2014;

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Luthar, 2006; Masten, 2014; National Scientific Council on the Developing Child, 2004). (For information on mechanisms of buffering the stress response, see Chapter 2.) Safe, stable, and nurturing relationships are also important for preventing maltreatment, as they have been identified as a moderator of intergenerational child maltreatment. A meta-analysis found that adult relationships and parent–child relationships had a protective effect against intergenerational child maltreatment when they were safe, stable, and nurturing (Schofield et al., 2013). A stable relationship with caring adults is also important in child care settings, which can affect social competence, behavioral issues, and thinking and reasoning skills. In a 2004 summary of key scientific findings on relationships, the National Scientific Council on the Developing Child identifies nurturing and stable relationships with caring adults as a cornerstone of healthy development. The Council identifies "serve and return" as a key mechanism of healthy interactions between parents and babies, by which the brain architecture is strengthened. This is also important in child care settings, which can affect social competence, behavioral issues, and thinking and reasoning skills. Finally, the Council concludes based on the science that secure and stable relationships can be protective from illness, chronic stress, exposure to toxicants, and preventable injuries.

HEALTH CARE

In addition to the social determinants of health (SDOH) described throughout this report, health care itself plays a major role in child development through the life course. From preconception through early childhood, there are several aspects of health care that are critically important from an equity perspective: access to quality care, coverage, provider bias (or lack thereof), and provider cultural/linguistic competency. The following section provides an introduction to these issues along the continuum of preconception, prenatal, and pediatric health care; Chapter 5 provides a more in-depth look into opportunities to enhance these systems. See Figure 3-4 for a visual of how this section ties to the report conceptual model.

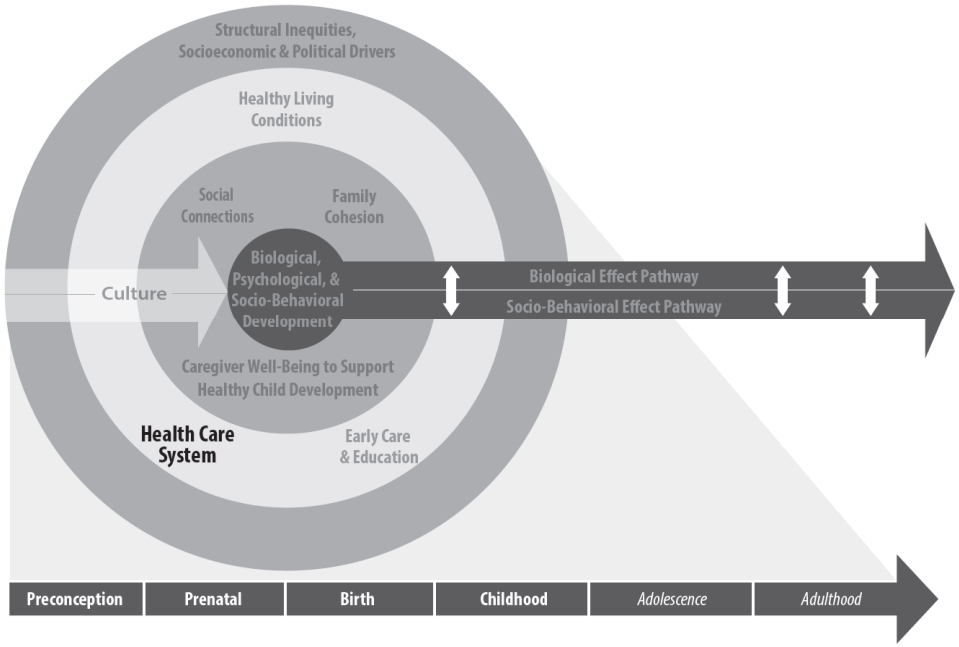

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FIGURE 3-4 Leveraging early opportunities to achieve health equity across the life course: A conceptual framework
NOTE: The element of focus in this section is reflected in the second outermost ring in the model: the health care system. This element is bolded for emphasis.

Preconception Care

The number of studies devoted to understanding preconception health and the interventions intended to improve health among women who may become pregnant have increased in recent years, but there is still a general paucity of data. According to a 2016 review of this research, results are mixed, and the studies themselves are generally of moderate to poor quality (Hussein et al., 2016). While preconception care may increase women's knowledge about certain health conditions and pregnancy (Callec et al., 2014; Chuang et al., 2010), it is not clear if this translates to healthier outcomes for infants and children. Across eight randomized controlled trials (RCTs) of preconception health care interventions, there was little evidence that interventions reduced adverse pregnancy outcomes (Hussein et al., 2016). This was true even though the interventions appeared to increase maternal knowledge and locus of control and reduce risky behaviors (Hussein et al., 2016).

It is also true that only a small minority of women receive preconception care, making it difficult to draw broad generalizations about its effectiveness (Hemsing et al., 2017). This is likely because access to preconception care is not widespread (Hemsing et al., 2017), and a recent review outlined several reasons why (Goossens, 2018). First, a lack of reimbursement mechanisms for administering preconception care reduces providers' incentive to deliver it. Lack of reimbursement, payment mechanisms, or insurance are also reasons that preventive services and health promotion services are not delivered more generally (Goossens, 2018). Limited time and heavy workloads were a second reason that health care providers do not engage in preconception care with widespread frequency. A third reason was simply a lack of knowledge on the part of clients themselves (i.e., they were not aware of preconception care or did not seek it out) (Goossens, 2018). A fourth reason was that providers themselves did not think it was their duty to provide preconception care; primary care providers, for instance, were likely to think it was the role of the obstetrician, and vice versa (Goossens, 2018). A final reason could be that, given the lack of strong evidence for improving birth outcomes (Hussein et al., 2016), women simply choose not to engage in preconception care because they do not think it will be effective.

Preconception care as currently conceptualized—typically a single medical visit initiated when one wants to become pregnant—is likely insufficient to address the accumulated life and medical risks up to that point for both parents. Thus, a broader, multisector view of preconception care seen through an SDOH lens is likely necessary to achieve the significant impact that would be expected when one mitigates potential risks prior to achieving pregnancy.

Prenatal Care

Delivered early and often, prenatal care increases the chances of delivering a healthy baby (Kilpatrick et al., 2017). Conversely, women who do not receive prenatal care are 3–4 times as likely to die from pregnancy-related issues compared to women who do receive such care (Bingham et al., 2011). There are well-documented negative effects on babies as well (Cox et al., 2011). Women who do not receive prenatal care are more likely to deliver babies that are underweight and preterm and more likely to die in infancy from respiratory problems, sudden

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infant death syndrome, and gastrointestinal issues; however, these outcomes are not solely due to lack of prenatal care (Association of Maternal & Child Health Programs, 2016).

In contrast to preconception care, prenatal care is one of the most common types of health care in the United States. In 2016, 77 percent of pregnant women initiated prenatal care in the first trimester of pregnancy (Osterman and Martin, 2018). Despite advances in prenatal care, there are significant disparities in coverage and access. Women who are young, are not white, have low educational attainment, or lack private insurance are more likely to have late or inadequate prenatal care (Osterman and Martin, 2018), and issues related to insurance coverage and access persist into postnatal life for the baby.

Pediatric Care

In general, the preponderance of evidence suggests that higher rates of health insurance have led to higher rates of pediatric health care access and better health outcomes for children (Leininger and Levy, 2015). Still, disparities in health care access remain, especially among those who do not have health insurance or have intermittent coverage (Leininger and Levy, 2015). While the number of children who are uninsured is low (~5 percent of children 0–18 years old), this varies from 1 to 11 percent based on state of residence (Kaiser Family Foundation, 2017). Immigrant children, including those who are undocumented, are less likely to be insured than children of the same age with U.S. citizenship (Leininger and Levy, 2015). Some studies suggest that intermittent losses of coverage—which is more common among families with income near the Medicaid eligibility line—decreases children's use of care (Leininger and Levy, 2015). For instance, one study found that even small gaps in insurance coverage decrease a visit to any doctor or a well-child visit by 4 and 9 percent, respectively (Leininger, 2009). Importantly, minority children are disproportionately likely to be from low-income families, particularly African American, AI/AN, and Hispanic children (Jiang et al., 2016). As a consequence of periodic bouts of being uninsured due to poverty (Leininger and Levy, 2015), these groups are less likely to receive routine pediatric care or treatment for medical conditions (Hodgkinson et al., 2017). These disparities in use are reflected in reported health statistics from parents: ethnic minority children are at least four times as likely to be described as in "fair or poor health" by their parents compared to white children. Similarly, children living in poverty are less likely to be described as in "excellent or very good health" by their parents (71 percent) compared to those not living in poverty (87 percent) (Kuo et al., 2012).

Barriers to Accessing Quality Health Care

There are a number of plausible reasons for the existing disparities in access to care, including lack of time and flexibility to seek services, long wait times, or the requirement of multiple appointments (Hodgkinson et al., 2017), and the decline of prenatal care in rural settings (Hung et al., 2017). These factors may make prioritizing pediatric care or treatment untenable for those living in poverty or enduring stress (Hodgkinson et al., 2017; Santiago et al., 2013).

It is important to note that, just as with other aspects of health care, bias and unequal treatment may play a role in women's preventative services and prenatal care (IOM, 2002). Stereotyping and bias on the part of health care providers may increase disparities in preconception care, as has been suggested of prenatal care (Kogan et al., 1994). In particular, messaging that places all the responsibility for the fetus' health on mothers may serve to

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exacerbate stigmas among those already facing significant societal disparities (Greaves et al., 2014; Sue, 2019).

Stigma, bias, or lack of training on the part of health care professionals may also exacerbate pediatric health care disparities (Sue, 2019). Some studies have shown that providers for lower-income families admit to these challenges, including a lack of understanding and training about the effects of poverty on children, confrontation of their own personal prejudice and biases, stigma that comes with working with low-income families and children, and difficulty applying typical diagnostic frameworks to children from low-income families (Smith et al., 2011, 2013).

As outlined in a 2016 National Academies report, *Parenting Matters: Supporting Parents of Children Ages 0–8*, medical providers (and service providers more generally) are met with increasing pressure to provide care in a culturally and linguistically sensitive way. Because of rapid changes in demographics, medical providers may face difficulty in adapting to language barriers or cultural norms, which may in turn reduce the willingness of ethnic minorities and families in poverty to access the health care system (NASEM, 2016a). Medical providers require the knowledge and tools to provide appropriate care to diverse populations in order to address the aforementioned health care disparities. Chapter 5 provides a more in-depth look into opportunities to enhance systems along the continuum of preconception, prenatal, and pediatric health care.

HEALTHY LIVING CONDITIONS

In the report's conceptual model, the committee identifies healthy living conditions as an important domain for study and intervention (see Figure 3-5 for a visual of how this section ties to the conceptual model). For the purposes of this report, healthy living conditions are the social, economic, cultural, and environmental factors that shape the odds for optimal child health and development. Specifically, these include economic security, nutrition and food security, neighborhood conditions, housing, and environmental exposures. These living conditions interact with multiple levels of the conceptual model. For example, adequate nutrition and food security encompass individual behaviors, such as breastfeeding, but they can also be shaped by the larger policy context, such as government programs (e.g., the Supplemental Nutrition Assistance Program [SNAP] and the Special Supplemental Nutrition Program for Women, Infants, and Children [WIC]). The following sections delve into the evidence on how healthy living conditions are critical for early development. Chapter 6 follows this thread and discusses the evidence for interventions (i.e., programs, policies, and systems) that show the most promise for promoting equitable healthy living conditions for all children.

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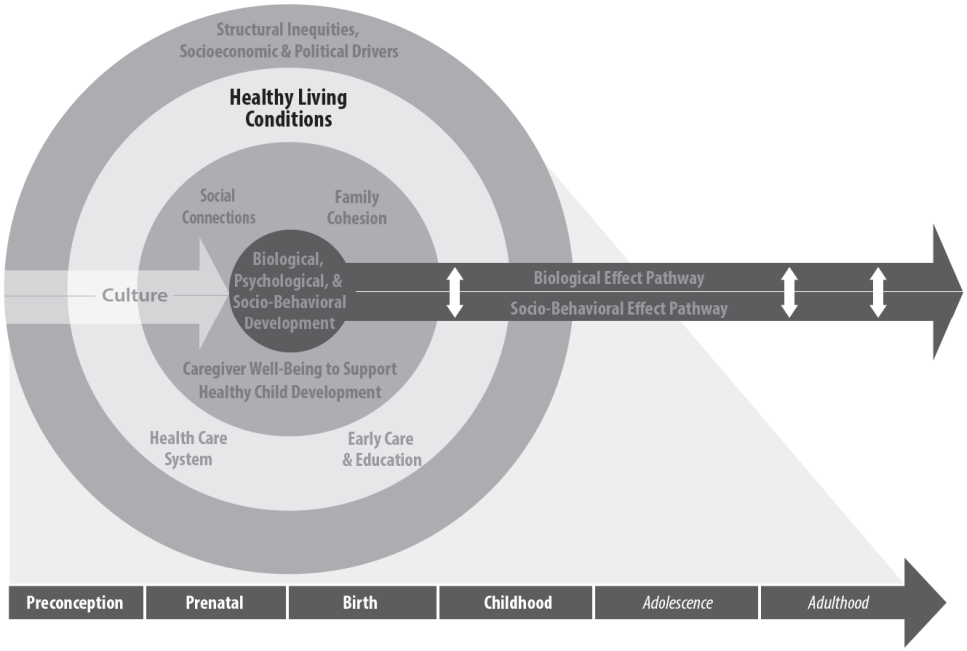

FIGURE 3-5 Leveraging early opportunities to achieve health equity across the life course: A conceptual framework
NOTES: The element of focus in this section is reflected in the second outermost ring in the model: healthy living conditions (i.e., economic security, nutrition and food security, housing, and environmental safety). This element is bolded for emphasis.

ECONOMIC SECURITY

Household Socioeconomic Status

Household socioeconomic status (SES), a construct often measured using income, occupation, and education, is an important social determinant of child health and well-being. For children, SES not only reflects household income but is also indicative of their neighborhood and housing conditions, food security, parental well-being and educational attainment, exposure to violence and other stressors, quality of ECE, and more. This section specifically explores how household SES as it relates to income, poverty, wealth, class, and education affects early developmental outcomes.

Poverty can be defined broadly as the lack of financial resources or material possessions to meet basic needs, such as shelter, food, and clothing. Poverty, income, and wealth are considered to be determinants of health, and there is a robust literature base that ties income and wealth to key health indicators, such as life expectancy, risk of chronic disease, and health-promoting behaviors (e.g., physical activity, healthy eating) (Chetty et al., 2016; NASEM, 2017a; Woolf et al., 2015). Poverty is a multidimensional construct that has many implications for the living conditions and environment that shape health and development.

For children, living in poverty is associated with negative health and developmental outcomes. The existing research suggests at least two pathways by which living in poverty can influence child outcomes (Evans, 2004). The first is through access to material resources and services (e.g., safe housing, nutrition, exercise, health care) (Braveman et al., 2018; Johnson et

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al., 2016; Woolf et al., 2015). The second is through psychosocial stress (e.g., parental job strain, exposure to violence, housing instability, food insecurity) (Braveman et al., 2018; Johnson et al., 2016; Lefmann and Combs-Orme, 2014). The convergence of these factors shapes early living conditions, which can lead to cumulative risk exposure for those living in poverty. Therefore, income and poverty can be conceptualized as upstream determinants of many of the social, economic, and environmental factors discussed in this chapter.

Trends and Disparities

While children under the age of 18 represent 23 percent of the nation's population, they make up 32 percent of people living in poverty (Koball and Jiang, 2018a). In 2016, 44 percent of young children (i.e., under the age of 9) lived in low-income households (i.e., below 200 percent of the federal poverty threshold). In addition, approximately 1 in 5 young children lived in poor households (i.e., below 100 percent of the federal poverty threshold) (Koball and Jiang, 2018b). It is important to note that child poverty, and poverty in general, can be measured in a variety of ways, which has implications for how trends are measured. For example, consumption-based measures of poverty—as opposed to those based on income alone—reflect permanent income and government benefits, and some argue that these are more likely to capture the effects of saving, ownership of goods (e.g., property), and access to credit (Meyer and Sullivan, 2009).

Socioeconomic disparities exist across race, ethnicity, age group, and geographic region. For example, black, AI/AN, and Hispanic children disproportionately live in low-income and poor households; they are also most likely to live in deep poverty (below 50 percent of the federal poverty threshold). In terms of upward economic mobility, there are differences among racial groups as well. In particular, research suggests that black and AI/AN children have the lowest rates of upward mobility, even when controlling for parental income (Chetty et al., 2018). When looking across age groups, younger children (up to 11 years old) are more likely to live in low-income or poor households. However, when stratified by racial or ethnic group, there are stark differences in the proportions (see Table 3-2).

TABLE 3-2 Percentage of Children (ages 0–11 years) living in high poverty areas from 2013–2017, by race and ethnicity

Race or Ethnicity	Percentage of Children (0–11 years) Living in Low-Income or Poor Households
American Indian/Alaska Native	28%
Asian and Pacific Islander	6%
Black or African American	28%
Hispanic or Latino	19%
Multiracial	10%
Non-Hispanic White	4%

NOTES: High poverty areas are defined as census tracts with poverty rates of 30 percent or more. These data are sourced from the Population Reference Bureau analysis of data from the 2013–2017 American Community Survey 5-year data.
SOURCE: Kids Count Data Center, 2019.

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Effects of Income and Poverty

Pregnancy and birth outcomes Blumenshine et al. (2010) conducted a systematic review examining the association between socioeconomic disadvantage and adverse birth outcomes (e.g., outcomes related to birth weight, gestational age, or growth restriction). The authors determined that 91 studies found significant associations between at least one measure of socioeconomic disadvantage and one adverse birth outcome. While differences in birth outcomes based on individual-level socioeconomic factors were almost always observed among non-Hispanic white women, the results were less consistent for African American and Hispanic women. For example, no studies included in this review found birth outcomes to be associated with individual-level socioeconomic factors among foreign-born Hispanic women. However, other studies show associations between neighborhood-level poverty and pregnancy and birth outcomes across racial and ethnic groups. The authors postulate that the cumulative effects of neighborhood conditions (e.g., poor housing, crime, pollution, stress), which are more common for communities of color, may have stronger health impacts than those associated with individual-level income or educational attainment, pointing to the need for community-level interventions (see the section on concentrated disadvantage more discussion of these studies at the neighborhood level).

Childhood outcomes Living in poverty during early childhood can have long-lasting implications for health and well-being. Family income has been found to be associated with risk of child maltreatment, where families with limited economic resources are at a higher risk (Berger et al., 2017; Cancian, 2010). Other research points to allostatic load as an important proximal outcome of living in an environment in the context of poverty. Blair et al. (2011) found that two aspects of the poverty environment were related to salivary cortisol (a measure of the stress response) in infancy: adult exits from the home and perceived economic insufficiency. In terms of neurodevelopmental and cognitive outcomes, the existing literature suggests that poverty affects the developing brain. This includes the structure and function of areas that regulate memory, emotion, cognitive functioning, and language and literacy (Johnson et al., 2016). There is also research that demonstrates the impact of family income on academic achievement, whereby a $1,000 increase in annual income could increase achievement by 5–6 percent of a standard deviation (Duncan et al., 2011; Wolf et al., 2017). Box 3-6 enumerates a few conclusions on the consequences of poverty from a recent report, *A Roadmap to Reducing Child Poverty*.

BOX 3-6
Conclusions from *A Roadmap to Reducing Child Poverty: The Consequences of Child Poverty*

The 2019 report *A Roadmap to Reducing Child Poverty* presents the current evidence on the health and well-being consequences of growing up in childhood poverty. Based on this evidence the authoring committee reached the following conclusions:

- The weight of the causal evidence indicates that income poverty itself causes negative child outcomes, especially when it begins in early childhood and/or persists throughout a large share of a child's life.
- Poverty alleviation can promote children's development, both because of the goods and services that parents can buy for their children and because it may promote a more responsive, less stressful environment in which more positive parent–child interactions can take place.

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See Chapter 6 in this report for a summary of the antipoverty packages evaluated in this report for their potential to reduce child poverty by half in 10 years.

SOURCE: NASEM, 2019

Implications of the Wealth Gap

Income is often the target of many policies and interventions and important for short-term outcomes, but a focus on the more encompassing construct of wealth is important for a health equity agenda because it has long-term and intergenerational implications. That is, income is necessary for building wealth, and wealth could buffer families from periods of income instability and low income. "Wealth, or economic assets accumulated over time, is calculated by subtracting outstanding debts and liabilities from the cash value of currently owned assets—such as houses, land, cars, savings accounts, pension plans, stocks and other financial investments, and businesses. Wealth measured at a single time period may provide a more complete picture than income of a person's economic resources" (NASEM, 2017a, p. 127).

U.S. Census data show that the median household wealth in 2013 was $1,700 for black families, $2,000 for Latinos, and an astonishing $116,800 for whites. The median household wealth in 1983 was $6,800 black families, $4,000 for Latinos, and $102,200 for whites (Asante-Muhammad et al., 2017). Using data from the Panel Study of Income Dynamics (PSID), Williams (2004) found that level of income was not a good indicator for black children's outcomes as it was for white children. Rather, having assets such as stocks or an IRA mattered more for black than for white children. Though focused on older individuals, Zhan and Sherraden (2010) found that household income was more associated with white children's college attendance and graduation, where wealth seemed to be more salient for black and Latino college students. This is also consistent with finding from Williams Shanks (2007) that family income is associated with white children's test scores but not those of black children. Kaushal and Nepomnyaschy (2009) find that although wealth, along with family sociodemographics and parental resources, accounts for the disparities in the black-white and Hispanic-white gap in children's participation in gifted programs, extracurricular activities, and grade retention, black children continued to face high risk of expulsion or suspension from school relative to white children regardless of wealth status. Furthermore, intergenerational transmission of wealth may not be as common, especially for black families (Chetty et al., 2018). While Latino families are moving up in their incomes across generation, black families have substantially lower rates of upward mobility and higher rates of downward mobility (Chetty et al., 2018). That is, black children from higher-income households are likely to become poor as adults compared to their white peers. This indicates that the pathway to wealth accumulation and maintenance differs across racial groups, which has implications for policies that are developed and implemented, especially if they do not address the continued loss of wealth for minority families and communities.

Parental Educational Attainment

Evidence shows that higher levels of caregiver/parental educational attainment are associated with positive child educational outcomes (e.g., school readiness, educational achievement), physical health outcomes (e.g., rates of LBW), and health behaviors (e.g., rates of smoking and binge drinking) (Child Trends, 2015). Research indicates that caregiver/parental

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educational attainment may also serve as a protective factor for outcomes in childhood, adolescence, and adulthood (Dubow et al., 2009).

Higher levels of maternal educational attainment, in particular, have been linked with improved health and well-being for children (Cutler and Lleras-Muney, 2006). Conversely, low levels of maternal educational attainment have been found to be significantly associated with negative infant and child outcomes. Infants whose mothers have not completed high school have been found to have twice the risk of dying before their first birthday compared to those of mothers who have completed college (Egerter et al., 2011b; Mathews and MacDorman, 2007).

Since 1974, the percent of parents (with children ages 6–18) with less than a high school diploma has decreased, while the percent of those with a bachelor's degree or higher has increased (Child Trends, 2015). However, disparities by race and ethnicity have persisted, particularly for African American and Hispanic mothers and fathers (see Figure 3-6).

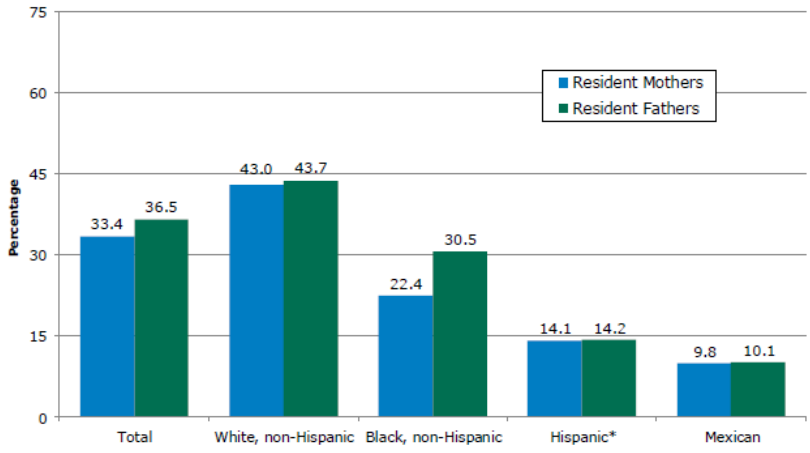

FIGURE 3-6 Percentage of school-age children whose parents have at least a bachelor's degree, by child's race and Hispanic origin and parent's gender, 2015
SOURCE: Child Trends, 2015
NOTES: * Hispanic includes Mexican. Data are sourced from Child Trends analysis of Current Population Survey, March Supplement.

Education is often touted as the gateway to prosperity or ensuring stability for families and children. However, accumulating evidence shows that there are racial differences in the impact of college attainment on income and wealth. Asante-Muhammad and colleagues of Prosperity Now show that "[w]hite families whose head of household holds a high school diploma have nearly enough wealth ($64,200) to be considered middle class (see Figure 3-7). A typical black or Latino family whose head of household has a college degree, however, owns just $37,600 and $32,600, respectively, in wealth. In fact, only black and Latino households at the median with an advanced degree have enough wealth to fit into [their] middle-class definition. By contrast, all white households except those who fail to attain a high school diploma could be considered middle class" (Asante-Muhammad et al., 2017, p. 10). Black and Latino families with college degrees have less than half the wealth of white families with a high school diploma. This differential impact of higher education is also seen with unemployment rates: black individuals with college degrees have a 19, 65, 31, and 31 percent greater chance of unemployment than Latinos, Hispanics, whites, and Asian Americans, respectively (Andolfatto, 2017). This is

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particularly concerning when many blacks and Latinos are less likely to have higher education degrees compared to their white peers. Thus, while education attainment is associated with better child outcomes, more research is needed to better uncover how racial disparities in education coupled with availability of wealth may be associated with the racial disparities in health and education.

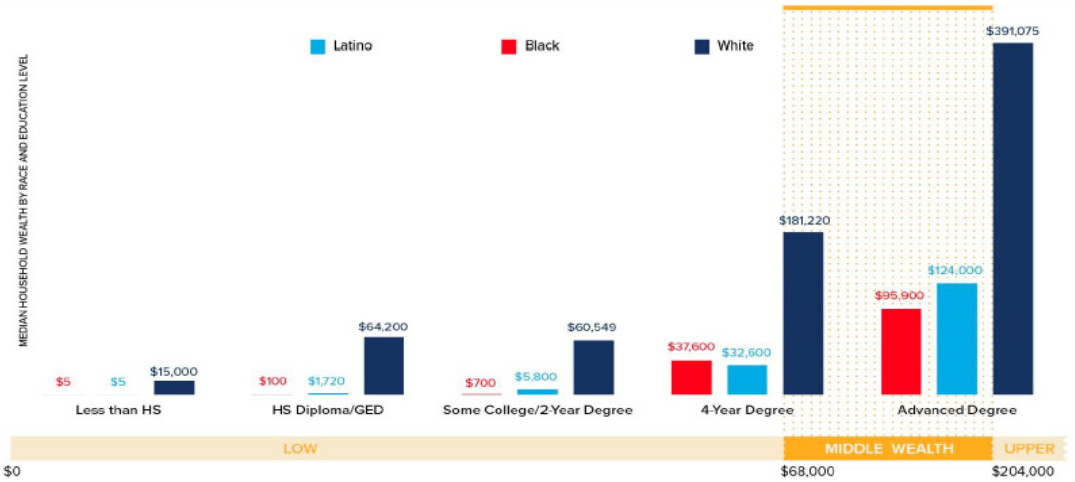

FIGURE 3-7 Having a college education is not enough to guarantee middle-class wealth for many black and Latino families.
NOTES: Wealth figures depicted above exclude durable goods. Data are sourced from the U.S. Census Bureau Survey of Income and Program Participation, 2014.
SOURCE: Asante-Muhammad et al., 2017.

NUTRITION AND FOOD SECURITY

Although access to adequate and nutritious foods affects development across the life course, it is particularly important for the preconception, prenatal, and early childhood periods. Further, the period from conception to 2 years of age (about the first 1,000 days) is especially critical, as the brain undergoes significant developmental changes during this period that may not be possible later in life (Schwarzenberg and Georgieff, 2018). In addition to macronutrients (protein, fat, and glucose), certain micronutrients (zinc, copper, iodine, iron, and selenium) and vitamins and cofactors (vitamin B_6, vitamin B_{12}, vitamin A, vitamin K, folate, and choline) are particularly important for healthy early brain development (American Pregnancy Association, 2017; Cusick and Georgieff, 2016; Georgieff et al., 2015; IOM, 1990; March of Dimes, 2018; Schwarzenberg et al., 2018). Malnutrition can take the form of undernutrition (inadequate intake of macro- and/or micronutrients) and/or overweight/obesity (excessive intake of calories but not necessarily adequate intake of macro- and micronutrients). Inadequate intake of macro- and micronutrients during critical or sensitive periods of development can result in negative developmental effects across the life course, with long-term effects being more likely the earlier such nutrient deficiencies occur (Georgieff et al., 2015; Schwarzenberg and Georgieff, 2018).

Access to healthy and nutritious foods for pregnant mothers is important to ensure healthy birth and developmental outcomes for their infants (Borge et al., 2017; Ramakrishnan et al., 2012; Thorne-Lyman and Fawzi, 2012; Veena et al., 2016; Yakoob and Lo, 2017; Zerfu and

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Ayele, 2013). In an overview of systematic reviews of international interventions to improve nutrition during pregnancy, infants of pregnant women who were provided oral supplements of vitamin A, low-dose calcium, zinc, and multiple micronutrients as well as nutrition education and preventive antimalarials were found to have a lower risk of LBW (da Silva Lopes et al., 2017).

Maternal nutrition during pregnancy can also affect overweight and obesity in children (Lau, 2014; Leonard, 2017). Tabacchi et al. (2007) describe that "the prenatal interactions between genetics and environment play an important role in determining the postnatal phenotype related to obesity" (see Chapter 2 for more detailed information on fetal programming and the interplay of genetics and environment). Evidence also suggests a correlation between maternal obesity and poor infant and birth outcomes (e.g., higher risk of preterm birth, being large for gestational age, fetal defects, perinatal death) (Aviram et al., 2011; Marchi et al., 2015), poor health outcomes in infancy and early childhood, including childhood obesity (Beckhaus et al., 2015; Mameli et al., 2016; Marques et al., 2013; Monasta et al., 2010; Nyaradi et al., 2013; Van Lieshout et al., 2011; Woo Baidal et al., 2016), and negative implications for disease and other poor health outcomes in adulthood (Langley-Evans, 2015; Poston et al., 2011).

Despite clear evidence linking maternal and infant nutrition with positive health and development outcomes, many communities, particularly low-income communities and communities of color, do not have access to affordable, high-quality, nutritious food (Harrington et al., 2009). The lack of access in these communities is compounded by greater exposure to unhealthy foods and beverages, including targeted advertising for such products (NASEM, 2017b). To achieve health equity for all children, it is critically important for all families to have access to healthy and nutritious foods. (For a discussion of the policies and programs in the United States that aim to decrease food insecurity and improve nutrition and healthy eating in the prenatal and early childhood periods (e.g., Temporary Assistance for Needy Families [TANF], SNAP, and WIC, see Chapter 6.) See Box 3-7 for statistics on food insecurity during childhood.

BOX 3-7
Statistics on Child Food Insecurity

In 2016:

- 13 million (one in six) children were food insecure.
- Fifty states and Washington, D. C. were home to food-insecure children. The number of food-insecure children living in Los Angeles and New York City was 800,000.
- The rate of food insecurity among households with children was 16.5 percent.
- The rate of food insecurity among households with children headed by a single woman was 31.6 percent, compared to 22.2 percent among households with children.
- In households headed by non-Hispanic blacks and Hispanics, the rate of food insecurity was 22.5 and 18.5 percent, respectively.
- 85 percent of counties with high child food insecurity were rural.
- In households with income less than 185 percent of the poverty threshold, the rate of food insecurity was 31.6 percent.

SOURCES: Coleman-Jensen et al., 2017; Medlin and McDonald, 2018.

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Breastfeeding

Exclusive breastfeeding[4] offers complete nutrition for infants during the first 6 months of life. Because of this, the World Health Organization (WHO) and the American Academy of Pediatrics (AAP) recommend that infants receive only breastmilk during this time to achieve optimal growth, development, and health. After that, WHO and AAP recommend continued breastfeeding combined with complementary foods up until at least 1 year of age.

Breastfeeding provides important health benefits for mothers and children, but it is important to note methodological difficulties in studying the health impact of breastfeeding. RCTs comparing breastfeeding to formula would be unethical, and observational study designs limit causal inferences. However, this is an area that has been studied for many decades, and there are several areas where the science has converged on the understanding of how breastfeeding promotes healthy outcomes for children and breastfeeding mothers.

There is evidence that exclusively breastfed infants have lower rates of infant mortality, otitis media, and gastrointestinal infections (Bowatte et al., 2015; Kramer and Kakuma, 2012; Sankar et al., 2015). The impact of breastfeeding on the child's health seems to endure, since breastfeeding (and longer breastfeeding duration) is protective against overweight and obesity in children, although reviews and meta-analyses describe such correlations as small (but consistent) and difficult to prove, with research needed to further elucidate the relationship (Arenz et al., 2004; Beyerlein and von Kries, 2011; Dewey, 2003; Harder et al., 2005; Horta et al., 2015; Lodge et al., 2015; Ryan, 2007; Wang et al., 2017). A growing body of research demonstrates important psychological effects of breastfeeding on children, including an impact on brain, cognitive, and socio-emotional development (Krol and Grossmann, 2018).

Breastfeeding may also be protective for mothers, with lower rates of breast and ovarian carcinoma, type 2 diabetes, and postpartum depression (Chowdhury et al., 2015; Dias and Figueiredo, 2015; Hahn-Holbrook et al., 2013; Ip et al., 2007). However, a study by Borra et al. (2015) showed that there is an increased risk of postpartum depression when women who had planned to breastfeed had not gone on to breastfeed, and it concluded that breastfeeding intentions during pregnancy and mothers' mental health during pregnancy both mediated the effect of breastfeeding on maternal depression. This aspect points to the need to provide not only "expert breastfeeding support to women who want to breastfeed but also … compassionate support for women who had intended to breastfeed, but who find themselves unable to."

Breastfeeding Disparities

In the United States, breastfeeding rates as a whole have been increasing (CDC, 2018a). According to the Centers for Disease Control and Prevention (CDC), of infants born in 2015 in the United States, 83.2 percent started to breastfeed, over half (57.6 percent) were breastfeeding at 6 months, and 35.9 percent were breastfeeding at 12 months. However, there are disparities in breastfeeding rates along a number of demographics.

Breastfeeding rates are lowest among African American mothers and mothers living in rural areas. From 2011 to 2015, the percentage of women who initiated breastfeeding was 64.3 percent for African Americans, 81.5 percent for whites, and 81.9 percent for Hispanics (Anstey

[4] Exclusive breastfeeding is defined by WHO as not giving the infant any other food or drink, including water, except needed vitamins and medications (WHO, n.d.-b). WHO recommends that infants start receiving complementary foods at 6 months of age, in addition to breast milk (WHO, n.d.-a).

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et al., 2017). Rates of breastfeeding are also lower in mothers who are young, have lower educational attainment, and have lower incomes (ASTHO, 2017). Roughly 80 percent of higher-income women were still breastfeeding at 1 year compared to 20 percent of lower-income women (ASTHO, 2017).

A multitude of factors impact a woman's decision to start and continue breastfeeding. Social factors, such as unsupportive cultural and social norms, concerns about milk supply, poor family and social support, and unsupportive work and child care arrangements, can make it difficult for many mothers to meet their breastfeeding goals (HHS, 2011). Black women disproportionally experience some of these barriers, such as returning to work soon after a child is born, lack of breastfeeding information from providers, and lack of access to professional breastfeeding support (Johnson, 2015). Black mothers also might not have consistent access to evidence-based supportive practices. A study by Chapman (2012) on hospital support for breastfeeding indicated that facilities in areas with higher proportions of black residents were less likely to fulfill supportive breastfeeding practices than those located in areas with lower percentages of black residents. Health care provider practices may also account for reduced rates of breastfeeding among black mothers, particularly with respect to breastfeeding encouragement and information (Jones et al., 2015). For mothers in rural communities, factors such as poverty and inadequate access to needed maternity and health services may serve as obstacles to breastfeeding (CDC, n.d.-b). Supportive programs and policies for mothers who plan to breastfeed are needed; see Chapter 6 for a discussion of some of these interventions (e.g., paid parental leave) (Pac et al., 2019).

NEIGHBORHOOD AND COMMUNITY CONDITIONS

A community is the place where a child lives, learns, and plays. By this notion, communities serve as the bedrock of health where experiences and behaviors are shaped by the conditions in which a child lives (NASEM, 2017a). The conditions that make up a community or neighborhood are critical for supporting healthy behaviors and providing safe environments for children. This section of the chapter explores the community conditions that can either promote or hinder optimal development during the prenatal through early childhood periods. These include factors that are deeply rooted in historical policies and structural inequities (as was introduced in the beginning of the chapter) and protective factors that can prevent or mitigate the manifestation of negative developmental outcomes among children. The following sections discuss disparities and scientific evidence related to developmental outcomes, including mechanisms across a number of neighborhood-level factors: concentrated disadvantage, physical environment, social environment, and exposure to violence. These factors are all interrelated, and despite the serious effects that some can have on early development, it is important to note that they are not immutable circumstances or effects (NASEM, 2017a).

Concentrated Disadvantage

Trends and Disparities

Beyond poverty at the household level, which was discussed in the previous section, nationally, 12 percent of children live in neighborhoods with concentrated poverty (i.e., Census tracts with poverty rates of 30 percent or more) (Kids Count Data Center, 2019). Concentrated disadvantage and neighborhood-level deprivation are important factors to examine because of

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the cumulative risk that is associated with these neighborhood conditions. Data from the Early Childhood Longitudinal Study indicate that the proportion of kindergarten children living in moderate and high-poverty neighborhoods increased from 1998–2010, which has important implications for school readiness (Wolf et al., 2017).

Neighborhood Disadvantage and Prenatal Through Childhood Outcomes

Individual-, household-, and neighborhood-level economic deprivation show a graded relationship to health outcomes (Pickett and Pearl, 2001). The Whitehall (Kumari et al., 2004) and GLOBE studies (van Lenthe et al., 2004) both offer strong evidence that individual and neighborhood-level economic conditions each contribute to health outcomes. Twin studies offer other compelling evidence; one such study of 3,738 same-sex twin pairs found that neighborhood deprivation had a significant within-pair association with depression after adjusting for individual socioeconomic indicators and other factors (Cohen-Cline et al., 2018). There is a strong body of associational evidence linking neighborhood economic environments to depression (Paczkowski and Galea, 2010) and other chronic diseases (Pickett and Pearl, 2001). Box 3-8 describes asthma disparities as an example of a health outcome in the context of neighborhood disadvantage.

Neighborhood-level economic deprivation has also been widely associated with pregnancy and birth outcomes, including reduced use of prenatal care (Schempf et al., 2009) and an increased risk of a range of adverse birth outcomes (Messer et al., 2006; O'Campo et al., 2008; Zeka et al., 2008). Studies have linked neighborhood-level economic deprivation indicators with birth weight (Buka et al., 2003; Morenoff, 2003; Pearl et al., 2001; Rich-Edwards et al., 2003; Subramanian et al., 2006), preterm birth (Ahern et al., 2003; Kaufman et al., 2003; O'Campo et al., 2008), and gestational age and fetal growth (Farley et al., 2006). However, this is not to suggest that neighborhood economic deprivation is the cause of poor health—Jokela's (2014, 2015) analyses of data from the British Household Panel Survey suggests that the observed associations may in fact reflect more fundamental social inequalities that shape health and hinder some people's ability to move to less deprived neighborhoods. Research has consistently shown an association between neighborhood deprivation and health, but a firm causal connection has not been established.

While there is evidence connecting neighborhood-level poverty and birth outcomes, the mechanism of this link remains unclear; one theory is allostatic load due to stress (Lefmann and Combs-Orme, 2014; Wallace et al., 2013). Wallace and colleagues (2013) found that while African American mothers who resided in poor neighborhoods had higher allostatic load than white mothers, allostatic load was not associated with preterm birth or LBW after accounting for race, neighborhood SES, maternal education, maternal age at time of birth, and smoking during pregnancy, among other things. This lack of link between allostatic load and pregnancy was also found in another study (Morrison et al., 2013). Nevertheless, African American women living in low poverty neighborhoods were more than five times as likely to have an LBW infant and to have given birth preterm than white women in low poverty neighborhoods. This racial disparity link between neighborhood disadvantage and LBW and preterm birth was also found in a systematic review and meta-analysis of population-based studies (Ncube et al., 2016). Margerison-Zilko and colleagues (2015) stress that it is not just about living in a poor neighborhood that is related to preterm birth but the length of time spent in a low-resourced, high-poverty neighborhood. In their systemic review and meta-analysis, examining the link between residential segregation and adverse birth outcomes, Mehra et al. (2017) found that black

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mothers living in segregated neighborhoods have the highest risk of LBW and multiple adverse birth outcomes.

BOX 3-8
Concentrated Poverty Case Study: Asthma

Childhood asthma disparities by neighborhoods are one example of the importance of neighborhood conditions. There are significant disparities in the rates of asthma between African American and white children. In 2016, the prevalence of asthma was 15.7 percent in African American children under 18 years old and 7.1 percent in white children (CDC, 2018b); the prevalence among African American children has remained about double that of white children since 2007 (Akinbami et al., 2009, 2014). Some evidence shows that this disparity in asthma prevalence may in fact stem from neighborhood conditions. For example, a study of children living in New Jersey between 2006 and 2010 found that in zip codes in which the highest fraction of the sample was African American, higher rates of asthma among LBW children were present regardless of race. As the authors state, "this distinction is important because unlike race, it is possible to change neighborhoods either by finding and remediating the hazards that are causing higher asthma prevalence or by helping vulnerable children to move" (Alexander and Currie, 2017). Furthermore, it is difficult to parse the effects of specific neighborhood factors (e.g., poor housing quality or exposure to environmental pollutants) in studies that examine zip codes or Census tracts because harmful neighborhood conditions or factors tend to cluster in the same areas.

Physical Environment

A number of studies have assessed how various physical characteristics of neighborhoods—the "built environment"—may shape health or birth outcomes, either through direct exposure or by limiting access to or the effectiveness of traditional health care strategies. Green infrastructure, such as vegetation or tree canopy, has been associated with reduced depression rates (Cohen-Cline et al., 2018; Fan et al., 2011), which may in turn impact birth outcomes (Accortt et al., 2015). The presence of active living attributes, such as walkability, mixed land use, and active transit options, has been associated with increased physical activity and reduced obesity in general (Feng et al., 2010; Saelens et al., 2003a,b). However, the evidence is unclear on whether this ultimately affects birth outcomes: the one study assessing built environments in terms of adverse birth outcomes did not yield clear evidence of impact (Vinikoor-Imler et al., 2011). It is possible that the condition of early built environments could be associated with "sleeper effects," whereby outcomes are observed later in life.

The importance of food security and nutrition has long been identified as a key determinant of prenatal and postnatal health outcomes. However, the *food environment* people live in—the accessibility of healthy food options in their neighborhoods—is an additional contextual factor that may shape their prenatal outcomes by facilitating or limiting their ability to follow prenatal nutritional advice. A few studies have found positive effects of healthy food environments on reducing obesity (Auchincloss et al., 2013), hypertension (Kaiser et al., 2016), and type 2 diabetes (Christine et al., 2015). However, a number of other studies of the food environment's impact on health outcomes have reported null results, leading to an overall mixed body of evidence (Cobb et al., 2015), and few such studies have moved beyond overall health outcomes to assess specific adverse birth outcomes. Two studies have found associational evidence that women living in neighborhoods with a high concentration of convenience stores versus healthier food sources (Ma et al., 2016) or in areas lacking proximity to supermarkets

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(Lane et al., 2008) had increased risk of some adverse birth outcomes. However, other studies have not found evidence that neighborhood food environments significantly impacted birth outcomes (Farley et al., 2006). Overall, the evidence in this area remains preliminary and somewhat mixed.

Social Neighborhood Environment

The social environment people live in can be as important as their physical environment. Social support has been hypothesized to benefit birth outcomes both by moderating or buffering the stress of pregnancy (or of other contextual challenges to a healthy pregnancy) and by exerting a direct positive influence on the health of a prospective mother (Cohen and Wills, 1985). Social isolation has been associated with whether women receive adequate prenatal care (Heaman et al., 2018) and with late antenatal presentation (i.e., delayed access to antenatal care) and poor fetal outcomes (Kapaya et al., 2015), while better social support has been associated with higher birth weight and Apgar scores at birth (Collins et al., 1993), reduced risk of maternal depression (Uebelacker et al., 2013), and improved fetal growth (Hoffman and Hatch, 1996). However, studies of interventions aimed at improving social support for pregnant women have not been uniformly shown to be effective (Lu et al., 2005), and the mechanisms by which social support may impact birth outcomes remains somewhat unclear.

Social support can refer to personal networks but also to neighborhood environmental context. These "neighborhood cohesion" measures have been associated with physical and mental health and well-being outcomes in some studies (Bures, 2003; Fone et al., 2007; Hutchinson et al., 2009), perhaps acting to help protect individuals from the harmful or stressful effects of neighborhood deprivation (Robinette et al., 2013). They have also been specifically associated with improved birth weight outcomes in several studies (Buka et al., 2003; Morenoff, 2003).

Exposure to Violence

Childhood exposure to violence can take on many forms, including primary exposure or direct victimization (e.g., maltreatment or abuse by an adult caregiver or bullying by peers) and secondary exposure (e.g., witnessing IPV between parents or community violence) (Gilbert et al., 2009). Regardless of the mechanism, the body of evidence on early exposure to violence shows that this is harmful to children's health and well-being (Egerter et al., 2011a; Moffitt and Tank, 2013; Shonkoff et al., 2012). Data also show that specific subgroups are at higher risk of being exposed to violence based on certain demographics, such as race and ethnicity, gender, SES, and parental education level. Research also indicates that children who are exposed to one incident of violent victimization are likely to be exposed to violence again, including other forms (Finkelhor et al., 2007, 2009, 2015).

Trends and Disparities

Data from the National Survey of Children's Exposure to Violence (2013–2014) including a nationally representative sample of children (n = 4,000; 0–17 years old) showed that 37.3 percent had experienced any physical assault in the past year, 51.4 percent had experienced physical assault in their lifetime, and 24.5 percent and 38.3 percent witnessed violence in the past year and in their lifetime, respectively (Finkelhor et al., 2015). For women, disparities in violence have been documented. For example, for 2003–2014, non-Hispanic black and AI/AN

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women experienced the highest homicide rates nationally, at 4.4 and 4.3 per 100,000 respectively (Petrosky, 2017). Furthermore, more than half of all female homicides (55.3 percent) were related to IPV, which is considered an ACE for children who are exposed to it. (See the section at the end of this chapter on Accumulation of Risk for more on ACEs.)

Neighborhood Violence and Child Well-Being

Research has linked neighborhood violence with negative health and well-being outcomes across the continuum of early development. For example, there is evidence to suggest that high rates of youth violence (e.g., assault, kidnapping, homicide, robbery, larceny, destruction of property) measured at the Census tract level are associated with elevated odds of preterm birth (Masho et al., 2017). The presence of community violence has also been shown to have a direct negative relationship with children's self-regulatory behavior and cognitive performance indicators (Sharkey, 2010; Sharkey et al., 2012). A systematic review of studies examined exposure to community violence and health outcomes in youth (0–18 years old) (Wright et al., 2017). Across the 28 studies included, the most consistent finding was that early exposure to community violence was associated with elevated blood pressure, asthma, and sleep disturbance, which the authors note can contribute to other health-related problems when it persists over an extended period. Among the studies in this review, one found that the level of collective efficacy[5] in a community moderated the effect of exposure to community violence on increased asthma risk, particularly among African Americans (Sternthal et al., 2010). In addition, social support has been identified as critical buffer for children against the negative effects of violence (Margolin and Gordis, 2004).

More recent studies have begun to examine the underlying pathways through which violence specifically influences health outcomes (Finegood et al., 2017; Theall et al., 2017). For example, a 2012–2013 study of African American children in New Orleans found that reports of domestic violence and rates of violent crime within a 500 meter radius of a child's home were associated with decreases in mean telomere length and the likelihood of reducing cortisol levels after a stress reactivity test (Theall et al., 2017). Some research suggests that underlying disadvantage (e.g., poverty, high unemployment, lower levels of educational attainment) associated with neighborhood violence partially explains the relationship between exposure to violence and negative cognitive and behavioral outcomes (Aizer, 2008). Egerter et al. (2011a) postulate that social and economic disadvantage increases the likelihood of exposure to violence, which can diminish levels of trust, social cohesion, and perceptions of safety in a community.

Limitations of the Evidence on Neighborhood Effects

Despite these associations, the relationship between neighborhood characteristics and health outcomes is complex, making causal connections difficult to establish in this literature. Associations may vary based on individual-level characteristics, such as age or sex (Meijer et al., 2013), and may be quite sensitive to the inclusion of other covariates in the models (Auchincloss et al., 2013; Blair et al., 2014; Paczkowski and Galea, 2010). As stated previously, it is also difficult to separate out the effects of factors that tend to cluster together (i.e., concentrated disadvantage) or are systematically present for specific populations, such as poor housing

[5] Collective efficacy can be defined as the level of trust among residents and their perceived willingness to engage in collective action (Sternthal et al., 2010).

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quality, poverty, and exposure to environmental toxicants. Results also vary widely depending on how a specific health domain is measured or how a neighborhood is defined (Paczkowski and Galea, 2010), and many studies capture only limited domains of neighborhood characteristics, leaving unanswered the question of which characteristics are most important or how those characteristics interplay with one another to shape outcomes. Last, few studies establish a causal connection between these endogenous neighborhood characteristics and health (Dohrenwend et al., 1992; Jokela, 2014; Ritsher et al., 2001).

HOUSING

Housing affordability, stability, and quality are well studied and documented SDOH across the life-span. This is also true, of course, during the prenatal and early childhood years. Quality, stability, affordability, and loss of housing in the prenatal and childhood periods can have significant effects on health, cognition, and neurodevelopment. As described previously, residential segregation has had persistent effects on communities, including limiting the availability of safe and affordable housing for some. In this section, housing instability, quality of housing, and affordability will be addressed in turn.

Housing Instability

According to a 2017 report from HUD, there were more than 550,000 individuals considered to be homeless on any given night in the United States in that year. More than 1.4 million people were homeless in sheltered locations (e.g., emergency shelter, transitional housing, safe haven) at some point in 2017 (Henry et al., 2017). Moreover, the risk of homelessness is high for many individuals: according to recent studies, nearly 3 million individuals renting are at risk of eviction, which indicates high rates of potential instability (Sandel and Desmond, 2017). The data indicate a vast imbalance in equitable access to stable housing, such that those experiencing homelessness are disproportionately likely to be black. Only 13 percent of the general population is black, compared to more than 40 percent of the homeless population (Olivet et al., 2018).

While housing instability and homelessness are notoriously difficult to measure accurately, families with children are estimated to make up 33 percent of homeless populations (Henry et al., 2017). A 2008 cross-sectional study of 12,746 children from low-income families suggests that nearly 30 percent of children from these households are in households with housing instability (Ma et al., 2008). This include frequent moves, difficulty paying bills or rent, spending a large proportion of income on rent, being evicted, or living in overcrowded conditions.

Housing instability—either chronic or intermittent homelessness—has well-documented negative health effects on developing children. A 2014 study of nearly 10,000 women found that mothers who were homeless while pregnant were more likely to deliver LBW children compared to those who were homeless after delivery (Cutts et al., 2015). Following birth, housing instability is associated with a wide range of cognitive, emotional, and behavioral outcomes. In a nationally representative, longitudinal study of children 2–21 years old, Coley and colleagues (2013) found that residential instability was associated with negative internalizing and externalizing behaviors—particularly for children with multiple housing moves. It was hypothesized that the negative effects of housing instability are due to changes in social

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structures, schooling, and the emotional health of the family (Coley et al., 2013). In a 2008 study, housing instability was associated with delays in seeking medical care or medications and increased emergency department (ED) visits among children from low-income families (Ma et al., 2008).

More broadly, it has been shown that homeless children—or children with frequent moves—are at increased risk for conduct problems at school, social difficulties, and low academic achievement (Buckner, 2008; Miller, 2011; NRC and IOM, 2010). These effects appear to be independent of low-income status; data on academic achievement and improvement on standardized tests suggest that students with frequent moves perform consistently worse in reading and math compared to other low-income students without frequent moves or housing instability (Masten et al., 2014; Obradović et al., 2009). Overall, the picture of housing instability is one of pervasive and sometimes long-term outcomes on children's health and well-being.

Housing Affordability

Housing is widely considered "affordable" if less than 30 percent of pretax income is required to own or rent (Schwartz and Wilson, n.d.; The Pew Charitable Trusts, 2018). By this definition, approximately 31.8 million U.S. households are living in housing that is unaffordable (Joint Center for Housing Studies of Harvard University, 2018). This population is generally low income. In 2015, 84 percent of renters in the lowest quintile of income were living in unaffordable housing; 70 percent of this group spent more than 50 percent of their income on housing costs (Fenelon et al., 2018). In 2016, 47 percent of all renters, and more than three quarters of families earning between $15,000 and $30,000, had unaffordable housing (Joint Center for Housing Studies of Harvard University, 2017).

According to the most recent data available, racial/ethnic minorities and single-parent families disproportionately experience high housing cost burden, with 55 and 54 percent of black and Hispanic renters in unaffordable housing, respectively, compared to 43 percent of white renters. Further, 63 percent of single-parent renters are in unaffordable homes, compared to 39 percent for married or partnered parents, and finding affordable housing in large metropolitan areas is particularly challenging (Joint Center for Housing Studies of Harvard University, 2017). Across the United States, there are only 37 available and affordable rentals for every 100 low-income households that require housing (National Low Income Housing Coalition, 2018).

There is mixed evidence for housing affordability on health and well-being outcomes among children. A 2010 critical review of housing characteristics and child development found that affordability was not well studied and was only marginally associated with children's health (Leventhal and Newman, 2010). These findings suggest that affordability, per se, is not the primary concern; rather, the amount invested in a child's health and enrichment may be a more robust predictor of developmental trajectories (Newman and Holupka, 2016). This includes spending in other domains that are important for optimal child development (Newman and Holupka, 2016), such as quality child care (Campbell et al., 2014), schools, and neighborhoods (Beyers et al., 2003; Knopf et al., 2016; Theall et al., 2017; Xue et al., 2005). Research suggests that a high housing cost burden is associated with lower spending on child enrichments (Newman and Holupka, 2016) and elevated maternal stress (Warren and Font, 2015), but its impact on maternal depression or anxiety is unclear (Harkness and Newman, 2005; Newman and Holupka, 2014). Families in unaffordable housing are more likely to miss rent payments (Warren, 2018) and therefore may be forced to move (Crowley, 2003; Desmond and Shollenberger, 2015). At the same time, research also suggests that by allocating a higher

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fraction of household income to housing, families may gain access to higher-quality homes (Kull and Coley, 2014), neighborhoods (Acevedo-Garcia et al., 2016b), and/or schools, and these contextual advantages of high housing costs may outweigh any negatives (Kull and Coley, 2014).

While high housing costs can be problematic, very low housing costs for low-income families may also present risks for healthy child development because they can be indicative of substandard or poor-quality housing conditions (Newman and Holupka, 2014, 2016). Accordingly, for low-income families, high or low housing cost burden could be harmful (Newman, 2008). Although prior research suggests risk associated with high and low housing cost burden (Newman and Holupka, 2014, 2016), we focus our discussion on high housing cost burden because this is the main problem for low-income families.

Economic and developmental theories used in child research (Becker and Tomes, 1986; Garner and Shonkoff, 2012; Shonkoff et al., 2012) and the scientific literature on the influence of poverty and child development (Brooks-Gunn and Duncan, 1997; Duncan, 2012; Duncan et al., 1994) suggest that understanding the role of housing affordability for child development requires considering (a) the multiple and interacting contexts in which children develop into adults (Bronfenbrenner, 1979; Bronfenbrenner and Evans, 2000; Bronfenbrenner and Morris, 2007), such as the home, school, and neighborhood; (b) the developmental timing and duration of unaffordable housing across childhood (Ben-Schlomo and Kuh, 2002; Ben-Schlomo et al., 2014; Brooks-Gunn and Duncan, 1997; Coley et al., 2013; Harkness and Newman, 2005; Hicks et al., 2018; Slopen et al., 2010); and (c) unequal health or educational benefits associated with a given level of parental income or education across racial/ethnic groups (Acevedo-Garcia et al., 2005; Assari, 2018; Assari et al., 2018; Chen et al., 2006; Shervin, 2018; Williams et al., 2010). For example, research suggests that youth are particularly influenced by socioeconomic variables during early childhood (Duncan et al., 1998; Ziol-Guest et al., 2009). Other research suggests that black and Hispanic children may be disproportionately harmed by unaffordable housing because investments in housing costs may have smaller returns for minority families (i.e., these investments may have diminished returns to school and neighborhood quality for minority children, relative to those for white children) (Assari, 2018; Shervin, 2018; Williams et al., 2010).

Housing Quality and Crowding

Families living in poverty are less likely to be able to afford high-quality housing, a factor clearly associated with child health (Rauh et al., 2008; Sandel et al., 2004). Although causal relationships are not always clear because most research is observational (Leventhal and Newman, 2010), many studies have explored the connection between child health and conditions in old and inadequately maintained residences, such as dampness, disrepair, poor ventilation, and lead paint. For instance, a study of low-income families waiting for housing vouchers found that poor-quality housing was associated with parents rating their child's health as "fair" or "poor" (Sharfstein et al., 2001). Overall, poor housing quality is consistently associated with "worse emotional and behavioral functioning and lower cognitive skills" (Coley et al., 2013). In addition, another common feature of low-income housing is overcrowding, which has been demonstrated to negatively impact children's well-being (Solari and Mare, 2012). The three subsections below focus on the relationship between child health and (1) exposure to allergens that cause asthma, (2) exposure to lead, and (3) overcrowding.

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Allergens and Asthma

According to the National Health Interview Survey, more than 6 million (8.4 percent) of U.S. children under the age of 18 had asthma in 2017 (Black and Benson, 2018). While the management of childhood asthma improved from 2001–2016, its prevalence remained relatively consistent (Zahran et al., 2018). This warrants an increased focus on the root causes and triggers, and especially on the housing conditions of children who are disparately impacted.

Poor housing conditions can lead to increased risk of asthma. Several studies have found connections between asthma and exposure to allergens, such as cockroaches (Rauh et al., 2008; Wu and Takaro, 2007). Other research has found a causal association between dust mite exposure and asthma (IOM, 2000). Dampness and mold are also associated with asthma and other respiratory conditions (Rauh et al., 2008; Wu and Takaro, 2007). One study found that density in housing code violations was associated with a greater likelihood of a revisit to an ED or a hospital readmission (Beck et al., 2014). Another study found that children living in improved public housing were less likely to experience repeat ED and urgent care visits for conditions unrelated to an initial visit (Kersten et al., 2014). Some researchers have estimated that eliminating specific asthma triggers in older children could reduce asthma prevalence by over 40 percent (Lanphear et al., 2001). Previous National Academies studies have explored these connections in detail (IOM, 2000, 2004).

Exposure to Lead

While there is no known safe amount of lead exposure, in 2012, CDC established a reference level of 5 micrograms per deciliter to identify children who have been exposed (Wheeler and Brown, 2013). The percentage of children estimated to have blood lead levels (BLLs) greater than this amount has decreased significantly over the past decades, from an estimated 8.6 percent for 1999–2002 to 2.6 percent for 2007–2010 (Wheeler and Brown, 2013). These reductions are attributed to the phased elimination of lead from gasoline from the mid-1970s to the mid-1980s (President's Task Force on Environmental Health Risks and Safety Risks to Children, 2018) and from paint in 1978 (Markowitz and Rosner, 2014). Lead paint was banned in 1978, but older homes, primarily in the Northeastern and Midwestern parts of the country, have a higher prevalence of lead paint (Rauh et al., 2008). When these homes are poorly maintained, this paint can chip or peel, and a child may ingest it or breathe in dust, resulting in elevated BLLs (Muller et al., 2018). In addition, lead remains in the soil in areas that were in close proximity to certain manufacturing sites (Muller et al., 2018).

Despite some improvements over time, current estimates suggest that more than 500,000 children ages 1–5 have blood lead levels (BLLs) over the reference level. Young black children are more than twice as likely (5.6 percent) than white children (2.4 percent) to have a BLL greater than 5 micrograms (Wheeler and Brown, 2013). Children who are living in poverty or who have Medicaid coverage are up to eight time more likely to have elevated BLLs (Wheeler and Brown, 2013).

Young children are particularly susceptible to absorption of lead (Lidsky and Schneider, 2003). Reviews of multiple studies have associated exposure to lead—even in low amounts—with IQ, test scores, impulsivity, and Attention Deficit Hyperactivity Disorder (ADHD) (Leventhal and Newman, 2010; Muller et al., 2018; Rauh et al., 2008). Some studies have linked perinatal or childhood lead exposure to adolescent BMI, impulsivity, anxiety, and depression (Kim and Williams, 2017; Winter and Sampson, 2017).

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Overcrowding

The U.S. Census collects information on the number of occupants per unit and defines crowding at more than one occupant per room within a single unit. By this metric, 4.3 percent of houses in the United States are overcrowded (U.S. Census Bureau, 2015). Units are most likely to be overcrowded in regions with disproportionately high rental prices. According to the U.S. Census from 2000, for instance, Los Angeles had more than four times the number of crowded units per capita compared to the United States as a whole (Solari and Mare, 2012).

Research has shown that living in a crowded home may have several detrimental effects on child development, although the overall picture is decidedly mixed. In 2012, Solari and Mare found that living in a crowded home negatively impacts children's academic achievement, externalizing behaviors, and physical health, even when controlling for several dimensions of SES. While this research is cross-sectional, making causal interpretations difficult, it does suggest several mechanisms by which crowding could impact child development. One hypothesis is that living in a crowded home reduces time spent studying and reading for school, thereby decreasing comprehension and test scores. Another is that overcrowding may disrupt a child's sleep, leading to mood changes and difficulty concentrating on academics. A final hypothesis is that overcrowding increases the chances that children will become sick, preventing them from attending school or concentrating while there. All of this, coupled with a lack of privacy, could impact a child's sociability and increase stress and behavioral problems. Each of these hypotheses need to be studied in more detail.

Other studies have shown mixed results. A 2010 study found that crowding during infancy was linked to negative cognitive and social skills, mediated primarily by mothers being less responsive to children in crowded homes (Evans et al., 2010). On the other hand, in 2012, Martin and colleagues found that overcrowding at age 2 was not a significant predictor of age 5 vocabulary, attention, or control of effortful behavior when accounting for other measures of household chaos (e.g., noise, family instability, lack of routine, and television watching) (Martin et al., 2012). This particular factor of housing requires more research, particularly because children growing up in crowded homes are more likely than their peers to end up in similar situations as adults (Leventhal and Newman, 2010), thereby "contributing to the intergenerational transmission of social inequality" (Solari and Mare, 2012, p. 3)

ENVIRONMENTAL EXPOSURES

While environmental exposures to toxicants and limited access to green spaces and healthy living areas have been found to be associated with poor health outcomes at multiple stages throughout the life course, the preconception, prenatal, neonatal, and early childhood periods represent several key developmental phases when humans may be particularly vulnerable to toxic environmental exposures. To best protect women, men, and children, it is important to understand how these substances can affect health and the methods to limit toxic exposures at these critical phases in development.

Environmental Toxicants

There are thousands of potential environmental toxicants that may be transmitted through the air we breathe, the water we use, and the soil and consumer products with which we, our food, and our water come into contact (Giudice et al., 2017). Many of these substances occur

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naturally in the environment (e.g., arsenic, radon, etc.), and many more are released through human-based processes (e.g., heavy metals, chemicals from plastic production and degradation, and particulates), such as through manufacturing by-products, fossil fuel use, mining, and disposal of waste (Di Renzo et al., 2015).

Several of these compounds are associated with poor preconception and prenatal outcomes. In particular, lead, methyl mercury, polybrominated biphenyls (PBBs), polychlorinated biphenyls (PCBs), and the pesticide chlorpyrifos have been shown to result in developmental neurotoxicity, leading experts in fetal development, obstetrics and gynecology, and fertility to conclude that there is no "safe" exposure. Further, endocrine disrupters, such as the pesticides and herbicides atrazine, glyphosate, and chlorpyrifos, are associated with cancer and neurodevelopmental disorders (Diamanti-Kandarakis et al., 2009; Gore et al., 2015). Animal models align with these human studies; endocrine disruptors in animal studies indicate that in utero exposure leads to obesity (Manikkam et al., 2013), abnormal sexual anatomy and sexual performance (Vandenbergh, 2004), and abnormal neuronal migration (Nakamura et al., 2012), similar to that seen in children with autism. Moreover, chemicals in personal products, such as phthalates, bisphenol A (BPA), and fluoroacetic acid, are related to reproductive outcomes (Giudice et al., 2017). However, hundreds of these chemicals remain untested, and researchers estimate that the markedly rapid increase in noncommunicable diseases, particularly those related to endocrine disruption, exceeds the expected natural occurrence of change due to evolution alone. Researchers conclude that these chemicals, on their own or in combination, are likely the drivers of this change (Di Renzo et al., 2015).

In addition, several substances are associated with poor early childhood development outcomes and long-term health problems. For example, particulate matter, ozone, and carbon monoxide are associated with poor child health outcomes (Giudice et al., 2017), including impaired lung function and neurodevelopment, as well as exacerbation of existing issues, such as asthma (Webb et al., 2016). Lead, mercury, and arsenic are associated with cognitive disorders and kidney disease among children (Weidemann et al., 2016). Certain groups in the United States are more likely than others to come into contact with these environmental toxicants or to be unable to access healthy living areas and green space. In particular, certain racial and ethnic groups and families and children living in poverty are disproportionally more likely to experience exposures and see poor health outcomes related to contact with environmental toxicants (ACOG Committee Opinion, 2013; Oberg et al., 2016). For example, a Chicago longitudinal study documented marked, persistent disparities in elevated BLLs of African American children compared to white children (Sampson and Winter, 2016). African American women exposed to higher levels of $PM_{2.5}$ were more likely to report higher depression severity during pregnancy and postpartum (Sheffield et al., 2018). In addition, African Americans and other people of color experienced higher $PM_{2.5}$ exposure compared to whites, as did those living in poverty (Mikati et al., 2018).

Specifically, the research literature has documented strong evidence of adverse pregnancy outcomes, such as increased risk of preterm delivery, related to exposure to certain metals, including lead, which may be transmitted to the fetus through maternal BLLs (Taylor et al., 2015). This study also documented an increased risk of reduced birth weight, smaller head circumference, and reduced crown to heel infant length related to exposure to lead from maternal BLLs. Similarly, higher maternal blood cadmium levels are associated with reduced birth weight, smaller head circumference, and reduced crown to heel length (Taylor et al., 2016), particularly for female infants. Mercury has long been shown to have negative effects on the

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developing fetus. One study demonstrated that levels greater than or equal to 10 ppm of mercury can be harmful to fetal development, causing brain pathologies that result in Minamata disease, microencephaly, seizures, intellectual disability, and stillbirth, as evidenced by methylmercury poisoning in Japan and Iraq, from industrial waste and the use of an imported grain treated with a fungicide containing methylmercury (Cox et al., 1989; Kalter, 2003).

Higher exposure to phthalates, chemicals added to plastics to increase flexibility and make other physical changes for use in consumer products, is associated with an increased risk of preterm delivery and spontaneous preterm delivery (Ferguson et al., 2014). In Swedish case-control study, higher levels of butyl benzyl phthalate (BBzP), bis(2-ethylhexyl) phthalate (DEHP), and polyvinyl chloride (PVC) were more likely to be found in children with a diagnosis of rhinitis and eczema; asthma; and asthma; eczema, rhinitis, and eczema, respectively (Bornehag et al., 2004). A Bulgarian case-control study found evidence of a higher risk of asthma among cases with high levels of DEHP (Kolarik et al., 2008). The presence of urinary concentrations of phthalate metabolites, particularly diethyl phthalate (DEP) and BBzP, was associated with a 6.6 percent and 8.7 percent increases, respectively, of fractional nitric oxide, a biomarker of airway inflammation. The BBzP association with fractional nitric oxide levels was particularly strong for children who wheeze (Just et al., 2012).

Phenols, particularly BPA, have been found to be associated with endocrine and neurodevelopmental problems. Prenatal BPA exposure among boys is significantly associated with a risk of decreased anogenital distance, a marker of testosterone production (Miao et al., 2011). The odds of preterm birth among infants with mothers who had higher plasma BPA levels was between 4.12 and 4.78 times higher than those without high BPA levels (Behnia et al., 2016). Each tenfold increase in maternal urinary BPA level is associated with an increased risk of anxious and depressed behavior and poorer emotional control among 3-year-old girls (Braun et al., 2017). There is evidence that exposure to composite fillings that leach BPA can have detrimental effects on early neurodevelopment, including a heightened risk of internalizing, problem, and delinquent behaviors in children, along with increased risk of anxiety, depression, social stress, and problems with interpersonal relationships (Bellinger et al., 2008). Heightened risk of asthma is associated with prenatal exposure to BPA, depending on the phase of fetal development in which the exposure took place (Spanier et al., 2012). Prenatal exposure among male infants is associated with a higher risk of high BMI, increased waist circumference, higher fat mass, and overweight/obesity (Gascon et al., 2015; Midoro-Horiuti et al., 2010; Spanier et al., 2012). Higher maternal blood BPA levels are associated with lower TSH levels in male infants, suggesting poorer thyroid function (Chevrier et al., 2013; Harley et al., 2013). Higher maternal blood levels of 2,4-DCP, 2,5-DCP, and triclosan (phenols other than BPA) are associated with lower weight among male infants. A higher maternal blood level of bisphenol-S is associated with lower weight among female infants (Ferguson et al., 2018). In the same study, Ferguson et al. (2018) also demonstrated that higher maternal blood levels of parabens and benxophenone-3 are associated with lower weight among male infants.

Among young children, postnatal exposure to BPA is associated with a heightened risk of asthma, depending on the phase of child development in which the exposure took place (Donohue et al., 2013). In studies of African American, Dominican American, and Chinese children, high BPA levels were associated with a higher risk of obesity (Donohue et al., 2013; Wang et al., 2012). Increased urinary BPA levels among children is also associated with an increased risk of albuminuria, which is an indicator of future health problems, including type 2 diabetes and cardiovascular disease (CVD) (Trasande et al., 2013).

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Limited and Developing Evidence

A number of studies have established a relationship between poor health or birth outcomes and living in places with close proximity to potential pollution challenges. Air quality through the gestational period has been associated with increased risk of LBW, preterm birth, and infant death (Padula et al., 2012). Exposure to metals such as cadmium and arsenic have been shown to increase incidence of LBW and preterm birth (Ahmad et al., 2001; Hopenhayn et al., 2003), as has other measures of poor water quality (ChangeLab Solutions, 2017). Maternal residences near power plants have been associated with high risk for preterm birth and LBW (Ha et al., 2015; Tsai et al., 2004), while women living in neighborhoods with high exposure to natural gas wells were 1.4 times more likely to have preterm birth outcomes (Casey et al., 2016). Living in areas with high traffic density, or near high traffic roadways, has also been associated in a number of studies with increased risk of preterm births (Currie and Walker, 2011; Fleisch et al., 2017; Harris et al., 2016; Miranda et al., 2013; Woodward et al., 2015), as has living near landfill sites (Elliott et al., 2001) or being exposed to pesticides (Wolff et al., 2007).

EARLY CARE AND EDUCATION

ECE[6] has significant effects on children's academic readiness and cognitive and socio-emotional development. ECE also impacts children's physical, emotional, and mental health and well-being, and access to high-quality ECE creates many pathways through which greater health equity can be achieved (Hahn et al., 2016). The following section describes disparities in access to ECE, including specific programs such as Head Start/Early Head Start and state-funded preschool programs. See Chapter 7 for a more detailed discussion of the link between ECE interventions and outcomes related to children's health, development, and well-being as well as recommendations to advance health equity through ECE. See Figure 3-8 for a visual of how this section ties to the report conceptual model.

[6] Early care and education can be defined as nonparental care that occurs outside the child's home. ECE services may be delivered in center-, school-, or home-based settings (NASEM, 2018).

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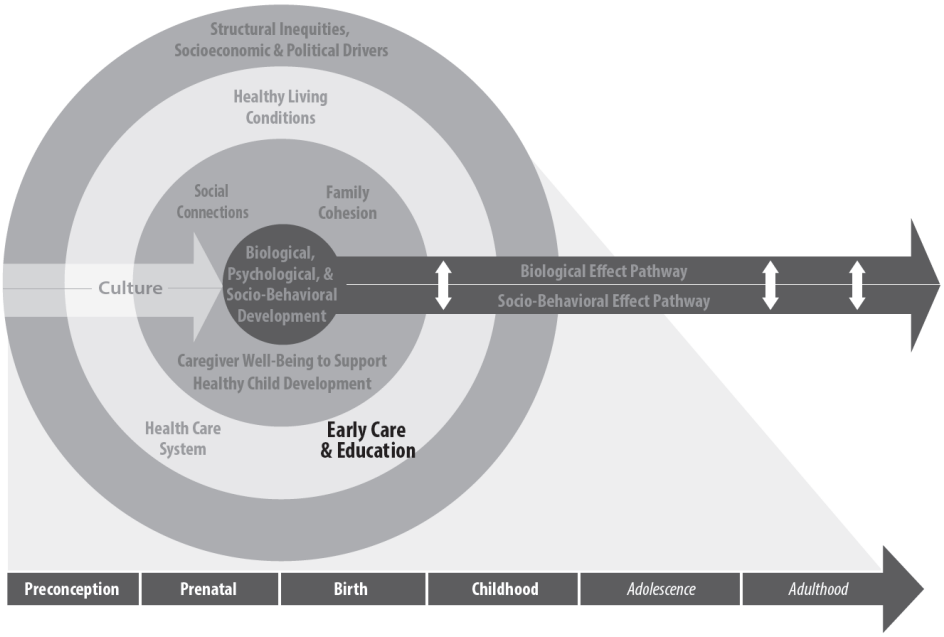

FIGURE 3-8 Leveraging early opportunities to achieve health equity across the life course: A conceptual framework
NOTE: The element of focus in this section is reflected in the second outermost ring in the model: healthy living conditions (i.e., economic security, nutrition and food security, housing, and environmental safety). This element is bolded for emphasis.

Access to Early Care and Education

Currently, about 60 percent of all children age 5 or younger who are not in kindergarten are enrolled in at least one nonparental care arrangement. The most common *primary* arrangement of early child care for all children was center-based care (29 percent), followed by home-based relative care (19 percent), home-based nonrelative care (10 percent), and multiple arrangements (2 percent) (de Brey et al., 2019). These percentages varied by race and ethnicity—see Figure 3-9 for the percentages of white, black, Hispanic, Asian, and two or more race children by type of early child care. In the United States, children under 2 years spend several more hours in informal care when compared to children in other Organisation for Economic Co-operation and Development (OECD) countries (25 hours per week compared to 3.5 hours per week) (Mathur, 2016).

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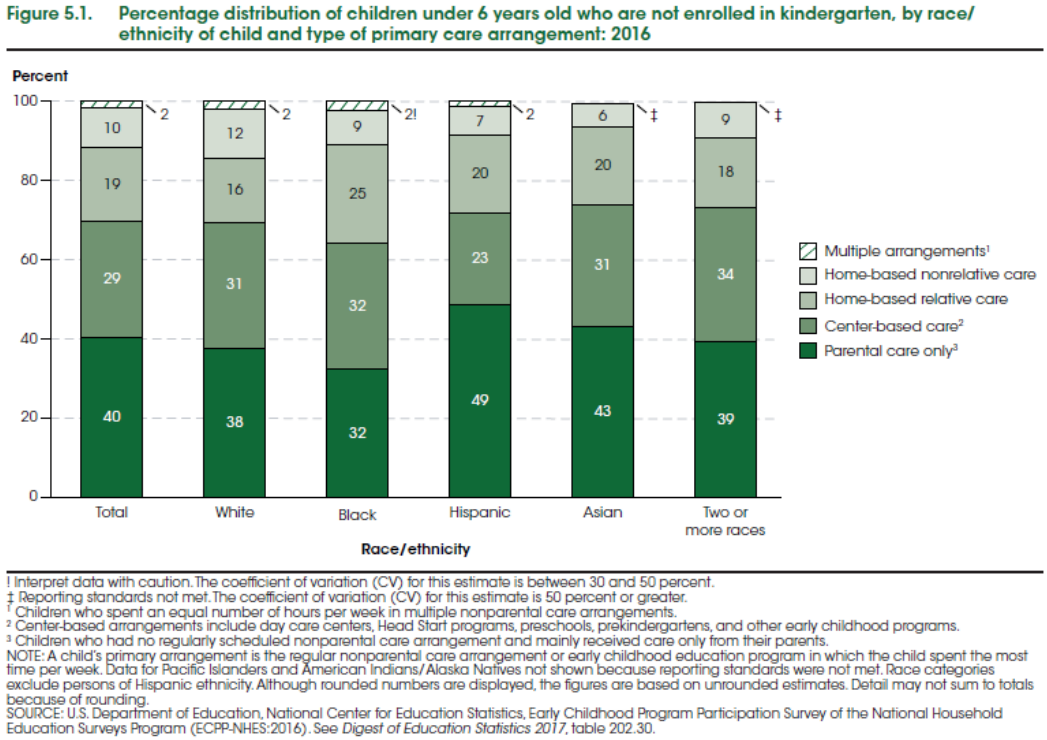

FIGURE 3-9 Percentage distribution of children under 6 years old who are not enrolled in kindergarten, by race/ethnicity of child and type of primary care arrangement, 2016
SOURCE: de Brey et al., 2019

Enrollment in ECE (including home-based care by a relative or a nonrelative and center/school-based arrangements) is associated with the following:

- Age. Preschool-aged children from 3 to 5 years old are more likely to participate than younger children.
- Race. White and black children are more likely to participate in one of these arrangements (62 percent and 68 percent of all young children, respectively) than Hispanic and Asian children (51 percent and 57 percent, respectively). However, white and Asian children are more likely to be placed in center/school-based programs (61 percent and 63 percent, respectively) than black and Hispanic children (57 percent and 52 percent, respectively).
- Home language. Children from homes where no parents or guardians speak English are less likely to participate in any ECE.
- Parental education. The lower the level of education of the parents, the less likely their children are to participate in ECE.
- Income. The lower the level of income of the parents, the less likely their children are to participate in ECE (see Figure 3-10). Fifty-four percent of poor children do not participate in any nonparental care arrangements, compared to 31 percent of nonpoor children (de Brey et al., 2019). While about three quarters of 4-year-olds from the highest income quintile attend preschool, only half of those from the lowest income quintile do (Cascio and Schanzenbach, 2016). However, the relationship is not completely linear. Children from households with income from $50,001–$75,000 are less likely to participate in center/school-based programs than those from the poorest ($20,000 or less)

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or highest ($100,001 or more) income groups. That said, poor children are much less likely than their nonpoor peers to be enrolled in center-based care or home-based nonrelative care as their *primary* form of ECE, defined as the arrangement where the child spends most of his or her time per week (Corcoran et al., 2019).

It is important to note that access to ECE does not guarantee access to quality care, especially for children from poor households and black and Hispanic children, who are more likely to be in lower-quality care than white and Asian children (Barnett et al., 2013; Valentino, 2018). Quality of access is also not available for AI/AN children.

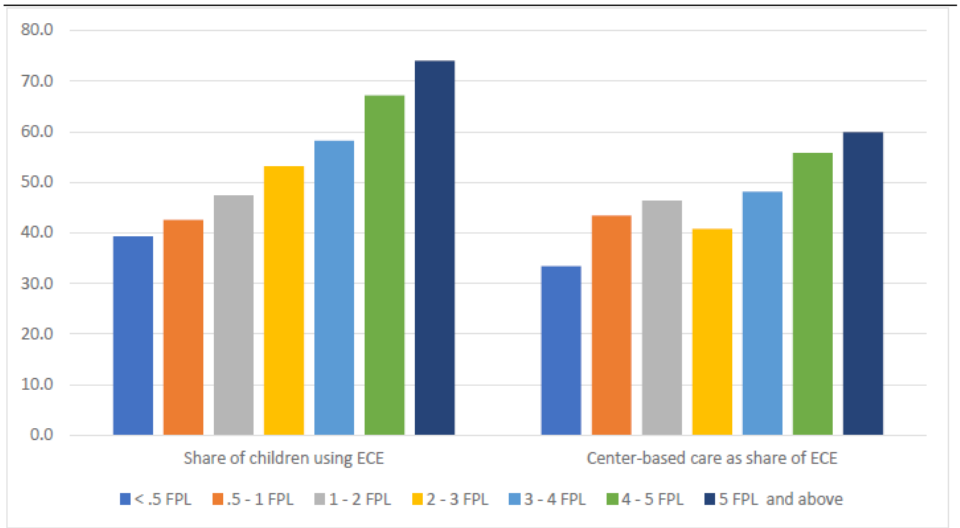

FIGURE 3-10 Patterns of ECE use by income category, all children age 0–5 years (not in kindergarten)
SOURCE: NASEM, 2018

The federal government subsidizes the cost of child care and provides early education programs for some low-income families through the Child Care Development Block Grant (CCDBG) and Head Start,[7] which may explain the greater access to ECE among low-income families, compared to those who earn slightly more. CCDBG provides child care subsidies to low-income families whose parents are working, going to school, or in a job training program. Children up to 13 years of age can be served in centers or family child care settings. (Two thirds of children served are younger than six [Office of Child Care, n.d.].) These funds are also used to improve the quality of care, coordinate resources, conduct research and evaluation, and provide technical assistance to grantees. There have been limited studies linking child care subsidies and child health and well-being, with mixed findings. In their econometric analyses drawing on the Early Childhood Longitudinal Study, Herbst (2010) found that subsidy use was associated with poor outcomes (i.e., reading, math, externalizing problems, approaches to learning, and interpersonal skills) for children compared to peers in the year prior to kindergarten entry. This was also found by Hawkinson, and colleagues (2013) for math outcomes using the same dataset. The authors hypothesize that these negative findings linking subsidy use and poor child

[7] Head Start is a program of the United States Department of Health and Human Services that provides comprehensive early childhood education, health, nutrition, and parent involvement services to low-income children and their families. See the following section and Box 7-3 for more information on Head Start.

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outcomes may be due to the quality of care that children with subsidies are likely to attend and the lack of availability of high-quality child care meeting the needs of low-resource families. In contrast, Krafft (2017), using a fixed effects approach that sought to adjust for selection bias, found that children receiving a child care subsidy experienced higher-quality care as reported by parents but there were no differences in the stability of arrangements or having multiple arrangements. The differences in these findings may be due to data used, methodology, analytical procedures, and how quality was defined and measured. However, these studies underscore the importance of ensuring that children from low-income families are able to access high-quality programs that meet their unique cognitive, emotional, social, and health needs.

Head Start is a federally funded preschool program for 3- and 4-year-olds from low-income households, created as part of President Lyndon Johnson's War on Poverty. In addition to focusing on children's learning and development, it also provides a range of services that address the holistic needs of children and families, such as health, nutrition, and parent education. Early Head Start was created in 1995 to focus on pregnant women and younger children from birth to age 3 (Office of Head Start, 2018a). Families receive services in homes, centers, or sometimes both settings (Office of Head Start, 2018b).

As of 2012, only about 15 percent of eligible families received CCDBG subsidies (Walker and Matthews, 2017), and Head Start serves less than half of all eligible 3- and 4-year-olds due to insufficient funding (Barnett and Friedman-Krauss, 2016). Early Head Start reaches about 5 percent of all eligible children (Schmit and Walker, 2016). Children from ages 2 to 4 and those living in deep poverty receive the highest rates of CCDBG subsidies (Chien, 2017). See Figure 3-11 for the percentage of children eligible for federal subsidies who receive such subsidies, by age and household income in 2013.

In terms of race and ethnicity, Latino, Asian, and AI/AN children tend to be underrepresented among eligible CCDBG recipients and Head Start children. In fiscal year 2016, 15 percent of eligible black children had access to CCDBG, but only 3 percent of eligible Asian children, 6 percent of eligible Hispanic children, and 7 percent of eligible AI/AN children were served (Ullrich et al., 2019).[8] As for Head Start, Asian and Latino children are somewhat underrepresented at 36 percent and 38 percent of eligible children served, respectively, while 54 percent of eligible African American children are enrolled in the program (Schmit and Walker, 2016). It is important to note that these patterns differ markedly across states.

Pre-K programs are funded and regulated by the state. Currently, 43 states and DC provide pre-K to 4-year-olds, with many also extending the program to 3-year-olds. Policies governing these programs are highly variable across states. Generally, unlike child care programs, state pre-K does not have any work or education requirements for the parents. Programs are also tend to be more focused on early learning and school readiness and less on other needs of children and families, although there are exceptions.

Access data for state pre-K programs are less available, since eligibility levels and data collection vary by state. Nationwide, 5 percent of 3-year-olds and 33 percent of 4-year-olds are served in a state-funded pre-K program. More than half of all programs target low-income children, with eligibility criteria ranging from 100 percent to 300 percent of the federal poverty line. About half the programs also take other risk factors or family backgrounds into consideration, including disabilities, abuse and neglect, homelessness, linguistic background, military duty, LBW, substance abuse, and teen parent status (Friedman-Krauss et al., 2018).

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[8] These data reflect federal eligibility parameters for CCDBG (Ullrich et al., 2019).

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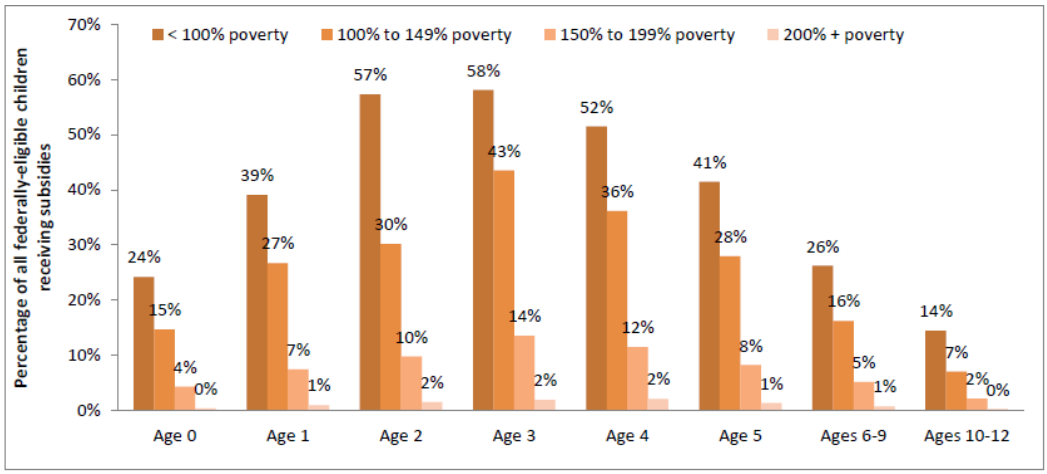
Enrollment varies greatly by state. For 4-year-olds, 12 states[9] enrolled less than 10 percent, 10 states enrolled more than 50 percent, and five states enrolled more than 70 percent. Only two (DC and Vermont) enrolled more than 50 percent of 3-year-olds. Seven states[10] continued to lack state-funded preschool programs. DC enrolled the highest percentages of 3- and 4-year-olds (66 percent and 88 percent, respectively) (Friedman-Krauss et al., 2018).



Note: Poverty figures are based on 2013 poverty thresholds published by the U.S. Census Bureau. For families with one adult and two children, 150 percent of poverty is $28,154 ($2,346 monthly).

FIGURE 3-11 Percentage of federally eligible children receiving subsidies by age and income
SOURCE: Chien, 2017

Head Start and state pre-K programs tend to have higher quality standards in terms of teacher qualifications and supports, learning standards for children, curriculum, group size, and adult-child ratios than child care programs outside those systems. To the extent that higher-quality programs are more likely to promote children's health and well-being, it is important for these programs to serve those with experiences of concentrated disadvantage, ACEs, and chronic stress. As mentioned earlier, while Head Start targets poor children, it currently serves only about half of all eligible children, and access by different racial groups is uneven.

Access to Special Education Services

About 2.9 percent of children birth through age 2 are served under Part C (the Program for Infants and Toddlers with Disabilities) of the Individuals with Disabilities Education Act (IDEA), and most were served at home in 2014 (Davis, 2016). Native Hawaiian or Other Pacific Islander and white infants and toddlers had risk ratios of 1.4 and 1.2, respectively, indicating that those in each of these racial/ethnic groups were slightly more likely than those in all other racial/ethnic groups combined to be served under Part C. In 2014, 6.1 percent of children ages 3 through 5 were served under Part B (Assistance for Education of All Children with Disabilities) of IDEA. In 2014, AI/AN, Native Hawaiian or Other Pacific Islander, and white children ages 3 through 5 had risk ratios above 1 (1.4, 1.5, and 1.2, respectively), indicating they were more likely to be served under Part B than were children ages 3 through 5 in all other racial/ethnic

[9] Rhode Island, Washington, Massachusetts, Delaware, Minnesota, Nevada, Arizona, Alaska, Mississippi, Missouri, Indiana, and Hawaii.
[10] Idaho, Montana, New Hampshire, North Dakota, South Dakota, Utah, and Wyoming.

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groups combined. In 2014, 8.7 percent of individuals ages 6 through 21 were served under Part B, with the most in specific learning disabilities. AI/AN, black or African American, and Native Hawaiian or Other Pacific Islander children ages 6–21 had risk ratios above 1 (1.7, 1.4, and 1.6, respectively), indicating that they were more likely to be served under Part B than children in all other racial/ethnic groups combined.

Unfortunately, access to early intervention and special education services is variable due to the wide discrepancy in eligibility criteria (Twardzik et al., 2017). Using data from the 2007 National Survey of Children's Health, Bethell and colleagues (2011) found that 19.5 percent of children received a parent-completed developmental screening. This varied across states and demographics. Screening was highest for children who were young, black, and publicly insured and lowest for children who were uninsured and had gaps in insurance coverage. They also found a significant gap between the developmental screening that is recommended by the American Academic of Pediatrics and what is reported nationally.

Public Education

Education attainment is correlated with health outcomes (NASEM, 2017a). Studies indicate that individuals with more education are more likely to have economic stability and better health outcomes (Heckman et al., 2018). Yet, disparities exist among income and racial groups in terms of both education inputs and outcomes. For example, recent data indicate increasing school segregation, with a growing trend of more schools that are high poverty and high minority, from 10 percent of schools in 2000 to 17 percent in 2013 (Chang, 2018). Other research examining longitudinal data established an association between court-ordered desegregation of schools (from 1954–1990) and subsequent adult outcomes for African Americans (e.g., educational attainment, adult earnings, probability of incarceration) through 2013 (Johnson, 2011). However, some contend that it is not segregation that is increasing but rather the number of nonminority students in public schools. Regardless, research also indicates that school districts serving the largest populations of minority (e.g., black, Latino, or AI/AN) students receive approximately $1,800, or 13 percent, less per student in state and local funding than those serving the fewest students of color. The United States spends approximately 7 percent—or $1,000—less per pupil on students in the highest poverty districts than those educated in the wealthiest (Morgan, 2018). Another estimate suggests a funding gap of $23 billion between school districts serving predominantly nonwhite students and those with mostly white students, despite serving the same number of children (EdBuild, 2019).

Funding inequities between rich and poor or majority-white and majority-minority schools are likely one factor contributing to the achievement gaps. Numerous studies have shown a relationship between school socioeconomic and racial/ethnic composition and student achievement outcomes (Kainz and Pan, 2014; Ready and Silander, 2011). Specifically, schools with large proportions of poor, African American, and Hispanic students have been shown to have lower rates of learning growth compared to schools serving larger proportions of white and high-income students. There are conflicting patterns of evidence regarding unique and overlapping effects of racial/ethnic and poverty concentrations on children's academic achievement. However, recent analyses of test data from the national public K–12 system have indicated that concentrations of poverty are the proximal explanation for observed negative associations between racial concentration and student achievement (Reardon, 2016). That is, racial segregation has a negative effect on education performance precisely because racial

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segregation is an indicator of students' exposure to poverty concentration and isolation and limited access to needed supports.

For example, data from the U.S. Department of Education, National Center for Education Statistics show that graduation rates collected from 2010–11 to 2015–16 increased from 79 percent to 84 percent (National Center for Education Statistics, n.d.). That is, more than 4 out of 5 students who were first-time 9th-graders in 2012–13 had completed high school by 2015–16 (within 4 years). There are some racial differences, with Asian/Pacific Islanders and white students graduating at higher rates, 91 and 88 percent, respectively, compared to Hispanic (79 percent), black (76 percent), and AI/AN (72 percent) students.

Similarly, the dropout rate has declined from 15 percent in 1970 to 6 percent in 2016. The youth dropout rate in 1972 was 12 percent for whites, 21 percent for blacks, and 34 percent for Hispanics. In 2016, the youth dropout rate for blacks was 6 percent (close to the national average) and 9 percent for Hispanics, compared to 5 percent for whites and 3 percent for Asians (Child Trends Databank, 2018). It is important to note that these estimates do not include those who were institutionalized, which is particularly pronounced for black and Hispanic male youth. There is a link between non-graduation and incarcerations (Kearney et al., 2014; Skiba et al., 2014).

There are many factors that lead to education attainment or lack of education (i.e., dropout). However, one predictor of high school graduation or dropout is reading proficiency at the end of third grade. In a study of nearly 4,000 students born between 1979 and 1989, Hernandez (2012) found that those who did not read proficiently by third grade were four times more likely to leave school without a diploma than proficient readers, with the rates increasing for those who could not master basic reading skills. According to the National Assessment of Education Progress, in 2017, while 47 percent of white and 59 percent of Asian 4th graders were proficient in reading, only 20 percent of African American and AI/AN students and 23 percent of Hispanic students achieved at that level (The Nation's Report Card, 2017). (A similar gap exists between low-income and nonpoor students.) Furthermore, the relationship between reading proficiency and dropout was particularly pronounced for students from poor households. Hernandez found that 31 percent of African American students and 33 percent of Hispanic students from poor households who were not proficient in third-grade reading did not graduate, compared to 22 percent for white children from poor households. The racial and education gaps disappear when students are proficient in third-grade reading and not living in poverty. However, the confound between poverty and race and ethnicity often makes it difficult to isolate the unique effects of each characteristic.

REVISITING CROSSCUTTING ELEMENTS

Racism and Discrimination

The 2017 report *Communities in Action: Pathways to Health Equity* describes racism as "an umbrella concept that encompasses specific mechanisms that operate at the intrapersonal, interpersonal, institutional, and systemic levels of a socioecological framework" (NASEM, 2017a, p. 104–105) (see Figure 3-12 for examples of how racism plays out at these levels). The report makes the distinction that racism is "not an attribute of minority groups; rather it is an aspect of the social context and is linked with differential power relations among racial and ethnic groups" (NASEM, 2017a, p. 105; see also Guess, 2006). Camara Jones' (2000) theoretic

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framework for racism uses an allegory of a garden to illustrate the relationships between these multiple levels and to provide a framework to guide interventions that would address these mechanisms of racism. It is also important to note that discrimination can affect a variety of racial and ethnic minority groups. For example, Asian Americans are often ascribed the stereotype of "model minority"; however, they can experience discrimination and its effects, as well as social or economic conditions that affect health (Chou and Feagin, 2015).

Reported racism[11] has been documented as a determinant of poor health outcomes across the life-span, particularly for mental health (e.g., depression, anxiety, psychological stress), with weaker evidence for physical health (Paradies et al., 2015). Bailey et al. (2017) highlight structural racism as an important but understudied driver of inequities, defining structural racism as "the totality of ways in which societies foster racial discrimination, via mutually reinforcing inequitable systems (e.g., in housing, education, employment, earnings, benefits, credit, media, health care, criminal justice, etc.) that in turn reinforce discriminatory beliefs, values, and distribution of resources, reflected in history, culture, and interconnected institutions" (p. 1,454). For more information on racism and discrimination as root causes of inequities, see Chapter 3 of *Communities in Action: Pathways to Health Equity* (NASEM, 2017a).

Childhood exposure to racism is mostly documented as experiences of discrimination. For example, the National Survey of Children's Health includes an item that asks parents if their child has ever been treated or judged unfairly because of their race or ethnic group. Data from 2016–2017 indicate that the overall prevalence of exposure to discrimination is low, at 3.7 percent nationwide. However, when these data are examined by subgroups, there are disparities. For instance, black and Other, non-Hispanic children were most likely to be treated or judged unfairly due to their race or ethnicity. (See Table 3-3 for a breakdown by race and ethnicity.)

There is an existing literature stream that draws connections between racism and early development and well-being outcomes. Specifically, negative prenatal and birth outcomes have been shown to be associated with discrimination or unfair treatment. An integrative review of 15 studies assessing adverse birth outcomes and discrimination found significant relationships between racial discrimination and LBW, preterm birth, and small size for gestational age (Alhusen et al., 2016). Studies in this review that examined mediating variables that are proximally related to birth outcomes (e.g., prenatal care, employment opportunities, neighborhood characteristics, or inflammatory markers) also found a significant association between these specific variables and racial discrimination. Another review found similar outcomes when investigating the relationship between racial discrimination and black-white disparities in birth outcomes, showing a consistent positive relationship between discrimination and preterm birth and LBW or very LBW (Giurgescu et al., 2011). The review findings also suggest that parental lifetime and childhood experiences of discrimination could have a stronger association with birth weight than experiences of discrimination reported during pregnancy.

[11] This meta-analyses defined reported racism as self-reported racism experienced directly in interpersonal contact; racism directed towards a group (e.g., based on ethnicity/race/nationality) of which the person is a member; vicarious experiences of racism (e.g., witnessing racism experienced by family members or friends); proxy reports of racism (e.g., a child's experiences of racism as reported by their parent); and internalized racism (i.e., the incorporation of racist attitudes and/or beliefs within an individual's worldview) (Paradies et al., 2015).

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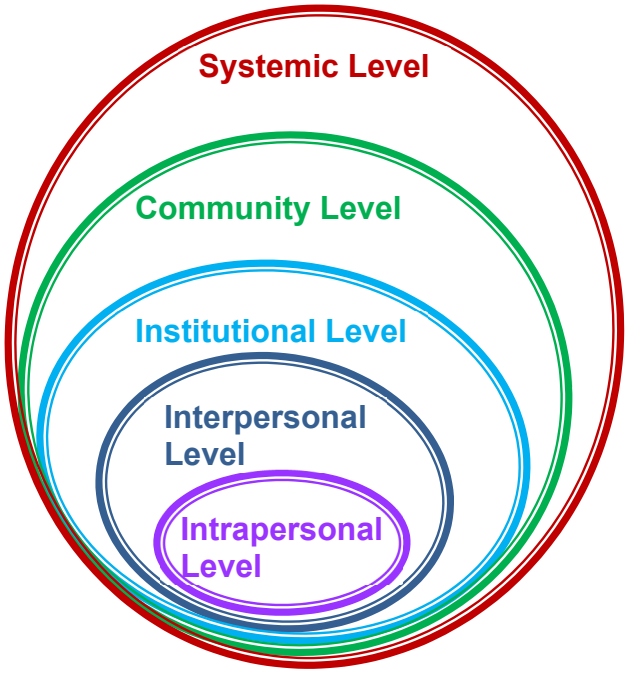

Systemic Level
- Immigration policies
- Incarceration policies
- Civil rights
- Predatory banking

Community Level
- Differential resource allocation
- Racially or class segregated schools

Institutional Level
- Hiring and promotion practices
- Under- or over-valuation of contributions

Interpersonal Level
- Overt discrimination
- Implicit bias

Intrapersonal Level
- Internalized racism
- Stereotype threat
- Embodying inequities

FIGURE 3-12 Socioecological framework with examples of racism constructs by level
NOTES: The mechanisms by which the social determinants of health operate differ with respect to the level. For the intrapersonal level, these mechanisms are individual knowledge, attitudes/beliefs, and skills. At the interpersonal level, they are families, friends, and social networks. At the institutional level, they are organizations and social institutions. At the community level, they are relationships among organizations. At the systemic level, the mechanisms are national, state, and local policies, laws, and regulations.
SOURCE: Adapted from NASEM, 2017a.

TABLE 3-3 Childhood Experiences of Discrimination by Race and Ethnicity

Race or Ethnicity	Percent of children who have ever been treated or judged unfairly because of their race or ethnic group.
Black, non-Hispanic	10.4
Hispanic	4.2
Other, non-Hispanic	7.4
White, non-Hispanic	1.0

SOURCE: National Survey of Children's Health, 2016–2017 from the Data Resource Center for Child & Adolescent Health, 2019

For childhood outcomes, the available evidence suggests that discrimination affects children through two pathways: (1) direct transmission to the child and (2) discrimination that affects the parent directly (sometimes called "vicarious racism"). A systematic review examining

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mostly cross-sectional studies synthesized the study findings related to racial discrimination and child health and well-being outcomes (Priest et al., 2013). Associations were most salient among mental health outcomes (e.g., depression and anxiety). The authors reported statistically significant associations between discrimination and 76 percent of the mental health outcomes studied. The findings also showed a consistent relationship between discrimination and reduced positive mental health outcomes, such as self-esteem, self-worth, and psychological adaption and adjustment. The authors reported weaker results for child physical health outcomes, but they also note that this likely reflects the extended period of time that it takes for physical health outcomes to manifest. With respect to vicarious racism effects, there is some evidence to suggest that there is a link between secondhand exposure to racism and socio-emotional (e.g., externalizing behavior) and mental health (e.g., depression) outcomes among children (Heard-Garris et al., 2018). Acevedo-Garcia et al. (2013) identify three directions for research on interpersonal and institutional discrimination and child health:

1. Incorporating a life course perspective into studies of discrimination and children's health,
2. Linking residential segregation with geography of opportunity[12] conceptual frameworks and measure, and
3. Considering residential segregation along with segregation in other contexts that influence children's health.

American Indian/Alaska Native and Children of Immigrant Background: A Closer Look

This section describes two populations of children and families, for whom the crosscutting-cutting elements in this chapter are particularly salient: AI/AN children and children of immigrant background. While there are many subgroups that have unique needs and historical contexts related to health equity, these are two that but the committee has identified these two as important to highlight here because of the lack of attention given to these groups, often times due to data collection or sampling issues (ASPE, 2007; NASEM, 2017a; National Congress of American Indians, n.d.). For these subgroups, racism and discrimination operate at multiple levels, including interpersonal, everyday experiences of discrimination and policies that obstruct access to health-promoting goods and services or induce historical trauma that can affect multiple generations. Their experiences across the domains discussed in this chapter (e.g., family cohesion, access to health care, neighborhood conditions) are amplified when examined with the additional lens of structural inequities.

American Indians and Alaska Natives

Health Outcomes across AI/AN Populations AI/AN populations experience stark disparities across a number of health indicators. Researchers hypothesize that a combination of factors may be related to these disparities, including the SDOH (i.e., poverty, unemployment, and education) (Elder et al., 2016), the effect of high rates of obesity and inactivity, alcohol and tobacco use, and a lack of access to health care (Bhaskar and O'Hara, 2017; Cobb et al., 2014; Rutman et al., 2016), particularly care that is culturally congruent (Lewis and Myhra, 2018; Rentner et al.,

[12] One tool for measuring opportunity by units of geography is the Childhood Opportunity Index, a data resource that compiles neighborhood-level indicators related to educational, health and environmental, and social and economic opportunity (Acevedo-Garcia et al., 2016a).

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2012). This stems in large part stems from the population's unique history in relation to U.S. policies, historical trauma, and the consequences for the health and well-being of generations (For more information on the historical and legal context of AI/ANs, see Appendix A in (NASEM, 2017). Moreover, health disparities and inequities vary by region, with some tribes experiencing high levels and intensity of alcohol use (Fortin et al., 2016), poor birth outcomes (Coughlin et al., 2013; Hwang et al., 2013; Kim et al., 2014), and poorer reproductive health (Rutman et al., 2012). AI/AN communities also experience have higher levels of racial misclassification, which reduces the reliability of existing health data for these populations (Jim et al., 2014). In addition, certain SDOH, such as violence against AI/AN women, is not consistently tracked across jurisdictions, making the justification for systemic changes challenging. For example, in some counties in the United States, AI/AN women are 10 times more likely to be murdered as compared to the national average, and are 3 times as likely to be raped in their lifetimes compared to any other racial or ethnic group, and the perpetrator of rape against AI/AN women is more likely to be white compared to other racial and ethnic groups. More research is needed to fully explicate the potential health impacts for AI/AN women living with the disproportionate level of violence specific to their race and gender in these contexts (Bachman et al., 2008).

Birth and Postpartum Outcomes While the research among on AI/AN populations remains limited, major disparities have been documented, particularly concerning birth outcomes. The mortality rate for AI/AN infants aged 0 to 1 year of age is 1.61 times higher than that of white children, with sudden infant death syndrome, unintentional injuries, and influenza/pneumonia being the top three causes of death (Wong et al., 2014). Concomitantly, AI/AN women experience some of the highest rates of diabetes and hypertension in the nation, as well as preeclampsia-eclampsia (Zamora-Kapoor et al., 2016), which are related to poor birth outcomes (Anderson et al., 2016; Dorfman et al., 2015). In a comparison between AI/AN, black, and Hispanic women, significantly lower rates of fetal and first day mortality were found among black and Hispanic women between for 2005–2008 as compared to 1995–1998 rates. However, among AI/AN, the rates did not decline significantly for AI/AN women.

Studies have found that preterm birth rates among AI/AN women (13.0 percent) are significantly higher as compared to the rate for all women in the United States (11.4 percent). Stress before and during pregnancy constitutes one of the biggest contributors to preterm birth. Research has demonstrated that AI/AN women experience the highest number of major stressors in the 12 months before becoming pregnant and in the 12 months before giving birth. Interpersonal violence rates, including rates of childhood physical abuse, rape, and multiple victimizations, are highest among AI/AN women. Access to prenatal care is limited for AI/AN women; they have, with twice the rate of AI/AN women accessing prenatal care late or receiving no prenatal care compared with other women (Raglan et al., 2016). In a study of nearly 300 AI/AN women, about 30 percent reported depressive symptoms 13–24 months postpartum. Nearly 70 percent of the sample reported financial events, 60 percent reported emotional events, 46 percent reported partner-related events, 38 percent reported traumatic events, and 19 percent reported IPV. Having had at least one partner-related or traumatic stressful life event or at least one traumatic stressful life event was significantly related to depression symptoms (Ness et al., 2017).

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Adverse Childhood Experiences In data from the 2011–2012 National Survey of Children's Health, AI/AN children were more likely to report 8 of 9 ACEs, including income deprivation, witnessing or experiencing violent victimization, racial/ethnic discrimination, household substance abuse, domestic violence, parental incarceration, divorce, and death of a parent. AI/AN children were also more likely to report more than one ACEs compared to non-Hispanic white children. After adjusting for sociodemographic factors, the difference between AI/AN and white children was no longer significant, which suggests that the heightened risk among for AI/AN children was explained by family and neighborhood factors. Among For AI/AN children reporting with one or more ACEs, parents reported an increasing prevalence of behavioral issues, including arguing, lack of emotional control, and school problems, and more provider-diagnosed behavioral disorders, such as depression, anxiety, and attention deficit and hyperactivity disorder (Kenney and Singh, 2016).

Health System Challenges Studies of health systems issues within AI/AN communities remain limited. The Indian Health Service provides national access to care statistics, but updated reports are lacking, and few focus specifically on the needs of early childhood. In one study of 59 emergency medical service agencies, or ambulance services, 46,761 annual emergency responses were reported, with 9,981 annual inter-facility transports. Pediatric emergency responses represented 15 percent (n = 7,190) of the total emergency responses, with an annual average of 180 pediatric responses per agency. Nine agencies (15 percent) reported that their agency did not have a medical director. Agencies with a medical director were more likely to report availability of pediatric medical direction than agencies without one. About 80 percent of agencies reported that their EMS providers needed pediatric continuing education for certification and recertification. About a quarter of agencies reported that all of their EMS providers received pediatric continuation education training, while six agencies reported no providers had received pediatric training. About half of agencies reported having a mass casualties plan, and of these, 15 had participated in a pediatric-focused mass casualties drill within 2 years of the survey. About 30 percent of all agencies had responded to a mass casualty incident involving a large number of pediatric patients that overwhelmed their service. Of these, about half of the agencies reported they did not have enough pediatric equipment available when they responded to the incident at that time (Genovesi et al., 2014).

Resilience Few studies examine factors contributing to resilience among AI/AN people;, however, one review found that, across eight relevant research studies examining resilience among AI/AN, resilience was an ongoing, dynamic process that responds to a changing environment, is evident within the life course, and can be accessed through culture. Cultural values, beliefs, and practices were identified as essential resources for AI/AN resilience along the life course and across generations. Three studies found culture could be best accessed through the use of narratives, lived experiences, and traditional stories (Ore et al., 2016).

Children of Immigrant Backgrounds

Children of immigrant backgrounds are one of the largest growing populations segments among the U.S. population of 0- to 10-year-olds. Nearly 90 percent of children in immigrant families are U.S. citizens, and about one quarter of them have an unauthorized immigrant parent (Koball et al., 2015). Thus, many children who live in immigrant families—irrespective of their own citizenship status—are greatly impacted by not only the household context but also the

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larger policy context for immigrant families (Filindra et al., 2011; Koball et al., 2015; Yoshikawa, 2011). In addition, research suggests that the United States will increasingly need to depend on immigrants and their children to maintain its workforce (Singer, 2012). To that end, supporting children of immigrant background and their families is essential to ensure not only individual healthy developmental trajectories, but also the future of the country. Much of the following discussion applies to documented immigrants, unless otherwise specified.

Many children of immigrant background face unique stressors that influence their development. For example, many parents who are first-generation immigrants have fled extreme poverty, political persecution, sexual violence, and other sources of oppression, which are traumatic experiences with long-term term consequences (Petrosky et al., 2017). In addition, many have experienced discrimination in housing and employment, as well as overt expressions of racism in their daily lives. These traumatic experiences can have implications for their parenting.

Residential segregation, among other factors, is one contributing factor as to why immigrants tending to end up living in concentrated poverty, which might lead to unhealthy levels of stress during pregnancy and in the infant's first years of life (NASEM, 2015b). Living in poverty in addition to an undocumented status may result in inadequate supports for caregiving: lack of high-quality quality day care, preschool, and elementary/middle school; and lack of access to adequate housing and health care (Yoshikawa and Kalil, 2011). There is growing evidence that documents the harm to children of parents with unauthorized status in terms of economic and socio-emotional hardship. For example, children fearing the deportation of a parent may show psychological distress by experiencing problems eating or sleeping, increases in headaches and stomachaches, anger or detachment, and depression and anxiety (Artiga and Ubri, 2017; Rojas-Flores et al., 2017).

The current policies toward immigrants in the United States have become another source of the toxic stress response for these families and their children. In particular, the dramatic recent increases in deportation of U.S. immigrants and the detention of parents at the borders seeking asylums have resulted in the separation of thousands of children from their parents (HHS Office of Inspector General, 2019). Research conducted since the 1950s shows that the scientific evidence is clear: separating children from families has long-term damaging health consequences for the children (Acker et al., 2019). These traumatic experiences can have cascading effects in multiple areas, even after reunification. Furthermore, recent immigration policies have had a documented effect on parents' willingness to enroll their children of immigrant background in public programs, including those in ECE settings (Cervantes et al., 2018). This could have serious implications because it increases the environmental barriers for these families; they refrain from using programs that could ameliorate or prevent educational and health problems (Koball et al., 2015; Yoshikawa, 2011). These barriers could be exacerbated as the increasing efforts to limit access to government services, such as education and health care, take effect at the federal or state levels.

However, research conducted both in the United States and internationally shows that if the right conditions are attainable and present, children of immigrant background can not only thrive, but also excel (García Coll et al., 2012; Motti-Stefanidi and García Coll, 2018). Many countries have, policies that support children of immigrant backgrounds exist at the state and national levels (Bachega, 2018; Line and Poon, 2013; OECD, 2018; Shinkman, 2018). Domestic and international studies show that policies providing support for immigrant families facilitate increases in positive outcomes, such as high school graduation rates. A U.S. national study

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showed that state family-supportive policies, such as allowing immigrant families to obtain driver licenses and, health care benefits, among others, lead to higher high school graduation rates (Filindra et al., 2011). An international study showed that immigrant- supportive public policies at the country level policies, also contribute also to more optimal health outcomes in children of immigrants (Marks et al., 2018).

At the community and neighborhood levels, research shows that school programs that support dual-language language learners (DLLs) facilitate positive developmental trajectories for children of immigrant background (NASEM, 2017b). In addition, interventions with a two-generation preventive focus also provide promising models and strategies to further support the healthy development of children of immigrant background (Valdez et al., 2013; Williamson et al., 2014).

ACCUMULATION OF RISK

This chapter has provided an overview of how family cohesion and social connections (Chapter 4), health care (Chapter 5), early living conditions (i.e., economic security, nutrition and food security, housing, and environmental exposures) (Chapter 6), ECE (Chapter 7), and racism converge through an accumulation of risk factors that influences a child's entire life course. Extensive evidence shows that exposure to multiple social risk factors or continuous exposure to a single risk factor is more harmful to children than one-time exposure to a single risk, across a broad range of outcomes (Evans et al., 2013; Sameroff, 1998). It is most common for researchers to calculate composite scores to represent the accumulation of risk factors by summing dichotomous indicators for each risk factor, although other, more complex methods exist (Evans et al., 2013). Across studies, there is variation in what is included within cumulative risk scores; in some studies, researchers focus on information about the family and residential context, whereas other studies may include information about the residential neighborhood context and/or school experiences.

Cumulative risk scores offer an easily interpreted measure and are strong predictors of child health (Evans et al., 2013). Notably, however, studies using cumulative risk scores typically do not consider the sequential timing or intensity of risk factors and are limited by the assumption that each type of risk is equally weighted and additive (i.e., not considering synergistic or interactive effects between risk and/or protective factors) (Dohrenwend, 2006; Evans et al., 2013). Longitudinal birth cohort studies with frequent assessments, such as the Dunedin Multidisciplinary Health and Development Study (Poulton et al., 2015) or the Avon Longitudinal Study of Parents and Children (Boyd et al., 2012; Fraser et al., 2012), provide an opportunity to investigate the chronicity of multiple risk factors over time (Danese et al., 2009; Dunn et al., 2018). The concluding section of this chapter explores the Adverse Childhood Experiences (ACEs) Study as an example of a composite approach to investigate the relationship between cumulative risk through childhood experiences and outcomes in adulthood (Felitti et al., 1998).

Adverse Childhood Experiences

As elaborated in Chapter 2, prenatal and early life environments have a profound role in shaping life course health, and even the health of the next generation, through a complex interplay of contextual and biological factors, including individual genetic characteristics, gene–

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environment interactions, family supports or stressors, environmental factors, and developmental experiences. This chapter looks more closely at the social and environmental contexts in which the aforementioned biological processes unfold and which structures promote the accumulation of risk or resilience. Perhaps the most well-known study of accumulated risk is the ACEs Study conducted by the Centers for Disease Control and Prevention (CDC) and Kaiser Permanente and published by Felitti and Anda in 1998 (see Box 3-9).

BOX 3-9
The Adverse Childhood Experiences Study

The phrase "adverse childhood experiences" was coined in 1998 following the publication of the Adverse Childhood Experiences Study (ACEs Study), a large epidemiological study conducted by the CDC and Kaiser Permanente, and refers specifically to the 10 categories of adverse events evaluated in the study, which include

1. Emotional abuse (recurrent),
2. Physical abuse (recurrent),
3. Sexual abuse (contact),
4. Physical neglect,
5. Emotional neglect,
6. Substance abuse in the household (e.g., living with an alcoholic or a person with a substance-abuse problem),
7. Mental illness in the household (e.g., living with someone who suffers from depression or mental illness or had attempted suicide),
8. Mother treated violently,
9. Divorce or parental separation, and
10. Criminal behavior in household (e.g., a household member going to prison).

The ACEs Study's findings were substantial and important, revealing that ACEs are common within the population (two thirds of study participants had experienced at least one ACE, and 12.6 percent had experienced 4 or more) and are associated with a broad array of long-term negative health outcomes in a dose-response relationship. Numerous epidemiological studies have supported these findings and demonstrated that those with an ACE Score of 4 or more are at dramatically increased odds for many of the leading causes of death in the United States (see Table 3-4).

SOURCES: CDC, n.d.-a; Felitti et al., 1998

Since the publication of the ACEs Study, subsequent research has demonstrated that the health impacts of ACEs are evident as early as infancy, and even in the prenatal and preconception periods. Parental ACEs are associated with increased risk of negative preconception and prenatal outcomes, including menstrual irregularity, reduced odds of conception, chronic health complications, preeclampsia, psychological risk (prenatal and postpartum depression), social risk (age at first pregnancy, education level, income, relational resilience, perceived stress, IPV, partner's drug abuse, maternal hostile behavior in infancy), behavioral risk (alcohol use, drug use, smoking), and negative pregnancy outcomes (miscarriage, premature delivery, reduced birth weight, shorter gestational age) (Ångerud et al., 2018; Brunton, 2013; Christiaens et al., 2015; Leeners et al., 2014; Li et al., 2018; Liu et al., 2018; Olsen, 2018; Racine et al., 2018; Smith et al., 2016).

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TABLE 3-4 Leading Causes of Death in 2015 and Their Relationship with the Experience of Four or More ACES

	Leading Causes of Death in the United States, 2015	Odds Ratio Associated with ≥4 ACEs[a]
1	**Heart Disease**	**2.1**
2	**Cancer**	**2.3**
3	**Chronic Lower Respiratory Disease**	**3.0**
4	Accidents	
5	**Stroke**	**2.4**
6	**Alzheimer's Disease**	**11.2**
7	**Diabetes**	**1.5**
8	Influenza and Pneumonia	
9	Kidney Disease	
10	**Suicide**	**30.1**

SOURCE: CDC, 2017
[a] Hughes et al., 2017 for all odds ratios except for stroke and Alzheimer's disease; Felitti et al., 1998 for stroke; Center for Youth Wellness, 2014 for Alzheimer's disease.

Outcomes related to prenatal and early childhood exposure to ACEs in children include altered neurodevelopment, neurocognitive function, cerebral processing, functional and structural brain connectivity involving the amygdala and (pre)frontal cortex, HPA axis, and autonomous nervous system (Brunton, 2013; McGowan and Matthews, 2018; Provençal and Binder, 2015; van den Bergh et al., 2018). Infants and children exposed to ACEs demonstrate increased risk of sleep disturbance, failure to thrive, growth and developmental delays, viral and bacterial infection, atopic disease (including asthma, allergies, and eczema), overweight and obesity, and learning and behavioral difficulties (Burke et al., 2011; Giordano, 2014; Kerker et al., 2015; Bjorkenstam et al., 2015; Matheson et al., 2016; Oh et al., 2017; Rhodes et al., 2012; Ryan et al., 2016; Shen et al., 2016; Thompson et al., 2017). Studies have also shown that ACEs are associated with increased risk-taking behaviors in adolescents, including early sexual initiation, teen pregnancy, teen paternity, substance use, and victimization (Hughes et al., 2017; Shin et al., 2009; Thompson et al., 2017).

Evidence points to several mechanisms by which ACEs and other early life stressors affect preconception, prenatal, and postnatal health:

1. Directly, through neuro-endocrine-immune dysregulation, metabolic regulation, and gene expression (as discussed in Chapter 2),
2. Through increased risk of health problems that may affect pregnancy outcomes, including obesity, diabetes, and autoimmune disease,
3. Through increased risk of parental mental health disorders,
4. Through increased risk behaviors, such as smoking, and substance use and risk exposures, such as victimization,
5. Through changes in epigenetic regulation that may be passed down via maternal or paternal germ lines, and

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6. Through difficulty with emotional regulation.

Beginning in 2009, many states began to systematically collect population-based representative data on ACEs, beginning with adults via the Behavioral Risk Factor Surveillance System (BRFSS). In 2011–2012, a modified ACE inventory was introduced into the National Survey of Children's Health. These surveillance activities provide scientists and policy makers with information on the prevalence of ACEs and data to study associations with health outcomes. According to the most recent published CDC data reporting from the BRFSS in 23 states, 62 percent of American adults have experienced at least one of the eight ACEs, and 15 percent have experienced 4 or more (Merrick et al., 2018). In 2012, The Institute for Safe Families formed the ACE Task Force to assess the prevalence of ACEs in Philadelphia in order to broaden the concept of ACEs to reflect the experiences of children of color in urban communities (Public Health Management Corporation, 2013).[13]

While the term refers specifically to the 10 categories identified in the ACEs Study, it is recognized that other forms of early life adversity, such as economic hardship, food and housing insecurity, unsafe toxic environments, and discrimination, also are also critical in shaping health and developmental outcomes. Cumulative exposure to negative experiences increases the risk of negative outcomes, but the converse is also true—supportive, nurturing, and buffering experiences also add up in a manner that is protective of health and neurodevelopmental outcomes. Thus, family, social, neighborhood, and structural environmental factors play an important role in shaping life course health and the health of future generations.

CONCLUSION

This chapter provides an overview of the risk and protective factors relevant to each level of the committee's conceptual model (see Figure 1-9). These factors are interrelated and complex, but they also demonstrate critical areas where there are opportunities for change and the science may be applied to inform interventions for children and families. In the chapters that follow, the committee takes the vast science base described in Chapters 2 and 3 and applies it to the necessary actions to advance child and family health outcomes and health equity across the domains discussed in this chapter (see Table 3-1 at the beginning of this chapter).

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[13] The original ACEs study sample was composed of primarily white, middle-class, and educated individuals (Felitti et al., 1998).

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4

Fostering Caregiver Well-Being Toward Healthy Child Development

INTRODUCTION

The focus of this chapter is on how to best foster children's healthy psychosocial development, emotional adjustment, and physical health using what science has shown about risk and resilience among children and families in high-risk contexts. With the goal of systematically addressing inequities, the broad question to address is, "For those children who are at risk for negative outcomes, what can be done—guided by science-based evidence—to expediently and effectively move them toward positive developmental trajectories?" In other words, how can the playing field be leveled, so that all children have the best possible opportunity to flourish and thrive in terms of their social-emotional, behavioral adjustment?

This chapter is structured in the following sections. At the outset is an overview of what developmental theory and research has demonstrated about "universals" in child development, beginning with a definition of core concepts in the phenomenon of resilience, followed by descriptions of risk and protective processes that are pervasive across diverse types of life adversities, with large and long-standing effects. Front and center among these universals are positive, supportive relationships and access to other basic and critical resources, such as good nutrition, health care, and early childhood education; these are all essential for developing children and for those charged with their care, and relevant theory and empirical evidence are provided on each. The primary focus is on fostering caregiver well-being and support, family cohesion, and social connections to promote healthy child development. This is reflected in the most proximal level to the individual child's development in the report conceptual model (see Figure 1-9). After discussions on these universals, the chapter continues with a consideration of some context-specific pathways to healthy child development: influences that are potent in and unique to particular subgroups of the population. As an example, issues of discrimination and segregation warrant attention for racial, ethnic, and sexual minorities. See Box 4-1 for an overview of this chapter.

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BOX 4-1
Chapter in Brief: Fostering Caregiver Well-being

Maximizing the well-being of young children and their primary caregivers within the context of the family, and with a focus on optimizing development in circumstances that place them at risk, is of utmost importance. The committee identified nine principles of psychological and behavioral human development. For example, for all children, the single most important factor in promoting positive psychosocial, emotional, and behavioral well-being is having a strong, secure attachment to their primary caregivers; in most instances, this is their biological mother. This chapter also examines the evidence on interventions that support strong relationships and highlights promising examples.

Chapter conclusions in brief:

- The primary caretakers of children—usually the mothers—are critical to fostering child well-being, especially in high-risk circumstances. Assuring the well-being of caretakers through ongoing support, extending beyond meeting basic needs, is necessary for child well-being.
- The most urgent parenting concern for families experiencing chronic stress and adversity is the potential for children's exposure to maltreatment.
- There is an urgent need to develop interventions well suited for fathers and other male caregivers; existing approaches that are developed for and tested with women cannot be assumed to generalize to other caregivers.
- Specific subgroups of children have unique needs and challenges when adjusting to adversity. Careful attention to potent subculture-specific processes be considered when working with subgroups well known to face serious inequities in relation to mental health.
- A tiered approach is needed for families known to be at risk due to their life adversities, with appropriate investment of resources to expand delivery of promising cost-effective programs and assessments via scientifically sound measures of critical constructs.

Chapter recommendations in brief:

- Implement programs that ensure mothers and families have access to high-quality, cost-effective community programs, including interventions to foster strong attachments and group-based supports in communities.
- Strengthen and expand evidence-based home visiting programs.
- Support research on:
    - the development of preventive interventions that target fathers and other male caregivers (with an emphasis on those that can be scaled up in the future); and
    - the development of culturally sensitive interventions that are tailored to meet the needs of subgroups of children known to be vulnerable.
- Routinely track levels of risk among mothers and children over time using periodic assessments.

The majority of this chapter is devoted to describing intervention programs that carry great promise in promoting positive outcomes among children in high-risk contexts. The focus is interventions that are well grounded in developmental theory and research; have demonstrated significant improvements with replications; have moderate or substantial effect sizes with a meaningful impact for the population; and, importantly, can feasibly be taken to large scale. Methods used to deliver the interventions, to ensure fidelity (to intervention strategies), and to assess outcomes are described in detail, to provide "blueprints" of science-based programs most likely to promote positive outcomes among young children and their caregivers who negotiate challenging life circumstances. For parameters and operational definitions used in this chapter, see Box 4-2.

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BOX 4-2
Maximizing the Well-being of Young Children and Their Primary Caregivers: Parameters and Operational Definitions

As described in Chapter 1, "inequities" are operationally defined for the purpose of this report, in part, as the *unequal likelihood of thriving or attaining positive adjustment outcomes over time because of differences in opportunity leading to unfair and avoidable differences in health outcomes* (NASEM, 2017).

The focus in this chapter is on behavioral/psychological/socio-emotional outcomes, rather than physical health or education; there are obviously some common "protective processes" involved in affecting all of these outcomes, but there are unique important issues for health (such as immunizations, primary care, and access to safe housing) or education (such as early Head Start and high-quality child care for early education) that are addressed in the following chapters.

The focus of this chapter is on discussing how to best maximize well-being among young children and families that are vulnerable at the outset. These are individuals whose life circumstances have rendered them statistically more likely—compared to national normative samples—to be on negative adjustment trajectories from early in life onward. In the field of risk and resilience, any group demonstrated to be at significantly higher odds for negative outcomes is described as being "at risk." For the purpose of this report, the term "children in high-risk contexts" will be used, as it is the contexts in which children live, grow, and play that puts them at risk.

The goal of this chapter is to describe how best to intervene to optimize developmental *trajectories across diverse groups of young children and their families in life circumstances that place them at risk for poor health outcomes*.

Resilience

Under the definition of inequities (see Box 4-2) as *unequal likelihood of thriving or attaining positive adjustment outcomes* over time, the construct of resilience from developmental science is directly relevant to discussions here, as it implies correcting what otherwise might have been negative trajectories, given major life stressors. Resilience can be defined as a dynamic process encompassing positive adaptation within the context of significant adversity (Cicchetti, 2010; Luthar et al., 2015; Masten, 2014). By this definition, there are two essential conditions that make up resilience: "a) exposure to significant threat, severe adversity, or trauma; and b) the achievement of positive adaptation despite major assaults on the developmental process" (Cicchetti, 2010, p. 145). With respect to a child's health, resilience is evident when physical, behavioral, and emotional well-being and development tip toward positive outcomes, even when a heavy accumulation of negative influences or risk factors exists.

Reviewing the body of accumulated scientific literature on resilience and socio-emotional functioning, Luthar et al. (2015) summarize the following core findings:

- Resilience happens; across risk and age groups; there will be a subset of individuals who show some positive adaptation.
- Resilience relies fundamentally on relationships, and this is true for children and adults.
- Children's resilience is shaped by multiple transactional systems (i.e., proximal and distal processes and environments affect the child, and the developing child in turn affects each of these).
- Resilience is not fixed over time but is dynamic; periods of positive adaptation can be interrupted by times of struggles.

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- Resilience is not a unidimensional construct; among those at risk, successful adaptation in some areas (such as overt behaviors) can co-occur with difficulties in others (e.g., covert anxiety or depression).

For information on physiologic resilience and variability in susceptibility, see Chapter 2.

As noted above, there are important considerations for *the context within which a child exists* and how that environment affects resilience. As is true for all children—facing adversities or not—the developing child's lived experience is shaped by the family, caregivers, community, and systems that the child interacts with on a daily basis. Reis et al. (2000) identify four propositions to describe this phenomenon: 1) from conception forward, individuals exist within the context of their social relationships; 2) each relationship is nested in a social and physical environment system; 3) each relationship is embedded in larger societal and cultural systems; and 4) all of these systems are continuously changing and interacting with each other over time. For these reasons, taking a multilevel, transactional, and relationship-centered approach to understanding resilience and its determinants for children is important (Luthar et al., 2015; Reis et al., 2000).

Of particular salience and proximity are family-level influences of relationships on resilience in childhood, as will be described in detail within discussions that follow. While risk factors such as exposure to significant adversity often unfold within the context of the family, so too do the protective factors that have been shown to foster resilience, such as parental warmth, responsiveness, and sensitivity (Hamoudi et al., 2015). Therefore, the parent or primary caregiver is a major target for interventions to promote resilience in children.

Across socioeconomic strata and in most parts of the United States (and the world), the primary caregiver is generally the mother or a mother figure. It is certainly true that in general, it is better for children to have two parents in the home, to share the responsibilities of child care and running the household. At the same time, even when there are two co-residing parents, one typically shoulders most of the child care, and this is generally the mother (e.g., see Ramey and Ramey, 2010). However, it should be clarified that in the discussions that follow, the term "mother" is used as a proxy for the primary caregiver who is a woman—whether that is the biological mother, grandmother, older sister, or other female relative. Considerations around men and fathers as primary caregivers are discussed separately later in this chapter, as the literature clearly indicates that there are distinct needs and issues that warrant specific, careful attention in interventions for men.

The critical importance of the parent–child relationship, specifically, is underscored in a 2017 special section published in *Child Development,* which highlights evidence-based interventions to maximize resilience in children and families. In their overview article, Luthar and Eisenberg (2017) outlined the key findings and recommendations that emerged from the collection of papers. Across these 11 papers, the single theme that most consistently recurred was the need to support the well-being of the primary caregiver—typically the mother. Contributing authors emphasized the importance of psychological and emotional support for mothers, particularly for single mothers (Taylor and Conger, 2017), those struggling with depression (Goodman and Garber, 2017), those at risk for maltreating their children (Valentino, 2017), and those living in poverty (Morris et al., 2017). Another, second, important target for intervention identified from this set of articles was specific types of parenting behaviors that most urgently warranted attention among at-risk families. It was particularly important to minimize harsh, rejecting, or neglectful parenting behaviors (e.g., Goodman and Garber, 2017; Harold et al.,

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2017; Reynolds et al., 2017), because the ill effects of maltreatment on children can be profound, pervasive across areas of adjustment, and long-standing, especially when it starts early in development and is prolonged. A third target area identified as beneficial for interventions was self-regulation and coping skills among adults—parents and teachers—as well as children (e.g., (Domitrovich et al., 2017; Modecki et al., 2017; Morris et al., 2017; Smith et al., 2017). Promoting self-regulation and coping for both the parent and child is essential because parents are likely to adjust their parenting behavior based on their children's behavior. Of course, for all parents living in challenging circumstances, there needs to be attention to not only their personal well-being and parenting but also the social and economic conditions that provoke the toxic stress response (discussed later in this chapter in the section on group difference).

HUMAN DEVELOPMENT IN THE CONTEXT OF RELATIONSHIPS

This chapter, and the associated conclusions and recommendations, are based on what is known from extant scientific literature about the major, critical building blocks to optimize the child's development from pregnancy through puberty (approximately 8–10 years of age). Sections that follow begin with a synthesis of this literature providing the basis for what are considered the universal principles of human child development; that is, principles that are largely generalizable to most children and families in the United States. Following this is a discussion of instances where children and families in discrete subcultural groups have important, unique needs, given the presence of potent risk or protective processes in their particular contexts. The remainder of this chapter is focused on a review of interventions that have been proven to be effective to promote relational supports and well-being of caregivers toward maximizing resilient adaptation.

Universals of Human Development

Universals: The interplay of biology and environment in human development

As described in Chapter 2, during prenatal development, genetic and environmental factors work together to build organ systems that promote the ability of the individual and species to thrive and survive (Center on the Developing Child at Harvard University, 2010; National Scientific Council on the Developing Child, 2010). It is an evolutionary conserved core principle that prenatal through early childhood development is both experience dependent and experience expectant.[1] Thus, the adaptability of the organism to future challenges is influenced substantially by genetic and epigenetic factors that contribute to homeostatic boundaries within which the brain and other organ systems operate over the life course and the experiences that occur during development that the organism anticipates as the "future" contexts in which it will need to operate to survive and thrive. This core principle relates to both physical survival, which Maslow's (1943) hierarchy of needs indicates is the most basic need for all humans (children and

[1] "Experience-dependent" describes the aspects of development that depend on the quality of the environmental input (for example, while babies are born with a capacity for language development, their ability to communicate verbally will depend on their exposure to language during early development). "Experience-expectant" describes the normal, generalized development when the environment provides infants with the needed input to develop neural connections that enable function in certain domains (such as vision and hearing). If input during these critical periods of development is not received, these functions will be lost permanently (Gallagher et al., 2003).

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their caregivers), and relationships that contribute to physical and behavioral well-being. See Figure 4-1 for Maslow's hierarchy pyramid.

All human circuits are built from the bottom up, with simple skills forming the foundation for the emergence of more complex skills (Center on the Developing Child at Harvard University, 2011). For sociability, the interaction between humans starts after birth, but the initial circuits needed to direct interactions that are fundamental to early development, such as caregiver–infant attachment, are established prenatally. Infants are ready to use their senses to attend to environmental stimuli—what they hear, see, smell, feel and taste—to receive the most important information that supports their immediate internal needs, such as hunger, thirst, and arousal (Hammock and Levitt, 2006). In essence, early experiences drive a process in which the motivation and attention circuits focus more and more on those stimuli that are most meaningful—a process known as contingent learning. Building these circuits occurs in a social context with the primary caregiver and others, providing key opportunities for the infant to associate stimuli with social meaning. Over time, as social context becomes more and more complex, the developing child engages cognitive and emotional regulatory processes to infer social meaning and ultimately, social responsiveness to cues—back and forth or "serve and return" interactions (National Scientific Council on the Developing Child, 2004). These skills develop over an extended period, through adolescence (Casey et al., 2016; Dahl, 2004).

There are extended discussions pertaining to the importance of all children having their basic needs met and the existing problems that place children and families at risk, including those related to access to adequate housing, good nutrition, and health care (along with the associated barriers, ranging from parents' inability to take time off of work and lack of transportation to logistical problems deriving from the spread of critical services across physical locations and delivery systems). These issues are discussed extensively in Chapters 3 and 6.

FIGURE 4-1: Maslow's hierarchy
SOURCE: McLeod, 2018
NOTE: Concepts from Maslow, 1943.

Universals: Development in the Context of Proximal Family Relationships

The focus of this chapter is the next most basic need in Maslow's (1943) hierarchy: belongingness and love. To set the foundation for the committee's discussions on this topic, the

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following section outlines nine committee-generated, universal principles of human development that are derived from the scientific literature on child development in the context of relationships. To reiterate, these principles all pertain to the broad domain encompassing children's psychological and behavioral adjustment. See Box 4-3 for a list of the nine universal principles of human development.

BOX 4-3
Nine Universal Principles of Psychological and Behavioral Human Development

1. For all children, the single most important factor in promoting positive psychosocial, emotional, and behavioral well-being is having a strong, secure attachment to their primary caregivers; in most instances, this is their biological mother.
2. This strong attachment presupposes effective parenting behaviors in everyday life, and "effective parenting" changes in nature and complexity and with development over time.
3. Effective parenting presupposes the caregivers' own well-being.
4. It is critical to ensure that children's mothers have the necessary supports for maintaining good mental health and psychological well-being.
5. If there is any concern about a parent "faltering," the single most important dimension of parenting in which parents need to be helped is maltreatment (by parent figures or others).
6. The early childhood years are critical in shaping subsequent child development—but at the same time, there remains much room to change course and impact these trajectories in a positive fashion.
7. The ill effects of toxic early environments (e.g., unhealthy relationships) can manifest in physical health problems in childhood and throughout life.
8. The ill effects of early childhood adversities (e.g., abuse or neglect by caregivers) can become most visibly apparent later in life (i.e., there can be latent or "sleeper" effects that are seen at later points in development).
9. Biology is not "destiny"; rather, while indicating relative risk rates, it sets "confidence limits" within which environmental influence can lead to relatively positive versus negative adaptation.

The first principle, deriving from over eight decades of research on resilience, is the following, mentioned earlier: *For all children, the single most important factor in promoting positive psychosocial, emotional, and behavioral well-being is having a strong, secure attachment to their primary caregivers*; *in most instances, this is their biological mother*. This conclusion makes good sense intuitively, as it is the primary caregiver who is the most constant and proximal person who has everyday contact with the young child and long-standing influence on the developing child from conception through late childhood. In most settings, this is typically the mother (relationships with fathers are obviously also extremely important, and are discussed below). Even in families where they are employed outside the home, across socioeconomic contexts, it is typically mothers who have the earliest, most constant, and most prolonged everyday contact with the child. However, it should be noted that even when the primary caregiver is another adult, the universals that are described here would largely generalize to these

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other main caregivers (with a few exceptions that are discussed explicitly). See Box 4-4 for discussion of gender specificity.

BOX 4-4
The Need for Gender Specificity in Referring to Parents

Developmental scientists have increasingly emphasized the need to avoid using gender-neutral terms in referring to parents in at-risk circumstances, and, instead, to describe patterns specific to fathers and mothers (see Cabrera et al., 2014; Luthar and Eisenberg, 2017; Phares et al., 2005; Rotheram-Borus et al., 2001; Yogman et al., 2016). By far, most of the existing developmental studies on parenting have focused on mothers, as it is mothers who are typically primary caregivers of young children and who usually enroll and participate in intervention programs (Pruett et al., 2017). Yet, there are many instances where fathers are primarily responsible for raising their children, and their unique needs require attention: *It cannot be presumed that what helps fathers do well as parents is the same as what most benefits mothers* (Cabrera et al., 2014; Luthar and Eisenberg, 2017). To that end, differences between mothers and fathers render it critical to better understand the unique factors that shape fatherhood and child development to inform interventions. To illustrate, fathering is more affected by the quality of the marital relationship (levels of satisfaction and support) than is mothering (Lamb, 2004; Lewis and Lamb, 2003). Similarly, programs based in supportive, close relationships among women (service providers and community-based "mentor mothers") can effectively recruit mothers (Luthar, 2015; Valentino, 2017) but are unlikely to attract men as strongly. For fathers, recruitment efforts in community settings where they are commonly found, including soccer games, employment centers, and shopping malls, are more effective (Pruett et al., 2017).

Following this first universal—that a strong attachment to primary caregivers is critical for the child's well-being—the second major universal is that *this strong attachment presupposes effective parenting behaviors in everyday life, and "effective parenting" changes in nature and complexity and with development over time.* In infancy, parenting is extremely demanding on physical energy, and, of course, on sleep; at a psychological level, it calls for sensitivity to the child's needs and appropriate responsiveness (e.g., soothing/comforting when the child is in discomfort or pain). Often, parents are physically exhausted and not sure about how to soothe a baby who might cry inconsolably for hours. As children grow into the toddler years, in addition to providing love and comfort, effective parenting calls for constant vigilance and appropriate limit-setting. During the toddler years, children are physically very mobile, oblivious to physical dangers (e.g., sharp objects or electrical outlets), increasingly and energetically willing to explore the world around them, rarely responsive to a simple "no" from adults, and prone to tantrums. The use of language as mediator of appropriate parent–child interactions becomes increasingly complex and important. In early childhood through the onset of kindergarten and formal schooling, in addition to the primary caregiver providing love and comfort and appropriate, consistent limit-setting at home, a new developmental task for the child—and thus for that caregiver—is adjustment to external influences. This subsumes the transition to child care and then to kindergarten with separation from the primary caregiver and learning to handle routines and a new set of socializing influences, including other adults plus children. During the elementary school years, failures and successes in various areas—academics, peer relations, sports or other specialized arenas—need to be processed at the socio-emotional level. Peers and others adults can provide support to children and parents as they navigate increasingly diverse and demanding contexts.

The third universal is that *effective parenting presupposes the caregivers' own well-being.* Being able to deal with the various tasks/demands of parenting is difficult at the best of

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times, and this becomes particularly challenging in the presence of high levels of life stress. Chronic exposure to poverty or prolonged periods of limited resources, as well as life events involving high levels of trauma, can lead to maternal depression, anxiety, and other mental health problems, which in turn can greatly compromise the quality of parenting. There are multiple studies showing that when mothers struggle with mental illness, this can significantly affect all their parenting behaviors (Knitzer, 2000; Luthar and Sexton, 2007; Rahman et al., 2013). In particular, maternal depression and high stress have been linked consistently with tendencies to neglectful or harsh parenting, increasing the likelihood of the child's exposure to ACES (Goodman and Garber, 2017; Plant et al., 2013; Toth et al., 2013; Valentino, 2017). Another important psychological element of effective parenting is parent agency or self-efficacy (i.e., the parent's perceived ability to influence the development of the child) (Jones and Prinz, 2005). The report *Parenting Matters: Supporting Parents of Children Ages 0–8* highlights parent self-efficacy as a contributing factor in shaping parenting competence and child self-regulation, social, and cognitive skills (NASEM, 2016). The report identifies one possible mechanism for this association as increased competence-promoting parenting practices (e.g., family routines or involvement with children in schools). And as the current report's conceptual model in Chapter 1 suggests (see Figure 1-9), caregivers and families live in the context of their social circumstances and living conditions; therefore, they need support in coping with related stressors or eliminating such sources of stress, in addition to efficacy-promoting supports.

The first three universals have been long cited in the developmental literature; the next has been more recently emphasized explicitly: from as early in a child's life as possible, *it is critical to ensure that the mother has the necessary supports for maintaining good mental health and psychological well-being*. Resilience researchers have clearly articulated that in order to achieve positive mental health and functioning, the same principles that apply for children apply to their mothers. Just as children need ongoing unconditional love from their mothers, so do the mothers need ongoing love and support from other adults to function well across their life roles (Luthar et al., 2015). In developmental science, for too many years, the tendency in studying families has been to discuss what parents should do and should not do, with little attention to what it is that allows parents themselves to function well and develop resilience at this daunting, decades long-task. In other words, what has been missing is mothers' well-being as the dependent variable.

In the last few years, this has changed (see, Balaji et al., 2007; Jolly et al., 2014; Luthar and Ciciolla, 2015; Taylor and Conger, 2017). There is now much more explicit emphasis on the need to understand what helps mothers do well over time, especially in highly stressful life circumstances. With accumulating evidence from both basic research and intervention trials, the answer is clear: the single most important need is dependable supports in their role as parents—emotional/intangible and tangible—from a dependable, reliable network of others in their lives (for a review on this topic, see Luthar and Eisenberg, 2017). The nature of these supports, furthermore, are both perceived (e.g., the expectation that comfort will be available when distressed or help will be forthcoming if needed for parenting/running the household) and instrumental (e.g., actual receipt of help with child care or aspects of running the home; see Balaji et al., 2007; Goodman and Garber, 2017 for more information). See Box 4-5 for more on supporting caregivers.

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BOX 4-5
Providing Supports for Caregivers

Recognizing the critical importance of supports for caregivers by no means implies parent-blaming or minimizing parents' agency. Rather, it is intended to draw much needed-attention to the great challenges of this life role and what even the most well-meaning caregiver needs, on an ongoing basis, to do this job well. As an analogy, consider the role of a surgeon. Ultimately, the physical manipulation of nerves, arteries, etc. is in his or her hands. Yet, it is common knowledge that for any surgery to go well, there needs to be high-level, ongoing supportive conditions—ranging from the assistants in the room to sterile operating room and appropriate medications. It would be illogical to turn to any "surgeon-blaming" if a procedure failed in the absence of these various essential supports. In the same way, the first order of business, in ensuring optimal child development, is to ensure that all primary caregivers have the necessary supports on an ongoing basis, if they are to sustain optimal parenting. Providing supports (e.g., in the form of services) does not assume any weakness on the part of the caregiver; instead, it promotes parents' agency to care for their children in the face of increasing demands as parents (e.g., work, child care) (Lino et al., 2017; NASEM, 2016; National Partnership for Women & Families, 2018).

A fifth universal, and also one more recently emphasized specifically in the literature, is that if there is any concern about a parent "faltering," *the single most important dimension of parenting in which parents need to be helped is maltreatment* (by parent figures or others).[2] In general, psychologists have shown that "bad is stronger than good" (Baumeister et al., 2001); that is, the ill effects of harsh behaviors are much stronger than the positive effects of praise (see also Fredrickson and Losada, 2005; Sparks and Baumeister, 2008). Additionally, the accumulated literature on resilience has shown that early and chronic maltreatment is the one category of adversity in which it is rare to find children who "do well" over time, showing little to no trouble across major adjustment domains; in fact, this has many serious long-term effects across salient aspects of child development (see Bolger and Patterson, 2003; Cicchetti, 2013).

Therefore, as a society, prevention of maltreatment needs to be a top priority in promoting the well-being of children in high-risk contexts. This again implies attentiveness when primary caregivers are stretched too thin, because they are at the most risk for maltreatment. As noted earlier, living with chronically high stress or mental illnesses, such as untreated depression, can lead the most well-meaning and loving parent to become angry and slip into neglectful or harsh parenting behaviors. If caregivers are believed to be at risk for abusing or neglecting their children—or of not being able to protect them from maltreatment by other adults—the situation needs to be treated as urgent and the appropriate help identified. Most importantly, the best help would be providing supports and/or mental treatment for the primary caregiver to address the root cause of maltreatment or neglect (Cicchetti, 2013; Cicchetti et al., 2006; Lieberman et al., 2005; Luthar et al., 2007; Toth et al., 2013). In the most extreme cases, continued high risk for maltreatment or actual maltreatment could entail removing the child from the home; however, there is evidence that suggests that this is not always the best course of action for the child (Toth et al., 2016).

Implicit within the above universals that the well-being of the young child inherently involves the well-being of the primary caregiver; in fact, Seifer (2003) explicitly pointed out that it does not make sense to think about resilience of infants; rather, we should consider the positive

[2] The Centers for Disease Control and Prevention (CDC) define child maltreatment as any type of abuse (e.g., physical, sexual, or emotional) and neglect of a child under 18 years of age by a parent, caregiver, or another person in a custodial role (CDC, 2014).

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functioning of the mother–child dyad. Additionally, in many countries other than the United States, people think about the needs of the child through the needs of the caretaker: if the child is to be helped, that help is most effectively offered if it goes through the caretaker (e.g., see Hrdy, 1999). In many cultural contexts, pregnancy, childbirth, and motherhood are embedded in a system of multiple generations of women who provide support and troubleshooting to the younger generations in all aspects of caretaking. Extended family systems and/or fictive kin are available and ready to help with any problem that arises; they affect the development and well-being of the child and also the mother (see Barlow and Chapin, 2010; Kopp, 2013).

To reiterate the caveat noted earlier—that in many instances, it may not be the mother but another family member (or caretaker) who serves as the primary caregiver—the principles described above are still the same; that is, whomever is charged with primary (or salient) responsibility of taking care of the young child needs to receive regular replenishment and support. Child care providers and educators also need ongoing support, including teachers from preschool onward through the K–12 system. These are all jobs that are among the lowest paid and with high levels of burnout and turnover, especially for early childhood educators. Once again, research has shown that these care and education professionals can serve critical protective functions, stepping in to provide the nurturance and support that complement or supplement what mothers, fathers, or other primary caregivers are able to offer. Yet, solutions rarely consider what it is that can and should be done to help replenish these professionals who are the primary adults during significant portions of children's daily lives. (For a more in-depth discussion of early care and education [ECE] professionals, see Chapter 7.)

Sixth, developmental research has clearly shown that relationships in *the early childhood years are critical in shaping subsequent child development—but at the same time, there remains much room to change course and impact these trajectories in a positive fashion*, with those starting out as negative changing to be more positive as well as the reverse. Without question, early attachments form the lens through which subsequent relationships are viewed (Lieberman et al., 2005; Sroufe et al., 2005; Yates et al., 2003; Zeanah and Zeanah, 1989), so that children with adverse early experiences are less likely to be open to or trusting of other nurturing people in their lives. Yet, there are certainly opportunities for reversals (e.g., at times of transition in life, including moves to new schools for children, and, later in life, marriage or employment changes). Many years ago, Rutter (1987) showed that women who were institutionalized as children were more likely to display positive parenting themselves if they married supportive spouses, and Sampson and Laub (2003) showed that for delinquent children followed over decades, both marriage and entry into the military conferred significant benefits to their well-being and adaptation. In short, even if subjected to early trauma, human beings are capable of recovering and healing.

The seventh universal principle is that *the ill effects of toxic early environments (e.g., unhealthy relationships) can manifest in physical health problems in childhood as well as throughout life.* Common health problems include asthma, headaches and backaches, and even elevated risk for heart disease and compromised immune functioning (e.g., Dong et al., 2004; Dube et al., 2009; Purewal et al., 2016; Wing et al., 2015; Kalmakis and Chandler, 2015). Therefore, once again, preventing exposure to such toxic experiences is of the essence, even if children appear to be adapting well behaviorally or psychologically to these adverse circumstances in their early years of childhood.

In a related vein is universal number eight: *The ill- effects of early childhood adversities (e.g., abuse or neglect by caregivers) can become most visibly apparent later in life (i.e., there*

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can be latent or "sleeper" effects that are seen at later points in development). (See Chapter 2 for more information on sleeper effects.) For example, attachment insecurities developed in infancy could manifest in severe separation anxiety from the parent on starting kindergarten or significant difficulties in relating well with peers (Cicchetti and Toth, 2015, 2016). Other examples include behaviors that become relevant or salient closer to puberty or adolescence, such as delinquent behaviors or abuse of drug and alcohol as a function of early relationship disturbances (Cicchetti and Toth, 2015, 2016). Perhaps the most obvious example is impaired parenting as adults; in general, there tends to be some intergenerational continuity of maltreatment (Adams et al., 2019; Cicchetti and Rizley, 1981; Kaufman and Ziegler, 1987; Pears and Capaldi, 2001; Rutter, 1987).

Universal nine—*biology is not "destiny"*; rather, while indicating relative risk rates, it sets "confidence limits" within which environmental influence can lead to relatively positive versus negative adaptation. Genetic factors, for example, can heighten children's vulnerability to disorders (physical and psychological) that afflict their parents. At the same time, intergenerational transmission of disorders is far from inevitable. As noted earlier, the notion of resilience encompasses the phenomenon wherein children who are at risk for psychopathology—or mothers who are at risk for negative parenting behaviors—can show healthy outcomes if they have the appropriate corrective or buffering influences in their lives. In other words, biological factors affect children's development, but they always operate in interaction with forces in the environment (Wise, 2009). The burgeoning field of epigenetics has shown that early life environments can lead to long-lasting changes in gene expression that "can leave indelible chemical marks on the brain and influence both physical and mental health later in life even when the initial trigger is long gone" (Murgatroyd and Spengler, 2011, p. 1; see also Dudley et al., 2011; Lerner and Overton, 2017; and Chapter 2 for more information).

To summarize, there are key take-home messages from the above discussion of universals to ensure positive psychological, behavioral, health, and social outcomes in early childhood. First, ensure the psychological well-being of the primary caregiver, as this will inevitably affect his/her parenting and the quality of attachment; this is best done by ensuring *ongoing* support for mothers in their everyday lives and communities (support during the child's early infancy will not protect the mother for the child's entire life). One population of particular importance for intervention is mothers under a high level of chronic stress and at risk for depression—as this is strongly related to increased risk for neglect and maltreatment, which, of all parenting behaviors, is the one that most urgently requires prevention. Finally, in considering whether a child or parent has done well in spite of early adversities, it should be noted that ill effects of these adversities can become manifest later in life. At the same time, significant healing can happen, even well past childhood and adolescence, so that mothers who experience suboptimal care themselves as children can be effective caregivers if they receive appropriate support and help as they negotiate the challenges of motherhood.

> *Conclusion 4-1: The primary caretakers of children—usually the mothers—are critical to fostering the well-being of children, especially in high-risk circumstances. Assuring the well-being of caretakers through ongoing support, extending beyond meeting basic needs, is necessary for child well-being.*

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Conclusion 4-2: The most urgent parenting concern for families experiencing chronic stress and adversity is the potential for children's exposure to maltreatment (by others within and outside the family).

Fathers

For the most part, this chapter has focused on mothers (or mother figures) because across cultures and socioeconomic settings, they are typically primary caregivers of young children (e.g., Hrdy, 2011; Luthar and Eisenberg, 2017); however, there are many instances where fathers are primarily responsible for raising children. This section reviews the importance of fathers for children's development. Following this, the extant evidence on interventions with fathers is reviewed and future needs are discussed.

While some research shows that fathers experience exclusion regarding parental involvement in child health and development (Plantin et al., 2011; Steen et al., 2012), there is empirical evidence that suggests they have an integral role in promoting the health and well-being of their children. Some evidence points to a direct relationship between fathers and specific childhood outcomes. For example, research suggests that father education and supportiveness shapes language and cognitive development in children (Cabrera et al., 2007). Furthermore, varying levels of father involvement or connectedness with children are associated with differential self-regulation and aggression in children (Vogel et al., 2006). One study has even found that positive father–child relationships, as reported by fathers, are protective from pediatric injury (Schwebel and Brezausek, 2010). In terms of father well-being, paternal depression has been associated with increased odds of child neglect (Lee et al., 2012).

Other evidence suggests an indirect pathway by which fathers can influence their children's well-being—through their mothers. Bloch et al. (2010) found that poor-quality relationships between fathers and mothers are associated with depressive symptoms, stress, substance use, and smoking among low-income mothers. Other research indicates that father involvement has a positive association with a mother's health behaviors (e.g., seeking prenatal care in the first trimester, abstaining from drinking, drug use, and smoking) (Teitler, 2001). Finally, fathers can support maternal well-being and birth outcomes. For mothers, a supportive relationship with the father can have buffering effects on maternal depression and preterm birth (Giurgescu et al., 2018).

Involving Fathers in Preventive Interventions

As noted earlier in this chapter, developmental scientists have increasingly emphasized that it cannot be presumed that what helps fathers do well as parents is the same as what most benefits mothers (e.g., Grusec and Davidov, 2010). To illustrate, Lamb and his colleagues have shown that fathering can be more affected by the quality of the marital relationship (levels of satisfaction and support) than is mothering (see Lamb, 2004; Lewis and Lamb, 2003). Accordingly, it is important to attend to the gender-specific needs of parents in designing strategies to recruit and retain them in prevention programs related to their families and children (Pruett et al., 2017).

Cross-national interventions further attest to the critical importance of attention to gender-specific needs of parents. Randomized trials with families affected by HIV indicated that across countries (United States, Thailand, and South Africa), men were more likely to respond to stress with flight-or-flight responses and women with tending and befriending (Rotheram-Borus et al., 2011). Accordingly, there have been efforts to design interventions to accommodate these

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gender differences in both delivery formats and activities. Community-based "mentor mothers," for example, were most beneficial for mothers seeking help via interventions, whereas men were more likely to be engaged via group activities, such as sports or vocational training. Similarly, Cabrera et al. (2015) note that fathers may be more willing to participate in a "game of basketball" than the "morning coffee" that many early childhood programs offer, and they recommend running support groups for "fathers" rather than just for "parents," sending the message that the role of a father is valued and is supported, independent of support to mothers.

In their review of existing programs for fathers, Pruett and colleagues (2017) indicate that whereas it is generally more difficult to engage fathers than mothers in preventive intervention programs, there are some avenues that seem promising. These include efforts to recruit fathers along with mothers via couples' work in early childhood (i.e., while the father is still involved with the mother and child, before any possible estrangement, as it is much more difficult to engage fathers when strife has entered the inter-parent relationship). Pruett et al. (2017) also suggest focused efforts to make the facilities themselves more "father friendly" (e.g., by addressing implicit biases of staff that fathers will intrinsically be uninvolved and portraying pictures of fathers with babies in the program facilities). At the most basic level, other approaches would be to highlight in the media those men who are positive role models of fatherhood early in life and even bring such individuals to high schools to talk with young men on this topic, well before they become fathers.

Davison et al. (2019) examine strategies for and barriers to father engagement in early obesity prevention during the child's first 1,000 days of life. The review finds that although obesity prevention programs have great potential to engage fathers, there are a number of micro- and macro-level barriers that hinder access and participation. Applying a policy, systems, and environmental perspective, the authors recommend the following strategies to enhance father engagement: integrate father-engagement standards into national practice guidelines (e.g., AAP, ACOG), develop educational and marketing materials (e.g., pamphlets) specific to fathers, and partner with local father advocacy organizations to identify new strategies to engage fathers. Other research has homed in on preconception (Kotelchuck and Lu, 2017) and pediatric care (Yogman and Garfield, 2016) as important platforms for engaging fathers.

What is clear is that speaking about promoting good parenting in gender-neutral terms will likely be a disservice to both fathers and mothers. The literature clearly delineates what mothers need most centrally to function well as mothers (again, aside from basic necessities for physical survival): authentic connections with and support from others in their everyday lives. Relationship-based strategies that are highly effective, such as those in home visiting programs for new mothers, will not necessarily generalize to new fathers, paving the way to their enhanced mental health, positive fathering behaviors, and attachment to the child. For example, in low-income African American families, mothers responded well to the tend and befriend approach; however the recruitment of biological fathers was more likely through opportunities for enhanced earnings to support their families (Jackson, 2015). Much more work is needed, therefore, on ascertaining how best to engage fathers and retain them in prevention and intervention programs.

It is important to note that NIH changed the policies about gender inclusion around the 1990s, when there was a concerted move to include women in intervention trials. The NIH Office of Research on Women's Health was formed in 1990 "in response to congressional, scientific and advocacy concerns that a lack of systemic and consistent inclusion of women in NIH-supported clinical research could result in clinical decisions being made about health care

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for women based solely on findings from studies of men—without any evidence that they were applicable to women" (Mazure and Jones, 2015). As researchers and practitioners think of how best to promote the well-being of young children via effective support for their primary caregivers, it would be prudent to avoid presuming that we can simply transfer program strategies and procedures from mothers to fathers. Although instances where a father is solely responsible for the child are not as common, this in no way obviates the need for understanding the most effective paths to reach and benefit these fathers. One evidence-based program that provides an avenue to reach fathers along with mothers is the Family Check-up (FCU) program (see Box 4-6 for a description of FCU as a promising model).

BOX 4-6
Family Check-up Program: Promising Model[a]

The FCU is a strengths-based intervention that reduces children's problem behaviors by improving parenting and family management practices. Stretching back over 20 years, the FCU integrates assessment with motivation-enhancement strategies to tailor intervention goals to meet the unique needs of each family and increase family engagement. It has been implemented in a variety of settings, including public school systems, American Indian tribal communities, publicly funded health centers, hospitals, and community mental health centers. Any professional who provides services to families with children can use the FCU.

Using a strengths-based, ecological, adaptive framework, the FCU focuses on children ages 2 through 17. It is based on a family-centered model derived from "basic research on social development and intervention science" (Dishion and Kavanagh, 2003, p. 3) and differs from traditional family-centered approaches by emphasizing methods to promote a family's motivation to change (Shaw et al., 2009). By providing parents with the tools that they need to manage their children's behaviors effectively and to build a strong and positive relationship with their children, FCU addresses the social environment as a social determinant of health. The program is multisector, engaging state research institutions, professional caregivers, mental health professionals and institutions, public schools, the Institute of Education Science, and NIH. Several studies have shown that the FCU is effective in achieving positive outcomes for families. For example, outcome data suggest that compared with the children of families who never participated in the FCU, children of families that did participate have fewer behavior and emotional problems, better emotion regulation, increased school readiness, and decreased risk for obesity (Brennan et al., 2013; Dishion et al., 2008, 2014; Lunkenheimer et al., 2008; McEachern et al., 2013; Shaw et al., 2006, 2009).

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

SOURCES: ASU Reach Institute, 2018; The Center for Parents and Children, n.d.

Reviews of past national-level efforts targeting fathers further attest to the importance of further work in this area. As Lu and colleagues (2010) have noted, many fathers in the United States face substantial barriers to staying involved in their children's lives, and this is especially the case for many low-income fathers and men of color. These barriers operate at multiple levels, including intrapersonal (e.g., poor knowledge, attitudes and behaviors toward parenting) and interpersonal (e.g., strained relationship with the mother or maternal grandmother), neighborhoods and communities (e.g., high unemployment and incarceration rates, which influence social norms), cultural or societal (e.g., popular cultural perceptions of black fathers as expendable and irresponsible, racial stratification and institutionalized racism), policy (e.g., the "marriage penalty" within Earned Income Tax Credit [EITC] and Temporary Assistance for Needy Families [TANF], child support enforcement), and even historic (slavery and forced

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family separation among African Americans and forced removal of American Indians from their land) (Lu et al., 2010). Life course factors, such as adverse childhood experiences (ACEs) or the absence of the father's father growing up, can also influence father involvement. As discussed in the next section, institutionalized racism against men of color leads to higher rates of incarceration that precludes many fathers from being present in their children's lives, establishing healthy relationships, and providing the necessary support to the mother.

In recognition of both the importance of fathers for child development, and the challenges fathers face, Congress has authorized and funded grants for fatherhood programs for more than a decade. Evaluation of four federally funded Responsible Fatherhood programs from 2011–2018 has yielded mixed results (Avellar et al., 2019). All four programs consisted primarily of group-based workshops covering topics such as the meaning of fatherhood, child development, co-parenting, and finding and retaining employment and even personal development topics, such as coping with stress, responding to discrimination, problem solving, self-sufficiency, and goal planning. Participation in the programs resulted in improved parenting, specifically their self-reported nurturing behavior and engagement in age-appropriate activities with children, but not in-person contact, financial support, co-parenting, earnings, or socio-emotional and mental well-being (Avellar et al., 2019).

One possible explanation for the mixed results is that most extant fatherhood programs still do too little, too late. Men's capacity to support and nurture others needs to be cultivated over their entire life course, not only after they become fathers (Lu et al., 2010). This includes their capacity to form stable, positive relationships; as discussed throughout this report, this depends on the presence of a stable, nurturing relationship with a caregiver during their own early childhood. Thus, future research on father involvement needs to take a more integrative, life course approach, studying how boys become men, men become fathers, and we as a society can better nurture the development of fatherhood across the life course.

> *Conclusion 4-3: There is an urgent need to develop interventions well suited for fathers and other male caregivers; existing approaches that are developed for and tested with women cannot be assumed to generalize to other caregivers, with equal effectiveness, in the successful recruitment, retention, and support of men and fathers who take care of young children in prevention programs.*

Future fatherhood programs also need to go beyond affecting individual knowledge, attitudes, and behaviors toward changing the family, neighborhood, social and cultural contexts of fatherhood. Public policy changes to remove barriers—such as the "marriage penalty" within TANF and EITC, making child support more "father friendly" by calculating payment based on actual earnings, adjusting and forgiving arrearages, and allowing a greater amount of the child support payment to be passed through to the child—can go a long way toward strengthening father involvement. Micro- and macroeconomic policies to reduce poverty and strengthen fathers' capacity to contribute financially to the family can also help. Most importantly, fatherhood initiatives, if they are going to succeed in advancing health equity, need to have a core objective that addresses institutionalized racism in employment, education, housing, health care, criminal justice and other systems that keep many low-income fathers and men of color from living out their true potentials in our society.

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Recommendation 4-1: Federal, state, and local agencies, along with private foundations and philanthropies that invest in research, should include in their portfolios research on the development of preventive interventions that target fathers and other male caregivers. Special attention should be given to the recruitment, retention, and support of men and fathers parenting young children from underserved populations.

The Role of Culture as Context

Anthropological perspectives on the study of children's development, pioneered by Margaret Mead, John and Beatrice Whiting, Robert A. Levine, and others, emphasize the importance of the cultural context that embeds children, families, and institutions that are important determinants of children's development. Subsequent elaborations by Super and Harkness (1986) labeled these influences the "developmental niche." The developmental niche consists of "(1) the physical and social settings in which the child lives [micro systems in Bronfenbrenner's (1979) work], (2) the culturally regulated practices of child care and child rearing [that exist in all microsystems, including family, education, and health care settings], and (3) the psychology of the caretakers" (p. 552). They expand as follows:

> "Regularities in the subsystems, as well as thematic continuities from one culturally defined developmental stage to the next, provide material from which the child abstracts the social, affective, and cognitive rules of the culture, much as the rules of grammar are abstracted from the regularities of the speech environment. The three components of the developmental niche form the cultural context of child development" (p. 552).

More recent formulations have refined these initial theoretical postulates to articulate more specifically how culture permeates the settings that children navigate and how they impinge on development. Vygotski (1978), Rogoff (2003), and Weisner (2002) describe culture as embedded in daily routines and interactions by which the child is enculturated into patterns of behavior, language/communication, and cognitions. A recent revision of Bronfenbrenner's bio-ecological model emphasizes how settings or microsystems through social interactions, daily routines, space, and other resource allocations enact on a daily basis cultural belief systems that may or may not align on behalf of the children's development (Vélez-Agosto et al., 2017). Service providers and caregivers need to be cognizant that practices that may be perceived or interpreted as harmful by other groups are embedded within a complex cultural system with its own deep embedded logic. (See the section on cultural considerations for subgroups for specific examples.) For that reason, cultural differences need to be accepted, incorporated, or renegotiated as part of interventions and educational efforts. Traditional cultural practices, as unfamiliar as they might seem for some, might be sources of strengths and resilience.

Subgroup Differences

Although there are universal processes of development that point out the importance of the quality of intimate family relationships for early child development, there can also be powerful influences that are largely unique to particular contexts; these potent *within-group* processes can

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affect children in both positive and negative ways. Considering the former, for example, specific processes have been identified as strengths or natural coping mechanisms that foster resilience within particular racial or ethnic groups, such as familism and respect for family obligations among Latinos and other collective-oriented cultures and effective family racial/ethnic socialization. Familism and family obligations provide the bases for family members to be involved in crisis management and the daily needs of mothers and children. Racial socialization provides children of color with an age-appropriate understanding of how racism operates and instills strengths and coping mechanisms based on positive and strong ethnic pride and identity. At the same time, these family contexts can place unique demands on children and families and lead to obstacles if they are not understood or accepted or are undermined by the majority culture; unintentionally, they might contribute to the emergence of health disparities very early in life.

This section describes subgroups in which there are specific, unique risk and protective factors that can affect the quality of parent–child relationships and the child's development; these need to be taken into account as interventions are designed to address health disparities. It should be emphasized here that ensuing discussions of subgroups does not imply that "all are the same" within them. There is invariably great heterogeneity between individuals and families within each subgroup, and it is important to recognize this, particularly in the design and implementation of interventions. Furthermore, while there may be unique risk and protective factors ascribed to these groups, the existing intervention research does not provide one-size-fits-all approaches to guide programs and policies working to serve these subgroups. These subgroups can differ from mainstream groups due to a variety of reasons, including

- Differential access to critical health-promoting resources (e.g., families not having access to safe housing and neighborhoods, preventive care, appropriate educational opportunities),
- Cultural differences due to language (e.g., non-English speakers in a monolingual English environment), religious background (e.g., Muslims) or recent migration,
- Chronic stress (e.g., groups in poverty struggling to meet basic needs, such as shelter and health care, and those facing excessive pressures to achieve, usually in well-educated affluent communities), and
- Discrimination (e.g., racial or ethnic discrimination, homophobia or discrimination against gender-atypical people [see Box 4-7 and the section on Lesbian, Gay, Bisexual, Transgender, Queer [LGBTQ] youth and families]).

BOX 4-7
Children of Same-sex Parents

In the last two decades, developmental researchers have provided a wealth of evidence that children of same-sex parents show patterns of adjustment that are comparable to those of heterosexual parents. Pioneered by Charlotte Patterson in the 1990s in response to heterosexist and homophobic questions in psychology, judicial opinions, and popular stereotypes, this burgeoning research in child development has shown that it is the quality of parent–child relationships, rather than having same- versus different-sex parents, that is critical for children's adjustment (e.g., Wainright and Patterson, 2008; Wainright et al., 2004). A meta-analysis of 19 studies confirmed that children raised by same-sex parents have patterns of adjustment that are just as healthy as those of their counterparts raised by heterosexual parents (Crowl et al., 2008).

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Children Living in Poverty: Meeting Basic Needs

As noted earlier, the most fundamental needs in Maslow's hierarchy are related to physical survival; when parents struggle to have their basic human needs met—for shelter, food, and health care—all in the family are at risk. Parents who are preoccupied by fulfilling basic needs for themselves and their children inevitably find it challenging to attend to their children's emotional, behavioral, and educational needs. Living in poverty is stressful on parents and can compromise the quality of their parenting behaviors, the evolving parent–child relationship, and the well-being of their developing children.

As discussed in Chapter 3, one of the most studied and documented factors that affects children adversely is living in poverty (Evans, 2004). Beyond physical survival, chronic poverty places children at risk because the physiological, safety, relational, self-esteem and self-actualization needs—including those of the parents and the child—are all jeopardized; all of these need to be minimally fulfilled to prevent health disparities. Low socioeconomic status (SES) and parental education are associated with limited access and use of critical health-promoting resources, and they contribute to psychological distress and physical health disparities. Research has consistently shown that poverty creates living conditions that negatively affects parenting and the mother–child relationship. The daily stresses associated with living in poverty lead to higher incidence of maternal depression, which, as noted earlier, is linked with a poorer quality of mother-infant interaction (Jackson, 2000; Taylor and Conger, 2017). Similarly, poverty has been linked to less capacity for parenting in a consistent and loving way, since it creates high levels of psychological distress that interfere with successful parenting (Conger and Conger, 2002; McLoyd, 1990). Families living in poverty with young children represent one of the most at-risk populations for health disparities, and programs and policies targeting them (see Chapter 6 for more on economic, food, and housing security) within a prevention and early intervention framework should be a priority for our nation.

Cultural Considerations for Subgroups

Other groups whose family relations might be at risk are those whose cultural background (i.e., lack of familiarity or congruency with the dominant North American culture and language) limits their access to critical health-promoting resources or can become an obstacle as they operate with different daily practices and routines that are different from that dominant culture. For example, a parent with a limited grasp of the English language might impede that parent's consideration of high-quality early intervention or preschool programs (Moss and Puma, 1995). They might also espouse different family constellations (e.g., where grandmothers are very involved in caretaking); these other adults might interfere with their family operations, as might external influences, such as educators or health care providers.

Differences in acculturation among family members can also be sources of risk and resilience for families of immigrant backgrounds or members of other nondominant cultures. It has been observed that as children acculturate more than their parents (i.e., become more familiar with dominant North American culture and language), there is more conflict between parents and children (Schwartz et al., 2012). In addition, children might assume parenting tasks, such as taking on adult roles, negotiating school systems, and communicating with other officials for services (drivers' licenses and bank accounts, for example) for their parents.

The evidence on the effects of these added responsibilities on the children's development, so-called family obligations or brokering, is mixed: in some contexts, they might be burdensome,

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while in others they seem to be protective (Dorner et al., 2008; Fuligni and Pedersen, 2002). When taken to extremes and in developmentally inappropriate ways, however, taking on these roles is unhealthy for children. In recent years, the increased number of parental deportations and separation of parents and children at the southern U.S. border has led to an increasing number of children of immigrant background being traumatized and pushed into developmentally inappropriate parenting roles, which will lead to increasing relationship disturbances and subsequent health disparities (Bouza et al., 2018).

In other instances, traditional cultural practices that are appropriate in their original context clash with mainstream North American institutions, and these cultural practices inadvertently place children in at-risk pathways and contribute to problems in family relationships. As an example, many societies (i.e., Korean, Japanese) do not use direct verbal communications with infants and young children. Yamamoto and Li (2011) have shown that some preschools make Asian American students feel inadequate because their quietness, a desirable trait in their home culture, is considered a "problem" in the preschool environment. By contrast, acknowledging and fostering bicultural competencies in families, children, and service providers offers a good foundation for fostering early development in diverse populations (Ho, 2015; Padilla, 2008).

Finally, as presented in Chapter 3, discriminatory practices by systems, institutions, and the population at large can affect family relationships. Cultural mistrust from years of oppression from dominant groups, such as those experienced by African Americans, American Indians/Alaska Natives (AI/ANs), and immigrants of various races and nationalities in this country, might interfere with family functioning. Cultural mistrust can be transmitted intergenerationally, as promoting mistrust has been identified as a particular type of ethnic racial socialization practice (Hughes et al., 2006). These attitudes permeate relationships with others, including with family members and health care providers (Moseley et al., 2007). Mothers and other main caregivers from other cultural backgrounds might not only have problems accessing supports from mainstream institutions but also experience family conflicts because the experiences of parents and children with mainstream institutions might differ. Although these factors can be listed individually, multiple factors are present in many individuals, families, neighborhoods, and regions (e.g., rural versus urban). This is why theoretical frameworks have emphasized the multiplicity of factors that are usually present in creating health disparities (Coll et al., 1996). Moreover, new theories of intersectionality refer to those processes as differential access to power and as exclusion from benefits and assets and the existence of oppression and privilege at many societal levels (Bowleg, 2012; Brown et al., 2016; Hankivsky and Christoffersen, 2008). Among the strongest factors associated with health disparities are gender, SES, and race (Becares and Priest, 2015; NASEM, 2017; Williams et al., 2010). In many instances, these factors are multiplicative, not additive, throughout childhood and adulthood.

At the same time, the protective factors that are unique for these populations need to be considered. The "immigrant paradox" refers to the findings that across physical and mental health and in educational and occupational outcomes, children and adults of immigrant backgrounds outperform native-born children, in spite of having been poorer and less acculturated to the dominant North American culture (García Coll and Marks, 2012). This phenomenon is poorly understood: it is unclear if motivational factors impel first and second generations to excel; if acculturation to poverty and its association with detrimental lifestyles and choices places subsequent generations at risk; or if discrimination and racism is more obvious to

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subsequent generations, and therefore its pernicious effects influence subsequent generations more.

Regardless, the traditional notion of rapid acculturation and complete assimilation as the successful model for incorporating immigrants has been questioned based on this evidence, and models of biculturalism and promotion of ethnic pride (Umaña-Taylor et al., 2014) are suggested as alternatives for successful incorporation. Finally, studies point out that state- and country-level policies that support families of immigrant background lead to fewer health disparities in youth (Filindra et al., 2011; Marks et al., 2018).

Children in Foster Care and Children with Incarcerated Parents

As discussed in the previous sections, a healthy (positive) relationship with main caregivers is one of the basic cornerstones of children's development and necessary to prevent many health disparities. There are populations that, for a variety of reasons, lack the stability and quality of parenting needed to promote their children's development. Children in foster care and whose parents have been incarcerated often have multiple stressful life events before removal from parental care (Leve et al., 2012; Murray et al., 2012). The actual parental incarceration or loss of child custody is usually preceded by involvement with illegal activities and/or parental substance abuse, unstable housing, domestic violence and/or child neglect or abuse. These children might be exposed to extreme stress, particularly during the prenatal period, when the in utero child should be especially targeted at the first indication of any family relational difficulties.

Children in Foster Care Children in foster care are overrepresented as a group in health disparities (Goemans et al., 2015, 2016; Leloux-Opmeer et al., 2016; Leve et al., 2012). In most instances, these children's parents have lost or given up their parental rights because they cannot provide their children stable, healthy relationships and other critical health-promoting resources needed for their well-being. However, children from certain racial and ethnic populations are disproportionately represented in the child welfare system. For example, AI/AN families are twice as likely to be investigated as the national population of families, and cases are twice as likely to be substantiated, with AI/AN children three times as likely to be placed in foster care as the national population of children (see Table 4-1 for data on this disproportionality) (Hill, 2007). Disruptions in attachment and other regulatory processes are consistently found in this population (Barone et al., 2017; Goemans et al., 2016; Leloux-Opmeer et al., 2016; Vasileva and Petermann, 2018). However, it has been found that foster care does not ameliorate these problems and actually might lead to more negative adaptive functioning after 1 year in such placement (Goemans et al., 2015).

TABLE 4-1 Disproportionate Foster Care of AI/AN Children: 15 States with the Highest Rates

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State	Disproportionality Rate[4] 2014	% of children who are AI/AN	% of children in foster care who are AI/AN
Minnesota	17	1.4%	23.9%
Nebraska	8.4	1.1%	9.3%
Idaho	5.2	1.2%	6.0%
Iowa	4.8	0.3%	1.7%
Wisconsin	4.8	1.1%	5.1%
Washington	4.3	1.5%	6.3%
Oregon	4.0	1.2%	4.9%
Montana	3.9	9.5%	36.9%
North Dakota	3.9	8.1%	31.4%
South Dakota	3.7	12.9%	47.9%
Alaska	2.6	17.8%	46.6%
Utah	2.5	0.9%	2.3%
New Hampshire	2.3	0.2%	0.5%
California	2.0	0.4%	0.8%
North Carolina	2.0	1.2%	2.4%
Massachusetts	1.2	0.2%	0.2%
Maine	1.1	0.8%	0.9%

SOURCE: NICWA, 2017.

Policies and practices of parental removal and replacement need to be carefully implemented. The stability of nurturing and supportive relationships, as described earlier in this chapter, are crucial considerations for these populations (Jones Harden, 2004). If parents are unavailable or incapable of providing such conditions, placement with other family members, foster parents, or adoption are advisable, but only if these substitute parents can, in turn, provide stability and responsive parenting. That means that the quality and timing of care provided by family substitutes, including foster and/or kinship care and/or adoption or the reunification with the original biological family, has to be carefully planned and implemented, to avoid re-traumatizing children through their interactions with these systems (Font et al., 2018).

In terms of supportive policies, the passage of the Family First Prevention Services Act in 2018 provides an opportunity to advance evidence-based child welfare policies through the investment of federal funds in children at risk for foster care, in foster family homes, or in qualified residential treatment programs or other special settings.[3] Most notably, the legislation provides support for programs and practices to keep children with their families safely (e.g., allowing states to use Title IV-E funds to prevent children's entry into foster care) (NCSL, 2019).

Children with Incarcerated Parents Incarceration is more prevalent in marginalized populations, adding to these populations' multiple risks for poor health outcomes. Analysis of incarceration rates shows much higher rates in low-income, minority populations (Acker et al., 2019; Wildeman and Wang, 2017). It has been widely documented that discriminatory policies and practices in the definitions of crimes, arrests, sentencing, and incarceration practices lead to an overrepresentation of low-income and minority populations, which has resulted in widespread incarceration within these communities (Acker et al., 2019; NRC, 2014). These higher rates not

[3] H.R.253—Family First Prevention Services Act of 2017.

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only affect the developing mother–child relationship but also disrupt the networks of support for parents frequently found in other family or neighborhood members.

Research clearly demonstrates the negative effects of parental incarceration on the child (Christian, 2009; Wildeman and Wang, 2017). Parental incarceration usually leads to family financial and housing instability and to more mental disorders and physical health problems in the partner (the parent who stays behind), which in turn is associated with higher incidence of child abuse and neglect (Acker, 2019). Various studies, including a meta-analysis, show that parental incarceration is associated with higher risks of children's antisocial behaviors, learning disabilities, and behavioral problems (Murray et al., 2012; Turney, 2014). Long-term educational, occupational, and physical and mental health outcomes as well as criminal behaviors are all related to parental incarceration (Acker et al., 2019).

Given the importance of attachment to the primary caregiver, maternal incarceration is believed to be more detrimental for young children than paternal incarceration and a serious risk factor for future psychopathology (Murray and Murray, 2010; Parke and Clarke-Stewart, 2001; Shlafer and Poehlmann, 2010). This is concerning, given that almost 80 percent of incarcerated women are mothers (McCambell, 2005). For that reason, policies and practices that ensure adequate access and visitation to the incarcerated parent and promote healthy attachments is seen as a fundamental need and right of the child(ren) left behind (UNM, 2011).

LGBTQ Parents and Children

The parents and children that do not conform to normative gender expression and sexual orientation schemas of society (the LGBTQ community) are also placed at risk by contextual circumstances. The current cultural, religious, and political climate in the United States is divided on the acceptance and approval of these behaviors. Although research indicates that same-sex parents have been found to raise children with the similar developmental outcomes as those of opposite-sex parents (see Box 4-7), about one third of the population still expect negative impacts on children (Drake, 2013).

From an equity perspective, the LGBTQ community has historically gained rights for families through legislation and court precedence, such as the Supreme Court decision to uphold same-sex marriage as a constitutional right in 2015.[4] This landmark court case was instrumental in securing marital benefits (e.g., financial, social) for same-sex couples and their children. As of 2016, 20 states and DC had enacted legislation that prohibits discrimination based on sexual orientation and gender identity or expression in employment, housing, and public accommodations—all are critical to promoting healthy development for children.

Even with such laws and policies, members of these populations are often subjected to discriminatory acts and microaggressions (IOM, 2011). Some have to "hide" their orientation and expressions from family, friends, and coworkers, which sometimes lead to the adoption of high-risk behaviors, such as unprotected sex, or internalizing disorders, such as depression, self-injurious behavior, and anxiety (IOM, 2011). Others may passively accept or actively repudiate discriminatory acts, exclusions, verbal and physical aggressions, and microaggressions as they express and adopt these less normative behaviors. These varied reactions to homophobia and discrimination can be associated with high levels of chronic stress and contribute to long-lasting health disparities (IOM, 2011).

[4] *Obergefell v. Hodges*, 576 U.S. (2015).

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While sexual orientation and gender identity may be more salient during adolescence, there is evidence to suggest that LGBTQ children demonstrate their inclinations to gender-atypical behaviors and same-sex orientation quite early (Rae et al., 2019). Many parents identify cross-dressing, toy preferences, and identification with heroes/heroines as displaying these preferences as early as the infancy and toddler years. As the typical puberty age has dropped, the issues that used to arise later in adolescence (i.e., identity, dating) are moving into childhood.

The LGBTQ population of children and youth is overrepresented in the adoption of risky behaviors and psychopathology. The IOM (2011) report on The Health of Lesbian, Gay, Bisexual, and Transgender People identified the following key findings from the literature about LGBTQ youth during childhood and adolescence:

- LGBTQ youth are at increased risk for suicidal ideation and attempts a depression.
- Rates of smoking, alcohol consumption, and substance use may be higher among LGBTQ than heterosexual youth.
- The homeless youth population contains a disproportionate number of LGBTQ youth. Some research suggests that young transgender women are also at significant risk for homelessness.
- LGBT youth report experiencing elevated levels of violence, victimization, and harassment compared with heterosexual and non-gender-variant youth.
- Families and schools appear to be two possible focal points for intervention research.

These findings are corroborated by data from the Youth Risk Behavior Survey, which identify LGBTQ youth as being at risk for bullying, intimate partner violence, and sexual assault (CDC, 2017).

Children in High-Achieving Schools

As noted at the outset of this section, the standouts in groups of children and families exposed to high levels of chronic stress include those at both extremes of SES. Evidence of the challenges facing children and families in poverty, summarized here, have been reviewed extensively in the literature, with studies burgeoning around publication of the landmark Special Issue of *Child Development* on poverty over 20 years ago (Brooks-Gunn and Duncan, 1997; Elder, 2018; Huston et al., 1994). By contrast, the evidence showing that youth in high-achieving contexts are an at-risk population is nascent, and the findings are counterintuitive for most. Extant evidence of adjustment difficulties among these children is presented here, following by consideration of why this subgroup might merit attention in this particular report, which is focused on family-based prevention for children known to be at risk given life adversities.

With regard to the existing evidence on children of well-educated, relatively affluent parents, studies using varied samples and methods have converged in indicating relatively high levels of adjustment problems, likely linked with long-standing, ubiquitous pressures to excel at academics and extracurricular activities. Studies of students attending high-achieving schools (HASs) (public and independent; day and boarding; in different parts of the country) have consistently determined that they show rates of clinically significant problems, notably depression, anxiety, rule-breaking, and substance use, at rates that are (a) much higher than national norms and (b) sometimes higher than those in urban poverty (the latter is particularly true for substance misuse; see (Luthar and Kumar, 2018; Luthar et al., 2013). Two large-scale analyses of national datasets in the United States and Norway (Coley et al., 2018; Lund et al., 2017) have confirmed a U-shaped association between community-level affluence and students'

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adjustment problems; in fact, students at the more affluent schools had the highest levels of substance use.

In a recent longitudinal study spanning 50 years, Gollner et al. (2018) showed that students at an HAS had *poorer* outcomes decades later than their counterparts at middle-achievement schools. The authors said that constant social comparisons among a population of largely talented students were likely implicated in this result, which led to lower educational expectations of the self. Similarly, a collection of three studies involving fifth- to tenth-graders in Germany (Pekrun et al., 2019) showed that in all three schools, being in a high-achieving context, had negative links with self-concept and emotions, and conversely, being in less-HASs was better for self-concept and emotion.

Over time, accumulating evidence on this population has led to global statements about the high levels of risk among these youth. Almost a decade ago, Koplewicz et al. (2009) referred to their problems as "an epidemic." A recent report on adolescent wellness, released by the RWJF, noted the following four as the most critical "high-risk" environments: exposure to poverty, trauma, discrimination/racism, and high pressure to achieve, usually found in relatively affluent communities (Geisz and Nakashian, 2018).

Importantly, risks can be further exacerbated for vulnerable populations in HAS settings. Recent research has shown that African American boys in relatively high SES communities reported significantly higher depression and substance abuse, probably as a result of greater discrimination/racism in their communities—likely dominated by white families (Assari and Caldwell, 2018; Assari et al., 2018a,b; Lewis and Van Dyke, 2018). There is value, therefore, in considering issues of racial socialization for children of color who have the "double jeopardy" of having few children who look like them in their schools, neighborhoods, and communities.

With regard to why this subgroup of children and families might be relevant for this report on early childhood and equities in adaptation, two factors are relevant; one has to do with "sleeper effects" and the other with the need for early prevention. With regard to the first issue (as has been discussed earlier), the charge of this committee is not just to prevent problems that are manifested in early childhood—temper tantrums or bed-wetting—but, equally, those whose seeds are sown in early childhood and then "incubate" and become evident by early adolescence. It is clear that the prevention of problems among HAS youth (and other youth who suffer poor outcomes due various types of adversity) cannot start when they are already teens; by then, a great deal of damage has already been done. Instead, what is needed is attention to these issues among parents when the children are still young. In fact, there are recent suggestions that the effects of high-achieving environments can be seen as early as preschool (Pekrun et al., 2019).

Second, there is a great deal of preventive potential in working with families in these settings toward raising awareness of the nature of risks and potential problems. Unlike low-income communities, where an influx of resources is often needed for prevention, in this case, dissemination of accumulated evidence from science, with associated recommendations (Luthar et al., 2013), can be helpful. It is particularly important for parents to (a) understand the level and seriousness of adjustment problems that researchers have recurrently documented in HASs and (b) be vigilant in their own homes, starting from early childhood, against being overly invested in the child's "resume-building" (Gollner et al., 2018; Luthar and Kumar, 2018 Pekrun et al., 2019).

It is also critical to note that just as children in high-achieving contexts can be vulnerable, so can their mothers (who are, again, as in most contexts, their primary caregivers). Panel studies over time have shown that well-educated mothers have shown substantial increases in hours

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spent on activities related to their children, with increases greater than those among well-educated fathers and less well-educated mothers or fathers (Kalil et al., 2012; Ramey and Ramey, 2010). In terms of sheer hours, therefore, this group can be especially stretched thin and chronically stressed. Cross-disciplinary studies have established other factors that can challenge their well-being, including the "invisible labor" involved in being responsible for overseeing schedules and activities of all in the household (see Ciciolla and Luthar, 2019; Lareau and Weininger, 2008; Yavorsky et al., 2015).

A recent study showing increases in depression over the last decade in the high-income groups, and among women in particular, provides additional testimony to the seriousness of these issues (Twenge et al., 2019). Changes over time were examined for four cohorts/generations and considering four groups of family incomes. The findings showed that cohort increases in serious psychological distress were the largest in the highest income group. Among adolescents, time period increases in rates of a major depressive episode were the largest in the highest income group and smallest in the lowest-income group; among adults, the cohort increase in suicidal ideation was largest and smallest in the highest-and lowest-income groups, respectively. The authors concluded that "with the exception of adult MDE and suicide attempts, the cohort increases were the largest and most consistently seen among those with the highest income" (Twenge et al., 2019, p. 10). Additionally, among adults, the increases in mood disorder indicators and all suicide related variables (suicidal thoughts, plans, and attempts) were consistently higher among women than men.

The evidence on youth and families in high-achieving contexts were reviewed above in some detail because this is a newly identified at-risk group, encompassing at least 20–25 percent of the U.S. population (e.g., Twenge et al., 2019). Additionally, despite the seriousness of problems that are increasingly documented, there remains reluctance to afford attention to them in research and policy (Geisz and Nakashian, 2018).

In addition to HAS families, there are other such groups that are increasingly being identified as showing high vulnerability relative to norms, such as subgroups based on sexual identity and preference. Questions on both these dimensions are now arising among children at younger and younger ages (often well before adolescence), creating stress and confusion for the child as well as parents. Children who identify as neither male nor female but rather as "nonbinary" show elevated rates of anxiety, depression, and other psychological difficulties, as do children who identify not as heterosexual but as lesbian, gay, bisexual, or questioning. Regarding transgender youth, considerations of gender reassignment are also now coming up in many families well before adolescence. All of these subgroups will need concerted attention in the years ahead, in terms of understanding how best to help both children and parents effectively negotiate these issues at home and outside it, such as in the children's peer groups.

Youth from Asian backgrounds are yet another illustration of a group that is potentially vulnerable but rarely included in discussions of children at risk. These children are often thought of as both privileged and "model minorities" (e.g., Lee et al., 2009; Ngo and Lee, 2007), but in fact, many U.S. families of Asian origin actually live in low-SES households. Further, issues of discrimination affect Asian youth, arguably, just as they do other racial and ethnic minorities. At the same time, Asians are not considered as an ethnic minority group in discussions of discrimination, prejudice, and stereotype (Wong and Halgin, 2006) the way that youth from African American, Latino, or AI/AN backgrounds are. It is important to emphasize that this is not a zero-sum game; research and policy attention to the needs and challenges of Asian

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Americans by no means implies detracting from the very real and well-documented challenges faced by other ethnic minority groups in the United States.

In addition to including such neglected subgroups in discussions of those needing attention to prevention, more attention to risk processes that were not an issue at the time that *From Neurons to Neighborhoods* (NRC and IOM, 2000) was released is also warranted, and social media is critical in this regard. In discussions of why there continue to be increases in children's depression since the early 2000s, social media is often noted as a major factor (Hoge et al., 2017). Yet, there remain several questions about which aspects of social media use are truly damaging to children's mental health. Possibilities range from the amount of time spent on social media to spreading rumors about others, being the victim of bullying, "sexting" and accessing pornography, and constant social comparisons or feeling "less than" others. In Chapter 3, there is a section devoted to the problem of bullying even among young children; clearly, there is a lot of work that still remains to reduce this major risk process—and, now, to also disentangle and address the new challenges introduced by the widespread use of social media from young ages onward. As Weisz and colleagues (2019) note, the nature of childhood disorders may be changing faster than our treatments are: "Threats to youth mental health are becoming more diverse and multiform…. Current threats encompass pressures to excel as well as new pressures to excel in increasingly competitive academic and social environments, images conveyed via advertising and social media that could make anyone feel inadequate, risks of harm via text messages and cyberbullying, and even fear of being gunned down at school" (p. 233).

To summarize, as policy makers and prevention scientists think about groups of young children and caregivers whose life circumstances are likely to place them at high risk for adjustment problems, the needs of all families need to be taken into consideration. Low levels of maternal education, coexisting with poverty and often single motherhood, clearly spell risks for adjustment problems and parenting difficulties among mothers, as noted earlier. At the same time, there are other groups that are increasingly being identified as requiring attention in future prevention efforts, including children and parents in extremely fast-paced, stressful, competitive communities, and sexual minority youth. Similarly, there are new risk processes that have arisen in the last couple of decades that potentially have strong negative effects on children and merit concerted attention in future research and policy when considering what should and should not be prioritized in preventive interventions.

In conclusion, psychologists, educators, and other health professionals need to work with groups of families that are especially vulnerable due to exposure to high levels of stress or adversity, with efforts beginning from preconception onward. These efforts will involve harnessing protective processes, and minimizing risk factors, in interventions that have a sound basis in developmental theory, science, and cumulative research evidence; have shown promising results in prior prevention studies; and are pragmatic (i.e., can feasibly be taken to large scale in the real world).

> *Conclusion 4-4: Specific subgroups of children have unique needs and challenges when adjusting to adversity. Careful attention to potent subculture-specific processes need to be considered in working with subgroups well known to face serious inequities in relation to mental health—including families experiencing chronic poverty, immigrants, LGBTQ children and LGBTQ parents, and those where children are separated from parents due to incarceration or foster care. Among the groups more recently identified as being at*

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high risk for chronic stress are children and parents, particularly mothers, in high-achieving, relatively affluent communities.

Recommendation 4-2: Federal, state, local, tribal, and territorial agencies, along with private foundations and philanthropies that invest in research, should include in their portfolios research on the development of interventions that are culturally sensitive and tailored to meet the needs of subgroups of children known to be vulnerable, such as those living in chronic poverty, children from immigrant backgrounds, children in foster care, and children with incarcerated parents.

In addition to addressing major goals relevant for children in general (e.g., fostering caregiver well-being and minimizing maltreatment), programs need to include components that specifically address unique, powerful risk and protective processes within these subgroups of children. As discussed above, it cannot be assumed that evidenced-based interventions and promising models developed in specific populations will work in other contexts. The validity and adaptability of interventions need to be empirically tested in groups that differ greatly in the sources of chronic stress and cultural backgrounds.

INTERVENTIONS TO PROMOTE RELATIONAL SUPPORTS AND CAREGIVER WELL-BEING

As developmental scientists have emphasized the critical importance of relational supports for the well-being and thus mothers' parenting, intervention trials have shown strong benefits of relational interventions. Indeed, the core components of several effective interventions suggest that over and above learning particular skills, it is improvements in mothers' overall well-being, within the context of supportive relationships, that is the most critical "engine" underlying the positive changes. To illustrate, the Incredible Years intervention by Webster-Stratton and colleagues (2001) targets children's school readiness and parenting skills while helping parents to cope with personal problems; parents are able to form valuable support networks with "buddies" from parent groups (Borden et al., 2010). Ammerman and colleagues' (2005) Every Child Succeeds program for depressed mothers provides in-home cognitive behavior therapy and a supportive home visiting component. A cardinal component in Dozier's Attachment and Biobehavioral Catch-up (ABC) is consistent respect, warmth, and support displayed by project staff or therapists to participating mothers (Dozier et al., 2005).

This section focuses on interventions with the strongest evidence base to foster healthy child development through supporting caregivers. First, there is a discussion of the evidence on home visiting, a well-established model for providing services to children and families, and the following section describes a range of interventions that have shown promise. The committee presents these as opportunities to use the science to drive action on caregiver and child well-being; however, the presentation of these interventions does not indicate prioritization of one over another. Different populations will require varying interventions based on their needs and context, in addition to specific tailoring to meet these unique needs depending on the population.

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Home Visiting Programs

One of the earliest program models built around the notion of supporting caregivers is home visiting. Home visiting programs in the United States focus on promoting child health and development by providing parenting support and other child-centered services in the home (e.g., developmental screenings and referrals to services and treatment) to at-risk families from the prenatal stage to through age of six (HRSA, n.d.). One of the primary goals of home visiting is to improve child health and educational outcomes. Home visiting programs have provided services to families in the United States since the late 1800s (Minkovitz et al., 2016), but the 2010 federal Maternal, Infant, and Early Childhood Home Visiting Program (MIECHV) led to the expansion and supported standardization of home visiting implementation and evaluation. In fiscal year 2018, the MIECHV program served approximately 150,000 parents and children in 76,000 families across all 50 states, DC, 5 territories, and 23 tribal and urban AI/AN communities, providing nearly one million home visits annually and 5.2 million home visits between 2012 and 2018 (HRSA, n.d.). By law, state and territory awardees have to spend the majority of their MIECHV program grants to implement evidence-based home visiting models, with up to 25 percent of funding available to implement promising approaches that will undergo rigorous evaluation.[5] Tribal home visiting programs are encouraged to select evidence-based home visiting curricula, but as the research evidence is limited, this is not a program requirement. Most tribal home visiting programs chose to implement evidence-based curricula with cultural adaptations developed with intensive community input and engagement, which enhanced family engagement and retention (Pratt and Chapman, n.d.). Home visiting services may also receive support from many other federal, state, and local public and private sources.

This section summarizes the evidence on home visiting, highlighting a few exemplar programs. Given the multidisciplinary and cross-sector nature of this topic, the chapters later in the report will discuss home visiting with respect to specific levers in certain domains to enhance or support such programs. (See Chapters 5 and 7 for more on the role of the health system and of ECE, respectively.)

Summary of the Evidence on Home Visiting Programs

A growing body of evidence suggests that home visiting by a nurse, a social worker, or an early educator during pregnancy and in the first years of a child's life improves a whole host of child and family outcomes, including promotion of maternal and child health, prevention of child abuse and neglect, positive parenting, child development, and school readiness. The positive impact continues well into adolescence and early adulthood. For example, in a 19-year follow-up of a randomized controlled trial (RCT), Eckenrode et al.(2010) found that relative to the comparison group, girls in the pregnancy and infancy nurse-visited group were 58 percent less likely to have been arrested and 43 percent less likely to have been convicted. Nurse-visited girls born to unmarried and low-income mothers also had fewer children and less Medicaid use.

One meta-analysis examined the impact of 18 implementation factors (including staff selection, training, supervision, and fidelity monitoring) and four study characteristics (publication type, target population, study design, comparison group) in predicting program outcomes (Casillas et al., 2016). The greatest program effects were found to be on increased

[5] See https://mchb.hrsa.gov/maternal-child-health-initiatives/home-visiting-overview for more information (accessed June 8, 2019).

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positive parenting and reduced likelihood of maltreatment; the former had the largest effects. The programs were significantly less effective for children's health and behavior. Further, provision of high-quality supervision and fidelity monitoring were key factors associated with beneficial outcomes. Programs serving families with one or more risk factors had significantly greater effect sizes compared to programs serving all families.

Avellar and Supplee (2013) reviewed the research on home visiting, and nearly 9 out of 12 programs (of the 32 reviewed) that met evidence-based criteria also showed favorable effects on children's social-emotional development or behavioral problems. Five programs showed one or more favorable effects on cognitive development, and five showed improvements in reductions in some aspect of child maltreatment (only six of the programs assessed maltreatment).

Another meta-analysis examined the effect of home visiting on six outcomes: birth outcomes, parenting behavior and skills, maternal life course, child cognitive outcomes, child physical health, and child maltreatment (Filene et al., 2013). Mean effect sizes were significant and positive for three of the six outcome domains: maternal life course outcomes, child cognitive outcomes, and parent behaviors and skills (Filene et al., 2013). Effect sizes for birth outcomes were significantly larger for programs that used nonprofessional home visitors, matched clients and home visitors on race and/or ethnicity, or included problem solving.

Effect sizes were larger for parent behaviors and skills programs that addressed parental substance use and taught parents developmental norms and appropriate expectations, discipline and behavior management techniques, and responsive and sensitive parenting practices. Improved child maltreatment outcomes were associated with problem solving and teaching parents how to select alternative caregivers for children. The effects on children's cognition were better in programs that taught parents responsive and sensitive parenting practices or "required parents to role-play or practice skills during home visits. Using professional home visitors was a significant predictor of better child physical health outcomes, as was teaching discipline and behavior management techniques" (Filene et al., 2013, p. S104).

Following a review of the research literature in 2018 (Sama-Miller et al., 2018), the Home Visiting Evidence of Effectiveness (HomVEE) project, established by the Administration for Children and Families (ACF) in the Department of Health and Human Services (DHHS), identified 20 home visiting models as meeting criteria of effectiveness to recommend for state-based home visitation programs (OPRE, 2018). Based on the available high- or moderate-quality studies, the review showed that most of these programs have multiple favorable effects with sustained impact, with Healthy Families America and Nurse-Family Partnership (NFP) showing favorable impact across the greatest breadth of outcomes. Most models had favorable impacts on primary measures of child development, school readiness, and positive parenting practices; none of the models, however, showed impacts on a primary measure of reductions in juvenile delinquency, family violence, and crime (Sama-Miller et al., 2018).

A national evaluation of the effectiveness of the MIECHV program from 2012 to 2017, based on random assignment of families, found statistically significant positive effects in 4 out of 12 confirmatory outcomes: the quality of the home environment, the frequency of psychological aggression toward the child, the number of Medicaid-paid child emergency department (ED) visits, and child behavior problems (Michalopoulos et al., 2019). Most estimated effects are similar to but somewhat smaller than the average found in past studies of individual home visiting models. Differences in effects among the evidence-based models are generally consistent with the models' focuses. For example, Parents as Teachers produced the largest increase in

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parental supportiveness, and NFP produced the largest reduction in ED visits for children (Michalopoulos et al., 2019).

Potential to Reduce Disparities

The field of home visiting research continues to expand, with researchers identifying measurement frameworks (West et al., 2018) and factors contributing to implementation and replication (Paulsell et al., 2014). The state and tribal MIECHV recipients are required to implement rigorous program evaluations, which also help inform best practices and identify model programs and approaches and support the development and implementation of cultural adaptations necessary to ensure program retention, engagement, and cultural fit (Denmark et al., 2018; Hiratsuka et al., 2018). While the evidence remains limited for significant improvements in the reduction of health disparities, some studies have demonstrated that home visiting resulted in improved outcomes and so offered savings in terms of a return on investment of $2–$4 for every dollar spent (Cannon et al., 2017).

Among nondepressed mothers or families without multiple Child Protective Services (CPS) reports prior to study enrollment, home visiting was associated with a significantly lower likelihood of CPS report recidivism. These results indicate potential for home visiting to prevent maltreatment recidivism but, importantly, suggest that home visiting approaches, with weekly in-home visits by a trained home visitor, is warranted for mothers exhibiting significant depressive symptoms or families with extensive CPS histories (Jonson-Reid et al., 2018). However, among 220 low-income home visiting enrollees in a regional setting, mothers who experienced maltreatment in childhood had a heightened risk for membership in the mild and moderate-severe depression subgroup trajectories (Teeters et al., 2016). Mothers reporting higher social support saw a significant reduction in the risk of experiencing either mild or moderate-severe depression subgroup trajectories, while mothers reporting an increased belief in having control over one's life (LOC) also saw a significantly reduced risk for assignment in either the mild or moderate-severe subgroups. Moreover, the depression score of mothers experiencing moderate to high depression was not significantly lower while enrolled in home visiting, though the depression scores for mothers at the minimal and mild levels of depression were significantly lower over time (Teeters et al., 2016). These results suggest the effect of home visiting may also depend on maternal depression level, demonstrating that tailoring home visiting curricula to best fit maternal risk factors may be warranted.

Exemplar Programs and Promising Practices

The NFP home visiting curriculum has demonstrated multiple positive outcomes, including a reduction in domestic violence reports (Eckenrode et al., 2017). Mothers receiving visits from nurse home visitors had fewer health care encounters for injuries and higher ratings of the home environment and parent's reports of caregiving (Kitzman et al., 1997; Macmillan et al., 2009; Olds et al., 1997; Zielinski et al., 2009). NFP is the only home visiting program to have statistically significant positive effects over time on substantiated cases of child maltreatment.

Healthy Families New York, a Healthy Families America–accredited home visiting program, has demonstrated positive effects for fathers' participation. Families with father participation were more than four times as likely to be retained, and engaged fathers were more likely to live at home with the child and to remain emotionally involved at 6 months (McGinnis et al., 2018).

The Family Spirit home visiting intervention is the only program that has demonstrated

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efficacy in American Indian communities. To date, it has only been implemented in three rural, tribal communities in the Southwest. Nevertheless, Family Spirit has seen that high-risk mothers had significantly greater parenting knowledge (effect size = 0.42) and parental locus of control (effect size = 0.17), fewer depressive symptoms (effect size = 0.16) and externalizing problems (effect size = 0.14), and lower past-month use of marijuana (OR = 0.65) and illegal drugs (OR = 0.67). Moreover, children in Family Spirit had fewer externalizing (effect size = 0.23), internalizing (effect size = 0.23), and dysregulation (effect size = 0.27) problems (Barlow et al., 2015).

Culture is embedded across many important components of parenting, and it has only recently been actively integrated into home visiting services. However, few culturally specific home visiting curricula have sufficient evidence to be deemed evidence-based practices. Some are considered promising, with some positive quasi-experimental support for program, family, and child outcomes. For example, *Kulia I Ka Nu'u* works with families with children 2.5–5 years old using a Montessori approach with Native Hawaiian culture to support parents in building their child's school readiness and school success (Yoshimoto et al., 2014). Parent educators help families learn more about Native Hawaiian values and practices and incorporate these cultural values into the curriculum, which includes literacy, math, art, social studies, and science. *Kulia I Ka Nu'u* is unique to the island of O'ahu. *Pulama I Na Keiki* (PINK) provides services to Native Hawaiian families prenatally to 3 years of age. PINK supports families in identifying and learning more about cultural traditions and how strengthening cultural knowledge contributes to family life. Parent educators provide culturally based information about prenatal health, childbirth, child development, and child rearing (Yoshimoto et al., 2014).

Expanding the Impact of Home Visiting

To maximize the impact of effective home visiting programs, there is a need to expand their reach by scaling them to larger and different populations; currently, only 40 percent of U.S. counties provide home visiting services, with 13 states offering home visiting in 75 percent of counties and 10 states providing it in 10 or fewer counties (NHVRC, 2017). In 2017, more than 300,000 families received home visiting services, though the National Home Visiting Research Center estimates that about 18 million additional families could benefit but were not being reached (NHVRC, 2018).The MIECHV program was reauthorized in 2018 at a funding level of $400 million annually through 2022 (Sandstrom, 2019). To address remaining unmet needs, the federal program needs to be funded at a much higher level in the next reauthorization. States can further expand support for evidence-based home visiting services through state general funds, Medicaid financing, or braiding of different funding streams (Johnson, 2019). Presently, 20 states are using Medicaid to expand state capacity and reduce unmet needs for home visiting services, including considering universal home visiting, such as in Oregon and Washington State. Many states are using a Medicaid State Plan Amendment to support home visiting services under targeted case management benefit; other states, such as Maryland and South Carolina, had made home visiting demonstration or pilot projects part of a larger Medicaid 1115 or 1915(b) waiver. South Carolina is braiding approximately $17 million in philanthropic investments with $13 million in Medicaid funding through a 1115 waiver to finance a pay-for-success approach. Under this approach, private (and public) funders provide up-front capital to expand social services through a "social impact bond;" the government pays for all or part of the program only if it measurably improves the lives of participants (Johnson, 2019). South Carolina has committed an additional $7.5 million in payment if independent evaluators find positive results

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from expanded home visiting services (Johnson, 2019).

Scaling up requires more than simple replication of an effective home visiting model to reproduce positive effects. It entails the careful consideration and balance of fidelity, spread, and tailoring models for local adaptation. To support fidelity in the home visiting field, Paulsell et al. (2014) argue that two conditions need to be met: 1) research on efficacy, effectiveness, and dissemination need to document challenges that threaten fidelity; and 2) the field needs to standardize measurement and improve understanding of valid and reliable fidelity measures within and across models. While model fidelity is an important mediator of program success, home visiting implementation requires flexibility to tailor the program to the needs and/or assets of the community or population being served. This is especially critical when applying an equity approach, so that all children and families receive the services that they need. One strategy may be for the MIECHV program to work with model developers in identifying core program elements across models, while allowing state grantees to choose additional elements from various models to enhance and tailor the core program to the differential needs and vulnerabilities of individual families and communities. Additionally, others have proposed proportionate universalism—a tiered approach where every new parent in a community receives at least one home visit, while the quantity and intensity of follow-up services are adjusted to the needs and assets of high-risk families, to improve program cost-effectiveness (Marmot et al., 2010). (See the section on Synthesizing the Evidence for Conclusion 4-5 on applying a tiered approach to services.) Most importantly, as the committee discusses in Chapter 5, home visiting must not become yet another siloed prenatal and early childhood service. Future evidence-based home visiting programs need to continue to partner with, coordinate, colocate with, and expand the reach and impact of other ECE programs and the future redesigned advanced medical home. Home visiting can thus serve as a linchpin between health care, social services, and early childhood education.

Lastly, the relatively modest impact of the MIECHV program demonstrated in the national evaluation has been observed in other early childhood programs that scaled up from RCTs to population-wide interventions (Center on the Developing Child at Harvard University, 2016). Studies that focus on average effects may mask wide variation in impacts if they do not or cannot account for the differential susceptibilities of individuals, families, or communities. Often, RCTs cannot easily be adjusted, adapted, or enhanced after randomization (see Chapter 1 for a discussion about the limitations of RCTs). Future home visiting and ECE research needs to move beyond searching for one-size-fits-all solutions to define more precisely which intervention works best for which patients or populations under what circumstances.

Researchers, program leaders, and policy makers should focus on expanding the concept of precision home visiting that advances which programs and activities are best for which family, in which communities, and for what outcomes.

Recommendation 4-3: To strengthen and expand the impact of evidence-based home visiting programs:

- **Federal policy makers should expand the Maternal, Infant, and Early Childhood Home Visiting Program.**
- **The Health Resources and Services Administration and the Administration for Children and Families should work with program developers to increase flexibility for states and communities, to tailor the program to the needs and/or assets of the community or population being served.**

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- **Federal, state**, local, tribal, and territorial **agencies overseeing program implementation should continue to strengthen programmatic coordination and policy alignment between home visiting, other early care and education programs, and medical home.**

State policy makers should further expand support for evidence-based home visiting services through general funds, Medicaid, and braiding of multiple funding streams. HRSA and ACF should continue to assure program effectiveness and accountability of the expanded program. Expansion of home visiting programs should be done in conjunction with the expansion of other public investments and services.

Given the resources needed to provide high-quality home visiting, cost is an important consideration. Yarnoff et al. (2019) developed and piloted a tool to standardize the collection of home visiting data, in addition to assessing program costs for 45 local implementing agencies that participate in MIECHV. The study reported an average cost per family served of $12,556 for agencies serving less than 100 families, versus $7,117 for agencies serving more than 100 families.[6] Burwick et al. (2014) conducted a cost analysis for 25 implementing agencies and found that the average estimated annual cost to operate a home visiting program was $580,972.[7] Costs varied drastically from program to program, ranging from $206,426 to $1,207,054. Estimating and comparing costs of home visiting programs—and by extension, of expanding them—can be difficult for a number of reasons. As noted above, program costs vary, and this could be attributable to the services offered, duration, transportation costs—particularly in rural and/or tribal communities—and family and agency characteristics (Burwick et al., 2014; Yarnoff et al., 2019). Further research is needed to understand how to expand home visiting programs cost effectively, and a tiered approach would provide tailored services based on family needs (see Conclusion 4-5 for more on a tiered approach). Finally, strengthening programmatic coordination and alignment across sectors, as indicated in the recommendation, could potentially reduce costs.

Other Evidence-based Programs Focusing on Mothers

Outside of home visiting, perhaps the first program to explicitly target mothers' well-being is Lieberman's Infant Parent Psychotherapy (IPP), designed with the recognition that a critical prerequisite in working with mothers at high risk for maltreating their infants is attention to the women's own psychological needs. The program has been expanded for work with mothers and toddlers (TPP), and therapeutic strategies combine cognitive behavioral, psychodynamic, and social learning theories. A cardinal ingredient, however, is high empathy and support for the mother, with recognition that "…the most effective interventions are not spoken but rather enacted through the therapist's empathic attitude and behavior toward the mother as well as the baby (Lieberman and Zeanah, 1999, p. 556). A number of studies have now documented the benefits of IPP and TPP, implemented with mothers across diverse settings (see Lieberman et al., 2011).

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[6] The study authors calculated cost of the program per family by dividing total program costs by the average monthly number of families served by the agency and then calculated weighted mean cost per family served as the sum of costs across all agencies divided by the sum of families served.

[7] The cost analysis was conducted using "the 'ingredient' or resource cost method, which involved itemizing the types of resources (or ingredients) needed to provide services, gathering information on the types and value of resources used by each agency during the study period, and aggregating costs to estimate total program costs" (Burwick et al., 2014, p. xii).

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Another program with a strong emphasis on respect, warmth, and support by intervention staff to participating mothers is Dozier's ABC. In this program, goals are to target insecure early attachments by intervention services to foster caregivers' positive qualities, including responsiveness, nurturance, and their own attachment states of mind (Dozier et al., 2002). The program has been refined and tested in RCTs across more than 20 years. Results show that those receiving the intervention showed significant gains compared to control groups, which included better quality of attachment between mothers and children and improved self-regulation in both (Bernard et al., 2015; Bick and Dozier, 2013; Dozier et al., 2002). See Box 4-8 for more on the ABC program as an example of a promising model.

BOX 4-8
Attachment and Biobehavioral Catch-up (ABC) Program: Promising Model[a]

Developed in 1994, the ABC Intervention is an internationally available training program for caregivers of infants and toddlers through early adolescence, including high-risk birth parents and caregivers of young children in foster care, kinship care, and adoptive care. Parent coaches conduct 10 weekly home visits, approximately 60 minutes each. The model is designed to help caregivers provide nurturance even when children do not appear to need it, mutually responsive interactions in which caregivers follow children's lead, and nonfrightening care. Parent coaches provide immediate feedback on the caregivers' interaction with the child to help the caregivers attend to the target behaviors. The model also incorporates homework and video feedback.

While the original version included children from 6–48 months old, the model has been updated to include two different versions: 1) infants (6–24 months) and 2) toddlers (24–48 months). The model is based on attachment theory and stress neurobiology, and it addresses the social and physical environments by helping caregivers sensitively respond to children's behavioral signals, enhancing children's behavioral and regulatory capacities, and fostering the development of secure attachments between children and their caregivers. The model was developed by academic researchers and continues to be evaluated by them, though it has expanded significantly since its origin. ABC has partnered with state agencies, community-based nonprofits, or religiously affiliated funds. NIH also funds many of their research projects. Infant ABC coaches are trained intensively and supervised for 1 year, while toddler ABC coaches have to receive additional training afterward. The project originated at the University of Delaware, and certified ABC parent coaches are currently available across the United States in communities in Delaware, Hawaii, Idaho, Kansas, Kentucky, Louisiana, Maryland, Michigan, Minnesota, New York, North Carolina, North Dakota, Ohio, Oklahoma, and Pennsylvania. Internationally, ABC coaches are available in Australia, Germany, Norway, Russia, South Africa, and Sweden.

The program was designed to address inequities in social environments that affect children exposed to trauma in early childhood. Positive interactive experiences can stimulate healthy development of the central nervous system, foster a positive sense of self, and help maltreated children learn to accurately interpret and respond to emotional, environmental, and social cues. Four independent RCTs support the efficacy of ABC and suggest that it is effective in improving attachment and helping regulate both biology and behavior in at-risk children (see, for example, Bernard et al., 2015; Lind et al., 2014). In addition to findings from these RCTs, researchers have also found evidence to support ABC's efficacy through dissemination studies with pre- and postintervention assessments.

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

SOURCES: ACF, 2018; Attachment Biobehavioral Catch-up, 2017

Yet another program with focused attention to the well-being of mothers is the Relational Psychotherapy Mothers Group intervention (RPMG) (Luthar and Suchman, 2000), developed for mothers at risk for maltreating their children—specifically, low-income women with histories of

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substance abuse. As with IPP and ABC, core therapeutic components in RPMG include a strong emphasis on empathy, genuineness, and warmth and using insight-oriented approaches to improve parenting skills and efficacy. There were two major differences between RPMG and programs for mother–child dyads, both related to potential for scaling up. The first was the choice of a group format rather than individual sessions, thus potentially reaching more women for an hour of clinicians' time. Second, RPMG was designed for mothers of children from birth through adolescence, with developmental issues addressed as part of the program, again allowing for inclusion of a wide range of women. Results of two clinical trials showed benefits of RPMG. Compared to treatment as usual in their methadone clinics, mothers who received the 24-week intervention showed lower risk for child maltreatment —by mothers' and children's reports— and better personal adjustment as well as both mothers' and children's reports of child maladjustment and on mothers' drug use via urinalyses (Luthar and Suchman, 2000). At 6 months posttreatment, RPMG recipients continued to be at an advantage, although the magnitude of group differences was reduced. In the second trial, RPMG was compared to relapse prevention therapy (RPT). RPMG mothers again fared better than RPT mothers at posttreatment per mothers' self-reports, children's reports, and urinalyses (Luthar et al., 2007). But again, at 6 months of follow-up, they lost many of their gains.

These findings collectively indicate that even after external interventions are completed, it is critical to ensure continuity in supports for mothers who are vulnerable to depression and other mental illness. Similar messages have emerged from Olds et al.'s (2007) widely used, efficacious program, the NFP, which provides regular home visits for low-income, single mothers by a warm, supportive nurse. These home visits begin in the prenatal months and continue through the babies' second birthdays. Yet, as Golden and colleagues (2011) underscore, mothers' needs for help are not necessarily resolved when children reach 2 years old. In fact, mothers who have received this program report widespread, continuing depression in their communities, with many women desiring and needing support from others who can be reliable sources of caring over time.

In view of this accumulated work, therefore, one critical challenge for those working with vulnerable mothers is to ascertain how these mothers might continue to receive support even when time-limited supportive interventions cease, and in this regard, peer-based interventions are promising. Specifically, there is potential value in groups run by "graduates" of supportive programs who receive subsequent training and supervision in facilitating groups for peers (see Bryan and Arkowitz, 2015). Reportedly, this strategy has been used successfully with mothers who themselves had experienced perinatal depression, who regularly met with other women who were currently depressed. Bryan and Arkowitz (2015) conducted a meta-analytic review and found that peer-administered interventions (PAIs) produced substantial reductions in symptoms of depression. The magnitude of beneficial changes over time was significantly greater than those in no treatment conditions and was also comparable to non-peer-administered interventions. Interestingly, PAIs that were administered purely by peers were more effective than those that also involved a professional in the treatment, in a secondary role.

In future efforts to use such peer-based groups for mothers, some have suggested drawing upon a type of model (not the intervention itself) that has been used extensively for decades and with minimal costs: the 12-step model (see Luthar et al., 2015). Community-based meetings in this program, specifically Alcoholics Anonymous, benefit more than 1.3 million Americans, and its benefits can be equal to or even greater than those of professional interventions (Kelly et al., 2012). A major "active ingredient" in this program, again, is ongoing access to dependable,

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authentic supports; members speak of unconditional acceptance with others sharing similar challenges and their ability to reach out to others when in distress. In the future, there could be value in using such community-based groups to promote resilience in at-risk mothers. Sustaining the type of supportive connections that mothers came to depend upon in the active phase of relational interventions might be possible with continued weekly meetings (Luthar, 2015).

This has been a recurrent theme in prevention and policy: scientists increasingly urge the use of networks that already exist in communities, bringing them to bear in interventions that become self-sustaining over time (Luthar, 2015). To illustrate, within affordable housing complexes, Antonucci and colleagues (2014) described programs focused on creating mutually supportive networks among residents. Kazdin and Rabbitt (2013) described several low-cost, feasible, and effective interventions using lay people (rather than professional therapists), including hair stylists in beauty salons, trained to assess depression and to provide appropriate referrals. Hoffman and Hayes (2019) underscored the potential value of training paraprofessionals in the specific procedures that highly trained clinicians have identified as being most promising in engendering improved mental health.

An alternative approach is to deliberately foster women's connections with reliable others in their everyday lives over the duration of a formal, relational intervention. This was the strategy used in an RCT of the Authentic Connections Groups (AC Groups) program at the Mayo Clinic, tested with mothers who were health providers—a professional group at high risk for depression, burnout, and even suicide, compared to norms. From the start of the 3-month program through the end, there was a focused, concerted effort to have women reach out to their "go-to committees" in their personal lives; thus, authentic connections were forged both within the groups and outside of them, in the mothers' everyday lives. In comparison with a control group (given 1 hour of freed time over 3 months to attend groups, as were intervention mothers), AC Groups women showed significantly greater gains across multiple adjustment indexes ranging from depression and self-compassion to lowered burnout at work and decreased levels of cortisol (Luthar et al., 2017). Moreover, effect sizes across outcomes were generally in the moderate range, and gains were still stronger 3 months after the intervention program was over (partial eta squares of 0.08–0.19; median, 0.16; according to Cohen (1988), partial eta squares correspond to the following effect sizes in the real world: .02 is small; .13 is medium; and .26 is large).

The success of these in-person AC Groups led to efforts to conduct them in a virtual format, via videoconferencing, to achieve potentially wider and easier dissemination (Luthar et al., in press). Results of this study indicated the successful completion of five different virtual groups (*n* 3–6 women each, for a total of 23), with the sample involving mostly mothers, most with advanced (graduate) degrees and almost all in high-pressure careers. As with the in-person version, the 3-month program had zero dropouts. Participants' mean ratings of the groups' effectiveness were 9.6 of 10, and the "Net Promoter Score" (percent of promoters subtracting percent of detractors) fell in the exceptionally high range (Luthar et al., in press). Open-ended responses emphasized participants' appreciation of access to robust, authentic connections with other caring women (which endured well after the program was completed). Given the low cost and ease of access—from mothers' own workplaces, homes, or even while traveling—such virtual groups might be considered a useful model to be explored further in future work with mothers (indeed, the use of virtual connections have been increasingly emphasized; see Hofman and Hayes, 2019; Weisz et al., 2019).

In concluding this section on interventions, the literature reviewed suggests the value of a

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tiered approach to psychological interventions for mothers at risk due to their life adversities, along with their children, and as early as possible—preferably prenatally. All approaches involve ensuring strong, dependable supports for the mother or other primary caregiver. At the broadest level of interventions, there is particular promise in those that bring together mothers in communities or virtually to provide such mutual, ongoing support. For women needing greater psychological help, time-limited, evidence-based group therapies, emphasizing connections and mutual support and making maximal use of existing resources, would be useful. The most intensive level would be one-one one interventions, such as those involving home visiting or work with mother–child dyads to enhance strong attachment quality. Given the critical need to support caregiver well-being and to prevent child maltreatment (see Conclusions 4-1 and 4-2), in addition to the review of high-quality interventions that are offered in community settings for families, the committee offers the following recommendation to support the development and implementation of such interventions.

Recommendation 4-4: Policy makers at the federal, state, local, territorial, and tribal levels and philanthropic organizations should support the creation and implementation of programs that ensure families have access to high-quality, cost-effective, local community-based programs that support the psychosocial well-being of the primary adult caregivers and contribute to building resilience and reducing family stress.

Measures to be Used

In concluding this section, there is consideration of measures that should be included in evaluating results of large-scale preventive interventions targeting young children and their mothers. At a minimum, there should be brief assessments of the mothers' depressive symptoms, stress, feelings of rejection to the child, any involvement with CPS, and intimate partner violence (IPV), as well as the degree to which they have positive, buffering relationships in their lives. Possible measures include the 20-item Zung Self-Rating Depression Scale (Zung, 1965), 10-item Perceived Stress Scale-10 (Cohen et al., 1983), and 10-item subscale on parental undifferentiated rejection from the Parental Acceptance-Rejection Questionnaire (PARQ; Rohner et al., 1980). For supportive relationships, potential measures might include the Satisfaction with Friends subscale (four items from the Inventory of Parent Experiences; Crnic, 1983) and two single items that have consistently shown significant, robust links with multiple aspects of mothers' well-being: "Do you feel seen and loved for the person you are, at your inner core?" and "When you are deeply distressed, do you feel comforted in the ways you need?" (Ciciolla and Luthar, 2019; Luthar and Ciciolla, 2015). Finally, if any studies propose additional expensive assessments to inform future interventions (such as, biomarkers as mediators, moderators, or outcomes), researchers need to explain how the findings from these will be applied not only in laboratory-based RCTs but also in large-scale interventions reaching vulnerable children and families in their own home, school, and community settings. Given the need to identify individuals at risk for early adversity and toxic stress, the committee offers the following recommendation to close a gap in the system. Furthermore, screening requires alignment with available services for children and families (see Chapters 5 and 8 for more on screening and alignment with services).

Recommendation 4-5: Health care providers who care for pregnant women and children should routinely track levels of individual health and social risk among

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mothers and children over time, using periodic assessments via a short set of scientifically validated measures.

SYNTHESIZING THE EVIDENCE

To summarize, major conclusions deriving from this chapter include the following. Promoting resilience in childhood rests, first, on ensuring the well-being of children's primary caregivers—usually, their mothers—and this requires the mothers' ongoing access to strong, supportive relationships, beyond meeting basic needs for survival. Maltreatment is of utmost importance among the various parenting behaviors that should be targeted in preventive interventions; minimizing neglect and abuse by caregivers and others needs to be a top priority. Gender specificity is advised in interventions for parents, with separate delineation of how mothers and fathers each are most effectively recruited, retained, and helped in interventions. When considering children who need preventive interventions, there are several subgroups whose unique challenges need to be considered. These include subgroups that have been extensively documented in past research to be at risk—such as children in chronic poverty, those separated from their parents, and recent immigrants—and those shown more recently as contending with high stress and distress, such as children and families in high-achieving communities and youth who are in the minority based on gender identification and sexual orientation.

As future interventions for mothers and children are taken to large scale, it will be useful to employ a tiered approach, with the intensiveness of services increasing with the level of the mother's distress. At a fundamental level, universal, supportive interventions for mothers from pregnancy onward are needed. This includes using existing resources for supportive group-based interventions (e.g., organized in their own communities, such as in preschools or schools) that are, where possible, led by peers and/or accessed virtually. Universal interventions also need to include at least one home visit after birth, to ascertain if mother and child have the supports they need, and, if difficulties are found, to provide connections to local resources for the family.

Conclusion 4-5: A tiered approach is needed for families known to be at risk due to their life adversities, with appropriate investment of resources to expand delivery of promising cost-effective programs and assessments via scientifically sound measures of critical constructs. This includes the need for:

- *Universal, supportive interventions for mothers and*
- *More resource-intensive programs.*

To promote these first-level, universal programs, federal, state, and local agencies overseeing program implementation could develop and monitor community-based support groups for mothers within existing programs, such as the Maternal, Infant, and Early Childhood Home Visitation program, Head Start, Early Head Start, and Women, Infants, and Children (WIC). Care providers within these programs require appropriate referral sources for mothers and children who need intensive, one-on-one care (e.g., through home visiting). Enhanced coverage by major insurance providers for programs offering group-based care, support, and therapeutic interventions for mothers is also necessary (see Chapter 5 for more in health care needs for this population).

To identify mothers who might need more intensive services, a brief battery of questionnaires should be used as a screening tool. Specifically, measures of maternal stress and

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depression (described in this chapter) should be routinely administered at prenatal and pediatric "well-baby" visits. Those mothers who manifest high distress will need referral to programs that offer evidence-based therapeutic interventions.

Intensive programs for mothers in distress can be group or individual. Programs that are short term and involve groups of women, such as AC Groups or the Mental health Outreach for Mothers (MOMS) Partnership, can help to maximize both reach and cost-effectiveness. (See Box 4-8 for information on the MOMS Partnership as a promising model.) Where necessary, mothers in serious distress need to be referred to more resource intensive, one-on-one programs, such as ABC and Infant/Toddler-Parent Psychotherapy. Home visiting, with concurrent referrals to available supports as needed—formal and informal—should exist within the community. Ideally, all types of intensive services would be coordinated with the pediatricians' offices, in a team-based approach to providing care (see more on team-based approaches in the health care setting in Chapter 5).

BOX 4-8
MOMS Partnership: Promising Model[a]

Based in New Haven, Connecticut, the MOMS Partnership is an evidence-based, community-driven initiative that offers mental health and trauma-related services and parenting and job readiness programs to single mothers in at-risk neighborhoods (Center on the Developing Child, 2019). The program recruits participants by "meeting" mothers at locations they frequent (i.e., grocery stores, laundromats, shelters, community colleges, libraries, and nail and hair salons) (White et al., 2018).The program's mission is "to reduce depressive symptoms and increase social and economic mobility among overburdened, under-resourced mothers, thereby strengthening generations of families to flourish and succeed" (MOMS Partnership, n.d.-c).

Beginning in 2010 and in partnership with mothers and community partners, the program has conducted surveys to identify and understand the needs of pregnant women, parents, other caregivers, and families in the community (Smith, 2018). Findings from more than 4,000 interviews have directly informed the design of the program. The program delivers a "bundle" of local services, including those to build skills (e.g., for stress management and work success), community mental health ambassadors, referral to local services and resources, and social networks and support (Smith, 2018).

The program has demonstrated promising initial outcomes, including findings from a 2012–2016 RCT in New Haven public housing and multigenerational outcomes from two quasi-experimental studies (Smith, 2018).Evaluation of the program has shown that at completion, 76 percent of participants have fewer depressive symptoms and participants experience a 67 percent decrease in parenting stress (MOMS Partnership, n.d.-a).

Beginning in 2019, the MOMS Partnership will replicate its program in Washington, DC, through a partnership with the DC Department of Human Services and via DC's TANF program (Clayton et al., 2018; MOMS Partnership, n.d.-b).The MOMS Partnership has also collaborated with Vermont Reach Up, that state's TANF program, to replicate the program in Vermont by 2020.

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

It will be critical to allocate resources to monitoring the quality of interventions delivered. All intervention programs need to be culturally sensitive, be adapted to community needs, have clear procedures for the selection and training of clinicians, and have mechanisms in place to ensure fidelity and quality of services. Support is needed for programs across tier levels of delivery, universal and intensive. Furthermore, taking an ecological perspective on promoting caregiver well-being and attachment would require supportive policies that facilitate healthy

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social connections and family cohesion during these critical years of early life. For example, paid parental leave following the birth of a child is one policy that has evidence for shaping child and family outcomes (AEI-Brookings Working Group on Paid Family Leave, 2017). (See Chapter 6 for a full discussion and Recommendation 6-1 on implementing paid leave.)

Finally, in evaluating the effectiveness of programs for at-risk mothers and their children, measures need to focus on those maternal dimensions that are known to strongly affect the mother–child relationship and hence child well-being. This would include measures of maternal depression and stress, feelings of rejection or hostility to the child, available support for mothers, and any contact with CPS. These assessments should be a routine part of prenatal, postnatal, and pediatric visits, along with other physical health indexes that are assessed and recorded. Careful, balanced consideration of all these aspects of intervention design, delivery, and measurement—with all the investment of resources called for by each aspect—is critical to move toward meaningfully increasing equity in well-being, considering America's most vulnerable children and families.

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5

Leveraging the Health Care System to Improve Outcomes and Promote Health Equity

INTRODUCTION

Significant differences in health status among women, infants, and children and inadequate response to the role of early life stressors on life course health indicate a need for an important transformation of the U.S. health care system and its delivery of preconception, prenatal, postpartum, and pediatric care, which are the focus of this chapter (see Box 5-1 for an overview of the chapter). Changes are urgently needed to better address health disparities, including those by race/ethnicity and socioeconomic status (SES) (see Chapter 1 for a more detailed discussion of disparities). For example, the most recent statistics on maternal and infant mortality and other birth outcomes (see Box 5-2) highlight significant disparities and also demonstrate the urgency of changes to the health care system to more rapidly and effectively address and reverse concerning trends. As the evidence in this report demonstrates, much chronic disease and disability among adults has origins in infancy and childhood. Furthermore, changing needs of the population, with growing diversity and greater understanding of the health impacts of social determinants of health (SDOH), also signal the need for major changes throughout the health care system.

BOX 5-1
Chapter in Brief: Leveraging the Health Care System

This chapter discusses opportunities to leverage the health care system in supporting healthy early childhood development, with a focus on improving access to health care services, improving quality of care, improving the organization and financing of care, and transforming the content of health care to address the social, economic, cultural, and environmental determinants of health. The chapter also addresses ways to apply a life course perspective and new knowledge about child development, including the effects of adversity, trauma, and the toxic stress response to address needs, challenges, and strategies for the health care system in the preconception, prenatal, postpartum, and early childhood periods at the practice, policy, and systems levels.

Chapter conclusions in brief:

- The current health care system focuses mainly on clinical goals and addresses the multiple determinants of health in fragmented and highly variable ways. Despite high-quality clinical care, the health status of America's children and young families is far worse than in comparable developed countries. U.S. health care provides only limited attention to integration of health care for the whole family across the life course, integration of mental and behavioral

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health with the rest of health care, or integration of healthcare within community systems to better support children and families.

- Although health insurance coverage has grown substantially in the past few decades, mainly through Medicaid expansions and other insurance enhancements, access remains a problem for many families with young children, who experience numerous barriers to obtaining health care services in addition to lack of health insurance coverage. Further efforts are needed to address financial and nonfinancial barriers to care and to ensure that all families have access to adequate preconception, prenatal, postpartum, and pediatric care.
- Promising strategies can improve quality of preconception, prenatal, and child health care, such as developing and implementing new measures, including for adversity and social determinants, along with efforts to strengthen the training of the health care workforce to better understand diversity and implicit bias and to address equity in health care.
- Programs that build on home visiting, referrals to community partners, and integrated community efforts have enhanced outcomes for children and families. New technologies have expanded care and access, increased understanding of the social determinants, and improved communication about health and chronic disease. New payment arrangements can accelerate the transformation of health services to programs to support families and population health.

Chapter recommendations in brief:

- Increase access to preconception, prenatal, postpartum, and pediatric health care.
- Expand accountability and improve quality of preconception through pediatric care.
- Adopt policies and practices that improve the organization, financing, and integration of care systems from preconception through pediatric care, with a focus on the caregiver and child together as the unit of care and collaboration with community-based services.
- Transform preconception, prenatal, postpartum, and pediatric health care to address the root causes of poor health and well-being, including social determinants.

As detailed in this chapter, the current health care system often delivers limited, episodic, inequitable, and fragmented services. It focuses on clinical medical care, which recognizes the myriad of social factors that affect health outcomes but often addresses such factors in fragmented and variable ways. The health care system serves as one platform, along with public health and other sectors, to address the social determinants that underlie many health inequities. To better address inequities, it is necessary to transform the organization, delivery, and financing of health care to incorporate community-focused teams and integration across multiple sectors to address the SDOH, poverty, mental and behavioral health (MBH), chronic disease, disparities, adversity, and family well-being. Achieving these goals will require assuring access to care, focusing and improving quality of care, changing the organization and financing of health care, and strengthening the content of health care.

Based on available evidence and existing resources, the committee identified three domains as important for focusing on preconception, prenatal, and early childhood interventions: health care systems and services (see Chapter 5), healthy living conditions (see Chapter 6), and early care and education (ECE) (see Chapter 7) (see the committee's conceptual model in Chapter 1, Figure 1-9). This chapter focuses on health care systems and services, including preventive care and clinical care delivery systems, and also emphasizes critical links and opportunities for alignment with other partners in the health care system and in other sectors in the community.

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BOX 5-2
Health and Health Services Disparities for Women, Infants, and Children

Maternal mortality—Maternal mortality in the United States increased from 7.2 deaths per 100,000 live births in 1987 to 17.2 deaths per 100,000 live births in 2015 (CDC, 2019c). There are large racial disparities as well—as of 2010, black women were three to four times more likely to die from a pregnancy-related complication than non-Hispanic white women (Creanga et al., 2015).

Infant mortality—In 2016, infant mortality rates were 11.4 percent for non-Hispanic black infants, 9.4 percent for American Indian/Alaska Native (AI/AN) infants, 7.4 percent for Native Hawaiian or other Pacific Islander infants, 5.0 percent for Hispanic infants, 4.9 percent for non-Hispanic white infants, and 3.6 percent for Asian infants (CDC, 2019b).

Low birth weight—In 2016, LBW rates in the United States were 13.7 percent for non-Hispanic black women and 7.0 percent for non-Hispanic white women (Martin et al., 2018). In the same year, rates among Hispanic subgroups ranged from 9.5 percent for Puerto Rican women to 6.9 percent for Mexican women (Martin et al., 2018). In 2013, rates among Asian and Pacific Islander subgroups were 10.6 percent for Asian Indian women, 9.4 percent for Filipino women, and 5.9 percent for Chinese women (Child Trends, 2015).

Preterm birth—In 2016, preterm birth rates were 13.8 percent among non-Hispanic black mothers and 8.63 percent among non-Hispanic Asian mothers. For Hispanic subgroups, rates ranged from 11.0 percent for Puerto Rican mothers to 9.1 percent for Central and South American mothers (Martin et al., 2018).

Prenatal care—In 2006, the rate of women who received prenatal care in the first trimester was 58 percent for black and Hispanic women and 76 percent for white women (69 percent of women overall received prenatal care) (Bryant et al., 2010).

Health insurance—In 2016, 55.4 percent of black children and 56.8 percent of Hispanic children were covered by public insurance, compared to 31.9 percent of non-Hispanic white children and 26.8 percent of Asian children (Child Trends, n.d.; U.S. Census Bureau, 2017). In the same year, black and Hispanic children made up 20 and 37 percent, respectively, of all children covered by Medicaid and the Children's Health Insurance Program (CHIP), yet they were 14 and 25 percent, respectively, of all children (Brooks and Wagnerman, 2018). In 2017, 4.9 percent of black children and 7.7 percent of Hispanic children were uninsured, compared with 4.3 percent of non-Hispanic white children and 4.6 percent of Asian children (Berchick et al., 2018).

Health status—In 2011–2012, 5.0 percent of Hispanic children interviewed in Spanish (3.6 percent of Hispanic children interviewed in English) and 3.6 percent of non-Hispanic black children had poor or fair health, compared to 2.7 percent of non-Hispanic white children (Pastor et al., 2015). In addition, children and youth receiving Supplemental Security Income (SSI) payments for severe disability are disproportionately black and Latino (NASEM, 2015).

The health care sector is positioned to play a crucial role in advancing health equity by providing care and services during the preconception, prenatal, postpartum, and early childhood periods. Preconception, prenatal, and pediatric care provide a point of entry into the health care system for women and men as well as children, especially those in the first few years of life. However, current health care system organizational fragmentation and the episodic delivery of health care services coupled with institutional and systemic disadvantage have resulted in significant disparities in access to and use of health care services. Health care that is well

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organized, accessible over time, high quality, universally available, and effectively integrated for all people could provide continuous access to a wide variety of resources and services and decrease disparities in use of health care services. Given that recognizing risk factors (biological, social, and environmental) as early as possible is fundamental to addressing health inequities, universal access to health care services is a critical component to decreasing and eliminating health inequities (Veugelers and Yip, 2003).

Although health care plays an integral role, the health care sector alone cannot meaningfully address health inequities, nor is it the primary actor or leader. Cross-sectoral and multidisciplinary collaboration is essential for decreasing health inequities. As discussed in Chapter 1, the United States spends more on health care than any other country yet has some of the worst outcomes and gravest disparities. Despite significant improvements in the last century, troubling trends have persisted and worsened in the current century (e.g., life expectancy has decreased, and maternal mortality rates have increased [see Box 5-2]) (CDC, 2019c; Murphy et al., 2018). Although children and youth represented about 45 percent of the Medicaid population in 2013, they received only about 19 percent of Medicaid expenditures (MACPAC, 2018). In addition, about 95 percent of U.S. spending on health is related to treatment and medical services; only about 5 percent is allocated to population-level health improvement and prevention (McGinnis et al., 2002). The United States spends the highest percentage of GDP on health care services among all nations but has poor health outcomes in many areas, including neonatal and maternal mortality, deaths from injuries, and rates of substance use. Improving health outcomes requires more rapid learning regarding interventions that work and those that do not, focusing investment in effective interventions and their deployment, and more equitable allocation of resources to other sectors outside of health care. As Steuerle and Isaacs (2014) document, federal spending on programs to support children and families has faced immense budgetary pressures as health care spending has increased.

The committee embraced the life course approach, which emphasizes the impact of an individual's experiences throughout a lifetime—and across generations—on health outcomes (see Chapter 3). Given the multigenerational impacts of toxic stress, food and housing instability, chronic disease, and parental ill health on the health of children, all family members need access to health care services across the life course. Thus, this chapter highlights the many ways in which our current system represents a patchwork of services offered at different times in life (e.g., little or no continuity, intermittent insurance coverage, poor access to providers, eligibility for services for short periods of time) and how U.S. health care needs to be redesigned and rebuilt on a firmer foundation of care across the life course with added boosters during life junctures most critical to a child's health. This chapter covers health care services delivered and received during the preconception, prenatal, postpartum, and early childhood periods—junctures at which well-timed services can boost the odds for good health across the life course.

This chapter generally focuses on health care provided by physicians, although there are many other practitioners that play an important role in the health care system during these life periods—such as nurses, nurse practitioners, nurse-midwives, doulas, social workers, specialized therapists, and mental health practitioners. However, given the scope of this report, the committee could not cover all of these in great detail. In addition, there are a number of forthcoming National Academies reports on topics related to this chapter, including the roles of other important practitioners. These includes studies on The Future of Nursing 2020–2030,

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which has a focus on reducing health disparities and producing a culture of health;[1] Assessing Health Outcomes by Birth Settings;[2] and Integrating Social Needs Care into the Delivery of Health Care to Improve the Nation's Health.[3] Finally, the 2019 National Academies report *The Promise of Adolescence: Realizing Opportunity for All Youth* (NASEM, 2019) covers topics not included here, such as sexual and reproductive health care for adolescents—including unintended pregnancy.

The health care system as a whole is robust and frequently interacts with children and their families by providing a nearly universal touchpoint with all women and children from the prenatal period to age 3, making it an important system through which to address health inequities. However, it is important to note that the health care system is not the main vehicle through which change should occur to address the SDOH, nor is it where additional funds should be funneled to do so. Rather, the health care system needs to be better leveraged to not only provide medical care but also address the SDOH (including barriers to access other than health insurance, such as lack of or inadequate transportation to medical visits, cost-sharing, and lack of culturally competent services) (Woolf, 2019). However, as noted above and described in detail below, the health care system does need to change to more systematically address the upstream causes of poor health and health inequities. To do so, the health care sector needs to engage and partner with other sectors to actively address the SDOH and find common solutions to meet the needs of children and families (see Recommendations 8-1 and 8-5 in Chapter 8 for more on the need for cross-sector approaches and integration of care to advance health equity).

This chapter includes an introduction to the history and current content of preconception, prenatal, postpartum, and pediatric health care and sections describing efforts to improve access, quality, and innovative delivery/financing of better health care during these critical and sensitive periods of life. As noted above, the life course perspective illustrates that health care from preconception through early childhood is a continuum of care that needs to take place across the life-span and take into account intergenerational effects. Thus, as this chapter discusses, an integrated health care system, which will require addressing a multitude of structural, professional, practical, and cultural barriers, is necessary to accelerate improvement in health care services, with the ultimate goal of improving health outcomes and decreasing health inequities.

BACKGROUND AND CURRENT CHARACTERISTICS OF HEALTH CARE

Preconception Care

A primary goal of preconception care is to improve the health of men and women during their reproductive years, especially shortly before conceiving a child. The Centers for Disease Control and Prevention (CDC) define preconception care as "a set of interventions that aim to

[1] For more information, see http://www.nationalacademies.org/hmd/Activities/Workforce/futureofnursing2030.aspx (accessed April 5, 2019).
[2] For more information, see https://sites.nationalacademies.org/DBASSE/BCYF/Research_Issues_in_the_Assessment_of_Birth_Settings/index.htm (accessed April 5, 2019).
[3] For more information, see http://nationalacademies.org/hmd/activities/healthservices/integratingsocialneedscareintothedeliveryofhealthcaretoimprovethenationshealth.aspx (accessed April 5, 2019).

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identify and modify biomedical, behavioral, and social risks to a woman's health or pregnancy outcome through prevention and management" (Johnson et al., 2006). As discussed in Chapter 2, preconception health is important not only for pregnancy outcomes but also for the lifelong health of children and even the health of the next generation (*The Lancet*, 2018). Disparities in preconception health can thus set up intergenerational transmission of health disparities. Using data from the Behavioral Risk Factor Surveillance System (BRFSS), 2013–2015, and Pregnancy Risk Assessment Monitoring System, 2013–2014, Robbins et al. (2018) found significant disparities in nine preconception health indicators by race-ethnicity, age, and insurance status (see Tables 5-1 and 5-2). They found that among older women (35–44 years), non-Hispanic black women, uninsured women, and those residing in southern states, prevalence estimates of risk factor indicators were generally highest and prevalence estimates of health-promoting indicators were generally lowest.

TABLE 5-1: Prevalence of Preconception Health Indicators Among Nonpregnant Reproductive-Aged Women (18–44 years), by Age Group, Race/Ethnicity, and Insurance—Behavioral Risk Factor Surveillance System, United States, 2013–2015

Characteristic	Depression† (2014–2015) % (95% CI)	Diabetes†,§ (2014–2015) % (95% CI)	Hypertension†,§,¶ (2013, 2015) % (95% CI)	Current cigarette smoking** (2014–2015) % (95% CI)	Normal weight†† (2014–2015) % (95% CI)	Recommended physical activity¶,§§ (2013, 2015) % (95% CI)
Age group (yrs) ¶¶						
18–24	19.2 (18.4–20.1)	1.0 (0.8–1.2)	5.0 (4.5–5.4)	13.4 (12.7–14.1)	57.0 (55.9–58.2)	53.3 (52.1–54.4)
25–34	22.6 (22.0–23.3)	2.4 (2.1–2.7)	9.2 (8.7–9.7)	19.5 (18.9–20.1)	42.7 (41.8–43.5)	49.7 (48.9–50.6)
35–44	23.1 (22.5–23.7)	5.3 (4.9–5.6)	17.0 (16.4–17.6)	16.8 (16.3–17.4)	37.9 (37.2–38.7)	49.0 (48.2–49.8)
Race/Ethnicity¶¶						
White	27.0 (26.5–27.6)	2.6 (2.4–2.8)	10.2 (9.8–10.5)	21.1 (20.6–21.6)	49.0 (48.3–49.6)	53.8 (53.2–54.4)
Black	16.2 (15.1–17.2)	4.5 (4.0–5.1)	18.3 (17.3–19.3)	15.6 (14.5–16.7)	30.0 (28.6–31.5)	42.8 (41.3–44.3)
Hispanic	15.5 (14.6–16.4)	3.6 (3.2–4.1)	9.5 (8.7–10.3)	8.9 (8.2–9.6)	37.2 (35.9–38.6)	46.0 (44.6–47.4)
Other	14.8 (13.6–16.1)	2.4 (1.9–2.8)	8.0 (7.1–9.0)	11.3 (10.3–12.4)	57.6 (55.6–59.6)	50.3 (48.2–52.4)
Insurance*††† **						
Yes	22.3 (21.8–22.7)	3.1 (2.9–3.2)	10.8 (10.5–11.2)	16.1 (15.7–16.5)	46.1 (45.6–46.7)	51.8 (51.2–52.4)
No	20.3 (19.2–21.3)	3.2 (2.8–3.6)	11.5 (10.8–12.2)	21.0 (20.0–22.0)	38.6 (37.2–40.0)	44.0 (42.7–45.3)
Overall	**21.9 (21.5–22.3)**	**3.1 (2.9–3.2)**	**10.9 (10.6–11.2)**	**16.9 (16.5–17.2)**	**44.9 (44.4–45.5)**	**50.4 (49.9–50.9)**

Abbreviation: CI = confidence interval.

* For indicators relying on annual standard core questions (i.e., questions that are asked annually by all states), estimates are based on 2014–2015 data. For indicators that are based on the biannual rotating core survey, CDC combined years 2013 and 2015; includes 50 U.S. states and the District of Columbia. Data self-reported by women aged 18–44 years.

† Self-report of ever having been told by a health care provider that they have the condition.

§ Excluded if occurring only during pregnancy.

¶ Hypertension and physical activity questions are included as part of the biannual rotating core that is administered in odd years; therefore, 2013 and 2015 data were used.

** Defined as smoking 100 or more cigarettes in a lifetime and currently smoking cigarettes every day or some days at the time of the interview.

†† Normal weight was defined as having a body mass index of 18.5–24.9 kg/m^2 as determined by self-reported weight and height.

§§ Participation in enough moderate and/or vigorous physical activity in a usual week was defined as meeting the U.S. Department of Health and Human Services recommended levels of aerobic physical activity. Respondents were classified as meeting recommendations if they reported at least 150 minutes per week of moderate-intensity activity, or at least 75 minutes per week of vigorous-intensity activity, or a combination of moderate-intensity and vigorous-intensity activity (where vigorous activity minutes are multiplied by two) totaling at least 150 minutes per week.

¶¶ In Chi-square tests, differences by age and by race/ethnicity are significant at $p<0.05$ for all indicators.

*** Defined as having any kind of health care coverage, including prepaid plans such as health maintenance organizations or government plans such as Medicare or Indian Health Service.

††† In Chi-square tests, differences by insurance are significant at $p<0.05$ for all indicators except diabetes and hypertension.

SOURCE: Robbins et al., 2018.

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TABLE 5-2: Prevalence of Preconception Health Indicators Among Reproductive-Aged Women (aged 18–44 years) with a Recent Live Birth, by Age Group, Race/Ethnicity, and Insurance—Pregnancy Risk Assessment Monitoring System, United States, 2013–2014.

	Recent unwanted pregnancy†	Prepregnancy multivitamin use§	Postpartum use of effective contraception¶
Characteristic	**% (95% CI)**	**% (95% CI)**	**% (95% CI)**
Age group (yrs)**			
18–24	6.4 (5.8–7.1)	17.9 (17.0–18.9)	64.9 (63.6–66.2)
25–34	4.9 (4.6–5.3)	37.4 (36.6–38.2)	55.1 (54.3–55.9)
35–44	9.8 (8.9–10.8)	45.4 (43.8–46.9)	50.6 (49.0–52.3)
Race/Ethnicity**			
White	5.0 (4.6–5.4)	37.8 (37.1–38.6)	56.8 (55.9–57.6)
Black	11.6 (10.4–12.8)	21.6 (20.2–23.2)	64.9 (63.1–66.7)
Hispanic	6.4 (5.6–7.3)	26.2 (24.8–27.7)	59.3 (57.5–61.0)
Other	6.0 (5.2–6.8)	31.7 (30.1–33.4)	44.6 (42.8–46.5)
Prepregnancy insurance††,§§			
Yes	5.8 (5.5–6.1)	37.4 (36.7–38.1)	56.7 (56.0–57.4)
No	7.3 (6.6–8.1)	17.1 (16.0–18.2)	57.9 (56.4–59.5)
Overall	**6.1 (5.8–6.4)**	**33.6 (33.0–34.2)**	**56.9 (56.3–57.6)**

Abbreviation: CI = 95% confidence interval.

* Includes Alabama, Alaska, Arkansas, Colorado, Delaware, Georgia, Hawaii, Illinois, Iowa, Maine, Maryland, Massachusetts, Michigan, Minnesota, Missouri, Nebraska, New Hampshire, New Jersey, New Mexico, New York, New York City, Oklahoma, Oregon, Pennsylvania, Rhode Island, Tennessee, Utah, Vermont, Washington, West Virginia, Wisconsin, and Wyoming. Data self-reported by women aged 18–44 years who recently had a live birth.

† Defined as a pregnancy among women who reported that just before they got pregnant with their most recent live-born infant, they did not want to be pregnant then or at any time in the future.

§ Defined as taking a multivitamin, prenatal vitamin, or folic acid supplement every day of the month before pregnancy.

¶ Includes male or female sterilization, implant, intrauterine device, injectable, pill, patch, or ring.

** In Chi-square tests, differences by age and by race/ethnicity are significant at $p<0.05$ for all indicators.

†† Defined as having private, Medicaid, other government plans such as TRICARE, military health care, IHS or tribal, and other kinds of health insurance during the month before pregnancy.

§§ In Chi-square tests, differences by insurance are significant at $p<0.05$ for all indicators except postpartum use of effective contraception.

SOURCE: Robbins et al., 2018.

Advancing health equity in birth and child health outcomes begins with reducing preconception health disparities. For decades, preconception care has been proposed as a key population-level strategy for improving birth outcomes. Clinical recommendations have been developed regarding the key components of preconception care, which include addressing primarily undiagnosed, untreated, or poorly controlled medical conditions; immunization history; medication and radiation exposure in early pregnancy; nutritional issues; family history and genetic risk; tobacco and substance use and other high-risk behaviors; occupational and environmental exposures; family planning and reproduction life plan; social issues; and mental health issues (Jack et al., 2008). Men could also benefit from preconception care, although the content of care is less well defined for them (Frey et al., 2008).

The evidence base supporting a range of services for preconception care as critical to child health has been well documented and includes folic acid supplementation; appropriate management of hyperglycemia; rubella, influenza, and hepatitis vaccination; low phenylalanine diet; and provision of antiretroviral medications to reduce the risk for mother-to-child HIV transmission (Johnson et al., 2006; Korenbrot et al., 2002). Yet, several recent reviews regarding the most effective health care structures to ensure translating this science into action have found mixed results. Burgess et al. reviewed nine studies and found that fertility intention screening was associated with improved knowledge related to healthier pregnancy but not increased provision of new contraception services for those not desiring pregnancy (Burgess et al., 2018). Lassi et al. (2014) reviewed 161 studies and found evidence of effectiveness for preconception care in improving outcomes for women with diabetes, epilepsy, PKU, and depression. Hemsing et al. (2017) reviewed 29 preconception interventions and found that the majority of interventions offered assessment or screening followed by brief intervention or counseling.

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Overall, these interventions demonstrated improvements in at least some of the outcomes measured (Hemsing et al., 2017). However, several other systematic reviews failed to find conclusive evidence of improved pregnancy outcomes associated with the following types of preconception care: routine prepregnancy health promotion (Whitworth and Dowswell, 2009), genetic risk assessment (Hussein et al., 2018), preconception care for diabetic women (Tieu et al., 2017b), interconception care for women with a history of gestational diabetes (Tieu et al., 2017a), preconception health and programs for women who are overweight or obese (Opray et al., 2015), preconception lifestyle advice for people with subfertility (Anderson et al., 2010), and preconception care in the primary care setting (Hussein et al., 2016). This could be largely due to a lack of randomized controlled trials (RCTs) or poor study quality, but this may also reflect the limits of preconception care services as they are currently conceived, organized and delivered.

Currently, the content of preconception care is too ill-defined and limited, its access is too restricted and episodic, its quality too disparate and inequitable, and its organization and delivery too fragmented and siloed to fully deliver on its promise as an intergenerational equalizer of population health (Verbiest et al., 2016). The timing of preconception care has been identified as a major limitation contributing to the health care system's failures in achieving optimal delivery of preconception care. Preconception care is commonly regarded as a single prepregnancy checkup a few months before the couple attempts to conceive. The timing of such an approach, however, could miss nearly half of all pregnancies, which are unplanned (Finer and Zolna, 2016), and approximately 37 percent of all births in the United States, which are unintended at the time of conception (Mosher et al., 2012). Moreover, such an approach may be appropriate to address certain risk factors (e.g., folic acid supplementation, low phenylalanine diet, or cessation of certain teratogenic exposures) but may be too late to address others. For example, preconception counseling 3 months before pregnancy may be timely to avoid ingesting methylmercury, which has a half-life of 50 days (CDC, 2016), but too late to reduce the bioaccumulation of dioxins and dioxin-like compounds (DLCs), which have a half-life of 7–8 years (IOM, 2003). DLCs are lipophilic and bioaccumulate in animal fat; hence, IOM recommends that girls and young women drink low-fat or skim milk instead of whole milk and eat foods lower in animal fat years before they become pregnant (IOM, 2003). Switching to low-fat milk and/or a low-fat diet a few months before pregnancy would do little to reduce fetal exposure to DLCs. Furthermore, to expect preconception care to reverse allostatic load (the cumulative physiologic toll from chronic stress) in a single visit may be asking too much. Thus, rather than considering preconception care as a single prepregnancy checkup, it needs to be re-conceptualized, tested and integrated into health care services delivered consistently, continuously, and comprehensively for women (and men) across the life course.

Another limit of preconception care, as commonly practiced, is its narrow clinical focus. While its benefits in reducing certain biomedical or behavioral risks (e.g., folate deficiency, PKU, smoking) have been well documented, these are often not the major drivers of disparities in birth and child health outcomes. Preconception care could have a greater impact in advancing health equity if it is better set up to optimize management of chronic conditions, such as hypertension, diabetes, and obesity, that disproportionately affect low-income women and women of color; yet, for many women, the lack of access to and the episodic nature of preconception care limit its effectiveness as a population-wide strategy for advancing equity in preconception health. Preconception care offers an important opportunity for addressing MBH issues, but in many underresourced communities, a positive screen is often not backed up by available referral services, such as cognitive behavioral therapy, alcohol rehabilitation, tobacco

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cessation counseling and referral, substance use treatment, or trauma-informed care (TIC). Most importantly, presently preconception care is ill equipped to address many social determinants of preconception health, such as food insecurity, housing instability, occupational or environmental exposures, or intimate partner violence. This is not a flaw of preconception care per se but of the larger U.S. health care system, which is poorly designed to tackle the SDOH; nonetheless, preconception care represents an important missed opportunity for advancing health equity, with its prevailing narrow focus on fixing biomedical and behavioral risks.

The evidence base regarding the effectiveness of preconception care is also limited by the relative dearth of research on what preconception interventions succeed in advancing health equity in birth and child health outcomes. For example, despite increasing recognition of the health impact of maternal allostatic load on not only birth and child health outcomes but also the developmental origins of health and disease, there has been a paucity of intervention research on what can be done during preconception care to reduce maternal allostatic load. Similarly, little is known on how preconception care might reduce the risk of aberrant placentation, epigenetic reprogramming, and neuroendocrine, immune-inflammatory, and metabolic dysregulation, which could contribute to disparities in birth outcomes and lifelong health.

Approaches to incorporate reproductive life planning discussions into routine visits for women of child-bearing age—e.g. asking every woman of reproductive age at every routine visit whether she would like to become pregnant at some point and, if so, when that might be (Callegari et al., 2017)—have led to increased delivery of preconception care services prior to pregnancy. However, these strategies have been criticized for focusing too much on reproductive *choice* and not enough on reproductive *justice*. The maternalism inherent in the traditional concept of preconception care (as narrowly defined by the current health care system) has been criticized by some for promoting the trope of "women as reproductive vessels" (Waggoner, 2013). Others have argued that preconception care (and contraceptive care) is limited by a primary focus on changing individual behaviors instead of the historical and social contexts of those behaviors and on promoting reproductive choice instead of broader reproductive access, especially for communities of color. This reality gave rise to the Reproductive Justice movement in the 1990s, which focused on a woman's right to *have* a child or *not* to have a child and explicitly recognized that, while pregnancy intention and choice are important, many women do not have the requisite resources to access the essential tools to control their own reproductive destiny. These include, but are not limited to, reliable birth control, adoption, abortion, and paid maternity leave. Reproductive justice activists have maintained that "reproductive safety and dignity depended on having the resources to get good medical care and decent housing, to have a job that pays a living wage, to live without police harassment, to live free of racism in a physically healthy environment—all of these (and other) conditions of life were fundamental conditions for reproductive dignity and safety—reproductive justice—along with legal contraception and abortion" (Ross and Solinger, 2017, p. 56).

Pregnancy intention is key to promoting good maternal, neonatal, and childhood health outcomes (Hall et al., 2017), but the decision to have a child does not occur in a vacuum. It is rooted in the environmental, socioeconomic, and political world in which a woman and her family live. A discussion of preconception care is incomplete if it is not centered in the broader context of U.S. history, which included concerted efforts to encourage some women to reproduce while going to great lengths to make sure other women did not. Such distinctions were often based on race, class, and/or immigration status, with a paternalistic presumption that some women were inherently fit to be mothers while others were not. Notable examples of public

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policies influencing the reproductive status of women in this country include the eugenics movement, forced sterilization campaigns, and welfare programs penalizing women for having a man in the home or having children (Kluchin, 2009; Ross and Solinger, 2017; Stern, 2005). As an example, in the 1970s, many low-income women and women of color, including Puerto Rican, African American, Chicano, and AI/AN women, experienced mass forced sterilization. AI/AN women suffered particularly serious abuse from federal policies that enabled AI/AN children to be taken from their families in addition to numerous violations of reproductive rights (Torpy, 2000). These examples underscore that there is much more at play than personal reproductive choice when a woman makes the decision of whether to have children and emphasize how the health care system can best support all women and families equitably in making this important decision.

Prenatal and Postpartum Care

For decades, the delivery of prenatal care has been a cornerstone of the U.S. strategy to reduce infant mortality and perinatal disparities (Alexander and Kotelchuck, 2001; Lu et al., 2003, 2010). The primary focus of prenatal care has shifted over time from focusing on medical intervention to providing more comprehensive intervention and prevention with public health approaches (Lu and Lu, 2008; Lu et al., 2010). Prenatal care originated from research conducted in early 20th-century England by John W. Ballantyne, who proposed that "to prevent fetal abnormalities and reduce maternal, fetal, and neonatal deaths, medical supervision for pregnant women should be provided throughout pregnancy rather than only during labor" (Lu and Lu, 2008, p. 592). In the United States, prenatal care began with a program of nurse home visiting to pregnant women by Mrs. William Lowell Putnam at the Boston Lying-In Hospital in 1901, which led to the establishment of an outpatient clinic in 1911 that provided prenatal visits consisting of history and physical examination, blood pressure measurement, and urinalysis (Lu and Lu, 2008). From the beginning, the content of prenatal care was influenced by concerns about toxemia (preeclampsia), which was diagnosed by high blood pressure and excess protein in the urine. Such concerns also contributed to establishing the timing and frequency of prenatal visits (Lu and Lu, 2008).

Several studies (Eisner et al., 1979; Gortmaker, 1979; Greenberg, 1983; IOM, 1973; Taffel, 1978) published in the 1970s found a significant association between no prenatal care and the incidence of low birth weight (LBW), a leading cause of infant mortality and perinatal disparities. Citing these studies, a 1985 IOM report concluded that the "overwhelming weight of the evidence is that prenatal care reduces low birthweight" (p. 146) and promoted prenatal care as a key population-wide public health intervention for improving birth outcomes in the United States (IOM, 1985). In 1986, the U.S. Public Health Service assembled an expert panel to assess the content of prenatal care. In its 1989 report, the expert panel identified three basic components of prenatal care: (1) early and continuing risk assessment, (2) health promotion, and (3) medical and psychosocial interventions and follow-up (NIH, 1989). Soon thereafter, Congress enacted a series of legislative initiatives that incrementally expanded Medicaid eligibility to low-income pregnant women and children independent of their welfare status. Many states then further expanded Medicaid eligibility and streamlined the process of enrollment into prenatal care (Handler et al., 2011). Arguments for expansion of access to prenatal care were bolstered by cost-effectiveness analyses, which suggested that savings could be achieved by reducing LBW, though the cost savings may have been overstated (Huntington and Connell, 1994).

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In part stemming from these national and state policies, the adoption of timely and adequate prenatal care has increased substantially over the past few decades (Kogan et al., 1998; Martin et al., 2002; Piper et al., 1994). This increase, however, did not lead to an immediate reduction in LBW or disparities in birth outcomes. While many reasons could have contributed to the persistent poor outcomes (Alexander and Slay, 2002), some began to question the effectiveness of prenatal care as a population-wide strategy for improving birth outcomes. Two reviews published in 1995 raised concerns regarding the validity of the evidence used to support the benefit of prenatal care (Alexander and Korenbrot, 1995; Fiscella, 1995). Citing problems with inconsistent results, insufficient adjustment for prematurity bias, and inadequate control for the effect of critical confounders and potential selection bias in earlier studies, Fiscella concluded that "current evidence does not satisfy the criteria necessary to establish that prenatal care definitely improves birth outcomes" (Fiscella, 1995, p. 475). Alexander and Korenbrot (1995) also concluded from their systematic review that "[t]here is little done during the standard prenatal care visit that could be expected to reduce low birth weight" (p. 113). Lu et al. (2003) concluded from a review of the *content* of prenatal care in 2003 that neither preterm birth nor intrauterine growth restriction (IUGR)—the twin constituents of LBW—can be effectively prevented by prenatal care in its present form. They contended that "[p]reventing LBW will require reconceptualization of prenatal care as part of a longitudinally and contextually integrated strategy to promote optimal development of women's reproductive health not only during pregnancy, but over the life course" (Lu et al., 2003, p. 362).

These critiques led to a dampening of enthusiasm for prenatal care and a search for alternative strategies, such as bolstering preconception care services, to improve birth and child health outcomes in the United States. It should be noted, however, that most extant studies examined prenatal care in a limited form, addressing primarily clinical risk factors for pregnancy complications rather than what truly matters to both maternal (and paternal) health and the developmental origins of the child's future health. They also focused on a few birth outcomes rather than examining a broader array of health and developmental outcomes for children and families (Lu et al., 2010). While there is great evidence that pregnancy is a critical life event and a sensitive period for healthy child development, experts still disagree about how the health care system should effectively organize and deliver prenatal care.

Later in this chapter, the committee calls for a redesign of prenatal care to improve access, content, quality, delivery, and financing. Access can be improved with outreach, care coordination, and technology. Content could be expanded to include more detailed assessment, education, and management of psychosocial and environmental risks. Quality improvement efforts could address implicit bias and unequal treatment. Most importantly, this committee calls for a transformation of the organization and delivery of prenatal care to achieve greater vertical, horizontal, and longitudinal integration, including greater linkages to community services and women's health across the life course. This report does not address health systems redesign to better support childbirth in the United States, as it is the subject of an ongoing study by another National Academies committee.[4]

[4] See https://sites.nationalacademies.org/DBASSE/BCYF/Research_Issues_in_the_Assessment_of_Birth_Settings/index.htm (accessed July 26, 2019).

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Postpartum Care

The postpartum period marks the time after delivery when maternal physiology returns to the non-pregnant state. This period, often referred to as the "fourth trimester," is generally considered to last 6–8 weeks (ACOG, 2018b). Many traditional cultures prescribe 30 to 40 days of rest and recovery (ACOG, 2018b), such as zuò yuè zi ("doing the month") in China and Taiwan (Pillsbury et al., 1978), a 21-day period of rest called sam chil il in Korea (Dennis et al., 2007; Park and Dimigen, 1995), la cuarentena (which also means quarantine and comes from cuarenta, the word for 40 in Spanish) in Latin America, and lying-in, which gave rise to the establishment of lying-in hospitals in England and the United States during the 20th century (Eberhard-Gran et al., 2010). In many cultures, the woman and her newborn are surrounded by family and community members who offer instrumental emotional support during this period. In the United States, however, many women have to navigate the postpartum transition on their own with little formal or informal support, wrestling with lack of sleep, fatigue, pain, stress, breastfeeding difficulties, and new onset or exacerbation of preexisting health and social issues, such as postpartum depression, substance dependence, intimate partner violence, and other concerns (ACOG, 2018b).

Postpartum care provides an opportunity to address these issues, but for many U.S. women, it is often limited to a single 6-week postpartum visit, and some women receive no postpartum care at all. In the Listening to Mothers III Survey, one-third of respondents reported attending one postpartum office visit, while 1 in 10 mothers reported not having a visit (Declercq et al., 2013). Among the latter group, "I felt fine and didn't need to go" (42 percent), "I felt that I had already completed all of my maternity care" (18 percent), "too hard to get to office" (12 percent), and "didn't have insurance" (7 percent) were the most common reasons given for not having a visit (Declercq et al., 2013, p. ix). Nonattendance is greater among certain groups, including low-income women, Medicaid insurance holders, and those with inadequate prenatal care (DiBari et al., 2014). One recent study of Medicaid deliveries in California found that only half of all women, and one third of African American women, attended a postpartum visit (Thiel de Bocanegra et al., 2017).

Even for women who receive postpartum care, the typical 6-week postpartum visit may be too late to address some early onset issues and too limited to address other late onset or persistent problems. For example, more than half of postpartum strokes occur within 10 days of discharge (Too et al., 2018) and 17.5 percent of pregnancy-related deaths occur between 43 and 365 days postpartum, often as a result of cardiomyopathy or mental health conditions (Building U.S. Capacity to Review and Prevent Maternal Deaths, 2018), which a 6-week postpartum visit may be ill equipped to prevent. Rather than an arbitrary "6-week" check, in 2018, the American College of Obstetricians and Gynecologists (ACOG) called for a reconceptualization of postpartum care as an ongoing process, the timing of which should be individualized and woman centered (ACOG, 2018b). ACOG recommends that all women have contact with their obstetric care providers within the first 3 weeks postpartum. Women with hypertensive disorders of pregnancy should be seen no later than 7–10 days postpartum, and women with severe hypertension should be seen within 72 hours (ACOG, 2018b). ACOG recommends that this initial assessment should be followed up with ongoing care as needed, concluding with a comprehensive postpartum visit no later than 12 weeks after birth.

ACOG also recommends that the comprehensive postpartum visit include a full assessment of physical, social, and psychological well-being, including the following domains: mood and emotional well-being; infant care and feeding; sexuality, contraception, and birth

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spacing; sleep and fatigue; physical recovery from birth; chronic disease management; and health maintenance (ACOG, 2018b). In practice, however, providing comprehensive postpartum care is disincentivized by prevailing health care financing practices, whereby the postpartum visit is bundled with the rest of obstetrical care, for which providers receive a global payment with no additional reimbursement for postpartum care. To deliver comprehensive postpartum care, ACOG also recommends an interprofessional postpartum care team, which consists of the primary maternity care provider, infant health care provider, primary care provider, and specialty consultants, as well as a care coordinator or case manager, lactation support provider, home visitor, and family and friends (ACOG, 2018b). Presently, the care settings where many women receive postpartum care preclude such interprofessional approaches, and transforming the organization and delivery of postpartum care in the absence of additional reimbursement will likely be challenging. Lastly, ACOG recommends that the postpartum care team help facilitate transition to ongoing well-woman care (ACOG, 2018b). This is especially important for women who have chronic health conditions or have experienced a pregnancy complication, given growing research suggesting that pregnancy may be a window to a woman's future health outlook. For example, women with pregnancies complicated by preterm birth, gestational diabetes, or hypertensive disorders of pregnancy have a higher lifetime risk of maternal cardiometabolic disease (Dassanayake et al., 2019). However, for many low-income women, especially in states without Medicaid expansion, access to ongoing well-woman care becomes limited upon termination of their Medicaid coverage at 60 days postpartum.

Recognizing the importance of the postpartum period as a critical time for a woman and her infant that sets the stage for their long-term health and well-being, this committee calls for a redesign of postpartum care to improve access, content, quality, delivery, and financing to better leverage its potential for advancing health equity. As will be discussed later in the chapter, access could be enhanced by colocation of maternal and infant services; greater use of home visiting, doula services, community health workers, and mHealth technology; and increased access to paid family and medical leave. Expanded care is needed to more holistically address not only clinical issues but also psychosocial and environmental concerns, with greater attention to social determinants of maternal, child, and family health and well-being. Quality could be improved by promoting quality measurement and continuous quality improvement (CQI), supporting workforce training, and, as discussed throughout the chapter, addressing implicit bias and unequal treatment along the care continuum. Organization and delivery of care could be strengthened through care coordination, systems integration, and interprofessional teamwork. To support this redesign, the committee calls for developing and testing innovative financing models, including pay-for-performance and pay-for-outcomes, unbundling postpartum care from global payment, and extending Medicaid coverage for 1 year postpartum.

Pediatric Care

The goal of pediatric care is to provide services to children and families that will improve their health status and functioning. Usual services include brief clinical encounters ("checkups") beginning in the hospital immediately after birth and continuing through childhood and adolescence, with decreasing visit frequency. Typical services have traditionally included providing regular immunizations, monitoring growth and development and nutrition, advising parents on common aspects of child development, and managing common illnesses and injuries. Child health care has a focus on prevention and includes regular screening for a wide range of

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conditions, including drowning risk, lead exposure, anemia, adversity, hunger, and infectious diseases, as well as child behavior and development.

Much of the work in pediatric well-child care grew from early efforts of the U.S. Children's Bureau and then the growth of public health departments in the states (Lesser, 1965). The Children's Bureau emphasized early nutrition and safe milk for babies and checked feeding and weight gain. With the development of immunizations to protect children (and communities) from dangerous infectious diseases, many state health departments developed immunization clinics. The 1921 Sheppard-Towner Act[5] led to substantial federal investment in well-child health programs. The American Medical Association (AMA) opposed the Act as government intrusion into health care, leading to the departure of pediatricians from the AMA and the formation of the American Academy of Pediatrics (AAP) (Baker, 1994). With the growth of the AAP, pediatricians (many of whom had worked in the public well-child clinics) increasingly integrated the elements of well-child care into their practices, ultimately overshadowing public service programs (Baker, 1994). Thus, much of pediatric well-child care grew from notions of improving child nutrition and weight gain and preventing serious infections (e.g., diphtheria, tetanus, polio). Many of those acute infections have disappeared (although immunizations remain critically important) and have been replaced with epidemics of chronic diseases in pediatric populations. Where less than 2 percent of children in 1960 had a serious chronic condition that interfered on a daily basis with their usual activities (e.g., school and play), by 2010, over 8 percent had such conditions, representing a 400 percent growth in these conditions (Halfon et al., 2012; Houtrow et al., 2014; Newacheck et al., 1986; Perrin et al., 2014). These high rates of chronic conditions mainly reflect greater numbers of children with obesity, asthma, neurodevelopmental conditions (especially autism spectrum disorders [ASDs]), and mental health conditions. While rates of mental health conditions increase with child age, 1 in 6 or more children ages 2–8 have diagnosed mental, behavioral, or developmental conditions (CDC, 2019a). Children with chronic medical conditions also have higher rates of mental and behavioral conditions than similar children without chronic conditions (Perrin et al., 2019).

These growing rates have led health care providers for children to recognize the importance of behavioral health in all aspects of pediatric care—the interaction of physical and mental/behavioral health and their common coexistence, along with the increasing rates of MBH conditions in children and adolescents. Initially codified in some of the Rochester Child Health Studies over a half century ago (Haggerty et al., 1975), this interest has grown into active work by several professional groups to build capacity and competence in behavioral health into the well-child experience (AAP, 2018; Foy and American Academy of Pediatrics Task Force on Mental Health, 2010). MBH accounts for a large proportion of visits in pediatric primary care and complicate care for most chronic conditions.

Screening is a key step in prevention and an integral part of pediatric care. Given the brief time available in a child health supervision visit, pediatric clinicians cannot do all recommended screening, so they make choices for their practices, based in part on the characteristics of their patient population. Generally, child health practitioners (e.g., pediatricians, family physicians, family nurse practitioners, physician's assistants) choose screening instruments that are brief, easily scored and interpreted, and able to identify conditions

[5] Also known as the National Maternity and Infancy Protection Act, the Sheppard-Towner Act provided states with federal funding to develop programs that would increase education of prenatal and infant care. The act was passed in an effort to decrease the high rates of infant mortality in the United Sates (https://embryo.asu.edu/pages/sheppard-towner-maternity-and-infancy-protection-act-1921 [accessed April 19, 2019]).

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that have moderate prevalence or saliency in their practice communities (AAP, 2016). Most child health professionals screen for growth and development, including behavioral issues and developmental delays, and for certain conditions that early treatment may ameliorate, including ASD. AAP has codified much of child and adolescent preventive care in its document, *Bright Futures*, and the Patient Protection and Affordable Care Act (ACA) included *Bright Futures* as the basis for pediatric preventive care (AAP, 2016).

The high rates and persistence of poverty among America's children have led many groups to address this problem. To address food insecurity, AAP and the Food Research & Action Center (FRAC) developed a toolkit for pediatricians that includes a validated, AAP-recommended two-question screening tool called The Hunger Vital Sign™ (AAP and FRAC, 2017). In addition, AAP, recognizing that poverty affects so many aspects of child health and development and essentially all types of clinical problems, joined these efforts by making poverty a major segment of its Agenda for Children (AAP, n.d.-a). All pediatricians face the consequences of child poverty in their practices. Children with leukemia and other serious conditions experience higher mortality, and poverty affects all children's ability to adhere to medication and treatment (Mishra et al., 2011). Low-income children have higher rates of most chronic conditions and typically have more severe cases (WHO, n.d.). Poverty affects parents' resources to care for their children's illnesses and limits access to many treatment services. AAP has educated pediatricians about poverty and what they can do in their practices (AAP Council on Community Pediatrics, 2016), and pediatricians increasingly screen for poverty and other SDOH, particularly checking for access to day care, home and neighborhood safety, hunger, and housing, especially along with partnering community agencies (Garg et al., 2015; Shekarchi et al., 2018).

Furthermore, pediatricians have provided much leadership in the development of life course sciences, theory, and practice, in part because of their perspective based in the dynamics of child development in the context of families and communities, along with their recognition of the early childhood antecedents of many long-term health and mental health conditions (Halfon, 2012; Halfon and Hochstein, 2002).

Over the past few decades, there has been significant growth in diversity in the pediatric population without similar diversification of the pediatrics workforce (AAP Committee on Pediatric Workforce, 2013). Recognition of disparities has driven pediatric efforts to address equity and assure equal access to services and treatment. Efforts to transform into team care and use telehealth mechanisms in part reflect the recognition that such changes will help pediatricians more effectively work on issues of poverty in their patients and communities. These changes acknowledge that enhancing pediatric practice with personnel knowledgeable and skillful in helping families access a breadth of community services will help address disparities.

Pediatric care in the United States is organized across a variety of small and large practices, federally qualified health centers (FQHCs), and hospital-based programs (for general and specialty care). Increasingly, smaller, community-based practices have merged to become larger, multisite practices, some limited to the care of children and youth, while others are multispecialty programs offering a larger range of services to a population of all ages. Many children's hospitals have hospital-based primary care programs and/or organized relationships or networks with primary care practices in their surrounding communities. Both safety net hospitals and FQHCs more often serve a large, low-income, Medicaid-insured population (Nath et al., 2016) with overrepresentation of black and Latino children and youth (Georgetown University Center for Children and Families, 2017, 2018). Pediatricians have pioneered the concept of the

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medical home, an organized and central place to coordinate all the health care needs of a patient or family. In recent years, this notion has spread among many other physician groups, especially family medicine and internal medicine. Expanded visions of the medical home increasingly embrace characteristics of community-based, comprehensive care (Ader et al., 2015; Bair-Merritt et al., 2015; Homer et al., 2008; Patient-Centered Primary Care Collaborative, n.d.; Stille et al., 2010). Medical home have helped efforts, especially in family medicine and pediatrics, to move to a whole-family, whole-child approach. Routine screening in pediatric care for parental mental health (especially postpartum depression) and other risks (e.g. smoking, firearms) and inclusion of parent training programs in pediatric programs are examples of ways that clinical care has become more family focused.

The distribution of subspecialists—physicians who care for more complex and usually chronic conditions—differs for children and adults. Adult-treating subspecialists are widely distributed, but subspecialists for children are mainly centralized in hospitals that care for large numbers of children and youth. This distribution leads to different problems with access for young children and families. These children's hospital programs, some freestanding and others part of larger general hospitals, provide the majority of health care for children with highly specialized needs—the groups with complex medical conditions and rarer childhood conditions. The substantial numbers of children and youth with chronic and complex health conditions, especially those with less common chronic conditions, need regular access to specialized pediatric care (e.g., specialized surgeons or pediatric cardiology).

> *Conclusion 5-1: The current health care system focuses mainly on clinical goals and addresses the multiple other determinants of health in fragmented and highly variable ways. Despite high-quality clinical care, the health status of America's children and young families is far worse than in comparable developed countries. U.S. health care provides only limited attention to integration of health care for the whole family, health care across the life course, or integration of mental and behavioral health with the rest of health care.*

The need for the integration of health care and the whole-family approach is discussed in more detail later in the chapter.

IMPROVING ACCESS TO HEALTH CARE SERVICES

To realize the full potential of health care services, people need access to regular primary and preventive care across the life course. The nation will be best served by a health care system that guarantees all people universal access to high-quality health care across the life course and in which preconception, prenatal, and pediatric care represent a series of well-timed, more intense encounters with a broader array of services ("boosters") during critical junctures in life. Cramming a life course worth of health care into a single preconception visit (or even a few visits) a few months before attempting to conceive will do little to advance equity in birth outcomes or children's health. Similarly, prenatal and pediatric care that is primarily based on episodic, short visits to a medical clinic or office for a narrow range of clinical services scheduled when convenient for health care systems and providers is not enough to reverse the trend of centuries of inequitable health care treatment and outcomes experienced by our nation's children.

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Health Insurance Is Necessary But Not Sufficient

Health insurance is a major facilitator to ensuring access to health care services; lack of insurance coverage is a significant barrier (Bailey et al., 2016; Choi et al., 2011; DeVoe et al., 2010, 2012a; Howell and Kenney, 2012; IOM, 2002; Sommers et al., 2015, 2017a; Tumin et al., 2019; Wallace and Sommers, 2016; Wherry et al., 2016). A Commonwealth Fund report found that insured women were significantly more likely than uninsured women to receive cancer screenings and other preventive services and to have a regular source of care (Gunja et al., 2017) (see Figure 5-1). Even for those with a usual source of care, health insurance improves access to more comprehensive services (DeVoe et al., 2008c, 2012a). For children, health insurance substantially improves health care access and use (IOM, 1998). The large majority of children and youth in the United States (about 95 percent) have health insurance coverage, with about 30 to 40 percent of it from public sources: Medicaid and CHIP (Alker and Pham, 2018; Cornachione et al., 2016). The small percentage of children and youth without health insurance use fewer health care services and fare much worse on measures of health and health care quality (DeVoe et al., 2008a, 2009b, 2010, 2012b).

Although studies of health insurance interventions are usually natural experiments, the Oregon Health Insurance Experiment, in which a 2008 lottery extended Medicaid to selected residents, represented a rare opportunity to assess the impact of insurance coverage through a randomized study design (James, 2015). Findings from the experiment confirm the well-documented associations between an individual's health insurance and access to health care and the causal link between a parent having access to insurance and a child gaining coverage (Bailey et al., 2016; DeVoe et al., 2015b,c; Gold et al., 2014; Hatch et al., 2016; Marino et al., 2016; O'Malley et al., 2016). Efforts to achieve expanded coverage in Massachusetts and several other states also led to similar landmark studies showing the importance of health insurance as a facilitator of health care access (Finkelstein et al., 2012; Smith et al., 2019).

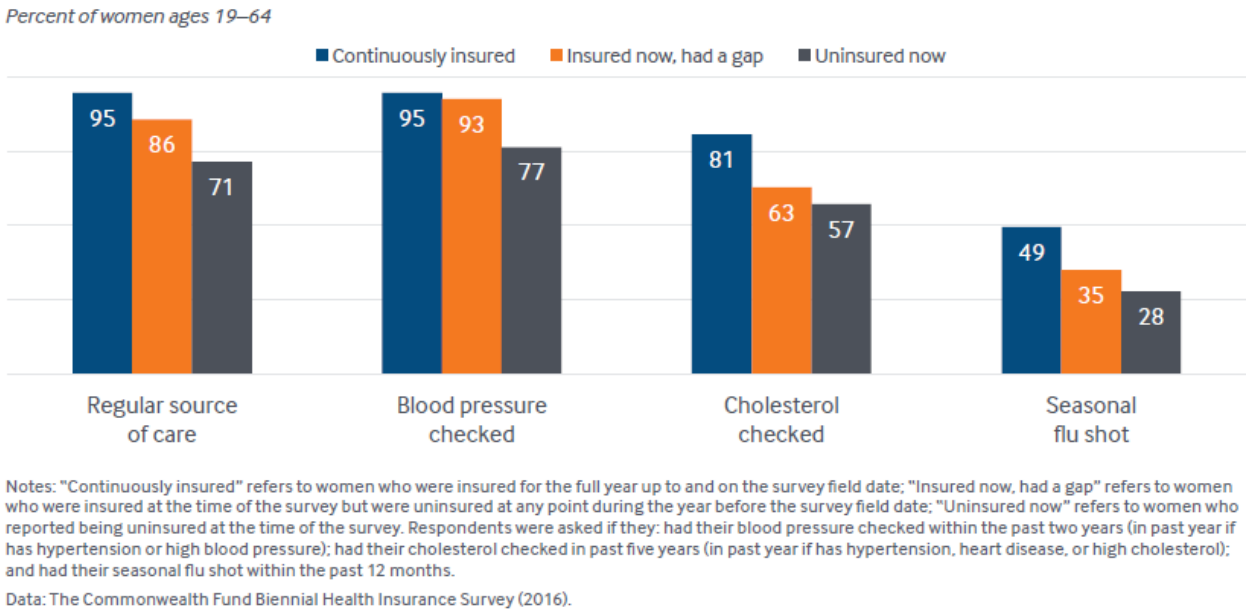

FIGURE 5-1: Women with a regular source of care and receiving preventive services by insurance status
SOURCES: Gunja et al., 2017, Commonwealth Fund

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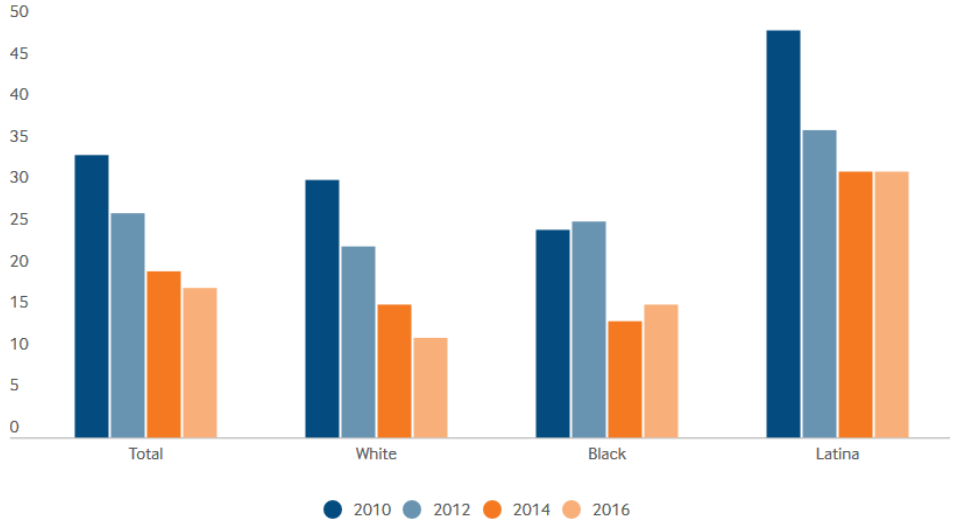
Shortly after Oregon's insurance expansions and related efforts to achieve expanded coverage in Massachusetts and elsewhere, the 2010 Patient Protection and Affordable Care Act (ACA) increased access to health insurance coverage nationally to millions of Americans through a combination of Medicaid expansions, private insurance reforms, and premium tax credits and subsidies through new "exchange" plans. After the ACA, the percentage of uninsured women decreased from 20 percent (19 million) in 2010 to 11 percent (11 million) in 2016 (Gunja et al., 2017). Low-income women and women of color have made particularly large gains. Women ages 19–64 earning less than 200 percent of the federal poverty level (FPL) who are uninsured fell from 25 percent in 2010 to 16 percent in 2016 for black women, 49 percent in 2010 to 32 percent in 2016 for Latina women, and 34 percent in 2010 to 18 percent in 2016 overall (see Figure 5-2). For children and youth, the ACA also led to the lowest rates of child uninsurance in U.S. history. While some of the insurance gains for pediatric populations reflected new eligibility under the ACA, most gains resulted from newly insured parents learning of their children's eligibility for existing programs (esp., Medicaid and CHIP) (Garrett and Gangopadhyaya, 2015), reconfirming that insurance is a family affair (DeVoe et al., 2008b, 2009b, 2011a, 2015a,c,d; Dubay and Kenney, 2003; IOM, 2002; Yamauchi et al., 2013).



Notes: FPL refers to federal poverty level. Income levels are for a family of four in 2016. Rates are for those uninsured at the time of the survey.
Data: The Commonwealth Fund Biennial Health Insurance Surveys (2010, 2012, 2014, 2016).

FIGURE 5-2: Percent of women ages 19–64 who are uninsured and early less than 200 percent of the FPL
SOURCE: Gunja et al., 2017, Commonwealth Fund

Prior to the ACA expansions, insurance coverage was not accessible to many low-income women of child-bearing age unless they became pregnant. Even after becoming pregnant, some women experienced long delays to obtain coverage, and some were unable to access care until the second or third trimester (see Table 5-3). CDC reported that in 2009, the percentage of women who were uninsured decreased from 23.4 percent in the month before pregnancy to 1.5 percent at the time of delivery, while Medicaid coverage increased from 16.6 percent in the month before pregnancy to 43.9 percent at delivery (D'Angelo et al., 2015). Even with the recent

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gains in access to coverage before and during a pregnancy, more than 1 in 5 (21.3 percent) American women who gave birth in 2016 began prenatal care after the first trimester; 4.6 percent began care in the third trimester, and 1.6 percent received no care at all. In all, approximately 15 percent of American women received inadequate prenatal care. Moreover, there are significant racial-ethnic, socioeconomic, and geographic disparities in prenatal care use (Osterman and Martin, 2018).

In addition to access to coverage, the ACA further improved access to essential care for women, especially preconception care, by mandating coverage for women's preventive services, including all FDA-approved contraceptive methods and counseling and at least one well-woman preventive care visit per year with no cost-sharing (HRSA, 2018; Women's Preventive Services Initiative, n.d.). Preconception care is also covered under well-woman preventive visits (Women's Preventive Services Initiative, n.d.). Additionally, the law mandated coverage for essential health benefits (EHBs), including maternity care and mental health services. The ACA prohibited gender rating (charging women a higher premium than men) and banned lifetime caps on benefits and exclusions based on preexisting conditions, which protected access for women with chronic conditions. A Commonwealth Fund report found that between 2010 and 2016, the percent of women ages 19–64 with health conditions who found it very difficult or impossible to find coverage they needed in the individual market decreased by nearly half, while those who said they were not getting care because of costs (did not fill prescription, skipped recommended test or treatment, had a medical problem but did not seek primary or specialty care) decreased from 48 percent to 38 percent (Gunja et al., 2017). Several other studies have also shown similar improvements (Angier et al., 2015, 2017, 2019a; Heintzman et al., 2017; Hoopes et al., 2016; Huguet et al., 2017, 2018; Sommers et al., 2016, 2017b).

Presently, it is unclear what effects recent efforts to deregulate the ACA, such as the repeal of the individual mandate, proposed extension of short-term coverage policies (which do not have to comply with many ACA consumer protections) (Keith, 2018), state waiver for EHBs, and religious and moral exemptions from contraceptive coverage mandates, will have on access to primary and preventive services for both men and women. It is likely that, as with prior policies that scaled back insurance access, these repeal efforts will negatively impact access to and use of recommended primary and preventive care services (Carlson et al., 2006; DeVoe et al., 2012a; Solotaroff et al., 2005; Tumin et al., 2019).

As demonstrated above, Medicaid is an important source of insurance coverage and facilitator of basic access to women's health care services, particularly for low-income women. In 2014, Medicaid provided more than 25 million low-income women with health and long-term care coverage (Henry Kaiser Family Foundation, 2017), approximately two thirds of them between ages 19 and 49. Medicaid is particularly important for low-income women and women of color. While overall 1 in 5 (20 percent) women of reproductive age were enrolled in Medicaid in 2015, 27 percent of Hispanic women, 31 percent of African American women, and nearly half (48 percent) of low-income women in the United States were enrolled (Sonfield, 2017). Expansion of Medicaid coverage for pregnant women in the 1990s led to significant increases in access to and use of prenatal care; today, Medicaid covers nearly half of all births in the United States (Kaiser Family Foundation, 2017). For nonpregnant women, the ACA legislation extended Medicaid eligibility to all individuals with household incomes up to 138 percent of the FPL; however, a 2012 Supreme Court ruling in *National Federation of Independent Business v. Sebelius* made Medicaid expansion optional for states, resulting in inconsistent coverage policies across the country (Henry Kaiser Family Foundation, 2017). As of May 2019, 36 states and DC

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have expanded Medicaid; 14 states have not (Henry Kaiser Family Foundation, 2019). Overall, the rate of uninsurance among low-income women of reproductive age decreased by 13.2 percentage points due to ACA Medicaid expansions (Johnston et al., 2018). A recent study found that from 2011 to 2016, states that expanded Medicaid showed significant improvements in black-white disparities in preterm, very preterm, LBW, and very LBW rates, compared to states that did not (Brown et al., 2019). Studies have shown that in states that expanded Medicaid, disparities by race and ethnicity in the rates of insurance coverage and access to care have narrowed more than in states that have not (Artiga et al., 2019; Hayes et al., 2017).

TABLE 5-3: Trimester that Prenatal Care Began, by Selected Characteristics: United States, 2016

			Timing of PNC		
				Late or no PNC[1]	
Selected characteristic	First trimester	Second trimester	Total	Late PNC[2]	No PNC
			Percent		
Total	77.1	16.7	6.2	4.6	1.6
Age of mother					
Under 20	61.2	27.6	11.2	8.3	2.9
Under 15	36.7	37.6	25.7	19.2	6.5
15–19	61.5	27.5	11.0	8.2	2.9
20–24	70.3	21.7	8.0	5.9	2.1
25–29	77.8	16.3	6.0	4.4	1.5
30–34	82.1	13.1	4.8	3.6	1.2
35–39	81.7	13.5	4.8	3.6	1.2
40 and over	78.4	16.0	5.6	4.1	1.5
Race and Hispanic origin					
Non-Hispanic, single-race:					
White	82.3	13.4	4.3	3.3	1.1
Black	66.5	23.5	10.0	7.0	3.0
American Indian or Alaska Native	63.0	24.5	12.5	9.2	3.3
Asian	80.6	14.0	5.4	4.6	0.8
Asian Indian	83.4	12.1	4.6	3.9	0.7
Chinese	81.2	11.4	7.4	6.9	0.5
Filipino	82.8	13.4	3.8	3.0	0.8
Japanese	85.5	10.5	4.0	3.2	0.8
Korean	85.3	10.6	4.1	3.4	0.7
Vietnamese	80.2	15.3	4.5	3.3	1.2
Other Asian	71.7	22.0	6.3	5.0	1.3
Native Hawaiian or Other Pacific Islander	51.9	28.9	19.2	14.2	5.0
Hawaiian	69.9	20.5	9.6	6.0	3.6
Guamanian	72.1	21.4	6.6	5.0	1.6
Samoan	56.9	29.1	14.0	10.4	3.6
Other Pacific Islander	43.8	31.3	24.8	18.4	6.4
Hispanic	72.0	20.3	7.7	5.6	2.1
Mexican	71.4	20.6	8.0	5.7	2.3

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Puerto Rican	76.2	18.1	5.7	4.3	1.4
Cuban	82.0	14.1	3.8	2.9	0.9
Central or South American	68.1	22.7	9.2	6.9	2.2
Other and unknown Hispanic	74.3	18.8	6.8	5.1	1.7
Live-birth order					
1st birth	79.0	15.3	5.7	4.4	1.3
2nd birth	80.1	14.8	5.1	3.9	1.2
3rd birth	75.8	18.0	6.2	4.6	1.6
4th birth or higher	66.2	23.8	10.0	6.7	3.3
Educational attainment[3,4]					
Less than high school	62.7	26.1	11.2	7.5	3.7
High school	73.4	19.5	7.2	5.0	2.2
Some college[5]	80.2	15.1	4.7	3.5	1.2
Bachelor's degree or higher	87.6	9.1	3.3	2.8	0.5
Source of payment for the delivery					
Medicaid.	68.1	23.3	8.6	6.4	2.2
Private insurance	87.0	10.3	2.7	2.1	0.6
Self-pay	54.8	25.4	19.8	13.2	6.6
Other[6]	75.0	16.8	8.2	5.8	2.3

[1]PNC that began in the third trimester and no PNC.

[2]PNC that began in the third trimester.

[3]Excludes women under age 25.

[4]Significantly increasing trend in first trimester PNC by educational attainment ($p < 0.05$).

[5]Includes associate's degree.

[6]Includes Indian Health Service, CHAMPUS (Civilian Health and Medical Program of the Uniformed Services) or TRICARE, other government (federal, state, or local), and charity. NOTES: PNC is prenatal care. Chi-squared test statistics for each variable by trimester prenatal care began were statistically significant ($p < 0.05$).

SOURCE: Osterman and Martin, 2018

Medicaid and CHIP finance health insurance for nearly 50 percent of U.S. children and youth. Children and youth with public insurance have about the same rates of use and quality of care as those with commercial insurance (DeVoe et al., 2011c,d), although those data do not account for the higher rates of chronic conditions and disability among publicly insured children. Although most U.S. children and youth currently have health insurance, children who are low income or from racial and ethnic minority populations face problems with lack of access to ongoing, comprehensive health care services (McCormick et al., 2001; Weinick and Krauss, 2000). Children and youth with public insurance (Medicaid and CHIP) disproportionately include black and Latino populations (Georgetown University Center for Children and Families, 2017, 2018).

Medicaid demonstration waivers have shown promise in allowing states to test new approaches in Medicaid that differ from federal program rules (Musumeci et al., 2018) in order to expand and broaden access to coverage for populations that have traditionally not been eligible for Medicaid. For example, 27 states have established limited-scope Medicaid family planning programs through waivers to extend access to family planning services to uninsured

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women who do not qualify for full Medicaid coverage. This includes low-income women whose incomes are not low enough or who have lost Medicaid eligibility after giving birth (Henry Kaiser Family Foundation, 2017). Additionally, Louisiana has added an interpregnancy component to their Medicaid waiver to provide interconception care for low-income, high-risk women who had an adverse birth outcome (ASTHO, 2013). Medicaid waivers have also helped children and youth with special health care needs access a number of treatment services in home and community settings. The original Medicaid waiver was for an Iowa child who had Medicaid coverage while hospitalized (because her parents' income was not considered for inpatient eligibility) but would lose it if she came home with multiple and complex needed services. The waiver allowed hospital care to be substituted with home care (Perrin et al., 1993).

Other Medicaid demonstration waivers have been proposed that could negatively impact eligibility, enrollment, and benefits, including work and reporting requirements, coverage lock-outs, premium cost-sharing, restrictions on presumptive eligibility and retroactive coverage, and time limits on coverage (Flowers and Accius, 2019). The impact of these waivers on access to preconception, prenatal, and pediatric health care services is still being studied; however, similar policies making Medicaid eligibility more restrictive in the past (e.g., proof of citizenship requirements) have had negative effects on access to care (Angus and Devoe, 2010; Bauer et al., 2011; Hatch et al., 2014). Arkansas has seen thousands of working-age adults, including many with dependent children at home, lose Medicaid benefits, along with no growth in job participation (Rudowitz et al., 2019; Sommers et al., 2019). However, federal courts struck down Arkansas's work requirement for expanded Medicaid coverage in March 2019. Recent reports indicate the first increase in rates of uninsurance among children in a decade, in part reflecting new restrictions on parents' access (Alker and Pham, 2018).

Many Factors Affect Access to Health Care Services

A complex array of factors beyond insurance coverage influence preconception, prenatal, and pediatric care use (Heaman et al., 2014; Kalmuss and Fennelly, 1990). Some of these factors include the complexity of household needs and social challenges (e.g., child care, transportation, addictions, lack of support), caregiver qualities (lack of time, negative behaviors), health system barriers (shortage of providers), and program/service characteristics (distance, long waits, short visits). Such barriers (particularly transportation to health care facilities, ability to pay for needed treatments, taking time from work for health care appointments, and costs related to housing and food) may take priority over health care for families.

Regarding access to prenatal care, a study of 246 African American women residing in Washington, DC identified psychosocial stress, substance use, child care problems, negative attitudes toward pregnancy and prenatal care, insurance/financial constraints, and nonparticipation in the Special Supplemental Nutrition Program for Women, Infants, and Children (WIC) program as key determinants of lower than recommended rates of prenatal care use (Johnson et al., 2007). Another study of low-income African American women in Milwaukee, Wisconsin identified structural barriers to prenatal care, such as transportation and insurance, negative or ambivalent attitudes toward prenatal care, perceived poor quality of care, and unintended pregnancy, and psychosocial stressors, such as overall life stress and chaos (Mazul et al., 2017).

Children and youth with special health care needs (SHCN) represent a population that experiences unique access challenges. A sizable percentage of U.S. children (10–20 percent or more, depending on the definition; about 11.4 percent of children ages 0–5 and 22.7 percent of

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children ages 6–11) have SHCN (HHS, 2015; NASEM, 2018) and their needs depend on the severity and prevalence of the condition. Most children and youth with more common conditions (asthma, obesity, mental health conditions, and neurodevelopmental conditions) receive the bulk of their care in community pediatric settings, including hospital outpatient clinics. Children and youth with more severe and less common conditions get a large part of their care from specialized pediatric hospitals (Perrin et al., 2014). Many households have difficulty gaining access to specialized care because most subspecialists are found in centralized children's hospitals, which may be distant from their homes. Children insured by Medicaid, disproportionately black or Latino (Georgetown University Center for Children and Families, 2017, 2018), particularly experience problems in accessing subspecialists (Bisgaier and Rhodes, 2011). Furthermore, narrow insurance networks may not include all needed specialists, creating barriers to needed subspecialty care.

Assuring access to quality health care requires addressing both financial and nonfinancial barriers. Lu et al. (2010) identified a number of promising strategies to increase financial and nonfinancial access to timely care, including policy initiatives to promote innovative care models, support outreach and care coordination, and increase provider participation in Medicaid-funded care. State Medicaid agencies have increasingly relied on managed care organizations (private-sector companies that contract with Medicaid to manage the program) to implement and manage Medicaid programs. Currently, about 80 percent of Medicaid recipients receive care through a managed care arrangement (MACPAC, 2016). Many managed care organizations have experimented with new programs or financing, although there are few consistent patterns (Institute for Medicaid Innovation, n.d.). It is also not yet apparent whether this trend has had a significant impact on reducing health disparities. Another example is Oregon's coordinated care organization (CCO) model, implemented in 2012 and designed to improve the coordination of care for the state's Medicaid beneficiaries. In the state, CCOs cover a distinct geographic area and have broad budgeting authority for Medicaid and CHIP funding within that area, along with incentives to improve quality and broaden attention to social determinants of care in part through active collaboration with schools and community agencies (Oregon Health Authority, 2018; Stecker, 2013). CCOs led to significant increases in early prenatal care initiation and a reduction in disparities across insurance types but no difference in overall prenatal care adequacy (Muoto et al., 2016).

Regarding access to postpartum care, colocating postpartum and well-baby care has been suggested as a strategy for improving access (Stuebe et al., 2019). Colocating care could reduce transportation, child care, medical leave, and other barriers and facilitate care coordination for issues that require joint assessment and management of both mother and infant, such as breastfeeding (Stuebe et al., 2019). Greater use of home visitors, doulas, and community health workers can also help improve access (Hans et al., 2018). Providing culturally congruent care to women of color can increase breastfeeding and reduce perinatal disparities (Kozhimannil et al., 2013), but only three states currently provide Medicaid coverage for doula services (Stuebe et al., 2019). Leveraging mHealth technology, such as SMS, remote blood pressure monitoring, and telehealth, can also improve postpartum follow-up (ACOG, 2018b; CMS Maternal & Infant Health Initiative, 2015). Presently, nearly one quarter of women return to work within 10 days postpartum, and nearly one half return to work within 40 days postpartum (Klerman et al., 2014); expanding paid family and medical leave could also increase access to and use of postpartum care (Rossin-Slater and Uniat, 2019) (see Chapter 6 for a discussion on paid family leave). Because many health and psychosocial issues persist or emerge beyond 60 days postpartum,

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some experts have called for extending Medicaid coverage for at least 12 months postpartum, especially in Medicaid nonexpansion states (Stuebe et al., 2019).

For nearly three decades, state Title V programs have played an important role in increasing access to and use of timely prenatal care and other family health care services through outreach and coordination by (1) facilitating partnerships among agencies that provide direct services to pregnant women; (2) helping to ensure that maternal and child health professionals in WIC, Head Start, and other public programs provide pregnant women with accurate and current information on coverage in their state; (3) increasing access to presumptive Medicaid eligibility, which provides pregnant women with access to immediate prenatal care; and (4) increasing continuity of coverage for low-income women who become pregnant (Association of Maternal & Child Health Programs, 2016). State and federal Maternal and Child Health Programs (Title V) have had even longer histories of involvement with very young children, following earlier Children's Bureau programs for safe milk and infant nutrition (Lesser, 1965). A t least 30 percent of federal support for state programs has been allocated for well-child programs, and at least 30 percent goes to programs for children with SHCN (formerly the Crippled Children's Service) (HRSA, 2019, n.d.). As with women's health, state programs have helped to facilitate partnerships across key agencies, improve knowledge of the health status of children and youth in the state, and link households with other agencies, especially Medicaid and the Social Security Administration (SSA) (for SSI coverage).

Recent growth of community acute care clinics—some operated by pharmacy chains and others by larger hospitals—have offered new community-based access to health care. These centers offer immediate and convenient care for (generally minor) acute conditions. The centers offer variable connections to children's and families' ongoing primary care; in several communities, such centers have worked to share medical records and referral information with ongoing primary care programs (Conners et al., 2017). Acute care centers, at times collaborating with telehealth companies, can offer immediate and convenient services. For example, a parent of a sick child can call from home (or work), reach a health care clinician, describe symptoms, and efficiently receive advice. Nonetheless, most innovative walk-in clinics, acute care centers, and commercial telehealth models have focused on middle-income communities and less on low-income areas. Thus, there is no evidence that they reduce disparities.

Improving Access to Needed Health Care

Based on its review of the evidence in the sections above, the committee concludes:

Conclusion 5-2: Although health insurance coverage has grown substantially in the past few decades, mainly through Medicaid expansions and other insurance enhancements, access remains a problem for many families with young children, who experience numerous barriers to obtaining health care services in addition to lack of health insurance coverage. Multiple agencies, especially the Maternal and Child Health Bureau, have worked to diminish barriers to care. Further efforts are needed to address financial and nonfinancial barriers to care and to ensure that all families have access to adequate preconception, prenatal, postpartum, and pediatric care.

To address the issues identified related to access to care, the committee recommends:

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Recommendation 5-1: The U.S. Department of Health and Human Services, state, tribal, and territorial Medicaid agencies, public and private payers, and state and federal policy makers should adopt policies and practices that ensure universal access to high-quality health care across the life course. This includes:

- **Increasing access to patient- and family-centered care,**
- **Ensuring access to preventive services and essential health benefits, and**
- **Increasing culturally and linguistically appropriate outreach and services.**

Achieving this recommendation requires:

- **Increasing access to patient- and family-centered care.** Specific actions include integrating services longitudinally, vertically, and horizontally to increase entry points to care for children and families; incorporating enabling services to facilitate access to care; expanding attention to whole family needs in clinical care; and overcoming household challenges, such as transportation and child care needs. (See the sections that follow for discussion on the need to integrate health care services across the life course.)
- **Supporting comprehensive access across the life course.** Expand comprehensive supports for health across the life course. Programs should increase awareness of and access to family planning services and general preventive health services that keep parents healthy and promote positive attachments essential to early life development.
- **Protecting access to benefits.** Assure continuous coverage and access for all men, women, and children. Coverage should include child and family preventive services and EHBs, with a prohibition against lifetime benefit caps and preexisting condition exclusions.
- **Actively promoting inclusion in coverage and care.** Promote culturally and linguistically appropriate outreach and services as well as increased diversity in the health care workforce.
- **Systematic application of measures:** For example, adoption of a measure to assess disparities in timely and adequate access to well-woman care, prenatal, and pediatric care as a national performance measure by the Title V Maternal and Child Health Block Grant.

PROMOTING QUALITY OF CARE

Quality health care is timely, equitable, safe, patient centered, efficient, and effective (IOM, 2001). Equitable care means care that does not vary in quality because of demographic characteristics, such as sex, ethnicity, geographic location, or SES (IOM, 2001). For populations who have access to health care services, the quality of the care they receive is not always safe and is often not equitable. Regarding preconception care, a survey of more than 800 women of reproductive age found that more than 1 in 4 (27 percent) were prescribed a medication with a potential teratogen (a potential cause of birth defects). Of these women, 43 percent received no counseling from their provider regarding the teratogenicity of the medication or the need for contraception (Schwarz et al., 2013). A recent review of 31 studies (Goossens, 2018) identified multiple barriers to high-quality preconception care at the provider level (unfavorable attitude

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and lack of knowledge of preconception care, not working in a medical discipline or clinical setting that provides maternity care, lack of clarity on the responsibility for providing preconception care) and the client level (not contacting a health care provider in the preconception stage, negative attitude, and lack of knowledge of preconception care). Limited resources (e.g., lack of time, tools, guidelines, and reimbursement) were frequently reported at the organizational and societal levels. Disparities are well documented in prenatal and pediatric care across similar domains.

Performance Measurement and Quality Improvement Efforts

Performance measurement is a common strategy currently being employed by state and local governments, payers, hospital accreditation bodies, and professional organizations to improve health care quality. It is often linked to change strategies—public reporting and pay-for-performance (Berwick et al., 2003; Chassin, 2002; Hibbard et al., 2005; IOM, 2006; Lindenauer et al., 2007; Millenson, 2004). Public reporting fosters interest in quality on the part of physicians and hospital leaders, perhaps by appealing to their professional ethos or creating market advantages (Marshall et al., 2000; Lindenauer et al., 2014). Pay-for-performance programs are intended to enhance the business case for quality improvement (QI) by rewarding excellence and reversing what have been described as perverse financial incentives that can deter providers and hospitals from investing in QI efforts (Dudley et al., 1998; Epstein et al., 2004; Millenson, 2004).

To date, the National Quality Forum has endorsed 18 perinatal and reproductive measures, none of which address ambulatory[6] preconception or prenatal care (National Quality Forum, 2016). The only prenatal and postpartum care performance measures currently in use are the two Healthcare Effectiveness Data and Information Set (HEDIS) measures on the rates of first trimester prenatal care and postpartum visits (National Committee for Quality Assurance, 2018).

There are more measures specific to the quality of child health care, which address some of the main aspects of well-child care (e.g., immunizations, regular visits, screening, developmental assessment). The reauthorization of the CHIP program in 2009 provided the first major support for quality measurement in public health insurance programs for children and youth and led to substantial growth in pediatric care measures and ongoing efforts to expand and refine these measures (AHRQ, 2018; Perrin, 2012). Measures have generally focused on children without chronic conditions, with the exception of the high-prevalence conditions (especially asthma), and quality-of-care measures for children with disability are much more limited (Perrin, 2012). Medicaid publishes and updates a quality measure set for children (Medicaid, n.d.). Several groups have catalogued children's quality measures (Beal et al., 2004), and the AAP and the National Initiative for Children's Health Quality, among others, have pursued consensus on best child health quality measures (Adirim, 2017).

The Collaborative Improvement and Innovation Network (CoIIN) is an example of a model aimed at improving maternal and child health that combines both CQI and collaborative learning to address infant mortality and perinatal disparities. The CoIIN was launched in 2012 in 13 southern U.S. states (Public Health Regions IV and VI), with early success leading to a national expansion of the program to other states. The CoIIN brought together state teams of clinical and public health leaders and policy makers to implement evidence-based and promising

[6] Ambulatory care is care provided by health care professionals in outpatient settings (AHRQ, n.d.-1).

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strategies, supported by virtual shared workspace, CQI experts, and a data dashboard that provided real-time data to drive real-time improvements (Ghandour et al., 2017). Between 2011 and 2014, early elective delivery at less than 39 weeks decreased by 22 percent versus 14 percent in other regions, smoking cessation during pregnancy increased by 7 percent versus 2 percent, and back sleep position increased by 5 percent versus 2 percent. Preterm birth decreased by 4 percent, twice that observed in other regions, but infant mortality reductions did not differ significantly (Hirai et al., 2018b). CoIIN is particularly notable in its application of CQI methodologies, which have largely been limited to clinical settings, to drive improvements in population-level perinatal outcomes.

While these efforts have improved adherence to standardized care processes, inequitable care remains pervasive throughout the system. Inequitable care means care that does not address the unique challenges and vulnerabilities made relevant by differential life experiences based on social characteristics, such as gender, ethnicity, language preference, geographic location, and SES (IOM, 2001). For example, a major component of preventive health care is culturally and linguistically sensitive and literacy-appropriate health education to promote behavioral changes. Thus, an office or clinic visit is only the prelude to the fully completing the care. To fully attain the effects of care, an individual or family needs to take that education to the home and follow through with it (e.g., eat more healthy foods). If the social environment in the community or home does not have the resources to support these behavioral changes (insufficient income or no neighborhood stores to purchase healthy foods), the care plan remains unfulfilled, and the effectiveness of the care remains limited, despite the practitioners' best intentions (Lu et al., 2010).

Multiple studies of the implementation of quality measures indicate substantial improvement in attention to specific processes in the delivery of health care services (e.g., increasing screening rates, testing for harmful side effects of treatment). Increasingly, payers have assessed quality of care (e.g., screening and immunization rates) and offer financial incentives to achieve quality thresholds. These incentives have improved processes linked to better health outcomes; further research is needed to determine their impact on health outcomes directly and on disparities. Often, these assessments are done at a population level and do not highlight disparities or account for practices caring for populations with different characteristics. Thus, in addition to traditional methods to assess performance and QI, there are a number of other areas that are receiving increased attention in efforts to attain equitable health care practices, including developing new metrics and measurement methods to account for child development and well-being in the context of an intersectional and multidimensional view of health and health equity and enhanced workforce education and training (including training to recognize and address implicit bias).

Development of New Metrics and Measurement Methods

Growing recognition of MBH concerns among children and youth has led to the development of newer measures of developmental and behavioral screening (including social and emotional screening), treatment, and outcomes, including measures of follow-up for medications used to treat some pediatric mental health conditions. Few measures routinely collected today address SDOH. Further, toxic stress and its precipitants, including many social determinants, such as adverse childhood experiences (ACEs), housing instability, food insecurity, or poverty and racism, are inadequately assessed in health care settings. Substantial evidence notes that much adult chronic disease has its origins in childhood and adolescence,

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often associated with the SDOH discussed throughout this report (Marmot et al., 2001). (See also the discussion of the Perry Preschool and Abcedarian Projects in Chapter 7.) The majority of adult mental health conditions originate in childhood or adolescence. An emerging body of research also suggests that some chronic childhood and adult diseases have a fetal origin (Barker, 1995, 2003, 2004; Calkins and Devaskar, 2011; de Boo and Harding, 2006; Kimm, 2004; Skogen and Overland, 2012) and that factors such as maternal and paternal stress, nutrition and physical activities, and occupational and environmental exposures may play a critical role in epigenetic modification and developmental programming of future health and disease. Information regarding these risks is not routinely collected, monitored, or reported during standard preconception or prenatal care visits, nor would this be feasible in a brief office visit. Until recently, pediatric care providers were not routinely screening for ACEs; however, the AAP released two policy statements and conducted substantial practitioner education, which bolstered rates of attention in community practice (Kerker et al., 2016; Szilagyi et al., 2016). Even having clear standards for screening does not assure that all child health practitioners screen for all relevant issues. A recent paper using parent reports found that, nationally, 37 percent of children received developmental screening with a validated tool (Hirai et al., 2018a). Another study, based on physician report, indicated that more than 80 percent of pediatricians reported using one or more formal tools to screen for autism, with 88 percent screening at 18 months and 74 percent screening at 24 months (Coury et al., 2017).

Several national initiatives have begun to advance primary care clinicians' understanding of ACEs and toxic stress and their capacities and competencies to identify and respond to risk factors for toxic stress and other adversity, such as the Trauma-Informed Primary Care Initiative, a partnership between the National Council for Behavioral Health and Kaiser Permanente;[7] the National Pediatric Practice Community on ACEs, an initiative of the Center for Youth Wellness;[8] the Trauma-Informed Care Implementation Resource Center,[9] developed by the Center for Health Care Strategies, and other initiatives (APA and AAP, 2017; APA Task Force on Childhood Poverty, 2013). The AAP has recommended regular screening for precipitants of toxic stress (Garner et al., 2012), and several professional organizations have compiled tools and resources to support clinicians in screening for social determinants (including risk factors for toxic stress), maternal depression, and early child development, such as the AAP Screening and Technical Assistance Resource (STAR) Center and the AAFP's Center for Diversity and Health Equity (AAP, 2010; Gleason et al., 2016; Sege and Amaya-Jackson, 2017).[10] Many other professional organizations are also issuing similar statements and recommendations; for example, the American Heart Association has called for upstream identification and mitigation of ACEs as a risk factor for cardiometabolic disease (NASEM, 2015; Suglia et al., 2018).

As the science advances regarding contributors to child health equity, new measures that capture SDOH, including indicators of cumulative adversity and family issues that may impact health and development, have become available. The development of these measures is well

[7] For more information, see https://www.thenationalcouncil.org/trauma-informed-primary-care-initiative-learning-community (accessed April 29, 2019).
[8] For more information, see https://nppcaces.org (accessed April 29, 2019).
[9] For more information, see https://www.chcs.org/resource/trauma-informed-care-implementation-resource-center (accessed July 17, 2019).
[10] For more information on the AAFP Center for Diversity and Health Equity, see https://www.aafp.org/patient-care/social-determinants-of-health/everyone-project/cdhe.html (accessed April 29, 2019). For information on AAFP's "EveryONE Project" to advance health equity in every community, see https://www.aafp.org/patient-care/social-determinants-of-health/everyone-project.html (accessed April 29, 2019).

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underway, addressing such issues as household hunger, lack of money for utilities, difficulty finding jobs, need for child care, or housing problems (AAP, n.d.-b; Arthur et al., 2018; Center for Youth Wellness, n.d.; Ellis, 2001; Garg et al., 2015; Gottlieb et al., 2016). The substantial increased attention to screening and identification of SDOH and adversity in early childhood has led to much consideration of how child and family health care services might prioritize screening centered on SDOH. However, many different screening tools exist for SDOH, with little consensus or guidelines on which ones are most appropriate or effective in different health care settings or contexts (Morone, 2017; Pai et al., 2016). Early studies show promising results with screening and intervention, and most of this work has been done in primary care practices, especially those that care for children and families (Angier et al., 2019b; Bazemore et al., 2016; Cottrell et al., 2018; DeVoe et al., 2016). There are a number of scientists actively engaged in building the evidence for SDOH screening and referral (e.g., which tools to use, how to screen, when to screen, how to incorporate technology). Providers will typically screen more when they can refer to programs that can address child and family needs. Where resources are limited, they are less likely to try to identify problems (Garg et al., 2016).

With the increased recognition of the need to expand risk assessment for ACEs, the development of evidence-based, effective strategies for early and continuing assessments of ACEs and other social and environmental determinants will be a key ingredient in transforming the delivery and measurement of preconception, prenatal, and pediatric care. In addition, there is an urgent need to develop biomedical measures to detect and treat toxic stress.

Interest has grown in measuring broader outcomes that may reflect more than health care, such as school readiness at age five (Jones et al., 2015) or a new composite measure of being healthy and ready to learn (Child Trends, 2018; Ghandour et al., 2018). These broader measures recognize the importance of cross-sector collaboration to keep children healthy and the health sector's role in helping children and their families access other needed resources (housing, food, and other supports) to help improve these outcomes.

As new metrics are developed, there is growing recognition that quality measurement is multidimensional and can be impacted by the clinical and social complexity of patients, families, and populations. For example, providers caring for patient populations in at-risk contexts are sometimes held accountable for achieving similar gains in quality metrics as providers in affluent communities, without adjustment for patient complexity, putting them at risk for financial loss. Misaligned financial incentives can perpetuate disparities in health care access for underserved populations and fail to recognize quality initiatives that are reducing disparities (or exacerbating them) for certain subgroups if all members of the population are not starting at the same baseline or improving at the same rate. Efforts are currently underway to create adequate adjustments for traditional measures (NCQA, n.d.), although developing adjustments for diverse child populations has been difficult (Kuhlthau et al., 2005).

Enhanced Workforce Education and Training

Workforce development and training is another important strategy for improving the quality of health care. The increased attention to identifying and addressing social, economic, and environmental factors adversely impacting the health of children and families has led to enhanced curricula in the training programs for child and family health care providers. These additional components strengthen the emphasis on learning about the social and community aspects of care for patients and families, implicit bias and unequal treatment, and how to more effectively collaborate with community organizations to improve care and outcomes. This

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education variably addresses adversity, biases, and disparities, with increasing attention to such issues as food and housing insecurity and the importance of screening for ACEs and recognizing microaggressions.

Advancing health equity requires sustained commitment and resources to increase the diversity and representativeness of our health care workforce. The next generation of health care providers also need to be better equipped with knowledge, skills, and tools to address MBH issues, ameliorate toxic stress, and provide TIC. This training requires explicit curricula on how to recognize and eliminate implicit bias and unequal treatment in health care. The future workforce needs to develop competencies in team-based care, working across disciplines and sectors to tackle the social and environmental determinants of health.

Transforming the health care system to provide culturally competent care is critical to advance health equity. It is crucial that the health care workforce receive education and training in cultural competence. Betancourt et al. (2003) define a culturally competent health care system as "one that acknowledges and incorporates—at all levels—the importance of culture, assessment of cross-cultural relations, vigilance toward the dynamics that result from cultural differences, expansion of cultural knowledge, and adaptation of services to meet culturally unique needs" (p. 294). The importance of culturally competent care is also relevant to efforts to increase patient- and family-centered care, as "cultural competence enhances the ability of health systems and providers to address individual patients' preferences and goals" (Saha et al., 2008). The wide diversity of household backgrounds—cultural, racial, ethnic, and linguistic—and the expected demographic changes that will result in successively more diverse future generations make it important that the health care system have capacity to understand and respond to cultural variations in health practices and understanding (see Box 5-3 on the role of doulas and Box 5-4 on the role of nurse-midwives and midwives in prenatal and postnatal care and culturally appropriate care). Although more providers entering the workforce have had cultural training and speak languages other than English than a generation ago, many households still face difficulties accessing culturally responsive and linguistically appropriate care. Access to such subspecialty care may be particularly challenging, especially diagnostic and long-term care services that address cultural and language needs. For example, young children with ASD often need specialized therapists who work intensively on communication with the child and between the child and family. A Spanish-speaking family, however, can find it difficult to identify a Spanish-speaking therapist (Dabney et al., 2015)

BOX 5-3
The Role of Doulas in Advancing Health Equity

A doula is "a trained professional who provides continuous physical, emotional, and informational support to a mother before, during, and shortly after childbirth to help her achieve the healthiest, most satisfying experience possible" (DONA International, n.d.). Doulas receive training on providing support during birth and/or the postpartum periods but do not perform medical or clinical tasks. For information on midwives, who receive medical and clinical training, see Box 5-4.

Evidence suggests that supportive care and services from doulas improves maternal and infant outcomes in the prenatal through postpartum periods. A joint consensus statement from the ACOG and the Society for Maternal-Fetal Medicine stated that "one of the most effective tools to improve labor and delivery outcomes is the continuous presence of support personnel, such as a doula" (Caughey et al., 2014). Similarly, a 2017 Cochrane systematic review of 26 studies found that women who received continuous support during childbirth, including from doulas, had improved maternal and infant outcomes (Bohren et al., 2017).

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Importantly, doulas of color can provide culturally appropriate support to women of color that is sensitive to the historical injustices experienced by people of color within the health care system. Doulas can help to bridge the divide between patients and their health care providers by assisting patients with health literacy and social support, thus improving both access to health care services and the quality of care (Kozhimannil et al., 2016). Providing such care is critically important to address the significant disparities in pregnancy outcomes in women of color, particularly African American women, in the United States.

Increasing access to culturally sensitive supports for women of color, such as doula care, is essential for decreasing health inequities. However, women of color are often least likely to have access to such care and services because doulas are rarely covered by insurance (Thich, 2016). An approach to increase access to doula care is a community-based doula program, which emerging evidence has found to be effective in improving health outcomes for women of color. Administered by HRSA and MCHB, the Community-Based Doula Program is a model that provides culturally appropriate peer-to-peer support based on the life course approach during the perinatal and early postpartum periods. The program serves women and families in communities with high levels of health and social needs (the majority of participants are black or Hispanic, and a small number are from tribal communities), and the program's community-based doulas "are of and from the communities being served" (HealthConnect One, 2014). Data findings of the program include rates of breastfeeding, including breastfeeding duration, that are higher than those documented by the CDC's Pregnancy Risk Assessment and Monitoring System (PRAMS) and breastfeeding goals of Healthy People 2020 as well as C-section rates that are lower than those documented by PRAMS.

Assuring equity calls for strategies to diminish cultural and linguistic barriers, through training and increased recruitment of health care providers from culturally diverse communities. It also requires explicit training on eliminating implicit bias and unequal treatment in health care. Unequal preconception, prenatal, and pediatric care based on race, ethnicity and SES has been well documented (Brett et al., 1994; Kogan et al., 1994; Kotelchuck et al., 1997). In a randomized trial of 524 providers who were shown videos depicting patients of varying sociodemographic characteristics, providers were more likely to recommend levonorgestrel intrauterine contraception for low-SES Latina and African American women than low SES white women (Dehlendorf et al., 2010). Patients from low-SES backgrounds were judged to be significantly more likely than patients from high-SES backgrounds to have a sexually transmitted infection (STI) and an unintended pregnancy and were also judged to be less knowledgeable. An analysis of data from the National Maternal and Infant Health Survey demonstrated that black women were less likely than white women to receive advice from their prenatal care providers about smoking cessation and alcohol use (Kogan et al., 1994). Stereotyping and implicit bias on the part of health care providers are factors that may play a role (ACOG, 2015). Advancing health equity in birth and child health outcomes will require addressing implicit bias and unequal treatment in health care (see Chapters 7 and 8 for more on implicit bias training).

BOX 5-4
The Role of Midwives in Advancing Health Equity

Midwives, defined here as certified nurse-midwives, certified midwives, and midwives whose education and licensure meet the International Confederation of Midwives Global Standards for Midwifery Education (AAP Committee on Fetus and Newborn and ACOG Committee on Obstetric Practice, 2017), are health care professionals who provide low-risk women with services that include primary care; gynecologic and family planning services; care during the preconception, pregnancy, childbirth, and postpartum periods; and care of the healthy newborn during the first 28 days of life (American College of Nurse-Midwives, 2012). Medicare and Medicaid reimburse for care provided by

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midwives in all U.S. states, and private insurance reimburses for their services in most states (American College of Nurse-Midwives, 2012).

The midwifery model of care tends to allow more time for patient interactions, with a holistic approach to health behavior education (Palmer et al., 2010; Vonderheid et al., 2007). In addition, African American women participating in CenteringPregnancy, a form of group prenatal care predominantly facilitated by midwives (see Box 5-5 for more information), have a significantly reduced preterm birth rate (Carter et al., 2016; Ickovics et al., 2007; Picklesimer et al., 2012).

Full integration of midwives into the health care system may provide an opportunity to both improve pregnancy outcomes and reduce health disparities. A recent study by Vedam et al. (2018) measured midwifery and health care integration with the Midwifery Integration Scoring System (MISS) and found an association between states with higher MISS scores and lower rates of infant mortality (including by race), cesarean section, preterm birth, and LBW infants than states with lower MISS scores.

Most existing training programs offer some, albeit limited, exposure to training in cultural diversity and how culture influences health care and behaviors. In a few settings, clinicians in training learn skills in team care, although integrated training programs are rare. Case Western Reserve University is starting a new program where medical students, nursing students, and others will train together in a new multischool training and research building. All of these areas (cultural and linguistic diversity, SDOH, MBH, and team experience) merit increased attention in medical training in general and in pediatrics specifically.

There is increased recognition that health care providers need to be not only well versed in medical interventions to treat illness but effective advocates in addressing the broader factors contributing to unequal access to services and treatment and impacting efforts to achieve health equity.

Improving Quality of Care

Based on its review of the evidence the committee concludes:

Conclusion 5-3: Strategies to improve the quality of preconception, prenatal, and child health care have included developing and implementing new measures, including for adversity and social determinants, along with efforts to strengthen the training of the health care workforce to better understand diversity and implicit bias and to address equity in health care.

To improve the quality of care provided in the preconception through early childhood periods, the committee recommends:

Recommendation 5-2: To expand accountability and improve the quality of preconception, prenatal, postpartum, and pediatric care:

- **Public and private payers should include new metrics of child and family health and well-being that assess quality using a holistic view of health and health equity. Federal, state, and other agencies, along with private foundations and philanthropies that invest in research, should support the development and implementation of new measures of accountability, including key drivers of health, such as social determinants, along with measuring variations by key subgroups to determine disparities;**

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- **Public and private payers, including the Health Resources and Services Administration's (HRSA) Bureau of Primary Care and Maternal and Child Health Bureau, Centers for Disease Control and Prevention, Centers for Medicare and Medicaid Services (CMS), and perinatal and pediatric quality collaboratives, should expand the use of continuous quality improvement, learning communities, payment for performance, and other strategies to enhance accountability; and**
- **Health-care-related workforce development entities should expand efforts to increase diversity, inclusion, and equity in the health care workforce, including diversity-intensive outreach, mentoring, networking, and leadership development for underrepresented faculty and trainees.**

Workforce development (bullet 3) will need to be addressed by several entities, including the Accreditation Council for Graduate Medical Education (ACGME) and specialty boards, professional schools, training programs, funders of graduate education in health professions (CMS, HRSA, and others), and teaching hospitals, including children's hospitals.
Metrics for accountability include:

- **Social determinants and risk measures:** Measures that reflect whether risks were identified early and whether families received needed help, with key drivers of health inequities that lie beyond traditional clinical purview but profoundly impact their health, such as housing instability, food security, and exposure to adversity or trauma.
- **Cross-sector developmental measures:** Measures that move beyond common indicators of child development, such as immunizations and management of acute infections or common chronic conditions, to address an expanded set of clinical indicators crucial for children and caretakers, including MBH. Measures should reflect healthy life course development, such as language at age three, school readiness at age five, reading proficiency at age eight, and high school graduation rates, as well as indicators of concern (need for special education, substance use, executive functioning, major behavior disorders).
- **Disparities as explicit measurement domains:** Measures that hold providers accountable not just for delivering services but also for improving outcomes and closing gaps in outcomes among key populations or subgroups. Adjusting quality measures for the social and clinical complexity of patient populations.

INNOVATIVE DELIVERY MODELS AND FINANCING CARE

Most preconception, prenatal, postpartum, and pediatric care is delivered through face-to-face office visits. For many families, but especially low-income families, this often requires taking unpaid leave from work, arranging child care and transportation, and waiting an hour or two for a 5- to 30-minute visit (Lewis et al., 2017). Preconception care is usually one short visit, with no consistency in what is accomplished during that visit, and a large percentage of families do not even receive this minimal level of care (Poels et al., 2016). Current prenatal care guidelines continue to recommend 14 or more prenatal visits during pregnancy, despite a lack of evidence supporting this number. Most visits consist of a spot blood pressure check and urine dip, a cursory auscultation of fetal heart tone and fundal height measurement, and a hurried conversation with a provider who, despite best intentions, often does not have sufficient time or

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training to educate patients about self-care let alone address their psychosocial concerns or occupational and environmental exposures (Lu, 2019). Similarly, postpartum care is usually one short visit (two after cesarean delivery), which consists of a cursory review of problems, a pelvic exam, and an often hurried discussion of contraceptive options, if the latter happens at all. Pediatric care also focuses on assessment of common issues, along with immunizations and other preventive services, in short office visits with limited time for extensive screening and counseling. Nonetheless, the multiple scheduled health supervision visits during the prenatal and early childhood periods can serve to connect families with trusted health advisors. For young children who have frequent pediatric care visits in the first years of life, health care provides a main entry point to health and many other services that can support the promotion of health for preschool children and their families (Garg et al., 2015; Patient-Centered Primary Care Collaborative, n.d.; Stille et al., 2010). In most communities, the current configuration of services does not take full advantage of this opportunity for maximizing collaborations across sectors and implementing concurrent interventions to strengthen children and families.

Concerns regarding increasing health care costs, health care provider availability, dissatisfaction with wait times, and the minimal opportunity for education and support associated with the individual care model have given rise to interest in alternative models of care. For children, a system of care designed to prevent and treat common infectious diseases faces a population whose health reflects infections less than noninfectious chronic conditions. Growing epidemics of nonfatal chronic conditions, especially MBH conditions (Halfon et al., 2012; Houtrow et al., 2014), much greater understanding of the SDOH, increasing diversity in the sociocultural makeup of the U.S. population, and the tremendous growth in the science of early childhood development, all call for new strategies and structures for delivering and financing preconception, prenatal, and pediatric care, including efforts to address inequities (Perrin and Dewitt, 2011). Social, economic, and environmental determinants of health generally affect the health of children and families more than usual medical treatments, and traditional ways of delivering health care have relatively limited impact on early childhood growth, development, and health. Some innovative ideas being tested in prenatal and well-baby care are group visits, multidisciplinary teams, strategies to link health care with other community resources, and increased use of new technologies (see section on the use of technology below).

Group Visits

Group visits and innovative designs that bring patients with similar needs together for health care encounters (face-to-face or virtually), increase the time available for the educational component of the encounter, improve efficiency, and reduce repetition (ACOG, 2018a). The group visit model is being tested for prenatal care because group visits are designed to enhance patient education while providing opportunities for social support and retaining the risk screening and physical assessment of individual care (ACOG, 2018a). In some settings, the groups continue meeting for postpartum and/or newborn care sessions. This model is especially promising in the reconceptualization of postpartum care, which would move away from a single clinical encounter toward more comprehensive, ongoing support for the postpartum transition.

While initial observational studies (Thielen, 2012) and a large RCT (Ickovics et al., 2007) found significant improvement in perinatal outcomes in group visit models, a more recent Cochrane review (Catling et al., 2015) and a meta-analysis of 10 observational studies and four RCTs (Carter et al., 2016) found no significant difference in preterm birth, LBW, breastfeeding, or neonatal intensive care (NICU) admissions. However, it should be noted that while there was

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no overall difference in outcomes between women in traditional and group prenatal care, black women in group prenatal care showed a 41 percent reduction in preterm births in the largest RCT (Ickovics et al., 2007) and the meta-analysis, respectively. CenteringPregnancy is a model of group prenatal care that has demonstrated some promising but mixed findings in improving maternal and birth outcomes, including decreasing disparities by race and ethnicity (see Box 5-5). CenteringParenting is a similar innovative model of group postpartum care that brings together a cohort of 6–7 mother-infant dyads for one year postpartum (Bloomfield and Rising, 2013); its impact on outcomes remains to be established.

Box 5-5
CenteringPregnancy: Promising Model[a]

CenteringPregnancy provides group prenatal care to women in a supportive, educational, and interactive environment. The model was developed and is implemented by the Centering Healthcare Institute, a Boston-based nonprofit started in the 1990s that works with health care providers to implement group care models (its three models are CenteringPregnancy, CenteringParenting, and CenteringDiabetes) in more than 585 locations, including large health systems, across the country.

The CenteringPregnancy group prenatal care model enables participants to spend more time with their provider and to learn, discuss with, and receive support from other pregnant women in a group environment (Centering Healthcare Institute, n.d.-2). In addition to each participant having individual time with their provider for physical health assessments, groups of 8–10 women due at about the same time participate in a curriculum of 10 90-minute or 2-hour sessions (in line with the recommended schedule of 10 prenatal visits) that includes provider- and staff-facilitated information sharing and discussion about health and nutrition, childbirth preparation, stress reduction, labor and delivery, breastfeeding, relationships, and parenting (Centering Healthcare Institute, n.d.-2). Participants are active in their own care by taking their own weight and blood pressure and recording their own health data, and they are women of varying age, race and ethnicity, and socioeconomic background (Centering Healthcare Institute, n.d.-2). The program aims for participants to form meaningful and supportive relationships with other participants, many of whom continue their participation with CenteringParenting's model of family-centered well-child care (Centering Healthcare Institute, n.d.-1).

Research shows that CenteringPregnancy has led to improved birth outcomes, including significantly lower risk of preterm birth, LBW or very LBW infant, infant small for gestational size, and fetal death (Chen et al., 2017; Cunningham et al., 2019; Gareau et al., 2016; Ickovics et al., 2016; Tanner-Smith et al., 2014). Ickovics et al. (2007) also found that participants were less likely to have poorer-quality prenatal care, were more knowledgeable and better prepared for childbirth, had greater satisfaction with their care, and were more likely to breastfeed than those receiving individual care (Ickovics et al., 2007). In addition, Gareau et al. (2016) estimate that South Carolina saved nearly $2.3 million with a $1.7 million investment in implementing the model, which achieved an average savings of $22,667 for every premature birth prevented, $29,627 for decreasing the rate of LBW, and $27,249 for decreasing the risk of a NICU stay.

The model has also been shown to decrease disparities in birth outcomes by race and ethnicity. Picklesimer et al. (2012) found decreased disparities in the risk of preterm birth for black women relative to white and Hispanic women who participated in the program (Picklesimer et al., 2012). In addition, Ikovics et al. (2007) found that black women who participated had the largest decreases in preterm birth (Ickovics et al., 2007).

Some studies and reviews have found no or mixed evidence of changes in the risk of preterm birth, LBW, prenatal care costs, and delivery between group and prenatal care (Carter et al., 2016; Catling et al., 2015; Ickovics et al., 2007; Tanner-Smith et al., 2014). However, the authors of two reviews note that further research is needed due to the small number of studies and participants and the lack of high-quality studies (Carter et al., 2016; Catling et al., 2015).

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

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Father and Partner Involvement

As discussed in Chapter 4, fathers can play an important role in their children's development, and the health system could do better in engaging fathers' involvement. Several authors have suggested innovative approaches to strengthen engagement through preconception (Kotelchuck and Lu, 2017), prenatal (CPIPO, 2010), postpartum, and pediatric care (Yogman and Garfield, 2016), though evidence of effectiveness is still lacking. Kotelchuck and Lu (2017) proposed a research agenda for advancing the father's role in preconception health, focusing on three priority domains: increasing the basic epidemiology and risk factor knowledge base; implementing and evaluating men's preconception health/fatherhood interventions (addressing clinical health care, psychological resiliency/maturation, and SDOH); and fostering more fatherhood health policy and advocacy research. The Commission on Paternal Involvement in Pregnancy Outcomes (CPIPO) proposed 40 research, practice, and policy recommendations for strengthening fathers' engagement, including the development of "father-friendly" hospital settings, practices, and policies (CPIPO, 2010). The Commission also called out the importance of developing more effective methods of recruitment and retention of men in communities with high levels of poor pregnancy outcomes in research. Similarly, Yogman and Garfield (2016) pointed out the important role of child health providers in supporting and encouraging father involvement, with special attention to fathers' involvement across childhood ages and the influence of fathers' physical and mental health on their children. Given the growing diversity of families, similar attention is needed to engaging partners of all types across the health system.

Multidisciplinary Teams

Another growing innovation is the sharp increase in multidisciplinary teams delivering care to children and families, including team members from sectors other than health care. Teams in primary care and obstetrical care settings take diverse forms, but they all reflect a dedicated move from care provided by physicians alone to much greater involvement of nonphysician providers in ongoing care (Halfon et al., 2014). Team care has grown from early models of physician–nurse practitioner collaboration dating back half a century. Many subspecialty programs, for adults and children, have long had teams providing care, with substantial documentation of their effectiveness (Katkin et al., 2017; Lahiri et al., 2016). Typical elements of a primary care team address (1) chronic care management (especially for the high-prevalence conditions other than mental health)—often with a nurse or nurse practitioner having main responsibilities, (2) mental and behavioral health—often through a colocated mental health professional (see below), (3) linking families with community resources (e.g., through a staff member knowledgeable about community benefits and resources or through medical-legal partnerships [MLPs]), and (4) helping families assess readiness for becoming parents and building their child-raising skills (e.g., programs to plan for parenthood, parenting programs in health care offices, connecting with home visiting programs, or encouraging families to read to children at an early age). Over the past few decades, nurses have played increasingly broad roles, including prevention and care management. Few teams include all these components, but the growth of team care has addressed all of them in different models.

Models that incorporate evidence supporting teams in health care settings to develop multidisciplinary care coordination programs involving families, social workers, paraprofessionals or peer workers, and community partners have evolved over time to help families of high-risk children be more proactive at managing health risks (Van Cleave et al.,

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2015). Programs such as the Parent-Focused Redesign for Encounters, Newborns to Toddlers (PARENT) have used health educator coaches for parents to deliver well-child care that incorporates social risk screening and referral, with developmental and behavioral assessments (Coker et al., 2016). Efforts to use team-based care (coupled with telehealth mechanisms) in part reflect the recognition that such changes will help to address more effectively the many factors contributing to the health of patients and communities. These changes recognize that enhancing health care practices with personnel knowledgeable and skillful in helping families access a breadth of community services will address disparities.

Integrating mental health and behavioral health services into primary care is a widespread innovation that usually involves having a mental health professional (e.g., a master's level psychologist, social worker, or psychiatric nurse specialist) colocated in the practice (Ader et al., 2015; Stancin and Perrin, 2014; Team Up for Children, n.d.). One such program embeds mental health workers in several community health center pediatric practices so that they can see patients jointly or transfer them easily and immediately.[11] These personnel both see patients directly and help to train the primary care practitioners to hone their own mental health skills. In another model that has experienced much growth (now in over 30 states), primary care clinicians can access telephone backup services that support their in-office care of mental health problems (Sarvet et al., 2010; Straus and Sarvet, 2014). Occasionally, backup providers will see patients directly for 1–2 visits, although the majority of services provided are either directly to the primary care provider or by referral to community resources for ongoing mental health care (e.g., community CBT providers).[12] Most studies of programs that integrate behavioral health care services into primary care to address the increased prevalence of mental health diagnoses, early childhood developmental conditions, and substance use disorders in families have shown substantial promise (AHRQ, n.d.-2; Balasubramanian et al., 2017; Kwan and Nease, 2013).

Embedded programs that directly address SDOH focus on having onsite professional social workers or other staff who provide in-person services or navigation to families (Fierman et al., 2016) or who help families find needed community resources. Several established programs do this kind of work, including the following (see also Box 5-6 for information on another initiative, Pediatrics Supporting Parents):

- The Health Leads program uses patient advocates to meet with families, guide them to community resources, and integrate SDOH care into the routines of clinical care (Garg et al., 2012) and has been shown to improve child health outcomes in a randomized trial (Gottlieb et al., 2016).
- Reach out and Read (ROR) is a practice embedded program in pediatric care settings encouraging parent–child interaction and literacy development and has been shown to result in higher language proficiency in at-risk children (Mendelsohn et al., 2001).
- The PARENT program uses coaches for parents to expand the capacity of providers to address family social risks. An RCT of the program showed improvements in use of developmental screening and other preventive care and reduced ED visits early in life (Coker et al., 2016).
- HealthySteps [with sites in more than 20 states, DC, and Puerto Rico (Zero to Three, n.d.-1)] combines practice-based services, using early child educators or nurses with early childhood training, with community linkages focused on newborn care, safety, and developmental issues. This program has shown some evidence for impacts on

[11] For more information, see https://www.teamupforchildren.org (accessed May 9, 2019).
[12] CURES Act and expansion of phone back-up programs

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parent–child communication (Minkovitz et al., 2007) (see Box 5-8 for more on HealthySteps).

- MLPs assist families with the legal challenges that often go hand-in-hand with unmet needs related to social determinants and have been shown to improve subjective well-being and positive impact health care use (Klein et al., 2013; Sandel et al., 2010) (see Chapter 6 for more information on MLPs).
- Help Me Grow is a coalition of 28 states, communities, and individuals invested in ambitious and resourceful early childhood systems that optimally serve all families and children. It is designed to help states and communities leverage existing resources to identify children in at-risk environments, link families to community-based services, and help families support healthy development of their children, including through child health provider outreach (Help Me Grow, 2017).
- Filming Interactions to Nurture Development (FIND) is a video coaching program that aims to strengthen positive interactions between caregivers and children to reinforce developmentally supportive interactions, or what's known as "serve and return." Early evaluation studies show participation in FIND Fathers project was associated with improvements in parenting stress, father involvement, and child behavior problems; other evaluations are ongoing (Center on the Developing Child at Harvard University, n.d.).

BOX 5-6
Identifying Strategies to Support Parents in Clinical Settings

Pediatrics Supporting Parents is a multiphase initiative supported by a consortium of foundations that is exploring opportunities in the context of pediatric well-child visits to promote children's healthy social and emotional development.

Phase 1: The Center for the Study of Social Policy (CSSP) has identified evidence-informed, scalable strategies to improve social and emotional health, the parent–child bond, and parental mental health (an important mediator of social-emotional development and the parent–child bond) during well-child visits. This includes a study of how strategies may differ with respect to pediatric practice or community setting.

Phase 2: The National Institute for Children's Health Quality (NICHQ) has assessed the findings from CSSP and uses them to inform a learning community of pediatric primary care practices who will pilot the strategies and make recommendations for scaling them up at a national level.

Based on the findings from the report *Promoting Young Children's (Ages 0–3)*, *Socioemotional Development in Primary Care* (NICHQ, 2016), this initiative has adopted the framework of four "design elements" of strategies to engage parents: (1) assessment, (2) education, (3) modeling, and (4) connection (NICHQ, 2016). The initiative's theory of change identifies these design elements as contributors to the primary outcomes of interest (i.e., social-emotional health, parent–child bond, parental mental health).

CSSP employed a multistep approach and applied a robust set of criteria (see Appendix C of CSSP, 2018 for more information) to identify 13 evidence-informed and promising programs to explore further through site visits. The organizing framework for these programs identified 10 important areas of strategy:

1. Anticipatory guidance
2. Screening, connection, and access
3. Health-related resources
4. Curriculum based courses for parents/caregivers

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5. Observations
6. Group well-child visits
7. Mental health consultation
8. Physician extenders
9. Home visiting
10. Trainings/CQI

SOURCE: CSSP, 2018.

The growth of teams also supports expanded attention to social, economic and environmental determinants of health among children and families. Team-based care helps practices encourage families to share information more effectively, as different team members focus on different aspects of a family's health and health determinants. Strong community knowledge and linkages help these efforts succeed, and some practices embed community health workers and other laypeople with lived experience, community expertise, and inherent trust among community members. Here too, the value of team-based community-oriented care has many advantages over traditional physician-centered practice. Community health workers know community resources (e.g., housing, food, employment) and can assist families in getting the help they need (e.g., resources to find improved housing for a child with asthma who wheezes because of mold in her apartment instead of repeated emergency department [ED] visits for nebulizer treatments).

Enhanced Services to Identify and Address Social, Economic, and Environmental Determinants of Health

There has been increased recognition of the impact of adverse social, economic, and environmental determinants on health outcomes over the past several decades. Federal and state public health efforts have moved to enhance care to better identify and address these factors (Lu et al., 2010). For prenatal care, enhanced care models have been designed to deliver coordinated, augmented, enabling, enriched, comprehensive, or "wraparound" prenatal care services—particularly for low-income populations. Enhanced prenatal care typically refers to routine prenatal care visits combined with ancillary services that may entail outreach efforts, counseling about the Special Supplemental Nutrition Program for WIC, case management, social work, psychosocial counseling, social support, health promotion/education, transportation, home visiting, and follow-up services to facilitate the ongoing use of the prenatal services offered (Alexander and Kotelchuck, 2001). The Comprehensive Perinatal Service Program (CPSP) enhances prenatal care with nutrition counseling, social services, and health education (Korenbrot et al., 1995). Most federally funded Healthy Start programs enhance prenatal care with care coordination, case management, and home visiting (Badura et al., 2008). In his systematic review of three types of enhanced prenatal care—home visiting programs, comprehensive-care programs, and preterm-prevention programs—Fiscella (1995) failed to find conclusive evidence of effectiveness of enhanced prenatal care for preventing adverse birth outcomes. It should be noted, however, that Fiscella examined the impact of enhanced prenatal care on only three immediate birth outcomes—perinatal death, LBW, and preterm birth; the impact of enhanced prenatal care on other short- and long-term health outcomes for children and families remains largely unexplored. A study of the Illinois Family case Management (FCM)

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program, another enhanced prenatal care program, did find that participation resulted in a lower LBW rate (Silva et al., 2006).

Health care providers have explored three basic approaches to the challenge of meeting social needs outside of their practices: (1) home visiting programs connected to the practice, (2) screening for risks and referring to community programs, and (3) community-level interventions.

1) Home visiting programs connected to the practice.

Home visiting has a long history of effective programs for young families (based on early experiments in Ithaca and Hawaii); nurses or other trained personnel make home visits for young families, in some cases during pregnancy and in all cases in the first few years of a child's life. For a more detailed discussion of home visiting, see Chapter 4. Some programs are closely integrated with health care providers; others work independently but share information. The Nurse-Family Partnership (NFP) program has shown evidence of reducing child abuse and neglect (Macmillan et al., 2009) with home visiting, for example. Some of the targeted programs that focus on specific needs, such as child abuse, child neglect, or LBW babies, have improved health outcomes in high-risk families (Avellar and Supplee, 2013; Radcliffe et al., 2013; Rushton et al., 2015). In the past few years, Congress has supported much growth in home visiting programs, allocating new funds to allow expanding the programs and the households covered. For an example of a community-based nurse home visiting program that has demonstrated promising findings in improving the health and well-being of children and their families, see Box 5-7.

BOX 5-7
Family Connects Durham: Promising Model[a]

Family Connects Durham (formerly Durham Connects) is a nurse home visiting program offered at no cost to families of newborns in Durham County, North Carolina. The program serves all families with newborns regardless of income or SES, and its mission is to "increase child well-being by bridging the gap between parent needs and community resources" (Center for Child and Family Health, n.d.). First implemented in 2008, the program was developed by the Duke Endowment and the founding director of the Duke Center for Child and Family Policy, Kenneth Dodge, in conjunction with community partners, with the goal to prevent child maltreatment and support all children and families in Durham, regardless of SES, with a model that could be replicated in other communities. The program maintains a commitment to community engagement through its Community Advisory Board, which includes representation from local agencies and is a collaborative effort of the Center for Child & Family Health, Duke Center for Child and Family Policy, Durham County Department of Health, and Durham County Department of Social Services. In addition to local grants, the program's funders include the Duke Endowment, Durham County Government, and United Way of the Greater Triangle.

A registered nurse with clinical and/community health experience visits, typically about 3 weeks (but up to 12 weeks) postpartum, to provide a weight and health check for the newborn and ensure that the mother is recovering from childbirth. Nurses may also provide information and community-based resources on topics such as breastfeeding, child care, postpartum depression, and social isolation. Families may be contacted 1 month after the home visit to ensure that the community resources and supports discussed during the visit were obtained. In addition, pediatricians, obstetricians, and family practitioners within the community partner with the program to share information and improve patient care. The program has Spanish-speaking nurses and matches families with nurses who speak their preferred language whenever possible. Nurses neither request nor report families' U.S. residency status.

In two RCTs, Dodge et al. found that by the time the infant was 6 or 12 months old, families who received a nurse home visit through the program had greater community connections, better use of higher-quality child care, higher-quality parenting behaviors, enhanced home environments,

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improved mother mental health, and reduced emergency medical care for infants (Dodge et al., 2013, 2014). Further, it is estimated that cities of similar size to Durham that average about 3,187 births per year could save about $6.7 million in community health care costs during the first 2 years of an infant's life if $2.2 million was invested each year in nurse home visiting (i.e., it is estimated that the program saves $3.02 in emergency health care costs for every $1 it spends) (Center for Child and Family Health, n.d.).

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

2) Screening in the practice and referral to a community partner.

WE CARE, a program based in pediatric primary care and serving low-income families, combines a screening tool and referrals to community resources for at-risk families who want assistance with social needs; the results from RCTs showed that families in the program were more likely to connect to social determinants resources, had fewer unaddressed needs, were more likely to be employed, and were less likely to live in a shelter at follow-up compared to usual (Garg et al., 2007, 2015). Other pediatric-based "screen and refer" programs, relying on either trained family specialists or volunteer community navigators, have shown similarly promising impacts on outcomes such as connection to social needs, increased immunization rates, and reduced early life ED use in randomized studies (Gottlieb et al., 2016; Sege et al., 2015). A range of more focused pediatric-based programs addressing specific social needs through screening and intervention have also shown promising results in high-quality studies, including programs focused on improving habitability for children with asthma (Krieger et al., 2005), the Safe Environment for Every Kid (SEEK) program focused on reducing intrafamily stress/violence and improving food security (Dubowitz et al., 2011, 2012; Feigelman et al., 2011), clinic-based referrals to Head Start (Silverstein et al., 2004), and StreetCred, which helps families get benefits they are eligible for (e.g., nutrition programs, EITC, SSI) (Marcil et al., 2018). Programs have also successfully deployed community health workers to do home assessments and education and reduced asthma triggers among children (Campbell et al., 2015; Williams et al., 2006). Other community collaboration models compile resource directories and connect people to publicly available benefits, including Temporary Assistance to Needy Families (TANF) or the Supplemental Nutrition Assistance Program (SNAP), as well as to community resources or private programs that can assist at-risk families (Henize et al., 2015).

Primary care providers have worked in this space for some time, with a growing body of evidence around effective programs and interventions. Several State Medicaid agencies have also begun to test promising models to incentivize providers in their efforts to address SDOH (including early detection of adversity and trauma experienced by children and their caregivers), greater integration with other community providers, and MBH integration (Van Buren, 2018). These innovative efforts together promise ways to strengthen preconception, prenatal, and pediatric care, help it move to team care and improve use of new technologies, and strengthen integration with other community services to enhance child health and well-being. North Carolina has developed an ambitious Early Childhood Action Plan, which has as goals healthy children who are safe and nurtured, learning, and ready to succeed (NCDHHS, n.d.). The plan builds on the science of early childhood and brain development and aims to address health equity. New York's First Thousand Days program includes statewide early home visiting,

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expansion of the CenteringPregnancy program (see Box 5-5), a requirement that managed care plans to have a child-specific quality agenda, and data system development to enhance cross-sector collaboration (United Hospital Fund, 2018).

3) Community-level interventions.

Lastly, addressing SDOH needs to encompass improved, collaborative systems for addressing medical and psychosocial risk factors at not only the individual child/family level but also the community level. Community-based parent support programs can provide resources through parent and child play groups, parenting information and support classes, and connecting families to medical or child care services (Trivette and Dunst, 2014). (For more information on supports for parents and caregivers, see Chapter 4.) The goal of these programs is to improve the health, well-being, and development of children by improving parents' caregiving skills and providing parents with adequate social supports and services (Goodson, 2014). Such programs are most effective when they are "family-centered as opposed to professionally-centered" and "capacity-building as opposed to dependency forming" (Trivette and Dunst, 2014). In pediatrics, family-centered care is care that is "based on the understanding that the family is the child's primary source of strength and support and that the child's and family's perspectives and information are important in clinical decision making" (AAP Committee on Hospital Care, 2003) (p. 691). Family-centered care can lead to improved child health and behavioral outcomes (Dunst and Trivette, 2009; Dunst et al., 2007; Kuo et al., 2012), and it is vital that community-based programs connect families to medical services where family-centered care is standard.

The 2016 report *Parenting Matters: Supporting Parents of Children Ages 0–8* describes specific elements of effective programs, which include (1) parents as partners, (2) tailoring interventions to parent and child needs, (3) service integration and interagency collaborative care, (4) peer support, (5) trauma-informed services, (6) cultural relevance, and (7) inclusion of fathers (NASEM, 2016). HealthySteps is a community-based pediatric primary care model that prioritizes the role of parents and caregivers as active participants in the care of their children (see Box 5-8 for more information on HealthySteps). Several initiatives have effectively coordinated health, social services, family support, and educational services, such as the Harlem Children's Zone (Harlem Children's Zone, n.d.) and the multisite Best Babies Zone initiative (Best Babies Zone, n.d.). A recent National Academies report, *Communities in Action: Pathways to Health Equity*, documents several such place-based, community-level initiatives (2017).

BOX 5-8
HealthySteps: Promising Model[a]

HealthySteps is a team-based, family-centered pediatric primary care model that aims to improve the health, well-being, and school readiness of infants and toddlers in low-income families. The cornerstone of the model is a child development professional, known as a HealthySteps Specialist (HSS), who connects with families during well-child visits as part of the primary care team. The HSS supports families by coordinating care and screenings, offering guidance and referrals to local agencies and programs (e.g., MBH services for maternal depression, food banks and legal clinics for food and housing insecurity), and providing on-demand aid, including through electronic communication and home visiting, between primary care visits. Infants are enrolled in the program at their newborn visit (or as early as possible before their 6-month visit), and families may continue in the program until the 3-year-old well-child visit at most sites and until the child is 5 years old at some sites (MacLaughlin et al., 2017). The program is a national network of more than 140 pediatric and family practices sites in 20 states, DC, and Puerto Rico (Zero to Three, n.d.-1) that has served more than 37,000 children ages 0–3, including refugee children (Buchholz et al., 2016).

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The program was analyzed through a 15-site national evaluation in 2003 (Guyer et al., 2003), and several single-site evaluations have also taken place. Findings from the evaluations suggest that the model can help to achieve improved outcomes in child health and development, breastfeeding and early nutrition, connections to resources, child safety, parenting knowledge and practices, parent and physician satisfaction, maternal depression, and early literary and school readiness (Zero to Three, 2017). For example, Minkovitz et al. (2007) conducted interviews with mothers when their children were 5.5 years of age and found that families who participated in the program were more satisfied with their care, more likely to remain at their original practice, less likely to use severe discipline, more likely to report concerns with their children's behavior, and more likely to have their children read books. Such findings indicate that the program's positive effects, while modest, may continue after the intervention has ended. Further evaluation of the program's recent innovations and for specific outcomes is ongoing (Zero to Three, n.d.-2).

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

In addition to knowing the community to better direct patients to resources, health care institutions can treat surrounding neighborhoods as "patients" and intervene more directly in the SDOH. In one such case study, the Healthy Neighborhoods, Healthy Families Initiative, a pediatric center invested in a multifaceted housing intervention in the surrounding neighborhood and significantly improved vacancy rates, though the health impacts on children in the area are still being evaluated (Kelleher et al., 2018). Similar, a community health center in Wisconsin partnered with urban planners to integrate health into sustainable land-use planning practices in an effort to shape overall community health outcomes (McAvoy et al., 2004). Health systems have also begun to participate in larger cross-sector efforts and partnerships predicated on the principles of collective impact, such as Accountable Communities of Health (ACH), which bring together a wide range of partners from across sectors to collectively address the SDOH. These efforts are nascent, however, and high-quality evidence on the health impacts of the ACH model or similar initiatives is not yet available. The Centers for Medicare and Medicaid Services (CMS) recently released a request for proposals to address similar opportunities at the child health level, with a strong emphasis on MBH and building community coalitions to improve outcomes based on social determinants criteria (CMS, 2019).

Another promising example is the redesign of the federal Healthy Start program in 2015 to place greater emphasis on improving women's health before and between pregnancies and across the life course, strengthening families, including increasing father engagement, and addressing SDOH through the collective impact model (whereby Healthy Start grantees serve as the backbone organizations in facilitating coordination and collaboration with social services, housing, economic and community development, and other nonhealth sectors to prevent infant mortality in the community). Results from more rigorous evaluation of the Healthy Start program are expected to be available in 2019 (National Institute for Children's Health Quality, n.d.).

Embracing New Technologies

Technological advances may help to transform the model of brief, episodic visits in a busy practice, especially in underresourced settings, by improving communication and care in several ways. A health care system redesign that better leverages eHealth technologies and social networking in innovative ways can enable more effective health promotion than current short

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visits. With many technological opportunities emerging to implement such a redesign, an important consideration is that machine learning algorithms may suffer from the same biases reflected in the data on which they are built, such that their use in health care may inadvertently perpetuate and even exacerbate existing health disparities (Char et al., 2018; Gianfrancesco et al., 2018). Research is needed to identify strategies to minimize such biases as new technologies are implemented more widely in health care and other sectors (Turner Lee, 2018).

Increasingly, health care providers are experimenting with telehealth strategies to augment their services and make them more accessible and convenient for families (Burke et al., 2015). Most used and studied in the area of providing mental health services remotely, telehealth has expanded substantially in the management of many chronic diseases of children and adults. New technologies allow better home/community monitoring of chronic disease and assessing symptoms and clinical signs over phone and video. Increased use of technological innovations might also improve access before and during pregnancy (Lu et al., 2010). Telehealth has been proposed as a way to help overcome many of the access barriers described earlier; women could be connected to their providers or specialists at anytime from anywhere. In addition, mHealth could make health promotion more accessible using simple mobile phone functions. Instead of bringing children and families to care, future research, practice, and policy initiatives to increase access should work on leveraging technological innovations to bring care to people in their homes and communities.

Technological innovations, data sciences, and design thinking could be leveraged to redesign care around the needs of children and families and not just provider or clinic schedules. Technologies such as wearables, sensors, and lab-on-a-chip hold potential if they are proven to reliably and more continuously collect high-quality data that leads to improved care, better patient experiences, and more equitable health outcomes. Such data, collected from the comfort of a woman's own home throughout her pregnancy, may include not only information on blood pressure or urine protein but also nutrition and physical activities, stress and sleep, and occupational, environmental, and other exposures that affect pregnancy outcomes and developmental origins of health and disease. With remote home monitoring, it is possible to continuously transmit data to the cloud, which, with the aid of AI and machine learning, could be used to improve predictive analytics. For preconception and prenatal care, this enhanced data might help triage women to different levels and components of care (e.g., routine follow-up, a call from a health educator, a home visitor, or an urgent appointment with a specialist). Instead of adhering to a uniform schedule, this approach might enable the frequency and content of preconception and prenatal visits to be determined by the specific changing needs and risks of each woman. Much work in childhood chronic disease, especially ASD and inflammatory bowel disease, similarly uses remote data to inform the need for office visits, rather than relying on routine follow-up periods. Linking these data with genomic, proteomic, metabolomic, psychosocial, and environmental data might help create a more precise risk profile that could inform the design of more personalized and precise interventions.

New health care technologies can also enhance other communication between health care providers and their patients and markedly change the character and components of regular care, if these are high quality and well focused on characteristics most valued by households (Olson et al., 2018). Texting has been used to encourage healthy behaviors or advise on routine care (e.g., developmentally specific infant care advice, such as text4baby) (Evans et al., 2012). Texting has also improved low-income mothers' adherence to immunizations for their children (Hofstetter et al., 2015; Stockwell et al., 2012).

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In addition, new and emerging technologies could play an important role in decreasing health inequities. Digital tools that leverage artificial intelligence and machine learning have the capacity to better identify social risk factors and improve systems of referral and follow-up for patients, when used with care and appropriate data sources (Padarthy et al., 2019). Advanced technological systems can help collect social risk screening data without relying so heavily on the point of care encounter, such as using patient-accessible electronic health records to pre-collect screening data in advance of clinic encounters. Indeed, not all screening need take place in clinical offices or visits at all—electronic practice gateways allow families to respond to questionnaires before or after visits, and texting can help encourage their participation. Head Start and other early childhood sites can also screen, and data sharing across communities can expedite care and response. In addition to ambulatory settings, some health systems have also implemented screening within EDs and trauma centers, especially around issues of violence and trauma, but evidence on the effectiveness within those settings is still preliminary (Juillard et al., 2016; Smith et al., 2013).

Health information technology (health IT) can be used in a variety of other ways to augment health care services (e.g., promote patient education, assist with care coordination). Health IT in health care settings can also be used to support provider decision making and reduce errors. For example, reminders generated by electronic medical records (EMRs) can be used to encourage prenatal providers to prescribe progesterone to eligible patients with a documented history of spontaneous preterm delivery, to tell pediatric care providers about overdue immunizations, or to prompt follow-up on abnormal lab results, which can sometimes be missed in busy, understaffed clinics, especially in in underresourced communities. Health educational materials are accessible on the internet and through smartphone apps. For example, pregnant patients can find information about self-care from the internet, on websites such as USDA's ChooseMyPlate.gov,[13] which provide useful tools for nutritional self-assessment and education to pregnant women. Parents can access a variety of parenting resources and guidance regarding health and wellness for early childhood. Health IT can also help link clients to needed services, such as the Healthy City website,[14] which maps community services in Los Angeles County down to the ZIP code and Census tract level using GIS technology.

Financing to Support Innovation

Payment arrangements for most health care services, from both public and private (mainly employer-sponsored health insurance) sources, rely on fee-for-service mechanisms, where payment reflects the number of services provided (e.g., health supervision visits, acute care visits, vaccinations) and cover only specified services. Fee-for-service arrangements provide few incentives for many of the changes that are critical to preparing health care providers to better deliver services to meet the needs of children, youth, and families, especially attention to mental/behavioral health, addressing SDOH, building community links, and incorporating telehealth. Traditional payment focuses on services, rather than on improving the health of populations. While providers recognize the many factors influencing health in prenatal and early childhood, traditional payment strategies maximize the numbers of patients per hour, often resulting in less time spent with each patient, without providing support for the longer visits

[13] For more information, see https://www.choosemyplate.gov/nutritional-needs-during-pregnancy (accessed May 9, 2019.

[14]For more information, see http://www.healthycity.org (accessed May 9, 2019).

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needed by some households (e.g., those facing a social determinant). Many health providers are already experimenting with new organizational structures to address changing needs and better respond to the main influences on health outcomes and well-being. Practices increasingly face the need to manage chronic conditions, address MBH, respond to cultural and linguistic diversity, and help with poverty and other SDOH, including efforts to address inequities in care and outcomes. These views reflect growing attention to child and family health in a holistic way. Health care providers are embracing team-based care models and new technologies (Katkin et al., 2017). Yet, current financing models prevent many health care providers from practicing in multidisciplinary teams; integrating health services with other community services; placing more emphasis on population health strategies; using technologies to enhance communication, assess risk, and extend care; and tailoring services to address equity and disparities. Optimizing care and support for postpartum families will also require policy changes. Presently, many insurers bundle reimbursement for prenatal care, delivery, and a single postpartum visit into one global fee, creating a disincentive for providers to provide comprehensive postpartum care or see patients more than once. Many women lose their pregnancy-related Medicaid coverage at 60 days postpartum. Payers often do not recognize the care provided to parents in pediatric and family medicine care settings. Thus, changes in the scope of postpartum care would require changes to reimbursement policies that support postpartum care as an ongoing process, rather than an isolated visit, such as unbundling from global obstetrics payment, pay-for-performance, and extension of Medicaid coverage for at least 12 months postpartum.

Public and private payers in the past few years have shown interest in moving to alternative payment mechanisms, where providers increasingly take (financial) responsibility for a specified population. These arrangements can provide very different incentives for the organization and provision of health care services. For example, they allow more care to take place out of office through the expanded use of telemedicine and lowering use of high-cost services of limited value (Berwick et al., 2008; Dzau et al., 2017; Wong et al., 2018). The growth of Accountable Care Organizations (ACOs) follows this interest in changing incentives to improve care. Several children's hospitals have developed ACOs (Makni et al., 2015; Perrin et al., 2017), although most of the growth in ACOs has come from large multispecialty programs with a main emphasis on practice transformation and cost savings for older populations.

As noted previously, Medicaid plays a major role in insuring children, youth, and pregnant women, and, increasingly, young parents. Its success and persistence are critical to the health of these populations, and Medicaid program enhancements generally implementing the changes in organization and financing described in this chapter will improve health and health inequities. As also noted above, children's hospitals (including organized children's health programs in general hospitals) provide most of the subspecialty care for children and youth with more complex and less common health conditions. Insofar as many children have public health insurance—with even higher rates among children with chronic health conditions—children's hospitals rely substantially on public financing. This reliance, however, puts these institutions at financial risk, as Medicaid generally pays much less than Medicare or private payers do for the same service. Although rates vary greatly among the states, on average, Medicaid pays at about two thirds of the Medicare rates (Biener and Selden, 2017).

Substantial moves to capitated or population health payments will greatly enhance the needed changes in health care arrangements. Here, too, several State Medicaid programs have innovated in their programs for children and youth. New York has focused on value-based payments, including efforts to define value measures for children and develop incentives to

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reward value improvements (NY Department of Health, 2017), based in large part on a careful analysis of value-based payment strategies for children (Bailit Health, 2016). Colorado has developed a Healthy Families Checklist, setting standards for Medicaid, strengthening eligibility opportunities for Medicaid, expanding benefits to include care coordination, and changing payment policy to support delivery system design (Ascend at the Aspen Institute, 2018). Massachusetts has reframed its Medicaid program as an ACO, emphasizing MBH integration, SDOH, and long-term supports and services (MA Executive Office of HHS, 2017). The committee supports health care payment reform efforts that promote value-based care, tie payment to population health outcomes rather than service delivery, and incentivize strategies that better address prevention and health equity.

BOX 5-9:
How Population Payments Can Enhance Use of New Technology

A 6-year-old girl in first grade complains of an earache. New technology allows a trained layperson to take a picture of the girl's ear membrane and send it to the child's health care provider. Paying physicians for the number of visits they carry out provides an incentive to have the child (with a parent) come for an office visit to determine whether she has an ear infection. Physicians who no longer receive compensation for each visit but are instead reimbursed based on the needs of the patient population can examine the eardrum remotely and decide whether the child has an infection—potentially saving a trip to the office for both child and parent.

Organization and Integration of Health Care Services

Based on its review of the evidence, the committee concludes:

Conclusion 5-4: Recent efforts to transform health care to address social determinants, early adversity, and mental and behavioral health integration and to develop community-based health care teams have increasingly addressed the changing needs of young families and children. Programs that build on home visiting, referrals to community partners, and integrated community efforts have enhanced outcomes for children and families. New technologies have expanded care and access, improved understanding of the social determinants, and improved communication about health and chronic disease. New payment arrangements can accelerate the transformation of health services to programs to support families and population health.

To advance the integration an organization of health care services, the committee recommends:

Recommendation 5-3: The Department of Health and Human Services, state, tribal, and territorial government Medicaid agencies, health systems leaders, and state and federal policy makers should adopt policies and practices that improve the organization and integration of care systems, including promoting multidisciplinary team-based care models that focus on integrating preconception, prenatal, and postpartum care with a whole-family focus, development of new practice and payment models that incentivize health creation and improve service delivery, and structures that more tangibly connect health care delivery systems to other partners outside of the health care sector.

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Achieving this recommendation requires the following:

- **Spread multidisciplinary team-based care models in community settings.** Promote the adoption and spread of multigenerational, team-based care models that support patients with a mix of traditional clinical professionals, such as doctors, nurses, social workers, and pharmacists, with mental health professionals, as well as community health workers or peer support specialists. Team activities include chronic disease management, integrated MBH, family support in early childhood, including access to parent training, and referral/connection to needed community services (housing, food, etc.).
- **Develop more integrated models for preconception, prenatal, and postpartum care delivery modes.** Models and interventions should allow women to engage in a continuum of services on their preferred terms, including culturally and linguistically appropriate service models, multigenerational care, approaches that employ home or community-based service delivery for women who prefer those settings, or programs that use new technologies and work to intentionally incorporate a woman's existing social support networks into her prenatal and postnatal care plan.
- **Adopt and spread integrated, whole-family and family-centered care models.** The best models give providers the ability to address the health of individuals and families comprehensively, including clinical health, integrated with MBH and health-related social determinants. Expanded child and family health models include assessment of family strengths and needs and strategies to address them, moving beyond individual care.
- **Develop and use new technologies that improve care and improve accessibility.** Advances include remote monitoring, as well as technologies to enhance ongoing communication, such as texting, virtual visits, and data sharing.
- **Align payment reform with health creation rather than service delivery.** Payment should promote value-based care and tie payment to population health outcomes rather than service delivery. Payment should incentivize strategies that address health creation and health equity and include comprehensive, coordinated, community-engaged care.
- **Develop cross-sector collaboration at systems levels to address the intersection of drivers across the health continuum.** Programs should seek collective impact or similar cross-sector efforts, such as ACH and other place-based initiatives, that aim to align health care, public health, social services, housing, education, and other sectors around aligned goals and common strategies. Shared governance structures should promote collaboration, including investment in administrative infrastructure and backbone organizations to manage collaboratives, assuring the flow of information and funding across sectors, as well as other strategies for sharing efforts and savings.

THE FUTURE OF PRECONCEPTION THROUGH PEDIATRIC CARE

Vision: To advance health equity, reduce health disparities, and improve birth and child health outcomes, the committee calls for a health care system that assures access for all to high-quality health care across the life course. Transformation of preconception, prenatal, postpartum, and pediatric care will address early childhood sensitive and key life periods by including attention to the root causes of poor health (for example, access to safe housing, high-quality education, food security), early adversity, and equity. The system will respond to the needs of children and their families holistically and through team-based care and by connecting them with community

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resources and integrating services across the life course. Assuring appropriate preconception, prenatal, postpartum, and pediatric care will have long-lasting effects on the health and well-being of our nation's children.

Recommendation 5-4: Transform preconception, prenatal, postpartum, and pediatric care to address the root causes of poor health and well-being—the social, economic, environmental, and cultural determinants of health and early adversity—and to align with the work of other sectors addressing health equity.

The Department of Health and Human Services should convene an expert panel to reconceptualize the content and delivery of care, identify the specific changes needed, develop a blueprint for this transformation, and implement a plan to monitor and revise the blueprint over time.

Implementation of this recommendation will require:

- **An update of clinical care guidelines and standards by the Women's Preventive Services Initiative, Bright Futures, American College of Obstetricians and Gynecologists, the American Academy of Pediatrics, the American Academy of Family Physicians, and others actively developing clinical care guidelines and standards to include this new content of care;**
- **Medical accreditation bodies, relevant programs, and agencies to develop performance monitoring and quality improvement based on this new content of care;**
- **Clinical care educational authorities, such as the Accreditation Council for Graduate Medical Education, to develop curricula, training, experiences, and competencies based on the updated guidelines; and**
- **Public and private payers to cover services reflecting this new content of care.**

This work should take place within a larger framework of social and reproductive justice and include diverse voices, especially from communities most affected by adverse birth and child health outcomes. To expand the content of preconception to pediatric care to address key drivers of health inequities better, specific actions include:

- **Recognize the impact of both adverse and enriching experiences across the life course and cumulative effects on health and well-being:** Address transitions between care providers and move from disjointed episodic care to an integrated continuum of longitudinal health care designed to optimize health production across the life course.
- **Include trauma assessment and response as an integral part of care:** Expand practice capabilities to screen for and respond to trauma and early life adversities as part of the standard of care for all families. Advance the biomedical detection and treatment of toxic stress in clinical practice, including the development of methods for early detection and implementation of evidence-based interventions such as connections to community resources designed to help address the effects of trauma. (See Recommendation 8-2 in Chapter 8 for more on screening and rapid-assessment).

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- **Change the content of clinical training to include social determinants of health, MBH integration, and early adversity:** Expand training, care protocols, and workflows to address the SDOH as a routine part of clinical best practices, especially in early life. To accomplish this, curricula and related training experiences need to be expanded to include competency-based training on screening and mitigation of early adversity, providing TIC, addressing the SDOH, and reducing implicit bias and unequal treatment in health care. Progress toward objectives and training outcomes should be benchmarked. (See Recommendation 8-3 in Chapter 8 on TIC.)
- **Implement an equitable whole-child, whole-family, multigenerational approach:** Expand clinical best practice to address the child and parents in an integrated, whole-family view of health that includes children, parents, and other caregivers. Train clinicians in ways that enhance the equitable delivery or care, including culturally competent caregiving and family-centered care that includes families and caregivers as partners in their own and their children's care. Clinicians and clinical staff should have ongoing training and accountability in areas of implicit bias and equity in evaluation and treatment.

CONCLUSION

Applying the science of early development to transform the preconception, prenatal, postpartum, and pediatric care has the potential to advance health equity. To better meet the needs of the populations receiving this care, the access, quality, and content of clinical care need to be addressed. This will require the health care system to be an active partner with other sectors and communities who are leading the way to address the root causes of health inequities—the social, economic, environmental, and cultural determinants of health.

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6

Creating Healthy Living Conditions for Early Development

MEETING FUNDAMENTAL NEEDS TO SUPPORT PRENATAL AND EARLY CHILDHOOD DEVELOPMENT

"I am trying to get gainful employment.... We all have that same thing in common of trying to do better for ourselves, trying to turn things around and do the right thing. But it's hard, it's hard because the resources that are available to us—we don't know about them, we are not aware of them, we don't know how to connect to the resources that are available to us." Parent on caregiver panel[1]

As described in earlier chapters, a child's most proximal influence in early development is the family unit, and specifically, the primary caregiver. Chapter 4 provides an overview of what children need from caregiver relationships and the critical role those relationships play for children to have the opportunity to flourish and thrive. However, these relationships do not exist in a vacuum, and neither do families—they are shaped by the social determinants of health (SDOH), as laid out in Chapter 3 (see also Figure 1-9). Families exist within the context of their communities, and all children need safe and healthy communities that promote optimal development. Healthy communities continuously create and improve physical and social environments and expand community resources that enable people to mutually support each other in daily life and in developing to their maximum potential (CDC, 2009). Chapter 4 also makes the case that it is essential to mitigate caregiver stress so that caregivers have the capacity and supports to care for their children and to serve as buffers against adversities. (See Chapter 2 for a brief discussion of the biological mechanisms of buffering and Chapter 3 for discussion of the importance of stable and nurturing relationships.) (See Box 6-1 for an overview of this chapter.)

[1] This quote is from a public meeting of the committee, held on October 1, 2018. The meeting webcast is available at www.national/academies.org/earlydevelopment.

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BOX 6-1
Chapter in Brief: Creating Healthy Living Conditions

Addressing the fundamental needs of families and children (i.e., economic stability, food security, and a safe and healthy living environment) is critical to achieving health and well-being during the prenatal through early childhood periods. This chapter identifies programs, policies, and systems changes that the committee concluded have the most evidence and promise for improving health and well-being outcomes for children and their caregivers, in addition to reducing disparities.

Chapter conclusions in brief:

- Increasing the economic resources families have available to meet basic needs when children are young (including prenatally) will improve children's health.
- Public programs that provide economic resources to families in the form of cash, tax credits, or in-kind benefits improve child health and development outcomes, which have long-lasting effects on health and educational outcomes.
- Income-support programs that are contingent on employment status or based on earned income have positive benefits for families yet may also have unintended consequences for child health and development outcomes through negative effects on attachment, breastfeeding, and caregiver stress. Thus, it is important to supplement work-support programs with basic support for families with young children that is not tied to employment.
- Given the importance of good nutrition for brain growth and development, providing resources to ensure families have access to sufficient and healthy foods can improve birth outcomes and child health outcomes.
- Child lead poisoning continues to be a pervasive problem in the United States. There are many effective programs and policies that, if implemented and funded, would prevent, or mitigate the impact of, lead poisoning prenatally and in early childhood.
- Healthy early development cannot occur without safe and stable housing. Lack of affordable housing and environmental hazards in housing disrupt healthy childhood development and parent/caregiver well-being.
- Lack of housing affordability and quality is an acute problem that disproportionately impacts people of color and contributes to health disparities among children.
- Current federal housing programs are not adequately funded, and there are not enough safe, affordable housing units in high-opportunity areas.
- Not all households experience the same level of risk of exposure to harmful environmental toxicants or pollutants. Poverty, substandard and/or unstable housing, race and ethnicity, and proximity to known sources of pollutants heighten pregnant women and children's risk of exposure and poor health and developmental outcomes.

Chapter recommendations in brief:

- Implement paid parental leave.
- Reduce barriers to participation to Women, Infants, and Children (WIC) and Supplemental Nutritional Assistance Program (SNAP) benefits; do not tie these benefits to parent employment for families with young children or for pregnant women.
- Increase the supply of high-quality affordable housing that is available to families.
- Develop a comprehensive plan to ensure access to stable, affordable, and safe housing in the prenatal through early childhood periods.
- Test new Medicaid payment models that engage providers and other community organizations in addressing housing safety concerns, especially focused on young children.
- Address the critical gaps between family resources and family needs through a combination of benefits that have the best evidence of advancing health equity, such as a combination of increased SNAP benefits, increased housing assistance, and a basic allowance for young children.
- Support and enforce efforts to prevent and mitigate the impact of environmental toxicants during the preconception through early childhood periods.

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This chapter addresses the fundamental needs of families and children that are critical to achieving health and well-being. In the report conceptual framework (Figure 1-9), these are the "healthy living conditions" situated in the second outermost circle, along with health systems and services (Chapter 5) and early care and education (ECE) (Chapter 7). Healthy living conditions are made up of the SDOH, or more specifically, the social, economic, environmental, and cultural drivers of health and well-being. These determinants are interdependent, and together, they create conditions that influence child health and the ability of a caregiver to fulfill a child's basic needs for healthy development.

Based on the core scientific findings in this report, this chapter seeks to address the challenges—for example, the barriers highlighted in the quote that opened this chapter—caregivers face with respect to securing economic stability and a safe and healthy living environment during the prenatal through early childhood periods. The committee reviewed promising community-level models and policy opportunities that focus on key neurobiological and socio-behavioral mechanisms needed for healthy development that yield the greatest impact to both mitigate and forestall the impacts of early life adversities on health.

Systems changes are needed to target multiple SDOH that shape early development and well-being. Systems that children interact with are most effective when they take into account developmental science and evidence when they are created, thereby meeting children's developmental needs. There are changes that could be made based on this science to existing policies that would make them more responsive to the needs of children. The recommendations in this chapter aim to provide predictability and security in the lives of children and their families through ensuring economic stability and a healthy and safe living environment. While the chapter takes a social determinants approach to addressing early living conditions, it should be noted that there are some important contextual factors that are not discussed here. For example, research shows that factors such as public transit, access to parks and green space, and mass incarceration all shape inequities for children and families (for example, see NASEM, 2017; Wildeman and Wang, 2017); however, these areas of programs, policies, and systems are not the focus of the solutions discussed in this report. The primary focus here are the programs, policies, and systems changes that the committee has identified as having the most evidence and promise for improving health and well-being outcomes for children and their caregivers, in addition to reducing disparities. The chapter includes discussions of the existing evidence and committee recommendations for solutions to address economic stability and security, food security and nutrition, housing, neighborhood conditions, and environmental exposures and exposure to toxicants.

ECONOMIC STABILITY AND SECURITY

Children's well-being and future health outcomes are strongly related to family income, and as the review in Chapter 3 shows, poverty is associated with significant detrimental effects on children's health, development, and well-being. A systematic review of the literature concluded that the evidence supports the conclusion that the link between income and child outcomes is causal; that is, "money makes a difference in children's outcomes" (Cooper and Stewart, 2013). The study also finds evidence that money in early childhood is important, particularly for cognitive outcomes. Thus, reductions in childhood experiences of poverty, and increasing the resources available to families to meet their basic needs, would be expected to improve children's health and developmental outcomes.

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Given the substantial evidence that money matters, an important factor in reducing health disparities in early childhood is to ensure that families with young children have sufficient resources. As the Council of Economic Advisors points out, current policies and public programs provide much less support for families when children are young compared to when they are school age, despite the needs and lower financial wherewithal, on average, of families with younger children (CEA, 2014). In this section, the committee reviews the evidence about U.S. safety net programs that are intended to increase financial resources of families with children through cash transfers or tax credits. In the following paragraphs, the committee assesses programs that provide targeted benefits to address food or housing shortfalls and programs to address neighborhood conditions. To retain a reasonable scope, the focus is on the largest safety net programs run by the federal and state governments that are offered to families with young children or pregnant women, while acknowledging that local governments, nonprofit organizations, and religious organizations also provide resources to help families in need.

Furthermore, the committee acknowledges the importance of providing parents and other caregivers with pathways to sustained economic security, such as educational opportunities and workforce development training. Chapter 3, for example, highlights the salience of parental educational attainment and household income as determinants of child health, well-being, and educational outcomes. Thus, an approach that enhances educational and economic opportunities and ultimately financial sustainability for caregivers would benefit children and families. Community-based programs that promote economic well-being for families are one promising avenue for advancing economic security. One such example is the Dudley Street Neighborhood Initiative's Fair Chance for Family Success—funded by the Boston Promise Initiative in partnership with the Family Independence Initiative. This is a peer-to-peer financial literacy and learning program, which reports improved outcomes for participating families in terms of amount of money in savings accounts, checking accounts, total assets, and subsidy income (NASEM, 2017). Similarly, programs that enable workforce participation or retention could also help families get on a path to economic security. WorkAdvance is one program that allows employers to place individuals with moderate job skills into training programs for specific sectors that have high demand for local workers (NASEM, 2019). Evaluation data for this program suggested large increases in workforce participation, training completion, and credential acquisition at a two-year follow-up (Hendra et al., 2016). While these types of programs are relevant to promoting healthy early development, the committee's approach in this report was to limit its scope to program, practice, and policy changes that had the strongest evidence for direct impacts on children and their well-being. Therefore, the committee did not include in-depth discussion of these types of economic and workforce support programs for caregivers in this chapter.

Policies and programs aimed at reducing the impact of poverty on children's health and well-being may provide cash benefits (directly, or indirectly through tax credits) or noncash or "in-kind" benefits, such as vouchers to buy food or housing. Alternatively, some programs directly provide food, housing, or education. In this section, the chapter first describes antipoverty programs designed to increase the level of (cash) resources families have, focusing on the two largest direct cash grant programs, Temporary Assistance for Needy Families (TANF) and Supplemental Security Income (SSI), and then on tax credits, focusing on the Earned Income Tax Credit (EITC) and child tax credits. Next, paid parental leave is discussed as another option for supporting families' needs when children are young. The committee examined the evidence on the extent to which these programs (i) increase cash resources and thereby reduce poverty, (ii)

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are associated with improved child health and development, including prenatal and birth outcomes, and (iii) are associated with longer-term health, educational, and economic outcomes. Throughout this section, the committee explores concerns about the strength of the evidence and the possibility of unintended consequences of these programs (which might directly or indirectly impact children's health). The committee's conclusions and recommendations build off those made in the 2019 National Academies report, *A Roadmap to Reducing Child Poverty* (NASEM, 2019), which provides a thorough analysis of the evidence for approaches to alleviate child poverty. Box 6-2 contains the conclusions from that report that are relevant to and informed this committee's conclusions and recommendations.

BOX 6-2
Relevant Conclusions from *A Roadmap to Reducing Child Poverty (2019)*[a]

1. Poverty alleviation can promote children's development, both because of the goods and services that parents can buy for their children and because it may promote a more responsive, less stressful environment in which more positive parent–child interactions can take place.
2. Some children are resilient to a number of the adverse impacts of poverty, but many studies show significant associations between poverty and child maltreatment, adverse childhood experiences (ACEs), increased material hardship, worse physical health, low birth weight (LBW), structural changes in brain development, mental health problems, decreased educational attainment, and increased risky behaviors, delinquency, and criminal behavior in adolescence and adulthood. As for the timing and severity of poverty, the literature documents that poverty in early childhood, prolonged poverty, and deep poverty are all associated with worse child and adult outcomes.
3. Periodic increases in the generosity of the EITC program have improved children's educational and health outcomes.
4. SNAP has been shown to improve birth outcomes and many important child and adult health outcomes.
5. The weight of the causal evidence indicates that income poverty itself causes negative child outcomes, especially when it begins in early childhood and/or persists throughout a large share of a child's life. Many programs that alleviate poverty either directly, by providing income transfers, or indirectly, by providing food, housing, or medical care, have been shown to improve child well-being.
6. Government tax and transfer programs modestly reduced the child poverty rate, defined by the Supplemental Poverty Measure (SPM), between 1967 and 1993 but became increasingly important after 1993 because of increases in government benefits targeted at low-income and nearly low-income people. Between 1993 and 2016, SPM poverty fell by 12.3 percentage points, from 27.9 to 15.6 percent, more than twice as much as market-income-based poverty.
7. A number of other program and policy options lead to substantial reductions in poverty and deep poverty. Two involve existing programs—SNAP and housing vouchers. The option of a 40 percent increase in EITC benefits would also reduce child poverty substantially.
8. The 20 program and policy options [that the committee considered] generate disparate impacts across population subgroups in our simulations[a] Although virtually all of them would reduce poverty across all the subgroups we considered, disproportionately large decreases in child poverty occur only for black children and children of mothers with low levels of education. Hispanic children and immigrant children would benefit relatively less.
9. Two program and policy packages developed by the [Poverty Roadmap] committee met its mandated 50 percent reduction in both child poverty (defined as 100 percent of SPM) and deep poverty (defined as 50 percent of SPM). The first of these packages combines work-oriented policy expansions with increases in benefit levels in the housing voucher and SNAP. The second package combines work-oriented expansions with a child allowance, a child support assurance program, and elimination of immigrant restrictions on benefits built into the 1996 welfare reforms. Both packages increase work and earnings, and both are estimated to cost between $90 and $111 billion per year.
10. The committee was unable to formulate an evidence-based employment-oriented package that would come close to meeting its mandate of reducing child poverty by 50 percent. The best package it could design combines expansions of the EITC, the Child and Dependent Care Tax Credit, a

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minimum-wage increase, and a promising career development program. Although this package is estimated to add more than a million workers to the labor force, generate $18 billion in additional earnings, and cost the government only $8.6–9.3 billion annually, its estimated reductions in child poverty are less than half of what is needed to meet the goal.

11. There is insufficient evidence to identify mandatory work policies that would reliably reduce child poverty, and it appears that work requirements are at least as likely to increase as to decrease poverty. The dearth of evidence also reflects underinvestment over the past two decades in methodologically strong evaluations of the impacts of alternative work programs.

[a] The charge of the National Academies committee that wrote the *A Roadmap to Reducing Child Poverty* report was to identify policies and programs that have the potential to reduce child poverty and deep poverty in the United States by half within 10 years. The committee examined 10 program and policy options. Four of them are tied to work, three of them modify existing safety net programs, two come from other countries, and the final one modifies existing provisions relating to immigrants. The committee then formulated two variations for each of the 10 options, yielding 20 scenarios in all.

SOURCE: NASEM, 2019

Cash Assistance Programs (TANF, SSI)

TANF provides cash assistance and sometimes other supports, such as job search support or child care subsidies, for eligible families with dependent children. Because TANF is a block grant, each state establishes its own eligibility rules, determines the type and amount of assistance to be provided, and sets other requirements and services (within broad federal guidelines). TANF participation is time limited and is intended to promote economic self-sufficiency, work, and marriage (HHS, 2012). In fiscal year 2018, 1.2 million families and nearly 2.4 million children received TANF assistance on average each month (ACF, 2019). TANF has been a shrinking component of the nation's social safety net for children since the passage in 1996 of the Personal Responsibility and Work Opportunity Reconciliation Act (PWRORA), when Aid to Families with Dependent Children (AFDC) was replaced by TANF. In 1996, 68 percent of low-income families with children received cash assistance through AFDC. In contrast, only 23 percent received TANF cash assistance in 2016 (CBPP, 2018a). In addition to reaching fewer low-income families, the size of grants has fallen in inflation-adjusted terms in most states. TANF benefits for a family of three in the median state were $447 per month in 2018 and have fallen by 20 percent in inflation-adjusted terms since 1996 (CBPP, 2018a). Overall, TANF is much less effective at reducing the severity of poverty than was the AFDC program: 18 percent compared to 56 percent of children moved out of deep poverty after receiving TANF versus AFDC grants (CBPP, 2018a).

Additional sources of direct cash support for families include the SSI for children and Social Security (SSA) programs. Children "with physical or mental condition(s) that very seriously limit his or her activities" and last over 1 year may qualify for SSI payments, which families can use to pay for basic needs, such as food and housing or medical care. To qualify, children need to meet the program's definition of eligibility and the family needs to have limited income and resources. In May 2019, about 1.1 million children under age 18 received an average monthly SSI payment of $674 (SSA, 2019). Nearly half (45 percent) of low-income families with an SSI recipient were lifted out of poverty by receiving SSI, according to a recent National Academies report (NASEM, 2015). Another 4 million children receive benefits through the Social Security program, as children of deceased workers (survivor benefits), children of workers with disabilities (through Disability Insurance), or children of retired workers. Summing across

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these groups, about $3.6 billion flowed to children through SSI and SSA programs (SSA, 2019). It is likely that the bulk of these monies goes to older children rather than those under age eight; nonetheless, for the families receiving these payments, the increase in income helps them to meet basic needs.

These programs (TANF, SSI, and SSA) all have the potential to improve children's health by raising family incomes; however, eligibility is limited, and the size of the assistance provided, especially for TANF, has not kept pace with rising costs of basic needs, particularly of housing. In terms of federal expenditures, the $12 billion of TANF spending on children and $12 billion in SSI for children with disabilities are only a small portion of federal spending on children in low-income families (Hoynes and Schanzenbach, 2018; Isaacs et al., 2017b). See Figure 6-1 for a breakdown of government spending on children by program from 1990–2015. The 2019 National Academies report *A Roadmap to Reducing Child Poverty* estimated that reductions in child poverty based on the current TANF program are small because of the low proportion of low-income children receiving TANF and the level of benefits. That report did not include expansion of TANF in the main strategies proposed to reduce child poverty, primarily due to the lack of evidence and the difficulties of assessing the effects of block grants on child outcomes when states have considerable flexibility in how the money is spent (NASEM, 2019). In contrast, they estimated that the child poverty rate (based on the supplemental poverty measure) would be 1.8 and 2.3 percentage points higher without SSI and SSA benefits, respectively (NASEM, 2019). While these programs (TANF, SSI, and SSA) provide some income support for young children, much more of the spending on children is through tax credits and in-kind assistance for food and housing, which we examine next.

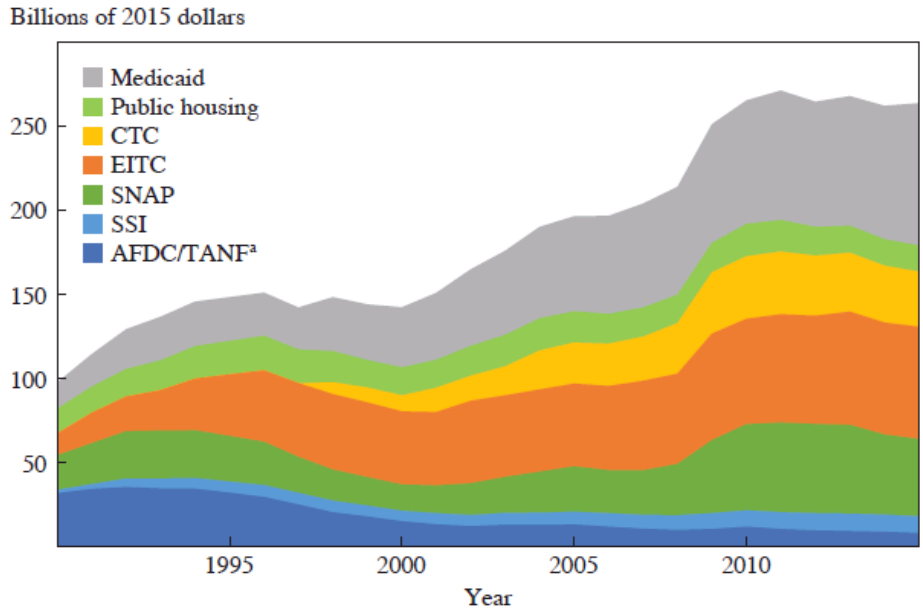

FIGURE 6-1: Government spending on children by program, 1990–2015
SOURCE: Hoynes and Schanzenbach, 2018

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Supporting Children Through Tax Credits

In contrast to cash assistance received on a monthly basis, tax credits represent an alternative financing mechanism for income support. When a family receives a tax credit, the taxes they owe to the government are reduced, and when the tax credit is "refundable," the family receives a payment if the credit exceeds the tax owed. In theory, a tax credit can provide the same amount of income assistance to a family as a direct cash transfer, although in practice, the amount and timing of the payments differ between the two types of mechanisms. Economists generally regard tax credits as having advantages over direct cash transfers in terms of the ease of administering the program (less bureaucracy) and lesser stigma from participation (Nichols and Rothstein, 2016). However, the extra income from a tax credit or refund is typically available only once per year and only if the family files a tax return. The two primary tax credits that apply to U.S. families with young children are the EITC and the Child Tax Credit (CTC), sometimes jointly referred to a working-family tax credits. While this section's focus is the federal level, many states offer similar tax credits to working families. In theory, working-family tax credits avoid the potential work disincentives of cash transfer programs, and research has demonstrated large increases in labor force participation, particularly for single mothers with children, as a result of the EITC (Nichols and Rothstein, 2016).

The EITC provides a refundable tax credit to eligible families based on earnings, number of children, and marital status. Initially implemented in 1975, EITC has seen its level and coverage expanded with bipartisan support several times over the past 45 years. The IRS and Census Bureau estimate that nationally, between 77 and 80 percent of eligible families claimed the EITC in 2015 (IRS, 2019). Although the take-up rate is high, outreach campaigns and use of tax preparation services or software can help increase the proportion of eligible families who receive the tax credit (Goldin, 2018). In 2016, the average EITC received by families with children was close to $3,200, which lifted an estimated 3 million children out of poverty and reduced the severity of poverty for nearly 7 million more children (CBPP, 2018c).

The CTC[2] is structured similarly to the EITC as a tax credit that is (partially) refundable. Low and moderate-income families can claim a tax credit for each of their children up to age 16. The "Tax Cuts and Jobs" Act of 2018 increased the CTC from $1,000 to $2,000 per child, with a maximum of $1,400 that is refundable. Over 90 percent of American families with children receive the CTC, and the average amount received in 2018 was $2,420 (per family) (TPC, n.d.). The average credit and share of families receiving the CTC is lower for families in the lowest income quintile, because some of these families will not have enough earnings to qualify and the CTC is not fully refundable. Nonetheless, the estimated effect of the CTC on poverty is notable: the Center on Budget and Policy Priorities estimates that the CTC lifted about 1.6 million children out of poverty and reduced the severity of poverty for 6.7 million children in 2017 (CBPP, 2018b). Most families in the second, third, and fourth income quintiles receive the CTC, while those in the lowest bracket benefit less (TPC, n.d.). The boost of $1,000–2,000 in the CTC is due to expire in 2025 (TPC, n.d.).

There is extensive research on the effects of the EITC on labor force participation (particularly of single mothers) and on health and educational outcomes for children in the United States. Less research has been conducted about the effects of the CTC, mostly because it was relatively small until recently. Given the similar structure of the CTC and EITC, many of the

[2] The refundable portion of the CTC is called the Additional Child Tax Credit (ACTC); here, both the ACTC and the CTC are included when referring to the CTC.

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effects are expected to be the same, with the important exception that the CTC is available to most moderate-income families, unlike the EITC, which is targeted at families with low incomes.

As noted above, the EITC and CTC together are effective in reducing child poverty: nearly 5 million children lived in families whose incomes were brought above the poverty level after including working-family tax credits, and over 7 million additional children experienced less severe poverty (CBPP, 2018c).[3] The poverty rate for children under 18 falls about 6 percentage points with the tax credits (16.4 percent compared to 22.8 percent without them) (Nichols and Rothstein, 2016). The reductions in child poverty are larger for these tax credits than other means-tested programs in the United States (Nichols and Rothstein, 2016). Given the well-established link between family income and health outcomes, by increasing family incomes, these tax credits are likely to lead to improved child health. One advantage of studying the link between EITC and outcomes is that researchers can use exogenous policy changes that increase families' income, avoiding the problem of endogenous income in studies of income–health links (Boyd-Swan et al., 2013).

While most studies have focused on the impacts of the EITC on labor force participation or reductions in poverty, a small but growing number of studies have examined the impacts of the EITC on health and education outcomes of children, both short and long term. Several studies found the EITC associated with higher birth weights and/or a reduction in the incidence of LBW (Baker, 2008; Hoynes et al., 2012; Komro et al., 2019; Strully et al., 2010). Using quasi-experimental methods, Hoynes et al. (2012) estimated a decline of 2–3 percent in LBW occurrence for a $1,000 increase in EITC, with larger effects for black than white women (among single women with a high school education or less). This study and one by Strully et al. (2010) also link EITC receipt with reduced rates of maternal smoking. In contrast, Baughman and Duchovny (2016) found that EITC receipt was not associated with improvements in parent-reported health status of children age birth to 5, although there were improvements for older children (age 6 to 14). Evidence linking the EITC to child cognitive or child development outcomes is limited (particularly for younger children). One study, by Hamad and Rehkopf (2016), used an instrumental variable approach to estimate the impact of the EITC on child development. They found "modest but meaningful improvements" in child behavior and home environments. The article notes that one mechanism through which the EITC impacts child development is through improved mental health of mothers (Evans and Garthwaite, 2010) and that it may lead to reductions in maltreatment (Berger et al., 2017). Studies focused on older children also find positive associations with EITC receipt on test scores (Dahl and Lochner, 2012; Chetty et al., 2011) and college attendance (Manoli and Turner, 2018).

It is important to note that effects of the EITC on health outcomes may be a result of the increase in family income; however, the EITC also strongly impacts work incentives, and impacts on child health and development may be due to changes in parent employment. Increases in maternal employment may increase the family's income but also reduce the amount of time mothers spend with their children. How these changes impact children's health and development likely depends on the quality of nonparental child care used and the stress experienced by parents, in addition to income changes (Hoynes and Schanzenbach, 2018).

Unlike the working-family tax credits, minimum-wage policies are not targeted specifically at low-income families with children. Nonetheless, some economists argue that minimum-wage policies play an important and complementary role in reducing child poverty along with the EITC (Nichols and Rothstein, 2016). Research summarized by Nichols and

[3] Note that these estimates are based on using the Census Bureau's SPM.

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Rothstein (2016) has demonstrated that the EITC provides a strong incentive for single mothers to work and has been a major factor underlying the rise in labor force participation of single mothers over the past two decades. An increase in the supply of labor, holding all else equal, could put downward pressure on wages, and the minimum wage provides a floor to keep wages from declining (Nichols and Rothstein, 2016). Parents who earn low wages benefit from a higher minimum wage in every paycheck along with receiving the tax credit when they file a tax return (CBPP, 2018c). A small number of studies also link increases in the minimum wage with health outcomes, though few examine children's health or birth outcomes. Two studies find small improvements in birth weight outcomes related to minimum-wage increases across states over time using quasi-experimental methods (Komro et al., 2016; Wehby et al., 2016). One study also found reductions in child maltreatment associated with higher minimum wages (Raissian and Bullinger, 2017). Given the evidence linking improved child health to higher family incomes, more research on the health effects of minimum-wage increases is needed to inform the policy debate.

Child Allowances

Many wealthy countries provide support to families through a child allowance or child benefit, which may be a cash grant or through the tax code. In the current U.S. tax code, the CTC and dependent child exemption act in many ways like a child allowance. The taxable income of families with dependent children is reduced, resulting in greater disposable income to meet the costs of raising children. A child allowance, distributed monthly, has two main advantages over the tax-code approach. First, it helps families with short-term needs for cash to meet expenses, compared to a once-per-year distribution through the tax system. Second, the lowest-income families often cannot take full advantage of tax credits and exemptions if their income is so low that they do not owe income tax. A child allowance paid to families on a monthly basis and not tied to earnings or employment would provide support for many of the lowest-income children in the United States whose parents do not work or have unstable and insufficient earnings.

A number of child allowance proposals have been proposed in recent years (NASEM, 2019; Shaefer et al., 2018). Schaefer et al. (2018) proposed a universal child allowance of $250 per child per month, possibly offering $300 for children under age 6 and slightly less for each additional child. Schaefer et al. noted that while this amount would not come close to covering the full cost of raising a child, based on their review of the research, it is large enough to have a meaningful impact on families and children and is comparable to child allowances in other countries. The National Academies *Child Poverty Roadmap* report considered two options for a child allowance: about $2,000 and $3,000 per year. The reductions in child poverty are larger for these two policies than any of the other individual policy changes considered by the report (although the costs are also higher).

A few key principles are important when considering the parameters of a child allowance that is intended to improve health outcomes for young children. First, targeting payments to families with the youngest children acknowledges that families with younger children have lower incomes on average than families with older children (see Schaefer et al., 2018). In addition, some costs (particularly child care) are higher for younger than for older children. The nation provides sizable public resources to children starting at about age 5 when they enter the public K–12 education system. A child allowance targeted at children under age five would help to balance public investments in different age groups.

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A universal program—in which all families with young children receive a child allowance—would reduce the stigma of participation relative to a means-tested program and may enhance social inclusion (NASEM, 2019, p. 5–31). While the costs of a universal program may exceed those of a targeted one, treating the allowance as taxable income would reduce the overall cost to the government. Alternatively, the child allowance could be phased out at 300 percent of the federal poverty level. Replacement of the current child tax credits with a monthly child allowance could provide families with a more regular source of cash income to support their children's needs (NASEM, 2019). Determining the specifics of a child allowance policy and funding mechanism requires additional research and modeling to compare the potential impacts on child health and health equity. In sum, based on the committee's review of the evidence and in accordance with the *Child Poverty Roadmap* report (NASEM , 2019), there is strong evidence that programs that provide direct income transfers or basic necessities such as food and housing lead to improvements in child health and well-being (p. 3–20). At the end of the chapter, the committee recommends expanding resources to support families with young children, with a child allowance as one important option to consider.

Paid Parental Leave

Maternal and paternal leave policies generally are intended to support mothers in recovering from childbirth and mothers and fathers in taking time off work to care for new infants. The United States remains one of the very few countries without a national paid guaranteed maternity leave policy. Across Organisation for Economic Co-operation and Development (OECD) countries, on average, mothers are entitled to 18 weeks of paid maternity leave (OECD, 2017). Although the United States is the only OECD country without a national-level policy of paid leave, California, New Jersey, New York, Rhode Island, Washington, Washington, D.C., and, most recently, Massachusetts have passed legislation to implement paid leave at the state and local levels (Mass.gov, 2019; Raub et al., 2018). See Box 6-3 for more information on California as an example of a state that has implemented a paid leave policy. A number of studies of paid maternity leave have found positive health effects, particularly lower infant and child mortality (Heymann et al., 2011; Nandi et al., 2018). Stearns (2015) estimated that paid maternity leave through temporary disability insurance in the United States reduced LBW by 3.2 percent and early births by 6.6 percent, with larger effects for certain subgroups, including black and unmarried mothers (Stearns, 2015). While improved maternal and child outcomes have been associated with paid leave policies in other countries, a review by Almond et al. (2018) showed mixed findings on child health impacts across studies, depending in part on the length of the leave. They concluded that "facilitating short maternity leaves is highly beneficial, but extended maternity leaves do not have a positive effect" on child outcomes (p. 1406). Rossin-Slater (2011) estimated that the Family and Medical Leave Act (FMLA), which allows for 12 weeks of unpaid maternity leave, resulted in lower infant mortality and slightly higher birth weight outcomes for college-educated women. Overall, however, based on her review of the literature, Rossin-Slater stated that "while extensions in existing paid leave policies have had little impact on children's well-being, the evidence suggests that the introduction of short paid and unpaid leave programs can improve children's short- and long-term outcomes" (p. 17).

Maternity leave policies have also been associated with higher rates of breastfeeding. As discussed in Chapter 3, breastfeeding provides important nutrition for developing infant brains and bodies; however, rates of exclusive breastfeeding for 6 months (as recommended by the

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World Health Organization and the American Academy of Pediatrics) are low in the United States, especially among black women. Further, the majority of mothers in the United States are not breastfeeding as long as they had planned (Mirkovic et al., 2014; Office of the Surgeon General et al., 2011). While the Affordable Care Act (ACA) has some protections for mothers who need to express milk while at work[4] (a critical component of successful breastfeeding for working mothers), many mothers are not given the time, appropriate space, and support needed to do so when in low-paying jobs (Murtagh and Moulton, 2011; Office of the Surgeon General et al., 2011). In a survey conducted by Declercq et al. (2013), 58 percent of women reported breastfeeding to be a challenge once they returned to work.

Paid leave "facilitates the initiation and continuation of breastfeeding" (Heymann et al., 2013). For example, a rigorous quasi-experimental study in California found that access to paid leave was associated with increased rates of exclusive and overall breastfeeding during the first 3, 6, and 9 months after birth (Huang and Yang, 2015). Studies (in both the United States and other developed countries) have found associations between maternity leave lasting at least 8 weeks and a higher probability of establishing breastfeeding (Guendelman et al., 2009; Ogbuanu et al., 2011; Skafida, 2012). Paid parental leave is hypothesized to increase mother–child attachment and give new mothers increased time to gain the skills and social support needed to maintain breastfeeding before returning to work.

The AEI-Brookings Working Group on Paid Family Leave published two reports, in 2017 and 2018, that focused on paid parental leave and paid family care and medical leave, respectively. Based on the extant literature on paid parental leave and its impact on family outcomes, the 2017 report puts forth a federal paid parental leave proposal. In addition to physical health and cognitive outcomes for children, the report cites improved labor force participation as a positive outcome associated with paid leave. For example, California and New Jersey's paid leave policies saw increases in labor force attachment among women in the months surrounding childbirth (Byker, 2016). This is important because continued workforce participation can help sustain household income and individual income, and other economic indicators, which have been linked to health and well-being (see, for example, NASEM, 2017; Woolf et al., 2015). The key elements of the AEI-Brookings federal paid leave proposal are benefits available to both mothers and fathers, wage replacement of 70 percent up to a maximum limit of $600 per week for eight weeks, and job protection for the individuals who take leave. The authors also suggest that such a federal paid leave program could be financed by a payroll tax levied on employees and/or savings in other areas of the budget (e.g., reduced tax expenditures in areas such as unemployment insurance (UI) or Social Security and disability programs) (AEI-Brookings Working Group on Paid Family Leave, 2017).

BOX 6-3
Paid Family Leave in California

California was the first U.S. state to implement a comprehensive Paid Family Leave (PFL) program in 2004, authorized by Senate Bill 1661 (Chapter 901, Statutes of 2002). The PFL program provides eligible employees up to 6 weeks of wage replacement leave (55 percent of regular weekly earnings) when they take leave from work to bond with a new child or to care for a seriously ill family member. The program is funded by a payroll tax levied on employees, so employers do not bear any direct costs. PFL benefit levels are indexed to inflation. California built the PFL program on the existing

[4] For example, the ACA updated the Fair Labor Standards Act to require U.S. firms with 50 or more employees to provide breastfeeding mothers with reasonable break time and space to express milk (DOL, 2018).

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State Disability Insurance system, so it is structured as an insurance benefit. California PFL was implemented as a virtually universal program—that is, almost all employees, excluding some self-employed persons, are eligible, regardless of the size of their employer.

The California Senate Office of Research reports that from 2004–2013, PFL applications increased overall, with almost a twofold increase in claims filed by men. This increase among men was reported to be driven almost exclusively by the number of men filing claims for caring for a new child. Applications for PFL for the purpose of caring for a new child make up about 88 percent of all PFL claims. Data from 2013 indicate that on average, women took 5.5 weeks of leave to care for a new child, while men took 4.5 weeks. One analysis used a differences-in-differences approach to examine how the PFL program affected leave-taking from 1999–2010. The authors found that the program doubled the use of maternity leave in California, with notable increases among non-college-educated mothers with infants (from 2.4 to 7.7 percent), unmarried mothers (from 1.9 to 9.3 percent), and African American mothers (from 2.0 to 13.7 percent) (Rossin-Slater et al., 2011). Other research has studied the California PFL program in relation to child and family outcomes. For example, the passage of the PFL program has been found to be associated with increased rates of breastfeeding (Huang and Yang, 2015).

SOURCES: Applebaum and Milkman, 2011; California Senate Office of Research, 2014; Huang and Yang, 2015; Rossin-Slater et al., 2011

Summary and Conclusions

There is considerable evidence that "income matters" for health outcomes, especially in early childhood. The *Child Poverty Roadmap* report concludes that "the weight of the causal evidence does indeed indicate that income poverty itself causes negative child outcomes, especially when poverty occurs in early childhood or persists throughout a large portion of childhood" (p. S-2). There is also strong evidence that the reverse is true: increasing family resources to meet basic needs supports the health and development of young children. Given the high rate of child poverty in this country compared to other wealthy nations, as well as large disparities across racial and ethnic groups in poverty rates, reducing childhood poverty is a critical, foundational step in reducing health disparities in early childhood.

> *Conclusion 6-1: Increasing the economic resources families have available to meet basic needs when children are young (including prenatally) will improve children's health and has the potential to reduce health and developmental disparities in early childhood.*

One way to increase the resources families have for basic needs is through social insurance and safety net programs that provide cash or tax credits to families. Studies demonstrate improved health outcomes when families receive assistance through government programs, such as the EITC and SSI. These programs are associated with improved birth, health, and educational outcomes for young children, which will set them on a better trajectory for lifelong health and well-being.

> *Conclusion 6-2: Public programs that provide economic resources to families in the form of cash, tax credits, or in-kind benefits improve child health and development outcomes, which have long-lasting effects on health and educational outcomes.*

Much of the support provided to families with children in the United States is in the form of "work supports," where eligibility and the level of benefits are closely tied to employment and

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earnings. These policies help to reduce poverty by both increasing resources and encouraging employment (which also can lead to higher family income in the future). However, the *Child Poverty Roadmap* report determined that a work-oriented package of programs and policies would be the least effective of the four packages they considered in reducing the number of children in poverty (see Table 6-1 for a summary of the components of each of the four packages). That report also concluded that mandatory "work requirements are at least as likely to increase as decrease poverty" (see Box 6-2).

TABLE 6-1: Components of the Four Packages and Their Estimated Costs and Impact on Poverty Reduction and Employment Change, from *A Roadmap to Reducing Child Poverty*

		1. Work-oriented package	2. Work-based and universal support package	3. Means-tested supports and work package	4. Universal supports and work package
Work-oriented programs and policy	Expand EITC	X	X	X	X
	Expand Child Care Tax Credit	X	X	X	X
	Increase the minimum wage	X			X
	Roll out WorkAdvance	X			
Income support-oriented programs and policies	Expand housing voucher program			X	
	Expand SNAP benefits			X	
	Begin a child allowance		X		X
	Begin child support assurance				X
	Eliminate 1996 immigration eligibility restrictions				X
	Percent reduction in the number of poor children	−18.8%	−35.6%	−50.7%	−52.3%
	Percent reduction I the number of children in deep poverty	−19.3%	−41.3%	−51.7%	−55.1%
	Change in number of low-income workers	+1,003,000	+568,000	+404,000	+611,000
	Annual cost, in billions	$8.7	$44.5	$90.7	$108.8

SOURCE: NASEM, 2019

In addition to a limited impact on reducing child poverty, further expansions of the work-oriented safety net programs may have unintended negative consequences for child health if

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parent employment results in lower rates of breastfeeding or disruptions to the attachment between infant and caregiver. Evidence of the importance of attachment and breastfeeding is discussed in Chapters 3 and 4. Work-oriented programs, such as the EITC, that increase families' incomes and increase employment are an important component of the social safety net. However, additional support for families with young children through paid parental leave or a child allowance that is not tied to parent employment would recognize the special needs of the earliest years, in which parent time and attention are critically important for children's health and development. Both paid parental leave and income support, such as a child allowance not tied to employment, may provide parents greater opportunity to take time out of the labor force to attend to their children's needs.

> *Conclusion 6-3: Income-support programs that are contingent on employment status or based on earned income have positive benefits for families yet may also have unintended consequences for child health and development outcomes through negative effects on attachment, breastfeeding, and caregiver stress. Thus, it is important to supplement work-support programs with basic support for families with young children that is not tied to employment.*

As noted above, additional income support for families with young children through paid parental leave would recognize the special needs of infants and their caregivers. Unpaid parental leave through FMLA does not cover all employees, and many families with low incomes are unable to afford to take an unpaid leave. Paid parental leave grants parents greater opportunity to take time out of the labor force to attend to their children's needs. Short, paid parental leave programs have been associated with positive health outcomes and higher rates of breastfeeding.

> **Recommendation 6-1: Federal, state, local, tribal, and territorial policy makers should implement paid parental leave. In partnership with researchers, policy makers should model variations in the level of benefits, length of leave, and funding mechanisms to determine alternatives that will have the largest impacts on improving child health outcomes and reducing health disparities.**

As of 2019, six states and D.C. have paid leave programs, and they finance these programs through employee payroll taxes (AEI-Brookings Working Group on Paid Family Leave, 2017). Some proposals for PFL follow a social insurance model, in which employees contribute through payroll taxes to a government-administered social insurance fund. Other financing options include an employer mandate, tax credits to encourage employers, or general funds (Isaacs et al., 2017a). Because there are a variety of options to implement, structure, and administer a paid leave policy, cost estimates for this program vary widely. In its 2018 report, the AEI-Brookings Working Group on Paid Family Leave offered three methods for assessing the cost of a hypothetical 8-week paid family medical leave program[5] had it been operational in 2016. The three methods use 1) national-level data, assuming uptake would be similar to private-sector participation under FMLA; 2) state paid leave data, assuming participation would mirror the rates of the states with operational programs; and 3) a simulation model to combine national- and state-level data. Because these methods differ with respect to data sources and assumptions

[5] The hypothetical program provides universal access to up to eight weeks of family and medical leave, including parental leave, with benefits paid at 70 percent of usual weekly wages up to a cap of $600 per week.

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on program use, the cost estimates vary widely and drawing comparisons can be difficult. Based on their analyses, the authors estimate that the program could be expected to cost from 0.10 percent of total wages or $7.65 million total benefits paid (based on New Jersey's state paid leave program) to 0.61 percent of total wages or $46.3 million total benefits paid (based on the FMLA national survey).

The committee did not study in depth other income-enhancement strategies to boost family resources that are not targeted particularly to health outcomes or early childhood, but these may be important for supporting the health and well-being of families and children. The National Academies *Child Poverty Roadmap* report details a number of additional strategies to reduce child poverty through, for example, increases in the minimum wage, job training programs, child care subsidies, and child support assurance, in addition to the policies discussed in this section. There is limited evidence of the impacts of these on child health, with the exception of the minimum wage (discussed earlier). The National Academies *Child Poverty Roadmap* report provides a careful assessment of a set of feasible strategies that could be used to reduce child poverty by half within 10 years (see Table 6-1). As discussed in Chapter 2, the scientific evidence amassed since *Neurons to Neighborhoods* has established that access to basic resources prenatally and in early childhood impact the developing child's brain and nervous system, immune function, and other organs (NRC and IOM, 2000). The toxic stress response of children living in poverty directly impacts behavioral and psychological well-being and substantially increases later-life risk for poor health and educational outcomes. Thus, reducing child poverty is a critically important, foundational strategy for improving child health outcomes and reducing health disparities in early childhood. Expansion of income-support programs that are not tied directly to parent earnings is likely to help those who need it most: children in deep poverty and the youngest children. Determining the specifics of a child allowance policy and funding mechanism requires additional research and modeling to compare the potential impacts on child health and health equity. At the end of the chapter, the committee recommends expanding programs to increase economic resources to support families with young children, with a child allowance as one important option to consider.

Policies that build family assets and wealth also deserve consideration in developing a national strategy to ensure all children have an equal opportunity to reach their full health and developmental potential. Individual Development Accounts (IDAs) and child savings accounts, for example, are typically targeted toward building savings for home ownership or postsecondary education. These strategies may have longer-term impacts on child and family well-being. Increasing education levels, particularly of mothers, also would likely lead to improved economic security for families. These policies support the broader goal of human capital development and long-term economic growth.

FOOD SECURITY AND NUTRITION

As described in Chapter 3, adequate and nutritious food are critically important for health outcomes during the preconception, prenatal, and early childhood periods. At times, adequacy of specific nutrients is crucial, such as folic acid during pregnancy. In each of these developmental periods, the overall adequacy and healthiness of food intake influence current health and development and have effects lasting into adulthood. Further, food insecurity may affect both children and parents through changes in eating habits and stress related to uncertainty and inadequacy of food availability. The neurobiological (and other) mechanisms underlying these

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effects were described in Chapters 2 and 3. In this section, we look at the programs and policies in the United States aimed at reducing food insecurity and improving nutrition and healthy eating, with a focus on the prenatal and early childhood periods.

Current Programs and Policies

Two major federal programs in the United States target the adequacy of food and nutrition for children living in households with limited resources: SNAP, formerly known as the Food Stamp Program, and the Special Supplemental Nutrition Program for WIC. In this section we examine the evidence on the effects of these two programs on children's health and development. Note that programs that operate primarily in schools and early education settings, such as the National School Lunch and Breakfast Program (SLB), are discussed in Chapter 7.

SNAP

SNAP provides assistance to eligible individuals and families to purchase food. Participants use an Electronic Benefit Transfer (EBT) card that functions like a debit card to purchase food from authorized retailers, which include supermarkets, grocery and convenience stores, and farmers' markets (CBPP, n.d.). Many participants enroll in SNAP for a short time—from 2009 to 2012, approximately 48 percent of participants received benefits for 24 months or less (Irving and Loveless, 2015; RWJF, 2018).

Participants need to meet requirements regarding income (gross[6] and net[7] monthly income), resources (such as cash, money in checking and savings accounts, and vehicles), and nonfinancial standards to be eligible to receive SNAP benefits (Cronquist and Lauffer, 2019). Undocumented noncitizens of the United States are not eligible for SNAP, but noncitizens who have lived in the United States for at least 5 years, receive disability-related assistance, or are less than 18 years of age are eligible (if they also meet the aforementioned income, resource, and nonfinancial eligibility requirements) (USDA, 2018b). The program expects that participating households will spend about 30 percent of their own financial resources purchasing food; thus, the amount in SNAP benefits received by each participating household is calculated by multiplying the household's net monthly income by 0.3 and subtracting the result from the maximum monthly allotment[8] for the household size.

Each month of fiscal year 2017, SNAP served 42.1 million individuals in 20.8 million households. Children were 44 percent of SNAP participants and received 43 percent of SNAP benefits. On average, the program provided assistance to 8.6 million households with children (42 percent of all households served by SNAP) each month. Of the total number of SNAP participants, 8 percent were children with U.S. citizen status living with noncitizen adults (Cronquist and Lauffer, 2019).

While SNAP benefits can be spent only on eligible food items, these benefits add to the total resources the family has to spend on all necessities. The average monthly benefit of $255 per household "represents a sizable income transfer to participants, and is expected to change the amount or quality of food purchased" (Hoynes and Schanzenbach, 2018, p. 13). A recent Urban Institute report estimates that the SNAP program reduced the number of children living in

[6] Includes a household's total, nonexcluded income, before any deductions have been made (USDA, 2018b).
[7] Gross income minus allowable deductions (USDA, 2018b).
[8] Maximum monthly allotments by household size are available at https://www.fns.usda.gov/snap/eligibility (accessed March 28, 2019).

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poverty by over a quarter and the number in deep poverty by nearly half (Wheaton and Tran, 2018). The report also found that SNAP reduced the poverty gap (defined as the additional income needed to lift all low-income families out of poverty) by 37 percent for families with children. Given the evidence on the links between health outcomes and income, one would expect these sizable reductions in poverty to lead to improved health outcomes. The evidence is growing that SNAP improves birth outcomes (Almond et al, 2011; East 2018), although, as discussed below, relatively few studies focus on the effects of SNAP on the health outcomes of young children.

A review of studies prior to 2003 concluded that SNAP participation increased household food expenditures (USDA, 2004), which suggests that SNAP would reduce food insecurity among recipient households. Because families experiencing greater hardship are more likely to participate in SNAP, however, some studies of SNAP's effect on food insecurity have found mixed and null results (Gibson-Davis & Foster 2006; Gunderson & Oliveira; Huffman & Jensen 2008; Wilde 2007; Wilde & Nord 2005). Gregory, et al. (2016) illustrate how estimates of the relationship between SNAP participation and food insecurity vary depending on statistical methods, demonstrating positive and negative estimates along with ones that were not significantly different from zero. They did conclude, however, that food insecurity was reduced by SNAP in a dose-response type model. Further, according to an Urban Institute report, "controlling for selection into SNAP is important for disentangling the effect of SNAP receipt on food insecurity" (Ratcliffe and McKernan, 2010) (p. 14). The authors found that the relationship between SNAP participation and food insecurity changed direction when they controlled for selection into SNAP using an instrumental variables approach. They concluded that SNAP participation reduced food insecurity by 16 percentage points (results for children not reported separately). Using methods to account for both selection and measurement error in reporting SNAP participation, Kreider et al. (2012) found a reduction of at least 8 percentage points in food insecurity for children, depending on the model assumptions. Deb and Gregory (2018) found that the effects of SNAP on food insecurity vary across the population; while it may have no effect for some, for those starting with low food security, it resulted in a much lower likelihood of food insecurity.

While SNAP increases household resources and reduces food insecurity for (at least) some families, studies of the impact of receiving food assistance on children's health outcomes are relatively rare. One study found the introduction of the Food Stamp Program in California was associated with a reduction in infant birth weight, particularly among first-time teen mothers for whom birth rates increased overall (Currie and Moretti, 2008). Potential mechanisms for this association could be related to fertility changes or the increased survival of low birth weight babies. The study findings also showed a small reduction in infant mortality for white babies in Los Angeles County. There is more recent evidence of a positive connection between receiving SNAP benefits (or food stamps) and improved birth outcomes. Almond et al. (2011) and East (2018) both found positive associations between food assistance during pregnancy and improved birth outcomes, using quasi-experimental methods. Almond et al. found larger improvements in birth weight outcomes for African American mothers and those living in high-poverty areas (Almond et al., 2011). They noted that these results occurred despite the fact that the Food Stamp Program was not designed to target pregnant women. See Figure 6-2 for data on the impact of in utero exposures to food stamps on likelihood of birth weight below selected cut-offs.

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Figure 2: Impact of In Utero Exposure to Food Stamps: Reduction in Likelihood of Birth Weight Below Selected Cut-Offs

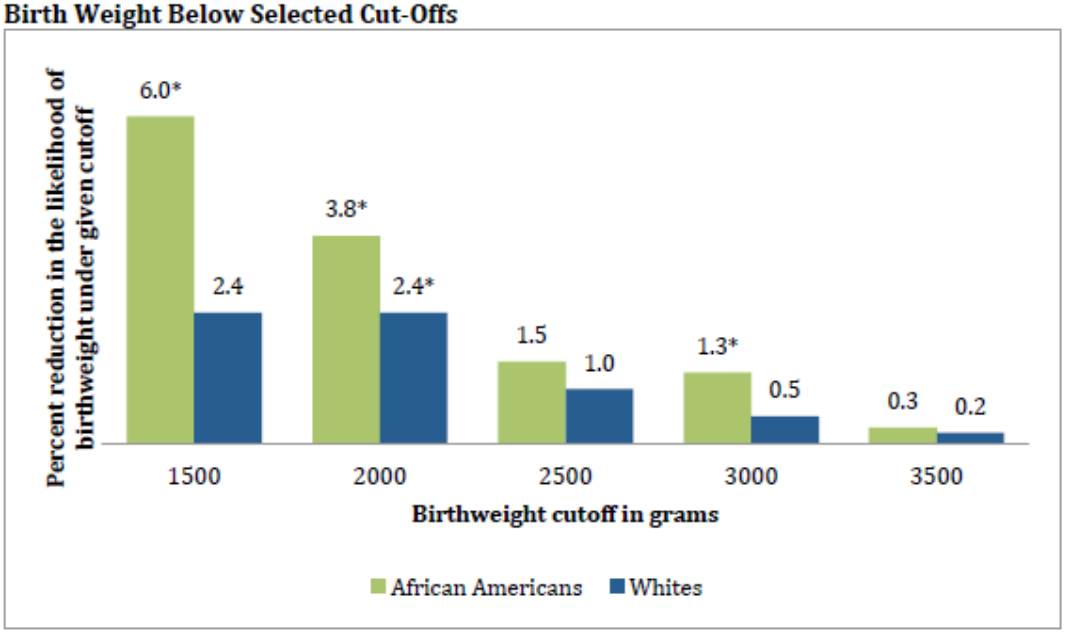

FIGURE 6-2: The impact of in utero exposure to food stamps on likelihood of birth weight under selected cut-offs.
SOURCE: Hoynes and Schanzenbach, 2016
NOTES: * denotes estimate statistically significantly different from zero. Data from Almond et al., 2011.

With respect to other health outcomes for children, there is limited causal evidence. Kreider et al. (2012) reported improvements in child health outcomes (along with reductions in child food insecurity); however the range of possible effect sizes is large. They accounted for both selection and underreporting of SNAP participation but did not specifically focus on young children. Most studies focus on adult health outcomes and found mixed results for adults (Kreider et al., 2012; see, for example, Gregory and Deb, 2015; Yen et al., 2012). Overall, there is limited evidence on the causal effects of SNAP on children's health because there have been few opportunities for random assignment[9] or limited variation in policies over time or in different places to exploit with experimental or quasi-experimental methods (East, 2018).

While studies of direct or contemporaneous effects on children's health are limited, one recent study demonstrated a link between receipt of SNAP in childhood and adult health outcomes. Hoynes et al. (2016) found that adult health measured by the metabolic syndrome index was significantly better for those whose childhood families had access to food stamps, particularly in early childhood (before age 5). Long-term positive health effects of SNAP—that is, adult health outcomes for those receiving SNAP as children—are consistent with short-term health improvements during childhood. Similarly, East (2018) found positive effects of SNAP participation before age 5 on health outcomes when children were age 6 to 16, using quasi-experimental methods for a sample of children born in the United States to immigrant parents. While conclusions about the immediate impacts of SNAP on young children's health are

[9] Random-assignment demonstration projects have been conducted recently and are underway in several sites as part of the Demonstration Projects to End Childhood Hunger project and the Healthy Incentives Pilot (Olsho et al., 2017; USDA, 2018a).

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provisional given the methodological challenges of estimating causality (Carlson and Keith-Jennings, 2018), the evidence of SNAP impacts on reduced food insecurity and later health outcomes suggests that children benefit in both the short and long run.

One of the concerns about the SNAP program has been the potential linkage between SNAP benefits and obesity, in both adults and children. Earlier studies that did not adequately control for selection into SNAP found positive correlations between SNAP and obesity, while other studies found reductions in obesity or no effects (Fan and Jin, 2015; Kreider et al., 2012). Myerhoefer and Yang (2011) concluded that "the balance of evidence points to a small positive impact of SNAP participation on obesity for women" but that the results for "childhood obesity are less consistent" (p. 313). For children, Kreider et al. (2012) concluded that the obesity rate was 5.3 percentage points lower due to SNAP.

In addition to reducing hunger and food insecurity, a second key objective of the nation's food assistance programs is to improve the healthfulness of American food consumption and provide nutrition education. Studies have examined the effect of SNAP on the quality or healthfulness of family's food consumption as a possible mechanism through which SNAP might affect obesity (and other health) outcomes. Most studies have focused on adult food intake, but Yen (2010) found no effect of the SNAP program on young children's nutrient intake. A number of demonstration projects have been conducted to evaluate ways to incentivize or influence consumption of healthful foods through SNAP. The Healthy Incentives Pilot (HIP) project provided a 30 percent rebate on purchases of a specified set of fruits and vegetables using SNAP benefits. Households receiving SNAP were randomly assigned to receive the rebate or not. The HIP evaluation reported that households receiving the rebates increased consumption of targeted fruits and vegetables by 26 percent, although some reported confusion and misunderstanding about how the rebate program worked and which vegetables and fruits were included (Olsho et al., 2017). The Summer EBT for Children pilot also was a random assignment design, but rather than targeting specific purchases, participants were provided (an extra) $60 per month per school-age child. The evaluation study found modest improvements in several child nutritional outcomes and null effects for others (Collins and Klerman, 2017) even though this program did not specifically target or incentivize healthful food purchases. While SNAP benefits can be used to purchase almost any food item from participating retailers, recent pilot projects found that increased benefits and incentives for purchasing specific healthful foods can modestly impact some health-related and dietary outcomes. Proposals to restrict SNAP purchases to prohibit less healthful foods, such as sugary beverages, or to incentivize purchases of fruits and vegetables, are highly controversial (Schwartz, 2017).

In summary, as discussed throughout this section, food assistance provided through SNAP is a major component of the nation's social safety net and reduces poverty and food insecurity for millions of families and children (Carlson and Keith-Jennings, 2018). While the evidence of SNAP's direct impact on children's health is limited, Hoynes, Schanzenbach & Almond (2016) found that adults who received SNAP as children experienced a reduction of 5 percent in heart disease and 16 percent reduction in obesity. Improvements in adult health outcomes related to receipt of SNAP in early childhood suggest that children are benefitting as well. Based on Hoynes and Schanzenbach's (2015) summary of the literature examining links between SNAP participation and health outcomes, the NASEM (2019) report on reducing child poverty concluded that "many (but not all) of the methodologically strongest studies show SNAP benefits having positive impacts on health" (p. 3-15).The report cites evidence that increasing SNAP benefits would substantially reduce child poverty and that current benefit levels do not

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account for food preparation time or geographic variation in food costs (Ziliak, 2016). Based on the evidence demonstrating the links between child health and family income or resources, increasing SNAP benefits would likely lead to improved child health and reduce health disparities.

WIC: The Special Supplemental Nutrition Program for Women, Infants, and Children

WIC provides assistance through breastfeeding support and education, healthy foods, nutrition education and counseling, screening referrals to other services, and vouchers to purchase fruits and vegetables from authorized farmers' markets (USDA, 2015). WIC services are provided in many locations, including county health departments, hospitals, mobile clinics, community centers, schools, public housing sites, migrant health centers and camps, and Indian Health Service facilities (USDA, n.d.).

To be eligible to receive benefits, WIC participants need to meet all four categories of requirements: categorical, residential, income, and nutrition risk (USDA, 2018c). Infants younger than 1 year; children younger than 5 five years; and women who are pregnant, postpartum (up to 6 months), or breastfeeding meet the WIC categorical requirement. To meet the residential requirement, participants need to reside in the state or local service area in which they apply but are not required to have lived in that area for a minimum amount of time. Participants also have to earn incomes at or below income standards that, set by state agencies, and fall within 100 and 185 percent of the federal poverty guidelines issued annually by HHS. Lastly, participants need to be determined by a health professional to have at least one medical (e.g., anemia, underweight) or dietary (e.g., poor diet) condition from a list of conditions indicating nutrition risk that is set by states (USDA, 2018c).

In fiscal year 2018, WIC served approximately 6.9 million people and cost the federal government about $5.3 billion (USDA, 2019). In 2016, it was estimated that 64 percent of individuals eligible to receive WIC benefits were children ages 1 to 4, 21 percent were pregnant and postpartum women, and 16 percent were infants, for a total 13.9 million individuals. Of these, 7.6 million (55 percent) received WIC benefits, with 86 percent of eligible infants receiving benefits but only 44 percent of children ages 1–4 (Trippe et al., 2019).

Numerous studies using varied methods have found evidence of improved birth outcomes to women participating in the WIC program (see, for example, Figlio et al., 2009; Fingar et al., 2017; Foster et al., 2010; Hoynes et al., 2011). These studies find reductions in the likelihood of LBW (Figlio et al., 2009; Hoynes et al., 2011) and reductions in infant mortality (Khanani et al., 2010). Figlio et al (2009) studied the effect of WIC in Florida between 1997 and 2001 by matching infant birth records with school records for older siblings to identify those who were marginally eligible and marginally ineligible in order to form groups for comparison. They estimated a significant reduction in the likelihood of LBW among WIC participants, although there was no significant effect on average birthweight or gestational age. Studies generally have found the effects of WIC on birth outcomes are stronger for women with lower education levels, those living in areas of high poverty, and African Americans (Hoynes et al., 2011; Khanani et al., 2010). Fingar et al. (2017) accounted for gestational age, which might bias estimates in other studies, and reported a significantly reduced risk of preterm birth, LBW, and prenatal death.

One of the potential channels through which WIC impacts birth outcomes is through changes in dietary quality and access to nutritional information and support for healthy behaviors, such as quitting smoking, for pregnant women. Participants can use their WIC vouchers only for specific foods, and changes in the approved foods after 2008 reflect dietary

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recommendations from the American Academy of Pediatrics (AAP) and IOM. The USDA published a final rule in 2014 that provides for more purchases of fruits and vegetables, whole-grain options, yogurt and soy options in place of milk, and more flexibility to tailor food packages to individuals (Carlson and Neuberger, 2018). While there has been concern about the possibility that providing infant formula to new mothers reduces breastfeeding, the changes to food packages and incentives after 2014 have encouraged breastfeeding (NASEM, 2016). The rate of breastfeeding among WIC participants has risen 45 percent over 12 years, reducing the difference between all women and WIC participants (Carlson and Neuberger, 2018).

There is solid evidence linking the WIC program to improved nutrient intake and eating more healthful food, although many of the studies report household-level consumption and do not focus specifically on children. The introduction of the improved food packages in WIC led to noticeable improvements in the percentage of families reporting that they eat more whole grains, drink lower-fat milk, and consume more fruits and vegetables (Andreyeva and Luedicke, 2013; Chiasson et al., 2013; Whaley et al., 2012). Thus, the additional resources to purchase specific food items provide by the WIC program is associated with changes in the types of food consumed by households (Whaley et al., 2012). In the case of the HIP, these changes may have been due in part to "promotional effects," whereby the incentive for certain foods provides information to participants about which foods are healthier (Olsho et al., 2017).

By providing access to healthier foods and nutrition information, WIC would be expected to improve the health and developmental outcomes of young children. WIC may also reduce food insecurity. One study found that WIC participation reduced the number of children experiencing food insecurity by 20 percent (Kreider et al., 2016). There is also some evidence about the impact of WIC on young children's cognitive and socio-emotional development. Jackson (2015) used matching and fixed effects estimation methods and found improvements in cognitive development at age 2 and reading and math scores at age 11 for children whose mothers participated in WIC prenatally. In contrast, Arons et al. (2016) found no significant improvement in socio-emotional development among young children receiving WIC; however, their sample size was small. Based on the current literature, the extent to which WIC supports cognitive and noncognitive development in young children is still uncertain.

Overall, the evidence is solid that WIC leads to improved birth outcomes and improved dietary intake for participants although there is less evidence that it directly improves children's health and development in the early years. Revisions to the food package and incentives for purchasing specific fruits and vegetables, have led to improvements in dietary quality. Evidence of savings on health costs, particularly postpartum, from the mid-1990s suggested that (back then) the cost savings far exceeded program costs (GAO, cited in Carlson and Neuberger, 2018, p. 24). Nearly two thirds of all infants and half of pregnant and postpartum women are eligible for WIC.[10] The broad reach of the WIC program has been viewed as a positive attribute, but some feel that it indicates that the program is not sufficiently targeted and resources could be spent more efficiently (Besharov and Call, 2009). However, many eligible families do not receive WIC benefits. In 2014, only half of eligible pregnant women participated, while 80 percent of eligible infants did (Johnson et al., 2017). While the WIC program is largely successful in supporting the health and nutrition of its recipients, investigating the barriers to participation and further studying heterogeneous effects on different subgroups are necessary to further understand its potential to reduce health disparities in early childhood. Furthermore, better coordination between the WIC program, ECE systems, and prenatal, postpartum, and

[10] See https://www.fns.usda.gov/wic/wic-eligibility-and-coverage-rates (accessed July 14, 2019).

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pediatric care would allow for a more integrated systems approach to addressing children's nutrition and developmental needs. (See Chapter 8 for more on applying a systems approach to promoting equitable healthy development.) Box 6-4 describes Healthy Mothers on the Move as an example of a promising model to improve nutrition and healthy lifestyles.

BOX 6-4
Healthy Mothers on the Move (Healthy MOMs)/
Madres Saludables en Movimiento (Madres Saludables): Promising Model[a]

Healthy Mothers on the Move (Healthy MOMs) / Madres Saludables en Movimiento (Madres Saludables) was a culturally tailored intervention that encouraged healthy lifestyle practices for Latina and African American women in southwest and eastside Detroit. Informed by community-based participatory research that identified the needs of pregnant and postpartum women in the community, the program aimed to decrease excessive weight gain during pregnancy and excessive postpartum weight retention in order to reduce risk factors for obesity, gestational diabetes, and type 2 diabetes (Detroit URC, n.d.; University of Michigan Prevention Research Center, n.d.).

The program consisted of a "healthy lifestyle intervention" and a "healthy pregnancy intervention" that served as a control group. The former consisted of culturally tailored programs, including curriculum-based education and home visiting, designed to equip participants with knowledge on pregnancy, childbirth, postpartum, and maternal and infant development as well as skills and supports to develop healthy lifestyles and manage stress. The programs were conducted in English and Spanish by community health workers. The control intervention provided the culturally tailored education on the same topics in English and Spanish (Detroit URC, n.d.).

The efficacy of the intervention was demonstrated through positive findings from longitudinal comparisons of the two interventions (Detroit URC, n.d.; Kieffer et al., 2014; Thornton et al., 2006). Participants were found to have decreased fat and sugar consumption, increased vegetable and fiber consumption, and decreased risk of depression (REACH Detroit, 2018).

The program's multisector partners included the Community Health and Social Services Center (CHASS) (a community-based nonprofit providing primary health care and support services to underserved residents of Detroit); the Detroit Department of Health and Wellness Promotion; Friends of Parkside (a community-based nonprofit); Harper-Gratiot Neighborhood Service Organization (a health and human services agency); Latino Family Services; the Michigan Department of Community Health; Southwest Solutions; St. John Health System; and the University of Michigan Schools of Social Work, Public Health, and Nursing. The program was funded by the National Institute of Diabetes and Digestive and Kidney Diseases from 2002 to 2008.

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

Summary

Both the SNAP and WIC programs have been studied extensively, and a large body of literature points to strong associations between program participation and positive outcomes, including less food insecurity, reductions in poverty, and greater consumption of healthy foods. The evidence is convincing that WIC, which is targeted to pregnant and postpartum women, infants, and young children, improves birth and postpartum outcomes. There is also strong evidence that SNAP improves birth outcomes and child health. In evaluating the impacts of both programs, however, confounding factors are important to consider: participants may be more disadvantaged than nonparticipants but also may self-select into the programs, and either factor may bias study estimates. A small but increasing number of studies use experimental and quasi-

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experimental methods to estimate causal effects, although few focus specifically on health outcomes for young children. In addition to providing additional resources to the family to meet their basic needs, both SNAP and WIC can increase the consumption of healthful foods through nutritional education and incentives.

> *Conclusion 6-4: Given the importance of good nutrition for brain growth and development (during the preconception, prenatal, and early childhood periods), providing resources to ensure families have access to sufficient and healthy foods can improve birth outcomes and child health outcomes.*

Because safety net programs, such as WIC and SNAP, have been shown to improve birth outcomes and to reduce food insecurity for young children, the committee recommends:

> **Recommendation 6-2: Federal, state, local, territorial, and tribal agencies should reduce barriers to participation to the Special Supplemental Nutrition Program for Women, Infants, and Children (WIC) program and Supplemental Nutrition Assistance Program (SNAP) benefits. Receipt of WIC and SNAP benefits should not be tied to parent employment for families with young children or for pregnant women, as work requirements are likely to reduce participation rates.**

As noted earlier, the National Academies *Child Poverty Roadmap* report concludes that the current level of SNAP benefits is inadequate. That report considers two options for increasing benefit levels: a 20 or 30 percent increase (along with a higher benefit amount for households with teenagers and a boost in summer benefits). The report also notes that SNAP has a larger effect on reducing deep poverty than other government assistance programs do. Given the strong evidence linking improved child food security and SNAP, along with evidence of longer-term positive outcomes, increases in SNAP benefit amounts are likely to reduce health disparities. This committee had insufficient evidence to compare the effects on health disparities of alternative means of increasing family resources, such as increasing SNAP benefits or providing a monthly child allowance. For the most part, a dollar increase in SNAP benefits will have the same effect as a cash dollar, although some families might increase food expenditures more with an increase in SNAP. Increasing family resources is a critical, foundational step to reduce child health disparities (see Conclusion 6-1). Careful study and modeling is needed to determine the most cost-effective way to do so, with particular attention to the potential impacts on child and caregiver health and well-being and maternal employment, attachment, and breastfeeding. At the end of the housing section in this chapter, the committee recommends expanding resources to support families with young children, with an increase in SNAP benefit levels as one important option to consider.

HOUSING

Housing Affordability and Child Health and Equity

Access to affordable housing is considered an “upstream” determinant of child development, as it has implications for housing quality (Evans et al., 2000), instability (Garboden et al., 2017; Jelleyman and Spencer, 2008), and loss of housing (Sandel et al.,

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2018)—all well-established determinants of child health (Leventhal and Newman, 2010). Unaffordable housing, or "high housing cost burden"—typically defined as housing costs above 30 percent of household income—is a critical social issue (Desmond, 2018) that has worsened during the past several decades (Joint Center for Housing Studies of Harvard University, 2017). In 2016, 47 percent of all renters, and more than three quarters of families earning less than $30,000, had unaffordable housing (Joint Center for Housing Studies of Harvard University, 2017). At the extreme, nearly 110,000 children are estimated to be homeless on any given night in the United States, and more than half of families who used shelters in 2016 identified as African American or black (U.S. Interagency Council on Homelessness, 2018).

The evidence discussed in Chapter 3 suggests that lack of affordable and quality housing, housing instability, and overcrowding have significantly detrimental effects to the health, well-being, and development of infants, children, and families. For more information on the evidence supporting the role of affordable housing in promoting positive outcomes for child health and development, see Chapter 3.

Improving Housing Affordability and Quality

Federal housing assistance is provided through a number of programs, including the Housing Choice Voucher Program, public housing, and the Low Income Housing Tax Credit. The Housing Choice Voucher Program is the largest federal housing assistance program for people with low incomes. Administered by the U.S. Department of Housing and Urban Development (HUD), this program provides funds to local public housing agencies (PHAs). PHAs have latitude in how the program is administered and what populations are prioritized. Eligibility is based on average median income in the geographic area and stratified by extremely low income, very low income, and low income. A PHA has to provide 75 percent of its vouchers to people with extremely low incomes (Eligibility.com, 2019). Although HUD provides housing assistance through the Housing Choice Voucher Program to more than 2 million families per year (CBPP, 2019)—which ensures that participating households contribute no more than 30 percent of their income to rent (CBPP, 2017)—only one quarter of all income-eligible households receive housing assistance (CBPP, 2017), and the average family will spend 26 months waiting for assistance (HUD, 2016). One analysis found that the percentage of families wih children receiving rental assistance decreased by 13 percent since 2004, while "the number of families that paid more than half their income for rent or lived in severely substandard housing rose by 53 percent between 2003 and 2013, to nearly 3 million" (Mazzara et al., 2016).

HUD housing assistance can help families to obtain improved housing quality and residential stability (social factors that are associated with child development and disparities) (Fischer, 2015; HUD, 2014, 2015). There is some evidence to suggest that housing assistance has a beneficial impact on child health (Slopen et al., 2018), although this is an underexplored area of research. Only a small number of studies have rigorously controlled for selection bias, thereby limiting interpretation of the results for many of the existing studies (Ahrens et al., 2016; Fenelon et al., 2018; Fertig, 2007; Jacob et al., 2015; Kimbro et al., 2011; Leech, 2012; Newman and Holupka, 2017; Slopen et al., 2018). According to an analysis by Chetty et al. (2016) of the Moving to Opportunity demonstration, the benefits of the voucher program may be greater for children who move when they are young (less than 13 years of age) and may "reduce the intergenerational persistence of poverty and ultimately save the government money" (p. 860). (See the section on Improving Neighborhood Conditions for more on the Moving to Opportunity study.)

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Although this program is designed to provide families with choices about residential location, new evidence suggests that the program falls short on multiple neighborhood characteristics for families with children (Mazzara and Knudsen, 2019). A 2019 study by the Center on Budget and Policy Priorities using HUD administrative data and Census survey data revealed that in the 50 largest metropolitan areas in the United States, voucher-assisted families with children are disproportionately clustered into high-poverty, low-opportunity, or minority-concentrated areas, relative to the distribution of voucher-affordable housing across the metropolitan area. For example, 33 percent of families with children using vouchers reside in high-poverty neighborhoods (Census-tract poverty rate at or above 30 percent) even though only 22 percent of voucher units are in high-poverty neighborhoods. Similarly, 61 percent of voucher-assisted families of color with children reside in "minority-concentrated" areas (Census-tract percent of people of color is at least 20 percentage points greater than the proportion in the entire metropolitan area), although only 32 percent of voucher units are allocated to minority-concentrated areas (Mazzara and Knudsen, 2019). Some programs, such as the Baltimore Mobility Program, include intensive counseling and require the use of vouchers in low-poverty areas for at least a year (Darrah and DeLuca, 2014). However, lack of affordable units in higher-opportunity neighborhoods remains a barrier (Misra, 2016). Other barriers include inflexible limits on search periods for families to find units and landlord resistance to voucher clients (Sard et al., 2018).

The 2019 report *A Roadmap to Reducing Child Poverty* modeled "expansions of voucher availability rather than other modifications, such as an increase in the level of housing subsidies, primarily because most experts agree that limited availability is currently the primary barrier preventing subsidized housing programs from having a larger impact on poverty reduction" (NASEM, 2019). That committee also noted that "there is as yet no consensus among researchers as to whether existing housing subsidy levels set by the government are sufficiently aligned with true market rents faced by low-income families."

As discussed in Chapter 3, housing quality is also a contributor to child health and development. A systematic review found strong evidence of effectiveness for home interventions focused on addressing asthma triggers, including multifaceted, in-home, tailored interventions (including mattress and pillow covers, HEPA vacuums and air filters, and cleaning), cockroach control through integrated pest management (including e-strategies, reducing access points, and using low-toxicity gel-bait pesticides), and combined elimination of leaks and removal of moldy items (Krieger et al., 2010). Several of the reviewed studies focused on children. A systematic review conducted by Crocker et al. (2011) found that home-based, multitrigger, multicomponent interventions reduced asthma symptoms and school absenteeism, as well as asthma acute symptoms, among children and adolescents. These assessments and interventions are often performed by home visitors or community health workers and have been found to be effective in both urban and rural areas (Chew et al., 2003; Crain et al., 2002; Levy et al., 2006; Morgan et al., 2004). The actions taken as a result of these assessments and interventions have been found to reduce disparities in asthma-related outcomes based on race/ethnicity and income (Postma, 2009).

Similarly, a comprehensive review by the Health Impact Project found residential remediation to be an effective primary prevention strategy to reduce childhood lead exposure (Health Impact Project, 2017). This remediation can range from complete removal or permanent containment of lead paint to scraping and painting over existing paint and covering contaminated soil. Secondary prevention through screening by pediatricians and other health care providers of

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young children and treatment for those with elevated blood lead levels (BLLs) is also critical for mitigating potential long-term harm.

Several states, including New York and Maryland, have undertaken state- and municipality-level lead prevention and mitigation efforts that have led to significant decreases in childhood lead exposure and poisoning. In New York, the City of Rochester implemented a Lead-Based Paint Poisoning Prevention Ordinance in 2006, which requires inspections for lead paint as part of existing inspections of most rental properties built before 1978. In addition, New York requires that all children undergo BLL testing at ages 1 and 2, which is overseen by the state's health department. The health department also provides educational and environmental interventions for children who are found to have elevated BLLs (City of Rochester NY, n.d.).

Enacted in 1994 and modified in 2012, Maryland's Reduction of Lead Risk in Housing Act has helped to make housing units safer for children by requiring owners of rental properties built before 1978 to ensure their properties comply with a lead paint risk reduction standard. The state has also invested in strong public enforcement of the act, which is coordinated by the Maryland Department of the Environment's Lead Poisoning Prevention Program.[11] Through partnerships with nonprofits that provide legal services, such as the Green & Healthy Homes Initiative,[12] the state has also increased compliance with the law via private enforcement (Trust for America's Health, n.d.). As of 2016, Maryland requires all children born in or after 2015 to undergo blood lead testing at ages one and two (Maryland Department of Health, 2016). The Maryland Department of Health and Mental Hygiene oversees the state's blood lead testing efforts and other services, including case management follow-up for children found to have elevated BLLs and community education for parents, tenants, rental property owners, homeowners, and health care providers (Trust for America's Health, n.d.). As a result of these initiatives, from 1993 to 2015, the number of Maryland children under 6 years old whose BLLs were 10 µg/dl or higher decreased from 23.9 percent (14,564 of 60,912 children tested) to 0.3 percent (377 of 110,217 children tested). In addition, from 2013 to 2014 alone, the number of rental properties that were treated and received certification for compliance with the lead paint risk reduction standard increased from 28,000 to 57,603.

While the United States has reduced the number of children at risk of lead poisoning, it has not completely eliminated lead hazards, and those most at risk for lead exposure are low-income and minority children.

Conclusion 6-5: Child lead poisoning continues to be a pervasive problem in the United States. There are many effective programs and policies that, if implemented and funded, would prevent, or mitigate the impact of, lead poisoning prenatally and in early childhood. Concerted efforts are needed to continue to ensure progress—through both policy and regulatory actions—on this preventable but serious problem.

Many high-quality reviews of lead poisoning prevention have been completed and contain important recommendations for remediation and prevention. For example, a report by the Health Impact Project (2017) includes the following recommendations and findings:

- **Reduce lead in drinking water in homes built before 1986 and other places children frequent.** Removing leaded drinking water service lines from the homes of children born

[11] For more information, see https://mde.maryland.gov/programs/Land/LeadPoisoningPrevention/Pages/index.aspx (accessed March 21, 2019).
[12] For more information, see https://www.greenandhealthyhomes.org/ (accessed March 21, 2019).

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in 2018 would protect more than 350,000 children and yield $2.7 billion in future benefits, or about $1.33 per dollar invested.

- **Remove lead paint hazards from low-income housing built before 1960 and other places children spend time.** Eradicating lead paint hazards from older homes of children from low-income families would provide $3.5 billion in future benefits, or approximately $1.39 per dollar invested, and protect more than 311,000 children.
- **Increase enforcement of the federal renovation, repair, and painting rule.** Ensuring that contractors comply with the Environmental Protection Agency (EPA) rule that requires lead-safe renovation, repair, and painting practices would protect about 211,000 children born in 2018 and provide future benefits of $4.5 billion, or about $3.10 per dollar spent.
- **Reduce air lead emissions.** Eliminating lead from airplane fuel would protect more than 226,000 children born in 2018 who live near airports, generate $262 million in future benefits, and remove roughly 450 tons of lead from the environment every year.
- **Clean up contaminated soil.**
- **Improve blood lead testing among children at high risk of exposure, and find and remediate the sources of their exposure.**
- **Ensure access to developmental and neuropsychological assessments and appropriate high-quality programs for children with elevated BLLs.**

Medical-Legal Partnerships

Medical-legal partnerships (MLPs) are multisector approaches to addressing legal issues, many of which are contributors to poor child health outcomes and disparities. Typically, a health care provider (or providers) partners with a legal aid entity to resolve a person or family's legal issues, including those related to housing, benefits, debt, or education. As discussed in more detail in *Communities in Action*, MLPs "play an important role in addressing the SDOH and are a relevant community-based solution for advancing health equity" (NASEM, 2017).

A systematic review conducted by Martinez et al. (2017) found that "researchers have established more findings regarding the capacity of MLPs to address legal outcomes than their capacity to address health outcomes" (p. 267); however, longer periods of study are likely needed to see improved health outcomes. Additional research is needed to identify child-specific health outcomes associated with MLPs.

MLPs represent a promising practice that has emerged over the last 10–15 years, and according to the National Center for Medical-Legal Partnerships,[13] there are now 333 MLPs in 46 states. See Box 6-5 for an example of a promising MLP between the Cincinnati Children's Hospital and the Legal Aid Society of Greater Cincinnati.

BOX 6-5
Cincinnati Child Health-Law Partnership (Child HeLP): Promising Model[a]

Cincinnati Child Health-Law Partnership (Child HeLP), which began in 2008, is an MLP between Cincinnati Children's Hospital and the Legal Aid Society of Greater Cincinnati. Child HeLP aims to support families experiencing legal and social issues that may be having detrimental effects on the health and well-being of their children. Physicians at Cincinnati Children's primary care clinics and social workers screen patient families for issues that may be affecting their children's health, such as food

[13] See https://medical-legalpartnership.org/.

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insecurity, inadequate housing, ACEs, and poor-quality education. Physicians receive training to better identify legal and social issues, which has been shown to successfully increase their comfort level with and knowledge of the SDOH as well as their familiarity with available community resources (Klein et al., 2011). Families identified as experiencing such issues are referred to Child HeLP to receive legal advice and assistance from the Legal Aid Society, which transmits information back to the family's provider to maintain open communication between the family's medical and legal teams (Cincinnati Children's, n.d.).

Child HeLP is a part of a more comprehensive, multisector approach to improving child health and well-being called the All Children Thrive Learning Network. The network includes a number of partners from the medical, public health, social services, legal, and education sectors as well as local agencies and community and faith-based organizations. Partnerships among these myriad stakeholders focus on four specific areas to improve child health and well-being: providing community-connected primary care and behavioral health services, decreasing preterm birth and infant mortality, strengthening neighborhood social influences, and improving access and quality of education to improve third-grade reading level outcomes (Cincinnati Children's, n.d.). Since 2008, there have been 6,600 referrals to the program, with 12,000 children and 6,100 adults helped.[b] One of Child HeLP's innovations is its practice of merging data sources and using GIS mapping to identify housing units with medium to high rates of housing code violations and high rates of pediatric asthma (Beck et al., 2014). Outreach to landlords and legal action has resulted in mitigation of asthma triggers (Beck et al., 2012).

Child HeLP has helped families improve the health and well-being of their children by addressing many issues related to the SDOH (Klein et al., 2013; Murphy et al., 2015; Sandel et al., 2010; Tyler, 2012). Families have received assistance in obtaining SNAP and WIC benefits, health insurance coverage, transportation to jobs, day care for young children, enrollment in school and special education services, and adequate housing to prevent homelessness. The program has also helped to prevent child maltreatment by resolving child custody disputes and obtaining relief for parents experiencing domestic violence (Cincinnati Children's, n.d.).

Dr. Kahn from Cincinnati Children's Hospital was quoted in a *New York Times opinion* piece as saying, "So much of child health is the result of poor social and physical living conditions for kids—food on the table, shelter, quality education. So much of what we do in pediatrics is driven by these broader well-being issues for the family. We do much better when we partner with groups that have that as a mission" (Rosenberg, 2014).

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

[b] Robert S. Kahn, Co-Director of the Cincinnati Child-Health Law Partnership, presented to the committee at its public information gathering session in August 2018. Presentation slides are available at: http://nationalacademies.org/hmd/~/media/Files/Agendas/Activity%20Files/Children/Prenatal-Early%20Childhood%20Development/2018-AUG-06/2-2%20Kahn%20-%20updated.pdf (accessed April 17, 2019).

Promising Tools

The following section describes tools that are available to communities and show promise in addressing poor child health outcomes and disparities through solutions to improve the affordability, quality, and stability of housing.

ChangeLab Solutions and Abt Associates have created a comprehensive and detailed toolkit that describes policies and programs to help preserve, protect, and expand the number of affordable rental units in neighborhoods where demand for housing is rising (Allbee et al., 2015). The authors state that "to ensure that people of all incomes, races, and ethnicities can continue to afford housing in neighborhoods experiencing rising rents, most communities will require a multifaceted strategy," one that includes a combination of the policies and programs described. Grouped into six areas of focus, these policies and programs are summarized in Table 6-2.

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TABLE 6-2: A Toolkit of Policies and Programs to Preserve, Protect, and Expand Affordable Housing

Area of Focus	Purpose	Policies and Programs
Preservation	Preserve the affordability of housing where low- and moderate- income renters already live	• Right of first refusal • Property tax incentives • Moving properties into subsidy programs • Preserving public housing through RAD*
Protection	Protect residents from the effects of rising rents or condo conversions by helping to reduce the risk of displacement or by helping them relocate to new units if necessary	• Good cause conviction policies • Condominium conversion protections • Rent stabilization
Inclusion	Ensure a share of new development is affordable to low- and moderate-income households	• Mandatory inclusionary zoning • Density bonuses and other voluntary inclusionary policies
Revenue Generation	Generate funding for affordable housing in neighborhoods experiencing rising rents and home prices by leveraging the development activity and economic growth associated with new development or redevelopment	• Tax increment financing • Linkage fees • Housing trust funds
Incentives	Offer a range of incentives to stimulate development of affordable housing in targeted areas	• Targeting of federal, state, and local housing resources • Local and state tax incentives • Parking incentives • Expedited permitting • Impact fees • Transfers of development rights
Property Acquisition	Gain control of desirable sites for development or redevelopment at affordable prices	• Using publicly owned land • Property acquisition funds

SOURCE: Allbee et al., 2015
*Rental Assistance Demonstration (RAD) is a federal program that "converts public housing subsidies into a form that can be used as the basis for securing private financing and can be combined more easily with other subsidies."

Another promising tool is the National Healthy Housing Standard,[14] which "provides health-based provisions to fill gaps where no property maintenance policy exists" and is "a complement to the International Property Maintenance Code and other policies already in use by local and state governments and federal agencies for the upkeep of existing homes" (National Center for Healthy Housing, 2014). The resource details minimum standards for healthy and safe homes, with information on each provision's public health rationale and further references and resources. Since the resource's release, the National Center for Healthy Housing also developed

[14] The National Healthy Housing Standard is available at https://nchh.org/resource/national-healthy-housing-standard-full-document/ (accessed March 21, 2019).

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an implementation tool to aid the adoption of the standard (National Center for Healthy Housing, 2017).[15]

Summary

Based on the evidence discussed in this chapter and presented in Chapter 3 on the effects of housing affordability and quality on health and developmental outcomes, the committee has reached the following conclusions about housing needs.

> *Conclusion 6-6: Healthy early development cannot occur without safe and stable housing. Lack of affordable housing and environmental hazards in housing disrupt healthy childhood development and parent/caregiver well-being. Children require affordable, quality, and stable living conditions to ensure that they can develop to their full potential.*
>
> *Conclusion 6-7: Housing affordability and quality is an acute problem that disproportionately impacts people of color and contributes to health disparities among children. Over half of black and Hispanic renters live in unaffordable housing, and health issues related to poor-quality housing, such as elevated blood lead levels and asthma, are more prevalent among these renters.*
>
> *Conclusion 6-8: Current federal housing programs are not adequately funded, and there are not enough safe, affordable housing units in high-opportunity areas. Additional funding for programs such as housing vouchers can move families out of poverty and allow families to reallocate money for other basic needs that support child health and development. Incentives and/or regulations, along with enhanced programming, can increase the supply of affordable housing.*

Given the evidence on the impact of housing for health and healthy child development, the committee recommends:

> **Recommendation 6-3: The U.S. Department of Housing and Urban Development, states, and local, territorial, and tribal public housing authorities should increase the supply of high-quality affordable housing that is available to families, especially those with young children.**

Increasing the supply of high-quality affordable housing will likely require additional federal funding to HUD and commitment from state and other local governments, as well as additional incentives or regulations to promote the development of new housing units.

> **Recommendation 6-4: The Secretary of the U.S. Department of Health and Human Services, in collaboration with the U.S. Department of Housing and Urban Development and other relevant agencies, should lead the development of a comprehensive plan to ensure access to stable, affordable, and safe housing in the**

[15] The National Healthy Housing Standard implementation tool is available at https://nchh.org/resource/national-healthy-housing-standard-implementation-tool (accessed March 21, 2019).

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prenatal through early childhood period. This strategy should particularly focus on priority populations who are disproportionately impacted by housing challenges and experience poor health outcomes.

Additional collaborators for this project include the agencies and organizations that are part of the United States Interagency Council on Homelessness.[16] The plan could include cross-sectoral initiatives that draw on resources from the health sector, such as a joint voucher program between HUD and HHS, MLPs to address housing problems within the clinical setting, and local investments in civil legal services and eviction prevention programs to help families stay in their homes during a short-term economic crisis.

Recommendation 6-5: The Center for Medicare & Medicaid Innovation should partner with states to test new Medicaid payment models that engage providers and other community organizations in addressing housing safety concerns, especially focused on young children. These demonstrations should evaluate impact on health, health disparities, and total cost of care.

Recognizing that reducing child poverty is a critically important, foundational strategy for improving child health outcomes and reducing health inequity, the committee recommends three key ways to ensure that families have the resources needed to meet children's basic needs. Founded in the review of the evidence of health impacts and the committee's expertise, the recommendation is based on increased resources for access to food and stable housing and income support in the form of a child allowance:

Recommendation 6-6: Federal, state, tribal, and territorial policy makers should address the critical gaps between family resources and family needs through a combination of benefits that have the best evidence of advancing health equity, such as increased Supplemental Nutrition Assistance Program benefits, increased housing assistance, and a basic income allowance for young children.

The costs of providing more resources available to young families by increasing SNAP benefits, housing assistance, or a child allowance depend on many parameters, including the size of the benefit per family and how many families are eligible and take up the benefit. Changes in these benefits may induce changes in parent employment, earnings, and receipt of other public benefits, affecting the total cost to the government. Full-scale analysis of the cost and behavioral changes of these proposals was beyond this committee's scope; however, the work of the National Aca Committee on Reducing Childhood Poverty does provide some illustrative examples of cost estimates. Using the Urban Institute's TRIM3 simulation model, the total change in annual government spending was estimated for a range of policy options (see the full report for the policy details and model assumptions). The change in government spending of increasing SNAP benefits by 20 percent was estimated at $26,414 million, including

[16] The collaborators in this council are HHS, Department of Education, Department of Labor, HUD, Veterans Affairs, Department of Agriculture, Department of Commerce, Department of Defense, Department of Energy, Department of Homeland Security, Department of Interior, Department of Justice, Department of Transportation, Corporation for National and Community Service, General Services Administration, Office of Management and Budget, Social Security Association, United States Postal Service, and the White House Faith and Opportunity Initiative. See https://www.usich.gov for more information (accessed June 15, 2019).

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employment and earnings adjustments (p. F-73). The cost of increasing housing vouchers by 50 percent was similar ($24,134 million) (p. F-78). In contrast, the estimate for a child allowance of $2,000 per year for families with children age 0 to 16 was $32,904 million (Page F-90).[17] A child allowance for families with children under age 6 would be considerably less costly. The actual costs of expanding SNAP or housing programs or creating a child allowance will depend on the size and scope of the program. Future research is needed to investigate how to implement these programs cost effectively and so that they will yield the greatest health benefits.

Increasing resources available for families with young children to meet their basic needs is of utmost urgency given the science-based connections between health and income for healthy development of young children and their later health and educational outcomes. While the National Academies Committee on Reducing Child Poverty recommended other important strategies, such as expanding EITC or CDCTC and increasing minimum wage, this committee's recommendation focuses on strategies that are likely to have particularly important impacts on health outcomes for young children. These include a child allowance targeted at families with young children not yet in K–12 school and increasing SNAP benefit levels and housing assistance. A child allowance would fill in some of the gaps in the current safety net and particularly benefit the lowest-income children and those most at risk of poor health outcomes. The key advantage of a child allowance (over, for example, tax credits) is that funds are available to families on an ongoing, monthly basis rather than once per year. In addition, under the current structure of the child and working-family tax credits, the lowest-income families receive few benefits. Children whose parents are in unstable employment or not employed suffer the short- and long-term health consequences of living in poverty. Reducing health disparities requires reaching these children during their earliest years, regardless of parental employment. Increased SNAP benefits and housing allowances would address current inadequacies in both of these programs and provide targeted support for the critical food and housing needs of young children. Paid parental leave would also recognize and support the special needs of the earliest years, in which parental time and attention are critically important for children's health and development.

Another way to increase resources to families with young children would be to provide more funding to subsidize child care. For example, the National Academies Committee on Reducing Child Poverty included expansion of child care subsidies as one of their options for reducing childhood poverty, and guaranteeing access to subsidies for families below 150 percent FPL would significantly increase resources available to low-income, working families. As noted in Chapter 3, only 15% of eligible families received CCDBG subsidies in 2012 (Walker and Matthews, 2017). However, these subsidies would flow only to families who meet the work requirements and use an eligible care provider, so these resources would be less likely to help families in deep poverty or those with intermittent employment. While child care subsidies are an important work support for low-income families, the committee focused on programs with more evidence of positive effects on children's health and those not dependent on parental employment. As noted earlier in this chapter, expansions of the work-oriented safety net programs may have unintended negative consequences for child health if parent employment results in lower rates of breastfeeding or disruptions to the attachment between infant and caregiver. In order to provide supports for all children, it is important to supplement work-support type programs with basic support for families with young children that is not tied to employment.

[17] It is important to note that these estimates are net costs to the government, and they are not suitable for comparison to other cost estimates cited in this chapter.

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In addition to material resources to meet their basic needs, children need a nurturing and healthy environment, responsive and sensitive caregiving free of maltreatment, and opportunities to develop the socio-emotional and cognitive skills to be healthy and resilient. Families with adequate resources may be better able to provide these important determinants of health, yet income alone may not be sufficient to ensure positive health outcomes. Thus, while addressing families' basic material needs is a critically important strategy for improving health outcomes of young children, strategies that focus only on reducing child poverty are unlikely to be sufficient to eliminate poor health outcomes and health disparities in early childhood. Specific recommendations with regard to the child's environment and the importance of relationships are discussed in other chapters of this report. The importance of confronting and eliminating structural racism in order to eliminate health disparities is discussed in Chapters 3 and 8.

NEIGHBORHOOD CONDITIONS

Neighborhoods play a critical role in the health, well-being, and development of children. Those that provide families with access to high-quality education, employment opportunities, safety, high-quality health care, and other essential services are essential to supporting children's healthy development. Persistent and increasing economic inequality, however, has contributed to neighborhoods facing increased economic segregation and concentrated poverty whose conditions and lack of resources can negatively affect healthy development and limit opportunities for children to flourish.

Neighborhood conditions can be defined broadly as the overall community context that is shaped by the natural and built environment (e.g., housing structures and conditions, availability of sidewalks and open/green space, presence of vacant lots, environmental exposures) and the availability or lack of health-promoting goods and services (e.g., access to healthy foods, affordable and safe transportation), in addition to the social environment (e.g., community cohesion, residential segregation, violence). This section addresses neighborhood conditions broadly; other sections in this chapter discuss the evidence related to specific neighborhood-level factors, such as housing, environmental exposures, food security, and economic security.

Concentrated Disadvantage

Residential segregation persists in the United States and has contributed to neighborhoods experiencing concentrated disadvantage where intergenerational cycles of poverty and adversity can derail children's healthy development. Data from the 2010–2014 American Community Survey shows that residential segregation by race and ethnicity has begun to decline slightly (Frey, 2015); however, economic residential segregation has increased (Rusk, 2017; Taylor and Fry, 2012).

Residential segregation, by race and ethnicity and by SES and income, increases racial health disparities and has pronounced effects on children's health outcomes. Its effects include socioeconomic disadvantages, such as limited opportunities for high-quality education and employment; increased exposure to crime, violence, and environmental toxicants; and limited access to transportation options, healthy food options, and health care services, which in turn leads to disparities in the quality of services and treatment (Acevedo-Garcia et al., 2008). For children in particular, segregation limits their access to high-quality schools and after-school

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programs and to neighborhoods in which they can play and exercise safely (Acevedo-Garcia et al., 2007, 2008).

Although policies that promote and increase segregation (including Jim Crow laws, redlining, and discriminatory banking and foreclosure practices) no longer exist overtly, their effects have persisted and remain widespread in communities experiencing intergenerational poverty and trauma. Segregation limits socioeconomic resources available to those living in concentrated poverty and has resulted in disparities in rates of disease, availability of high-quality health care providers, and opportunities to engage in health-promoting behaviors. Its effects have also included access to a higher density of alcohol, tobacco, and fast food outlets, increased risk of exposure to environmental hazards and toxicants, and higher risk of exposure to violence. For more information on the historical and contemporary effects of segregation policies, see Chapter 3 of NASEM, 2017 or Reskin, 2012.

Exposure to Violence

Exposure to neighborhood crime and violence can significantly affect children's cognitive (Burdick-Will, 2018; Sharkey, 2010), emotional and behavioral (Kim et al., 2014; McCoy et al., 2016; Sharkey et al., 2012), and health outcomes. Improving neighborhood conditions to diminish exposure to crime and violence plays an important role in reducing children's biological stress and improving their health outcomes. For more on exposure to violence as an ACE, see Chapter 3.

In addition to adversity faced at the individual level, repeated exposure to crime and violence can contribute to community trauma. A report by the Prevention Institute describes that "community trauma is not just the aggregate of individuals in a neighborhood who have experienced trauma from exposures to violence"; rather, symptoms of community trauma "are present in the closed-cultural environment, the physical/built environment, and the economic environment" (Pinderhughes et al., 2015). The report describes several community strategies to implement within each of these three environments to mitigate the negative effects of community trauma, build more resilient communities, and improve health and well-being outcomes (see Figure 6-3). The authors note that across all environments, approaches that are most effective will "build on indigenous knowledge, expertise, and leadership to produce strategies that are culturally relevant and appropriate" (Pinderhughes et al., 2015, p. 5).

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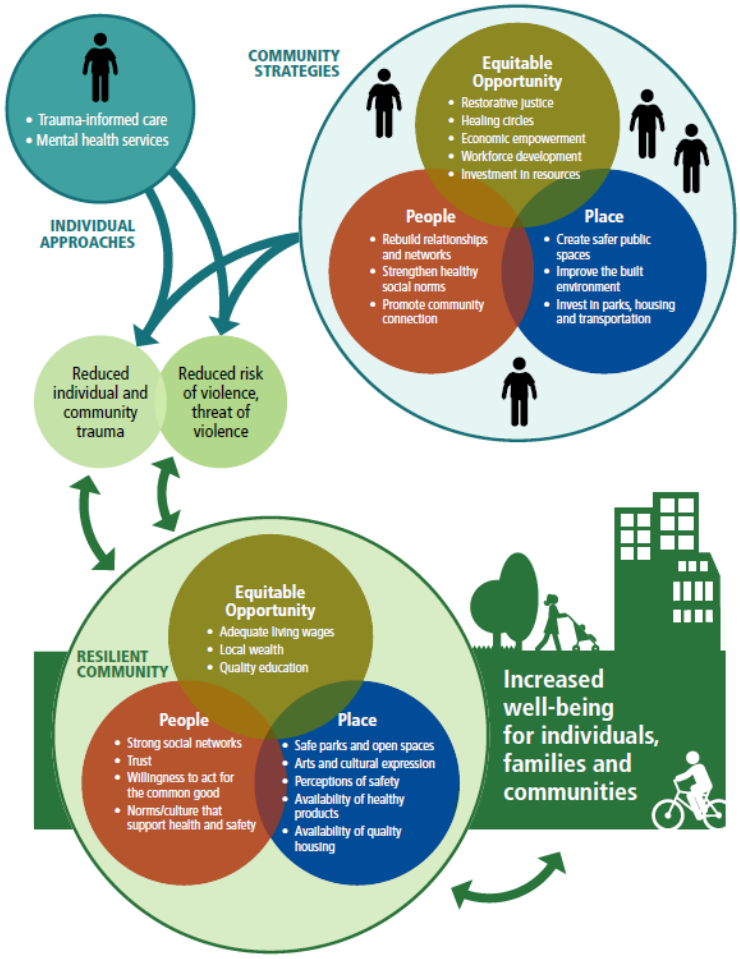

Figure 6-3: Promoting community resilience: from trauma to well-being
SOURCE: Pinderhughes et al., 2015
NOTE: Figure provided by Prevention Institute, www.preventioninstitute.org.

Improving Neighborhood Conditions

HUD's Moving to Opportunity (MTO) study provides clear evidence that better neighborhood conditions are strongly associated with improved health and development outcomes (Kling et al., 2007; Ludwig et al., 2013). MTO randomized assistance designed to help low-income families move to less impoverished neighborhoods; after 10–15 years, adults chosen to receive that assistance had lower prevalence of obesity and diabetes and better subjective health outcomes than adults who were selected not to receive it (Ludwig et al., 2011). Despite these positive findings, the MTO experiment has also yielded mixed results. Early MTO studies resulted in findings that ran counter to the hypothesized outcomes; for example, Sanbonmatsu et al (2006) found no evidence of improvement in academic outcomes for any age group four to seven years after randomization. As noted previously in the chapter, Chetty et al. (2016) found strong evidence to suggest that children who moved to lower-poverty neighborhoods when they were young (i.e., under 13 years of age) were more likely to experience positive outcomes later in life, such as attending college and earning higher incomes. However, the study also showed contrary outcomes for those who moved when they were older. Other research indicates that

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while girls in the MTO experiment exhibited positive mental health outcomes (e.g., decreased rates of major depression and conduct disorder), boys experienced the opposite (e.g., increased rates of PTSD, major depression, and conduct disorder) (Kessler, 2014). In terms of long-term outcomes (i.e., 10 – 15 years), Ludwig et al (2013) concluded that moving to lower-poverty neighborhoods during childhood leads to improvements in adult physical and mental health, but not economic self-sufficiency.

MTO established that places matter for health but not what it is about them that matters most: it was not designed to tease apart which aspects of places drove those health improvements, nor did it specifically address prenatal or birth outcomes. Other studies have attempted to further illuminate which aspects of a neighborhood—for example, better economic environments, built environment features that promote healthy lifestyles, or improved social cohesion—might represent the most important levers for improving health and birth outcomes. Other than MTO, however, randomized studies on place and its impacts on health are extraordinarily rare, so the evidence is almost entirely limited to observational studies.

The Washington Center for Equitable Growth describes two approaches to addressing segregation within neighborhoods: 1) invest in neighborhoods to make the consequences of segregation less severe and 2) reduce the level of segregation in neighborhoods directly (Sharkley, 2016).

Approaches to increase and shift investments to low-income communities include

1. Providing work supports (including supplemental wages and guaranteed public service jobs, as in the New Hope program, and services to residents of public housing developments and rent incentives to encourage work, as in the Job Plus program) for individuals and families in high-poverty communities;
2. Investing in evidence-based programs for young people, such as those that include tutoring or mentoring, sports activities, cognitive behavioral therapy, and summer jobs, which randomized trials have shown lead to increases in academic achievement and decreases in involvement in crime and violence;
3. Identifying and/or establishing a single institution in every low-income neighborhood "that takes ownership over the community and takes responsibility for all the residents within it, […] so that everyone within that neighborhood knows that there is going to be an institution serving them for the long haul and will have resources sufficient to bring about long-term change" (Sharkley, 2016).

Addressing the lack of affordable housing is crucial to reducing segregation in neighborhoods directly (Sharkley, 2016). To improve housing affordability, the Washington Center for Economic Growth provides the following approaches:

1. Expanding the supply of housing vouchers. (See the following section on Housing Affordability for more on housing vouchers.)
2. Providing support to allow families to access opportunity neighborhoods.
3. Providing incentives and regulations to preserve and expand affordable housing in exclusive markets.
4. Establishing a long-range mobility bank (Sharkley, 2016).

In developing approaches to enable families to access opportunity neighborhoods, an important consideration is ensuring that adequate supports are available to families to buffer the negative effects of stress associated with moving to other communities. For an example of a

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promising initiative that aims to support and strengthen families by providing community-based supportive services, see Box 6-6.

BOX 6-6:
SHIELDS for Families: Promising Model[a]

SHIELDS, started in 1991, provides multipronged community-based services, operating 34 different programs. SHIELDS' goals are to promote family reunification and support families remaining intact in the community, strengthen families by providing comprehensive and collaborative services, improve the general well-being of families through comprehensive health programs and preventative social services, and encourage self-sufficiency and economic independence by assisting individuals with obtaining employment and advancing education (SHIELDS For Families, 2011). SHIELDS consists of five divisions: child welfare, children and youth, mental health, substance abuse, and supportive services.

SHIELDS supports high-risk families in South Los Angeles, with several programs specifically for pregnant and parenting women and their children ages 0–5. SHIELDS' wraparound programs address employment, health services, housing, income and wealth, public safety, the social environment, and transportation. SHIELDS engages multiple sectors and is funded by private foundations, LA County, and federal grants. For wraparound support programs, SHIELDS has partnered with private hospitals, the Los Angeles Department of Children and Family Services, the Los Angeles Department of Probation, the Los Angeles Homeless Services Authority, Special Services for Groups, California's Post-release Community Supervision jurisdiction, the California DOJ, training and vocational schools, and many more organizations and agencies.[b]

SHIELDS reports that 4,308 families were linked to over 15,000 services through their family resource center; 362 families received mental health treatment; 78 percent of families successfully completed treatment; and 69 percent of families reunited with children (SHIELDS For Families, 2011).

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.Caregivers and representatives from SHIELDS for Families presented to the Committee at its public information gathering session in October 2018.
[b] For more information, see https://www.shieldsforfamilies.org (accessed April 17, 2019).

ENVIRONMENTAL EXPOSURES AND EXPOSURE TO TOXICANTS

Each year, Americans are exposed to chemicals in the environment at increasingly greater levels. EPA tracks new chemical substances in its Toxic Substances Control Act Chemical Substance Inventory, which is updated every 6 months (EPA, n.d.-3). Currently, more than 85,000 chemical substances are listed in the TSCA inventory, with companies introducing about 700 new substances each year (Tollefson, 2016; EPA, n.d.-1). As described in Chapter 3, many of these substances are related to poor health outcomes for mothers, infants, and children. In its review of the literature, the committee found that children (from fetal development through early childhood) are at greater risk than adults from adverse health effects of environmental exposures due to their smaller size and proportionally large intake of food, air, and water to body weight and are subject to rapid developmental processes that may be influenced and disrupted by chemicals and toxicants. Parents' exposures to environmental toxicants in the preconception phase also present a risk to early child development. Further, as a result of exposure to toxicants, children are more likely than adults to suffer from developmental problems, chronic conditions, and death. The committee also concludes that:

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> *Conclusion 6-9: Not all households experience the same level of risk of exposure to harmful environmental toxicants or pollutants. Poverty, substandard and/or unstable housing, race and ethnicity, and proximity to known sources of pollutants heighten pregnant women and children's risk of exposure and poor health and developmental outcomes.*

The following section provides a review of the limited research currently available that specifically examines policies and practices to reduce the risk of environmental exposures among preconception, prenatal, and postnatal populations.

Reducing the Risk of Environmental Exposures in the Home and in Child Care Settings

The Center for Medicaid and CHIP Services (CMCS) serves millions of pregnant women, children, and families by providing health care access to low-income families and families who would not otherwise qualify for health insurance. CMCS also establishes reimbursement criteria for these health services, including environmental exposure screening, testing, and case management. How these criteria are implemented varies by state, though some aspects of coverage are mandatory. For example, blood lead screening is mandated for Medicaid recipients at 12 and 24 months of age. However, states vary in how this mandate is implemented (CMS, n.d.). Some states provide these screenings only to Medicaid-enrolled children, while others offer universal screenings; one state, Arizona, has chosen to implement a targeted screening approach (Arizona Department of Health Services, 2017; CMS, 2016), while still others do not currently have a lead screening program (Dickman 2017; NASHP, 2018). Moreover, some states require reporting the results to state health departments or the state lead registry, which may result in follow-up care or case management. Other states do not have a registry or other tracking system, and still others do not provide any form of follow-up to children with elevated BLLs.

Medicaid can cover home investigations of lead exposures, case management for exposed children, and support to states to implement education, screening, and outreach efforts to areas at high risk of lead exposure (CMS, 2017), though it is unclear how these programs are applied at local levels. In addition, greater collaboration with tribal nations is needed to improve state screening and registry efforts (President's Task Force on Environmental Health Risks and Safety Risks to Children, 2018), though some federally funded research studies have led to tribal law and policy updates to improve lead screening among children (Petersen et al., 2007). No other screening programs aimed at environmental exposures receive Medicaid support; lead exposure is an exception because of the estimated magnitude, frequency, and duration of exposure, the significance of the health impacts, and the potential for prevention (NASHP, 2017). However, several states have asserted regulatory authority over other environmental toxicants, including bisphenol a (BPA), cleaning agents in schools, and flame retardants (NCSL, 2017).

Child care settings represent another source of potential environmental exposures for children. With more than 13 million preschoolers in child care every day, including 6 million infants and toddlers (Amoah et al., 2016) (see Chapter 3 for more information on the number of children enrolled in various ECE programs), addressing potential environmental exposures in child care settings could reduce the risk of related health issues for a large proportion of children under the age of six. Taking steps to address potential exposures can present challenges, as child care facilities may be located within individual homes, community centers, and office buildings,

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which may not be consistently assessed or regulated to reduce the risk of environmental exposures. Moreover, licensing guidelines vary, with limited requirements addressing child care provider training on mitigation of early childhood exposures. Reviewing existing state policies on the regulation of environmental exposures in child care facilities offers policy makers an opportunity to compare policy approaches and consider nonregulatory approaches to effect change (Environmental Law Institute, 2015).

In addition to reviewing existing policies, evaluating the effectiveness of training programs can also inform decision making around environmental exposure risk reduction. In a study of 60 child care centers in Washington, D.C., an environmental risk assessment training was developed and tested that covered the following areas: (a) air quality, (b) arsenic, (c) asbestos, (d) built environment, (e) chemicals in art supplies, (f) lead, (g) mercury, (h) mold, (i) noise pollution, (j) pesticides and integrated pesticide management, (k) physical education and nutrition, (l) plastics, (m) radon, and (n) safe cleaning alternatives (Amoah et al., 2016). More than 580 child care workers received the training and pre-/post-assessments, and knowledge scores increased by about 20 percent, on average, between the two assessments. Nearly 70 percent of participating child care centers reduced their environmental risk assessment scores. This training is an important approach to reducing the environmental risk for children receiving child care outside the home.

Educational approaches are also effective at the level of the individual household. In a meta-analysis of seven community health worker–delivered interventions focused on reducing asthma symptoms, most of these significantly decreased asthma symptoms, lessened daytime activity limitations, and reduced emergency and urgent care use (Postma et al., 2009). Programs that included higher intensity and frequency of home visiting reported the most positive health outcomes. Similarly, in a systematic review of 20 studies focused on reducing the risk of asthma morbidity among children, most of them were effective in significantly reducing the number of days with asthma symptoms, school days missed, and asthma acute care visits (Crocker et al., 2011). The review examined studies expressly focused on informing multitrigger, multicomponent, home-based environmental interventions, which addresses the complex nature of reducing asthma risk within the home due to the many potential sources of risk. A randomized controlled trial (RCT) aimed at educating pregnant women on reducing exposure to hazardous air pollutants resulted in a significant increase in preventive behaviors (Marzieh et al., 2017), demonstrating that interventions aimed at parents can also be effective in reducing the risk of toxic exposures for infants and children.

Reducing the Risk of Environmental Exposures in the Community

In a 2018 study, researchers linked 2011 National Emissions Inventory data with block groups from the 2009–2013 American Community Survey data (Mikati et al., 2018). For particulate matter of 2.5 micrometers or less in diameter ($PM_{2.5}$), Americans living in poverty and people of color experienced significantly higher levels of exposure compared to white Americans. Racial disparities, particularly for black Americans, were greater than for those living in poverty alone (Mikati et al., 2018). In a subsequent 2019 study, researchers found that, on average, non-Hispanic whites experience a “pollution advantage” (Tessum et al., 2019). In other words, they experience 17 percent less air pollution exposure, comparing the amount to which they are exposed and the amount that they are responsible for due to their consumption levels. In comparison, black and Hispanic Americans, on average, experience 56 percent and 63 percent excess exposure, respectively, relative to the exposure caused by their consumption

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(Tessum et al., 2019). While $PM_{2.5}$ exposure for all groups fell about 50 percent between 2002 to 2015, due to increased regulation and population density reductions near polluted areas, the inequities remain above and beyond this reduction. Scientists suggest addressing these inequities will likely require multilevel approaches, including reducing consumption levels, improving manufacturing and other processes to minimize pollution, and implementing more comprehensive measures, such as evaluating plans for construction and urban development with the purpose of reducing dependence on automobile transportation to manage and lessen current inequities, along with meaningfully involving communities in assessment processes to effect broader policy change (Schulz et al., 2016).

For example, researchers collaborated with the Rural Empowerment Association for Community Help in North Carolina to train middle schoolers to assess asthma indicators, lung function, and air pollution (Guidry et al., 2014). School administrators and students reported positive perspectives on the project, which offers an example for community-based research aimed to improve air pollution in rural settings (Guidry et al., 2014). In another example, the Community Action to Fight Asthma Initiative in California used an environmental justice approach to reduce risk factors for asthma in school-aged children. Statewide coalitions, which included local residents and technical assistance experts, supported policies to reduce children's exposures to environmental triggers. Technical assistance, community involvement, and multilevel strategies led to policy change that addressed reduction of environmental inequities (Kreger et al., 2011).

The BREATHE Project, based in Barcelona, which aimed to assess traffic-related air pollutant exposure among schoolchildren, is a promising example of using complex measures and modeling to reduce pollutant exposure (Rivas et al., 2018). Using advanced measurement and statistical techniques, researchers identified eight factors/sources of pollutants, including minerals, traffic, road dust, secondary sulfate and organics, secondary nitrate, sea spray, heavy oil combustion, metallurgy, and organic/textile/chalk, and were able to characterize the air quality within and near 39 urban schools. This novel approach allows for detailed mapping of exposure risk, which supports positive policy change through data-driven decision making.

Community Infrastructure

Sustainable infrastructure is defined as “systems that have the capacity to endure over a long period of time; enabling the human-built environment to thrive and providing an opportunity for human society to improve its quality of life, without compromising the integrity and availability of natural, economic, and social assets for future generations” (Hendricks et al., 2018, p. 2). Properly managing infrastructure (i.e., replacing lead drinking water pipes, mitigating flood zones to reduce damage to homes and schools, and updating technology to ensure timely notification of emergent hazards), through the lens of assessing and maximizing sustainability, therefore has important downstream implications for public health. This is particularly the case with respect to ensuring that the necessary environment and resources are equitably available to support early childhood development and minimizing exposures to harms that arise from poor infrastructure, lack of sustainable practices, and the concomitant effects on the environment. Moreover, reducing these risks can lead to cost savings. For example, the annual U.S. cost of environment-related pediatric disease in 2008 was estimated to be $76.6 billion, or 3.5 percent of total health care costs (Trasande and Liu, 2011). These costs are limited to pediatric diseases due to environmental exposures. Including costs of pediatric diseases resulting from severe climate conditions’ impact on aged or damaged infrastructure, such as

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heat-related deaths and illnesses and increased exposures to lead and other toxicants, mold and poor water quality from flooded and damaged dwellings, and poor air quality from increased forest fire smoke and smog, could support prioritization and decision making to best address failing infrastructure.

The United States currently faces a critical infrastructure crisis (Maxwell et al., 2018), a result of decades of deferrals at every governmental level and lack of effective accountability on the side of private institutions to ensure essential infrastructure received necessary upgrades, including improvements that would minimize negative environmental impacts and ensure long-term sustainability (American Society of Civil Engineers, 2017). Using strategies such as the framework put forth by Koehler et al. (2018) for improving community health through better environment decision making could provide important approaches to mitigate future harms and reduce costs. In addition, existing models do not specifically examine risks to early childhood development and ensuing costs. Additional research and environmental assessments are needed to ensure harms are minimized, especially for populations who are disproportionately impacted by harmful environmental exposures. Investing in the rebuilding of the nation's infrastructure at every level—waste and water management, walkable streets and parks, cleanup of toxic waste sites, and so on—along with strengthening capacity for community engagement and civil rights actions around environmental justice, can provide a cross-sector approach that strengthens infrastructure in the United States, reducing potential harms to infants and children and ensuring lower medical costs for pediatric injury and disease associated with severe climate effects

The Role of Civil Rights

Civil rights strategies are an important yet underused tool to promote clean and healthy communities for developing children. This was acknowledged as a community-driven solution to promote health equity in the 2017 National Academies report *Communities in Action: Pathways to Health Equity*. The report asserts that "civil rights, health, and environmental justice laws and policies provide a framework to promote equal access to publicly funded resources and prohibit discrimination based on race, color, national origin, income, gender, disability, and other factors" (p. 351). Federal laws and civil rights legislation have historically been a codified source of rights to protect individuals and groups from harmful environmental exposures (see, for example, Title VI of the Civil Rights Act of 1964,[18] the Environmental Policy Act,[19] and the Clean Water Act[20]). In 1994, an executive order was issued on Federal Actions to Address Environmental Justice in Minority Populations and Low-Income Populations.[21] Civil rights laws and their enforcement not only aim to protect populations in at-risk contexts—thereby reducing disparities—but also allow for a crosscutting approach that can apply to many of the determinants of health and development discussed in this report (e.g., housing, environmental exposures, education). (See Hahn et al., 2018 for a brief summary of civil rights history and discussion on the relationship between civil rights laws and the determinants of health.) Specifically, civil rights strategies can be used to mitigate discriminatory burdens, lower or remove barriers to community participation in decision making, and improve access to health

[18] Title VI of the Civil Rights Act of 1964 (42 U.S.C. § 2000d to 2000d-7): "No person in the United States shall, on the ground of race, color, or national origin, be excluded from participation in, be denied the benefits of, or be subjected to discrimination under any program or activity receiving Federal financial assistance" (GPO, 2010).
[19] 42 U.S.C. § 4321 et seq.
[20] 33 U.S.C. §1251 et seq.
[21] E.O. 12898. 59 FR 7629; February 16, 1994.

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and environmental benefits that are foundational to safe and healthy communities (NASEM, 2017; USDA, 2012).

At the state level, California's EPA operates an Environmental Justice Task Force, which develops initiatives in communities where compliance and enforcement can help mitigate or stop the harmful effects of exposure to pollution. At the community level, there are tools that can be employed to ensure that these rights are protected to engender equitable outcomes for children and families. For example, public health, civil rights, and environmental justice practitioners use a five-step planning framework by which to assess proposed or current environmental policies or practices for their potential to harm or benefit communities (Environmental Justice Leadership Forum on Climate Change, 2016; NASEM, 2017; The City Project, 2016):[22]

1. Describe what is planned in terms that are understandable to the community,
2. Analyze the benefits and burdens on all people,
3. Analyze alternatives to what is being considered,
4. Include people of color, low-income people, and other stakeholders in every step in the decision-making process, and
5. Develop an implementation plan to distribute benefits and burdens fairly and avoid discrimination.

(See NASEM, 2017 for a more in-depth discussion about the planning process, in addition to a few examples of communities that have successfully applied the process.)

The *Communities in Action: Pathways to Health Equity* report (NASEM, 2017) also outlines some guidance for civil rights attorneys, public health professionals, community groups, public agencies, recipients of public funding, foundations, and other stakeholders to advance equity using civil rights tools (see Box 6-7 for a listing of these guiding strategies).

Finally, based on the committee's expertise and its review of civil rights legislation, literature, and past action to address equitable access to the health-promoting social determinants, the authoring committee of *Communities in Action: Pathways to Health Equity* reached the following conclusions in its report:

> Civil rights approaches have helped mitigate the negative impacts of many forms of social and health discrimination. Continuing this work is needed to overcome discrimination and the structural barriers that affect health.
>
> Using civil rights approaches in devising and implementing community solutions to promote health equity can guard against unjustified and unnecessary discriminatory impacts, as well as against intentional discrimination in programs that affect health. For example, those implementing community solutions can employ methods and data in ways that include full and fair participation by diverse communities (NASEM, 2017, p. 362).

Given the critical role of civil rights strategies in advancing health equity in communities, the report also included a recommendation to foundations and other funders to expand their community support beyond their traditional roles by supporting education, compliance, and enforcement related to civil rights law and other areas (see Recommendation 7-1 in NASEM,

[22] This planning framework is based on Title VI, Executive Order 12898, case law, and best practices by federal agencies (NASEM, 2017).

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2017 for the full recommendation). The report acknowledges the barriers associated with federal tax laws that may preclude foundations from supporting such activities, but it also suggests the use of general operating funds as opposed to program-specific funds (NASEM, 2017). The alignment between this report and *Communities in Action* with respect to health equity and the applicability of civil rights strategies for addressing environmental exposures in communities warrant serious consideration of these findings and the abovementioned recommendation for promoting healthy and equitable early development.

BOX 6-7
Guidance for Communities to Advance Equity Using Civil Rights Tools

- Communities and other stakeholders can work together on compliance and equity plans for programs or activities by recipients of public funding that use the civil rights framework by describing what is to be done, analyzing the impact on all communities, analyzing alternatives, including full and fair participation by diverse communities, and promoting health equity.
- Compliance and equity plans can be used to guard against unjustified and unnecessary discriminatory impacts, as well as against intentional discrimination, in health and wellness programs and activities.
- Communities, when appropriate, can work with civil rights attorneys to use problem-solving strategies, including coalition building, planning, data collection and analysis, media, negotiation, policy and legal advocacy out of court, and access to justice through the courts.
- Communities can work with attorneys and public health experts together to promote a better understanding of the civil rights dimension of the challenge of health disparities and to show how to address these civil rights concerns for their communities to ensure that civil rights laws against discrimination in health and other publicly funded programs and activities are strengthened and not rolled back.

SOURCE: Excerpted from NASEM, 2017

Leveraging Clinical Settings to Reduce Environmental Exposure Risk

Recent evidence confirms that preconception and prenatal exposures can adversely impact fetal development, which may result in long-lasting health effects (Grandjean et al., 2015). Training tools for health practitioners offer a potentially promising prevention approach to reduce harmful environmental exposures and the risk of future adverse health effects for prenatal and preconception patients (Sathyanarayana et al., 2012). In a national online survey of over 2,500 obstetricians, nearly 80 percent agreed that they can reduce patient exposures to environmental health hazards by counseling patients. However, half of respondents reported that they rarely take an environmental health history, and less than 20 percent reported routinely asking about environmental exposures commonly found for pregnant women in the United States. Moreover, only 1 in 15 reported any training on how to assess environmental exposures among patients (Stotland et al., 2014), demonstrating the need for additional supports for providers, particularly given the complexity of the assessment process. There are several online tools for providers to triangulate exposure risk, yet the sheer number of possible chemical exposures, the relatively limited data available on possible risks, and the expertise and time commitment needed to complete an assessment may pose significant challenges for primary care or specialty providers (Koehrn et al., 2017).

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Medical Education on Environmental Health

Education on the effects of preconception and prenatal exposure to environmental toxicants is largely insufficient in medical education and curricula (Gehle et al., 2011; Tinney et al., 2015). Pediatric practice, which has historically recognized the influence of environmental health exposures (Tinney et al., 2015), is an important setting to screen for exposure to environmental toxicants. However, there are also earlier opportunities to identify and ensure timely treatment for harmful exposures during the preconception and prenatal periods (Tinney et al., 2015).Many professional organizations have emphasized the urgency and importance of addressing exposure to environmental toxicants. In a joint committee opinion from the ACOG Committee on Health Care for Underserved Women and the American Society for Reproductive Medicine (ASRM) Practice Committee, ACOG and ASRM call for "timely action to identify and reduce exposure to toxic environmental agents while addressing the consequences of such exposure" (ACOG, 2013). The International Federation of Gynecology and Obstetrics (FIGO), a professional organization of obstetrical and gynecological associations from 125 countries that includes ACOG, states that "obstetricians, gynecologists, midwives, women's health nurse practitioners, nurses, and other health professionals should … make environment health part of health care" and that "exposure to toxic environmental chemicals during pregnancy and breastfeeding is ubiquitous and is a threat to healthy human reproduction" (Di Renzo et al., 2015). However, despite broad consensus on the importance of addressing exposure to environmental toxicants studies show that many obstetricians and gynecologists do not screen their patients for harmful environmental exposures (Grindler et al., 2018; Stotland et al., 2014).

While qualitative data suggest that pediatricians express an interest and need for more training on environmental medicine topics (e.g., environmental history taking) (Kilpatrick et al., 2002; Trasande et al., 2006), this type of education is not routinely included in pediatric curricula (Roberts and Gitterman, 2003). The National Environmental Education and Training Foundation and the Children's Environmental Health Network formed two working groups with expertise in medical and nursing education to explore opportunities to incorporate environmental health into pediatric education (McCurdy et al., 2004). After reviewing the transition from undergraduate student to professional status, the medical education working group recommended improving education about children's environmental health in medical school curricula, residence, training, and continuing medical education, in addition to expanding fellowship training in children's environmental health. The nursing working group similarly recommended enhancing content on children's environmental health at the undergraduate, graduate, and continuing nursing education levels. Both working groups identified leverage points (i.e., key organizations and stakeholders) that could facilitate these changes.

There are several existing resources and strategies to accelerate the integration of environmental health into existing medical education and curricula. A work group of faculty members and residents[23] formed at a meeting convened by the Mid-Atlantic Center for Children's Health and the Environment and The George Washington University's Department of Obstetrics and Gynecology identified several approaches to better incorporate environmental health into medical education, including

[23] Faculty members and residents who participated in the meeting represented 16 academic obstetrics-gynecology programs from 5 states (Delaware, Maryland, Pennsylvania, Virginia, and West Virginia) and DC.

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1. Integrating environmental health into basic science courses and the organ systems approach (Gehle et al., 2011) into medical school and residency curricula[24],
2. Implementing clinical training programs on environmental health, such as the model established by the University of California, San Francisco, Program on Reproductive Health and Environment (PRHE)[25],
3. Adding questions on environmental health on resident training and board-certifying exams[26] to ensure that programs will include environmental health in their curricula,
4. Training leaders in medical education and faculty members on reproductive environmental health through the Association of Professors of Gynecology and Obstetrics, and
5. Requiring continuing education in environmental health[27] to ensure that practicing obstetricians and gynecologists receive introductory education to environmental health (Tinney et al., 2015).

There are also resources to support providers in offering their patients counseling and health education on environmental exposures. Located across the United States and Canada, Pediatric Environmental Health Specialty Units (PEHSUs) are a national network of experts in reproductive and children's environmental health who are located at academic medical centers and provide medical information and advice and respond to requests for information on the prevention, diagnosis, management, and treatment of the effects of environmental exposures on children and adults of child-bearing age (PEHSU, n.d.). The AAP publishes a guide for pediatricians, Pediatric Environmental Health, which is meant to help identify, prevent, and treat pediatric environmental health problems (AAP Council on Environmental Health, 2012).

Similarly, the Organization of Teratology Information Specialists (OTIS) provides an online service known as MotherToBaby that provides evidence-based information on the safety of medications and other exposures during pregnancy and while breastfeeding to health care professionals and the general public (MotherToBaby, n.d.). Lastly, a collaborative initiative[28] known as SafetyNEST provides health care professionals and the general public with "accurate, evidence-based, and personalized information about the effects of toxic chemicals on prenatal and early childhood health" through an educational online platform (SafetyNEST, 2018).

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[24] The George Washington University's Department of Obstetrics and Gynecology has a medical school and residency curriculum on environmental health specifically for obstetricians and gynecologists (Tinney et al., 2015).
[25] PHRE includes rotations programs for obstetrics-gynecology clinical trainees and maternal-fetal medicine fellows as well as resources, such as educational materials for patients and environmental health history forms for clinical practice, for reproductive health professionals (Tinney et al., 2015).
[26] Such as those administered by the Council on Residency Education in Obstetrics and Gynecology, the American Board of Obstetrics and Gynecology, and the American Osteopathic Board of Obstetrics and Gynecology (Tinney et al., 2015).
[27] The Agency for Toxic Substances and Disease Registry has created online resources on environmental health, including case studies on environmental medicine, for health professionals completing continuing education (Gehle et al., 2011; Tinney et al., 2015).
[28] Partners include the UCSF Program on Reproductive Health and the Environment, the Icahn School of Medicine at Mount Sinai, the American Medical Women's Association, and Universidade Federal do Paraná in Brazil (SafetyNEST, 2018).

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Research, Monitoring, and Testing of Environmental Exposures

Existing conceptual frameworks fail to encapsulate the complexities inherent in early childhood development. The Public Health Eposome Conceptual Model attempts to capture the changing nature of these exposures, along with the life course considerations needed to accurately assess risk (Tulve, 2016). In particular, children may experience chemical exposures from their environments at every life stage throughout the life course. Children are exposed to chemicals in foods, water, air, and surfaces they touch, such as chemical residues on surfaces or in dust and soil. Revising existing conceptual models aims to improve evaluation and assessment of environmental exposures to improve accuracy, enhance the overall fit of statistical modeling, and ensure improved predictive power in research studies examining environmental exposures and effects on early childhood development.

In another approach, a conceptual framework relying on a holistic approach to maternal and child health research includes both psychosocial stressors and environmental hazards to better explain factors that influence poor health outcomes for populations that experience higher risks of environmental exposure. This approach makes it possible to include chronic stressors and environmental hazard exposures to better understand health inequities evident across income levels and racial and ethnic groups in the United States (Morello-Frosch et al., 2006).

Given the effect that environmental exposures have on some child mental health problems, researchers suggest "broadening outcomes to include dimensional measures of autism spectrum disorders (ASDs), attention deficit hyperactivity disorder (ADHD), and child learning capacity, as well as direct assessment of brain function" (Rauh and Margolis, 2016, p. 1). Longitudinal studies examining these outcomes may inform how key exposures result in child mental health problems, which may better address or prevent these issues (Rauh and Margolis, 2016).

Existing Regulatory Approaches

The federal government plays an essential role in enhancing the resources and technical assistance available to states, tribes, and local agencies through programs, policy development, and implementation by the Centers for Disease Control and Prevention (CDC), EPA, Food and Drug Administration (FDA), U.S. Consumer Product Safety Commission, and other relevant rule- and policy-making entities focusing on environmental exposures and children's health. In addition, many of these agencies collaborate on joint initiatives to address environmental exposures among American children and families. For example, CDC monitors environmental chemicals in children's blood, reports on trends over time, and establishes key recommendations for safe levels among children (CDC, 2019a). CDC also provides training and other resources to state, local, and tribal governments to support environmental health services, including tools focused on ensuring food safety and safe water (CDC, 2019b).

EPA also provides critical resources and supports to states, local, and tribal governments to reduce the risk of environmental exposures among children and to mitigate, if necessary, environments with high levels of contaminants. For example, through the *America's Children and the Environment* report, EPA reports on environmental health data, specifically concerning children, and establishes indicators to minimize potential adverse health outcomes related to environmental exposures. Areas of focus include criteria for air pollutants, drinking water contaminants, lead, mercury, cotinine, perfluorochemicals, phthalates, BPA, perchlorate, and certain health outcomes, such as respiratory diseases, childhood cancer, and neurodevelopmental

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disorders (EPA, n.d.-2). However, as EPA has noted, several scientific challenges exist with regard to children's environmental health, including (1) the dispersion of data and information across systems that are difficult to access, (2) the need to assess environmental health from a systems perspective, (3) the need for new and more complex methods and models to evaluate risks specific to the early life stage of development and how exposures could result in health outcomes in later life stages, and (4) the need for translational research to better support community action and decision making (EPA, 2015). FDA regulates products and ensures the safety of food, including infant formula, the levels of BPA in food, and ingredients in cosmetics. For example, in 2014, the FDA BPA Joint Emerging Science Working Group reviewed the scientific literature on BPA effects on humans and found existing margins of safety to be adequate for exposure to BPA due to food contact (e.g., BPA exposures from coatings used in food containers) (Bisphenol A (BPA): Use in food contact application, 2018). Similarly, the U.S. Consumer Product Safety Commission regulates products that are manufactured for children and that may be used by children. For example, the CPSC limits the levels of phthalates in toys (GAO, 2017).

In addition to federal regulations, state environmental and public health agencies engage in efforts to prevent environmental exposures and to translate existing data and research findings into actionable policies and practices. For example, Washington State has enacted the Washington Children's Safe Products Act to limit levels of lead, cadmium, phthalates, and certain flame retardants in children's products (Department of Ecology, 2008). California (DTSC, n.d.) and Oregon (Oregon Health Authority, 2017) have implemented similar efforts to reduce the levels of toxic chemicals in consumer products. In all, there are at least 72 laws in 25 states addressing BPA, green chemistry, decabromodiphenyl ether, biomonitoring, and chemical safety (NCSL, 2014). However, the development and implementation of state regulations vary widely from state to state.

Frank R. Lautenberg Chemical Safety for the 21st Century Act

The Frank R. Lautenberg Chemical Safety for the 21st Century Act was enacted in 2016, with overwhelming bipartisan support, and gives EPA the authority to regulate or ban new and existing chemicals that pose a risk to human health or health of the environment. Prior to this law, which amended the Toxic Substances Control Act, EPA bore the burden of proof to demonstrate that a chemical posed an "unreasonable risk" to public health or the environment. In addition, the law also required EPA to choose the "least burdensome" regulation with consideration of both public health and the impact on the manufacturer (Gerlach, 2016). The law included new requirements for premarket testing of new chemicals ["EPA has to make an affirmative finding on the safety of a new chemical or significant new use of an existing chemical before it is allowed into the marketplace" (Camacho-Ramos, 2016)]; ongoing risk evaluation of the toxicity of chemicals already in commerce ("EPA must develop a screening process for all existing chemicals that ranks them according to their level of risk, then develop a risk management strategy for chemicals that raise concerns" (Gerlach 2016), and it is no longer required to select the "least burdensome" option); and a user fee (up to $25 million) charged to companies to cover the costs of risk evaluation. Importantly, for the first time, the law required EPA to specifically assess health threats to children, pregnant women, and other vulnerable populations (EDF, 2016). As of June 25, 2019, 60 new chemical reviews had been completed in May 2019. More than 2,300 new chemical reviews have been completed since enactment (EPA, 2019).

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The President's Task Force on Environmental Health Risks and Safety Risks to Children

The President's Task Force on Environmental Health Risks and Safety Risks to Children was established by Executive Order 13045 in 1997.[29] The Task Force was created with multidisciplinary and cross-sector membership, including representation from the Departments of Education, Labor, Energy, Justice, HUD, Agriculture, and Transportation. According to the executive order, the mission of the Task Force is to

> Recommend to the president federal strategies for children's environmental health and safety, within the limits of the administration's budget, to include the following elements:
>
> a. statements of principles, general policy, and targeted annual priorities to guide the federal approach to achieving the goals of this order;
> b. a coordinated research agenda for the federal government, including steps to implement the review of research databases described in Section 4 of the executive order;
> c. recommendations for appropriate partnerships among federal, state, local, and tribal governments and the private, academic, and nonprofit sectors;
> d. proposals to enhance public outreach and communication to assist families in evaluating risks to children and in making informed consumer choices;
> e. an identification of high-priority initiatives that the federal government has undertaken or will undertake in advancing protection of children's environmental health and safety; and
> f. a statement regarding the desirability of new legislation to fulfill or promote the purposes of this order (Executive Order 13045, 1997).

Over the years, the Task Force has developed and published guidance on federal strategies to protect children from environmental health and safety risks. These publications include information on strategies and programs to address topics such as childhood asthma (President's Task Force on Environmental Health Risks and Safety Risks to Children, n.d.), healthy housing (Federal Healthy Homes Work Group, 2013), and lead exposures (President's Task Force on Environmental Health Risks and Safety Risks to Children, 2016). The Task Force's 2016 work plan identifies three areas of priority: 1) reducing lead exposures by addressing sources of lead (e.g., paint, drinking water, and consumer products), 2) protecting health in a changing climate by preparing families and communities to understand and mitigate the effects of climate change on children's health, and 3) reducing the burden of environment-related illnesses by addressing the environmental origins of diseases to promote health and reduce health disparities (PTFCEH, 2016). The work plan also delineates short- and long-term strategies for making progress on these three priority areas. The strategies offer a multipronged approach by identifying opportunities for communication and engagement, educational innovation, regulation and policy, and research.

National Research Approaches to Establish the Effects of Environmental Exposure

Fifteen years ago, spending on federal maternal and child health programs neared $57.5 billion dollars. Most spending fell within the U.S. Department of Health and Human Services

[29] For more information, see https://ptfceh.niehs.nih.gov (accessed April 8, 2019).

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(HHS), but it included several HHS agencies, the Departments of Defense, Education, Agriculture, HUD, and EPA. A 2012 study examining these spending trends noted that supporting a continued focus on data sharing and integration of funding streams could yield synergistic effects and economies of scale. With a goal of realizing a coordinated, integrative MCH system, researchers recommended that federal, state, and local partners eliminate existing silos and increase community and consumer involvement (Kenney et al., 2012). It is unclear whether the federal government has attained these goals, and more research would support a re-evaluation of whether the implementation of federal MCH funding has resulted in improvements in outcomes, data sharing, or improved community participation. In particular, it is unclear how federal spending furthers goals related to the elimination or mitigation of environmental exposures from preconception through age eight. Additional research examining federal funding linked to environmental exposures for these populations could support decision making at multiple levels and across agencies charged with regulation of environmental exposures.

In 2016, the NIH introduced the Environmental influences on Child Health Outcomes (ECHO) Program, a group of over 70 cohort studies aimed at understanding the relationship between environmental exposures and five main domains: upper and lower airways; obesity; pre-, peri-, and postnatal outcomes; neurodevelopment; and positive health outcomes (ECHO, 2019a,b). ECHO has enrolled a combined sample of more than 50,000 children (ECHO, 2019a,b). Exposures are measured from before birth to 5 years of age, and outcome data are collected through adolescence (Forrest et al., 2018). Nearly 150 cohort-focused articles have been published in the first 3 years of the project. In February 2019, the project received single institutional review board approval to begin the ECHO-wide cohort data collection protocol (ECHO, 2019a). While the multiple measures over time on the same participants in longitudinal research provide important information on possible cause and effect relationships, pooled cohort studies offer the advantage of being able to include a variety of different population groups and multiple variables at the same time.

National survey data also continue to play a vital role in our understanding of environmental exposures among children. Biomonitoring data in NHANES among pre-school-aged children are limited (Calafat et al., 2017) but offer an important key to establishing exposure trends among children as young as 3 years old, using both parental report and analysis of biomarkers within urine samples. With these methods, researchers detected over 37 chemicals from children aged 3–5 years, including plasticizers, combustion products, personal-care product chemicals, and pesticides (Calafat et al., 2017). NHANES data also offer the opportunity to establish benchmarks for environmental exposures among children, which enables programs and community-based efforts to improve interventions to reduce the risk of environmental exposures. For example, in a recent study of farmworkers, researchers measured a significant reduction in the levels of pesticide metabolites among parents and children who received an educational intervention on steps to reduce transmission of pesticides in the home (Griffith et al., 2018). Moreover, NHANES data were key in establishing that children exposed to higher levels of one chemical were also exposed to higher levels of other chemicals (Hendryx and Luo, 2018), which may place them at higher risk for poor health outcomes.

Conclusions and Recommendations to Reduce Exposure to Environmental Toxins

Given the importance of safe early learning environments, as discussed above, the committee recommends:

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Recommendation 6-7: The Administration for Children and Families, Maternal and Child Health Bureau, and federal and state regulators should strengthen environmental protection in early care and education settings through expanded workforce training, program monitoring, and regulations.

Training professionals who regularly interact with children and their families on how to prevent and mitigate exposure to environmental toxicants during early development is an important opportunity to advance health equity; therefore, the committee recommends:

Recommendation 6-8: Professional societies, training programs, and accrediting bodies should support expanded or innovative models training of prenatal and childhood health care providers on screening, counseling, and interventions to prevent or mitigate toxic environmental exposures.

Given the need to continually monitor, prevent, and mitigate the impact of environmental toxicants in the preconception through early childhood period, it is critical that federal, state, local, tribal, and territorial governments continue their efforts in this area.

Recommendation 6-9: Federal, state, local, tribal, and territorial governments should support and enforce efforts to prevent and mitigate the impact of environmental toxicants during the preconception through early childhood period. This strategy should particularly focus on priority populations who are disproportionately impacted by harmful environmental exposures. This includes:

- **Environmental Protection Agency fully exercising the authorities provided by Congress to safeguard children's environmental health under the Toxic Substance and Control Act as amended by the Frank R. Lautenberg Chemical Safety for the 21st Century Act.**
- **Continued allocation of resources and technical assistance from the federal government through Center for Disease Control (CDC), EPA, Food and Drug Administration (FDA), and the U.S. Consumer Product Safety Commission to translate existing data and research findings into actionable policies and practices.**
- **Ongoing review and updating of environmental exposure levels by federal agencies to reflect health and safety standards specific to the unique vulnerability of children (from fetal development through early development).**

Other ongoing governmental activities that should be continued include regularly examining children and adolescents in the NHANES and other national surveys for conditions related to environmental exposures, to track and monitor national and regional trends; development of policies and regulations requiring the remediation of persistent environmental exposure risks, including but not limited to children's exposures to lead, mercury, arsenic, and chemicals and by-products of manufacturing, and the continuation of environmental evaluations; and case management of children with demonstrated risk of environmental exposures, particularly within

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communities experiencing higher risk levels, such as those with disproportionate poverty rates, substandard housing, higher concentrations of air pollutants, and other risky exposures.

In summary, while the approaches discussed in this chapter offer a few promising practices or approaches, several unknowns continue to pose challenges to a comprehensive exposure reduction effort. For example, more research and monitoring is needed to assess the effects of multiple long-term chemical exposures, paternal exposure burden and effects, and potential interactions between chemical exposures and gene expression. Evaluation of new, systemic approaches to policy development and implementation with demonstrable effects of reducing or mitigating chemical exposures remains key (Wang et al., 2016). One possible example is genetic studies examining gene variations that may increase susceptibility to environmental exposures. Combining these data with prospective, longitudinal studies could expand our knowledge of the role of critical developmental stages in the etiology of childhood diseases related to chemical exposures and human development (Wright and Christiani, 2010) and lead to novel approaches to further reduce environmental exposure risk.

CONCLUSION

Creating healthy living conditions to promote optimal development requires a multipronged approach to support caregivers and families so that they may meet the fundamental needs of their children. This chapter builds on the evidence presented in Chapter 3 on early life influences and discusses the evidence on programs, policies, and systems changes to ensure that all children have access to healthy living conditions. Among the critical needs discussed in this chapter are access to nutrition, safe and stable housing, protection from environmental exposures, and importantly, economic resources to ensure these basic needs are met.

Based on its review of the extent literature and committee expertise, the committee offers program and policy recommendations to support families in promoting optimal development from the prenatal through early childhood periods. The conclusions and recommendations in this chapter emphasize the importance of public programs to provide access to economic resources, healthy foods, and stable housing. Furthermore, the recommendations highlight the many roles of government at all levels in shaping healthy early living conditions (e.g., removing barriers to accessing public nutrition programs, creating a comprehensive plan for healthy housing, passing legislation to authorize and fund paid parental leave, strengthening environmental protections in ECE settings). Implementation of the program and policy solutions in this chapter would enable families and caregivers to care for young children in healthy communities.

This chapter takes a systems perspective by delving into the multiple, interactive systems that shape the social, economic, and environmental determinants of health for children (see Chapter 8 for more on a systems approach to promote health equity during the prenatal through early childhood periods). The following chapter focuses on the ECE system and the various programs and policies that can be leveraged to promote health equity among children.

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7

Promoting Health Equity Through Early Care and Education

INTRODUCTION

Several important threads are evident throughout this report: the importance of intervening early, preferably before adversity occurs, but if not, soon after; the inextricable interplay between genes and the child's environment in producing health; the need to support caregivers of children—those who spend significant time with children and therefore have an important impact on children's growth and development; and the need to create healthy supportive environments. Knowing that most young children participate in some type of nonparental care on a regular basis (formal or informal arrangements), the early care and education (ECE) platform is a significant opportunity for health promotion and advancing health equity (see the committee's conceptual model, Figure 1-9, in Chapter 1). ECE is defined here as nonparental care that occurs outside the child's home. ECE services may be delivered in center-based settings, school-based settings, or home-based settings (i.e., a setting other than a child's home) (NASEM, 2018); however, this chapter also discusses programs that support parents, such as home visiting. Education itself is incredibly important when it comes to health (García, 2015). Because educational attainment positively correlates with health outcomes, investments in ECE are critical to decreasing disparities to set the stage for future success (Barnett, 2013; NASEM, 2017a). In this chapter, the committee discusses how to apply the important learnings from early development to the ECE system, including the importance of a properly supported and trained ECE workforce, access to quality ECE, and resources to support these needs. At the end of the chapter, the committee provides recommendations detailing the specific actions needed to ensure that ECE meets its potential to promote child health and well-being. See Box 7-1 for an overview of the chapter.

While ECE has primarily focused on whether it improves children's cognitive and socio-emotional development, as well as academic readiness, there is some indication that ECE may influence child (and even adult) health outcomes, including physical, emotional, and mental health (Campbell et al., 2014; D'Onise et al., 2010; Muennig et al., 2011). What is also of critical importance is *how* ECE is related to children's cognitive development, socio-emotional development, academic readiness and achievement, and health and well-being, as well as how it can lead to health equity. Hahn and colleagues (2016) postulate that ECE advances health equity through several interrelated systems (see Figure 7-1).

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BOX 7-1
Chapter in Brief: Early Care And Education

This chapter discusses the role of ECE in ensuring that children are healthy and ready to learn, with a focus on incorporating health and health equity into a comprehensive approach to school readiness and success. The chapter examines the evidence linking ECE to health and health equity outcomes through leveraging such programs and systems as platforms to deliver health-related services, social-emotional and behavioral curricula, and interventions, supporting parents and the home environment as well as educators and other members of the ECE workforce. The chapter also emphasizes the importance of improving the quality, access, and affordability of ECE programs, especially for underserved populations.

Chapter conclusions in brief:

- For ECE programs to contribute significantly to a health promotion and equity strategy, there is a need to intentionally, cohesively, and simultaneously address adequate funding that supports comprehensive, evidence-based standards and practices that promote health equity in the ECE system, an adequately compensated and competent workforce, a connection to community resources and support, continuous quality improvement, and a systematic examination of effectiveness at multiple levels.
- Policies and systems that prepare and support early childhood educators and program leaders, including those in public schools, need to incorporate the latest evidence about how educators can better support children's school readiness and success by fostering their health and well-being. This would entail providing comprehensive supports and resources to degree granting institutions and preparation programs, including the development of curricula, textbooks, practicum experiences, toolkits, and fact sheets, with an emphasis on equitable practices that address the diverse experiences and needs of children and families.
- Maximizing the impact of ECE on positive childhood development and health and well-being at the community or population levels will require increasing public funds for ECE programs. Currently, eligibility for ECE programs is limited, and among eligible families, access is low due to lack of funding and availability of programs and services. Therefore, even if existing publicly funded programs have the resources to provide robust supports that improve young children's health and well-being, they will not reach most children, especially those who live in low-income households or experience adverse experiences and toxic stressors.

Chapter recommendations in brief:

- Develop a comprehensive approach to school readiness that explicitly incorporates health outcomes, standards, and practices and leverages ECE systems and programs, including home visiting.
- Develop and strengthen coursework, practicums, and ongoing professional learning opportunities that focus on competencies of educators, principals, and ECE program directors that are critical to children's health, school readiness, and life success.
- Develop and implement a strategic plan to (1) improve the quality of ECE programs by adopting the health-promoting standards and (2) expand access to comprehensive high-quality and affordable ECE programs across multiple settings.

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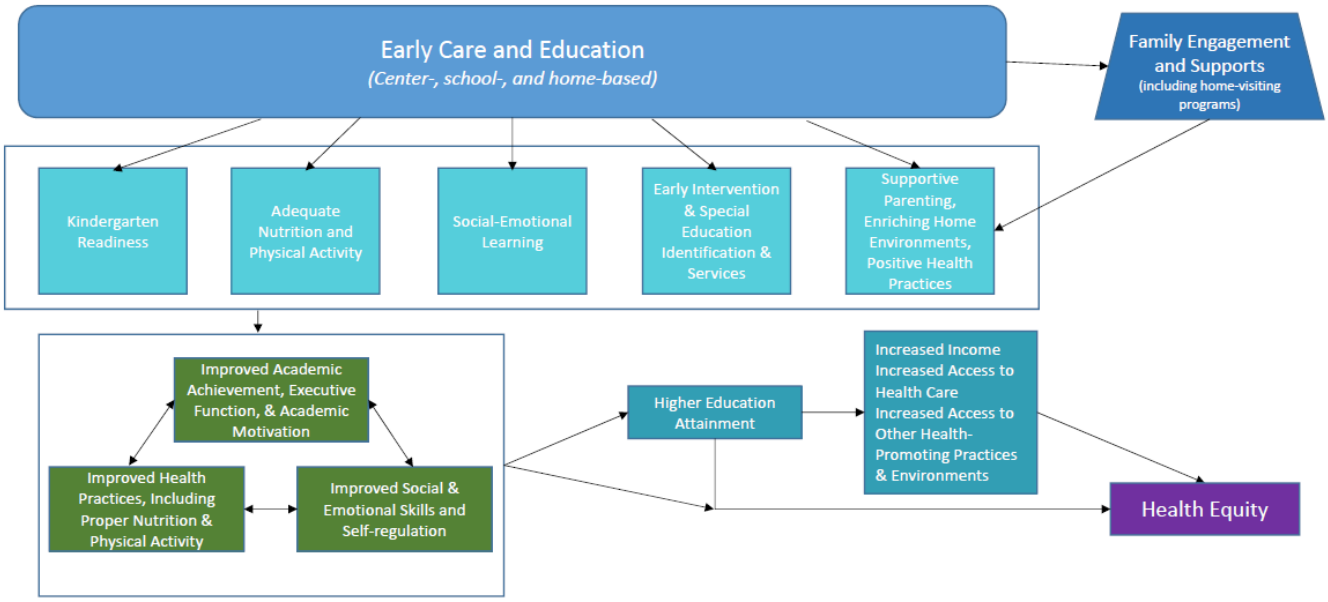

FIGURE 7-1: Conceptual framework: How ECE is linked to health promotion and health equity
SOURCE: Informed by Hahn et al., 2016.

ECE programs increase children's cognitive, social, and health outcomes through enhancing children's motivation for school and readiness to learn and identifying problems that impede learning. This, in turn, helps children improve their cognitive ability and social and emotional competence, while increasing their use of preventive health care. There is also evidence that participation in a high-quality early learning program is associated with children's self-regulation; approaches to learning, such as their motivation and persistence; and executive function (EF) skills, which are domain-general skills that transfer to many areas of development, including learning to read, making friends, and dealing with new challenges (Pianta et al., 2009; Holliday et al., 2014; Yoshikawa et al., 2013). These short-term outcomes of ECE are then expected to lead to lower risk of dropping out of school, greater school engagement, and subsequently better educational attainment, which results in increased income and health care, decreased social and health risk, and improved health equity. Traditionally, ECE is thought of as being confined to a specific age range. For this report, the committee discusses ECE in the context of birth through 8–10 years of age. Another National Academies report, *The Promise of Adolescence: Realizing Opportunity for All Youth* (NASEM 2019), picks up from here, discussing health and development from the onset of puberty into adolescence and early adulthood.

DIRECT LINKS BETWEEN ECE AND HEALTH EQUITY

Health, social-emotional, and other health-related behavioral outcomes are some of the most commonly reported from evaluations of ECE programs, aside from the often-cited cognitive outcomes (Cannon et al., 2017; Carney et al., 2015; Fisher et al., 2014; Rossin-Slater, 2015). ECE programs have been shown to reduce externalizing and internalizing behaviors (Carney et al., 2015), improve social-emotional skills (D'Onise et al., 2010; Hahn et al., 2016), reduce substance use (Cannon et al., 2017; Jones et al., 2015), and improve physical health or well-being (D'Onise et al., 2010; Rossin-Slater, 2015; Sabol and Hoyt, 2017). ECE can produce

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this range of results through different pathways. It can provide services to children and/or their parents directly that impact their health outcomes and related skills and behaviors; implement evidence-based curricula or interventions to improve children's social-emotional skills, which are associated with both short and long-term health and cognitive effects; and support the training and well-being of early childhood educators.

While reviews of the research literature suggest that ECE programs can be a promising lever for improving health outcomes and equity, they also show instances where ECE programs have weak, nonexistent, or even negative impact on children's behavior and health (Cannon et al., 2017; D'Onise et al., 2010; Hahn et al., 2016; Herbst and Tekin, 2011; Rossin-Slater, 2015). These mixed and negative findings could reflect factors such as lack of program quality, poor fidelity to the program model, and limited program duration. The following section takes a deeper dive into specific ECE programs or interventions that have produced significant results on health and discusses their characteristics.

Links Between ECE and Health Outcomes

A number of studies and reviews of the literature have found positive relationships between participation in ECE programs and physical health indicators and outcomes (Hahn et al., 2016; Kay and Pennucci, 2014). Most of the associated benefits tend to be related to obesity, access to health care, and early screenings and detection. For example, low-income preschoolers enrolled in a center-based program are less likely to experience food insecurity than if they were cared for by parents exclusively or by an unrelated adult in a home setting (Gundersen and Ziliak, 2014). Using a quasi-experimental methodology on data from the National Institute of Child Health and Human Development Study of Early Child Care and Youth Development (NICHD SECCYD), Sabol and Hoyt (2017) found that 4-year-olds who attended center-based ECE programs had lower blood pressure when they were 15 years old than those who were in a home-based environment, whether that was with a parent, a relative, or nonrelative.

Hahn et al. (2016) sought to examine the impact of ECE on fostering the health equity outcomes of low-income and racial and ethnic minority children through a meta-analysis. They focused on state and district programs, the federal Head Start program, and foundational model programs, such as the HighScope Perry Preschool (PPP) and Abecedarian programs. They included studies that were for children aged 3 or 4 years; primarily focused on low-income or racial and ethnic minority populations; not conducted only in the summer; based on behavioral interventions; included assessment of effects on children's health and health-related or academic outcomes; and had a control or comparison population and provided enough data for analysts to calculate effect size and adjust for confounding. Findings were included for the following outcomes: standardized achievement (an effect was found across program types); high school graduation (effect found only for Head Start); grade retention (effect found across program types); assignment to special education (Head Start was not evaluated on this measure; effect found for all other programs); and crime (effect found across all program types) (see Table 7-1 for more information). Additional analyses examining the persistence of effect of programs on academic achievement and cognitive ability showed a rapid decrease of effects after the program ended, then a gradual decline over time. Higher program quality based on observational data and having teachers with a bachelor's degree or higher had greater effects on student standardized achievement. There was insufficient data to examine impact of class size, hours, duration, or benefits of additional components, such as family engagement or health access. In sum, there was consistent evidence that center-based ECE programs improved educational and health-

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related outcomes for low-income and ethnic minority preschool-age children, with some indication of long-term outcomes. Hahn et al. (2016) further note that the fade-out of center-based ECE effects for cognitive and achievement outcomes could likely be because many low-income and ethnic minority children are likely to attend low-resourced (i.e., lower-quality) elementary schools and have teachers with fewer credentials. Others, such as Duncan and Magnuson (2013), have postulated that "preschool programs may affect something other than basic achievement and cognitive test scores, and perhaps these other program impacts, unlike achievement and cognitive impacts, persist over time" (p. 120). That is, looking at discrete and constrained skills, such as letter naming, may not be good predictors, whereas focusing on unconstrained skills, such as self-regulation and expressive language, would be more appropriate. Thus, there is more to understand about the fade-out effect, or arguably the catch-up effect, especially in the changing landscape where more children are in out-of-home settings.

TABLE 7-1 Effects of Center-based Early Childhood Education Programs on Education, Social, and Health-related Outcomes (data for all program types combined).

Outcome (Number of Studies; Program Types Included)	Mean Age at Follow-up, y	Standardized Mean Difference (95% CI)	Effect Meaningful?	Consistent Across Body of Evidence?
Test scores (27 studies; all types)	3.7	0.29 (0.23-0.34)	Yes	Yes
High school graduation (7 studies; all types)	20.0	0.20 (0.07-0.33)	Yes	Yes
Grade retention (12 studies; all types)	17.0	– 0.23 (–0.43 to –0.02)	Yes	Yes
Assignment to special education (6 studies; state and district and model programs)	15.5	– 0.28 (–0.49 to –0.08)	Yes	Yes
Crime (5 studies; all types)	25.0	– 0.23 (–0.45 to 0.05)	Yes	No
Teen birth (3 studies; Head Start and model programs)	18.0	– 0.46 (–0.92 to 0.0)	Yes	No
Self-regulation (5 studies; state and district and Head Start programs)	18.0	0.21 (0.14-0.28)	Yes	Yes
Emotional development (7 studies; state and district and Head Start programs)	4.0	0.04 (–0.05 to 0.12)	No	No

SOURCE: Hahn et al., 2016.

The research is equivocal, however, especially for health, social, and emotional outcomes. Herbst and Tekin (2011) found that 4-year-old children of single mothers who were enrolled in nonparental care through a child care subsidy were more likely to be obese or overweight when they were kindergartners than those who stayed home. As with a study by Hawkinson and colleagues (2013), this study also found an association between subsidy use and poor cognitive outcomes. A review of 37 studies by D'Onise et al. (2010) found "generally null effects of preschool interventions across a range of health outcomes" (p. 1432), leading the authors to caution against relying on a "flimsy evidence base" to inform policy (p. 1432). However, the study did find "a general trend toward beneficial effects, with particularly beneficial effects for overweight and obesity, mental health, social competency, and crime prevention" (D'Onise et al., 2010, p. 1432). They also found that across the studies, half of the comparisons related to immunization and general health yielded positive impacts (and none produced adverse effects).

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Foundational Research in ECE

The foundational studies of the HighScope Perry Preschool Program (PPP) and Carolina Abecedarian (ABC) project provide the most robust findings regarding the link between ECE and health equity throughout the life course. As described below, these two programs—which occurred in two different states and were conducted by two different teams in two different decades—have similar short- and long-term outcomes. They also shared some common characteristics: they focused on children with the greatest needs and employed an educated and responsive teacher, low child-teacher ratio, active and language-rich learning opportunities, child assessment, and home visiting and family support activities.

Some caution, however, should be taken in generalizing these findings due to the limitations of these studies. They occurred more than fifty years ago, when most children in poverty did not have access to early education services and programs. The samples, primarily African American children, are not representative of the general population. They were also small and continued to decrease over time. Finally, these controlled programs have not been adequately replicated at a large scale. It is also critical to note that not all children in the treatment groups performed at the highest level and in the end, did not surpass their more economically advantaged peers. For example, in the PPP study almost a third of children from the treatment groups were arrested five or more times by age 40, almost a third did not graduate high school, and almost two-thirds of children from the treatment group required public assistance as adults (Gomby et al., 1995). Thus, these programs did not equalize the outcomes for children from low-incomes households in comparison to their higher income peers.

While PPP and ABC provide a blueprint to build from to support children's school readiness, achievement, and health equity throughout the life course, policymakers and practitioners alike need to base their decisions on lessons beyond those from these studies, such as more contemporary ECE programs and interventions discussed in this chapter, to ensure that all children, especially children with the greatest needs, have the same opportunity to thrive and lead healthy lives.

HighScope Perry Preschool Project (PPP)

The HighScope PPP started in 1962 with a focus on serving 3- and 4-year olds (it was 1 to 2 years long) with a home visiting component. The program is aims to promote the social and cognitive development in children placed at risk due to poverty. Schweinhart and Weikart (1997) showed that students enrolled in the program in 1986 had more positive behavior and attitudes than students in the control group (Schweinhart and Weikart, 1997). In addition, experimental evaluations of study participants in their teens and 20s showed that even years later, when study participants were in their teens and 20s, students formerly enrolled in the program "had higher academic grades and earnings, higher rates of high school graduation, fewer arrests and out-of-wedlock births, and lower levels of welfare receipt than their peers who were not in a preschool program" (Child Trends, 2012). Furthermore, children in the intervention group had higher rates of safety-belt use and engaged in fewer risky health behaviors, such as smoking and illicit substance use, in adulthood compared to those in the control group. At age 27, former PPP students were more likely to be employed and had higher earnings than students in the control group (Child Trends, 2012; Schweinhart et al., 2005). This continued into age 40, with former PPP students having higher earnings, committing fewer crimes, and being more likely to hold a job and to have graduated from high school than adults who did not participate in PPP. In 2019, Heckman and Karapakula presented findings from the HighScope PPP Age 55 Study (Heckman

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and Karapakula, 2019a). They indicated that the program kept parent engagement active longer, which resulted in more warmth and less authoritarian parenting. They also found that at age 55, the female participants in the early childhood intervention group had lower cortisol (39.01 versus 89.29 pg/mg) compared to the control group, and the male participants were less likely to have high cholesterol levels (mean differences in high cholesterol[1] were 0.71 in the treatment group versus 0.94 in the control group). Female participants in the intervention group were also less likely to be uninsured for a prolonged period compared to the control group (Heckman and Karapakula, 2019b). In addition, they reported intergenerational effects for children of intervention participants: completion of high school, good health status, stable employment, and a history of never having been suspended and arrested (Heckman and Karapakula, 2019a).

Carolina Abecedarian (ABC) Study

The Carolina ABC Study is a center-based intervention that enrolled families between 1972 and 1977 based on a high-risk index. During recruitment, 111 infants were matched on high-risk scores and then assigned to preschool treatment or control status. Fifty-seven infants were assigned to the experimental group and 54 to the control group (Campbell et al., 2002). The families in the study were mostly African American, with young mothers, with less than a high school education, unmarried, living in multigenerational homes, and reporting no earned income (Campbell et al., 2002). The service delivery model for the experimental group was a 5-year, full-day, year-round, center-based program with a comprehensive curriculum (LearningGames®) (Sparling and Lewis, 1979) focused on educational games addressing children's cognition, language, and adaptive behavior. The program also emphasized health care and family support programs. Activities were individualized for the child's needs, with more conceptual and group-oriented activities as children got older. Families in both the experimental and control groups received supportive social services. Findings showed that children in the 0–5 intervention group had better cognitive, academic, and emotional outcomes (Ramey and Ramey, 2004). This also had persistent effects when children were in their 20s, with children in the treatment group having better intellectual test performance and reading and mathematics test scores, more years of education, and a greater likelihood of being enrolled in college (Campbell et al., 2012; FPG Child Development Institute, 2012). By age 30, the treatment group were more likely to have completed a bachelor's degree, have consistent employment, not use public assistance, and have delayed parenthood. Regarding health outcomes, Campbell et al. (2014) found through biomedical data that children in the intervention group at age 35 had significantly lower risk factors for cardiovascular and metabolic disease, especially for male participants (i.e., mean systolic blood pressure for the control group was 143 versus 126 for boys in the intervention group). "One in four males in the control group was affected by metabolic syndrome, while none in the treatment group were" (Ramey, 2018, p. 539).

Looking across the body of research, Head Start appears to be particularly effective at promoting young children's physical health (see Box 7-2 for more information on Head Start). The Head Start Impact Study showed that children enrolled in the program had better access to dental care while they were in the program and to health insurance when they were in kindergarten (Puma et al., 2012). Broader reviews of research on Head Start show positive impacts on obesity, immunization, screening for hearing and vision problems, and even child mortality (Belfield and Kelly, 2013; Rossin-Slater, 2015; Yoshikawa et al., 2013). In addition, in

[1] High total cholesterol indicates whether total cholesterol concentration in mg/dL is 220 or higher.

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their review of ECLS-B data, Belfield and Kelly (2013) found that Head Start provided its participants "protective effects against . . . asthma, respiratory ailments, allergies, and being on medication" (p. 322). Lee et al. (2013) used data from ECLS-B to analyze low-income children's nutrition, weight, and health care receipt at kindergarten entry. They compared (a) Head Start participants and all nonparticipants and (b) Head Start participants and children in pre-K, other center-based care, other nonparental care, or only parental care using propensity-score-weighted regressions. They found Head Start effects were larger compared to informal child care settings rather than center-based settings. Specifically, Head Start children had lower BMI scores and probability of being overweight compared to children in home-based settings; had better healthy eating habits than children in center- and home-based settings; and were also more likely to have dental care checkups compared to children in any other type of setting, including pre-K and center-based settings. Furthermore, dosage appears to matter. Frisvold and Lumeng (2011) analyzed administrative data from more than 1,500 children from Head Start programs in Michigan from 2001 to 2006 and found that children who participated in full-day Head Start were 25 percent less likely to be obese at the end of an academic year than those who enrolled in half-day programs. The effect seems to be more pronounced for boys and African American children.

BOX 7-2
About Head Start

Head Start is a program of the U.S. Department of Health and Human Services that provides comprehensive early childhood education, health, nutrition, and parent-involvement services to low-income children and their families.

Program quality in Head Start and Early Head Start programs is defined by the Head Start Program Performance Standards (HSPPS), which include provisions related to teaching and the learning environment, curricula, child screenings and assessment, oral health practices, child nutrition, child mental health and social-emotional well-being, and family engagement. In addition, the HSPPS includes standards of practice for special populations, including tribal communities, dual-language learners (DLLs), children with disabilities, and pregnant women.

The federal government assesses the extent to which programs meet HSPPS through the Designation Renewal System. Programs are evaluated on a regular basis, and if deficiencies are found, programs are required to recompete for their federal grant. If the program does not achieve a high level of classroom and teacher-child interaction quality, as measured by the Classroom Assessment Scoring System (CLASS), it will need to recompete. Generally, Head Start programs score well on the Emotional Support and Classroom Organization domains of the CLASS assessment (about a 6 out of 7), but the average score for Instructional Support is only a 2.9. "In only two states (Kentucky and Vermont) were scores statistically significantly above 3 such that we can be confident the state average exceeds the threshold" (Barnett and Friedman-Krauss, 2016, p. 5).

Head Start's standards stand out among the major ECE programs for the attention to the holistic needs of children and families (including health care) and value placed on engaging families as partners and leaders. Programs are required to help children and families access health services (including insurance), promote oral health, meet children's nutritional needs, and work with mental health consultants. Programs also need to develop strategies to communicate with parents; implement intake and assessment processes to identify families' strengths and needs; connect them to resources that "support family well-being, including family safety, health, and economic stability"; increase their capacity to support their children's learning and development; and include them in the governance of the program. Finally, Head Start programs have to establish relationships with other programs that serve families in their communities, such as health care providers, schools, child care, libraries and museums, housing agencies, and other social services agencies.

In 2016, the federal government released updated standards for Head Start (previously not updated since the authorization of the Improving Head Start for School Readiness Act of 2007) to reflect

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the latest research on child development and program quality. More research is needed to elucidate how these changes have affected program quality and outcomes.

SOURCES: HHS, 2017; Rose, 2010.

One reason that Head Start stands out among ECE programs with respect to impact on physical health outcomes may be that the program design includes a robust health component (such as requiring programs to provide diverse nutrition and health services, helping families receive physical examinations by scheduling screening appointments or offering screenings directly onsite, assisting families in applying for age-appropriate health care services, providing health promotion activities directly onsite, and tracking each child's health progress) (Lee et al., 2013). In their review of the literature, Yoshikawa et al. (2013) concluded that "in contrast to the literature on Head Start and health outcomes, there are almost no studies of the effects of public prekindergarten on children's health" (p. 5) because pre-K programs typically do not provide health-related services.

However, pre-K programs can incorporate such services and potentially achieve similar outcomes. A recent evaluation of the Universal Pre-K (UPK) program in New York City found that the expansion of the program "led to increases in rates of diagnosis of asthma and vision problems, to increased rates of screening for immunization or infectious disease, and to increased rates of treatment of hearing and vision problems" among Medicaid recipients who were eligible for UPK (Hong et al., 2017, p. 3). The researchers attribute some of these findings to UPK's program requirements, which include immunizations and developmental screenings for all enrolled students. In other words, by incorporating direct services[2] into their program design, Head Start and New York City's UPK initiative have become opportunities to effect broad improvements in young children's health outcomes.

Head Start and the UPK program in New York City are just two examples of large-scale, publicly funded ECE programs that have demonstrated generally positive results for preschool children. Others include state-funded pre-K programs in Georgia, New Jersey, North Carolina, and Oklahoma and district-run programs, like that of Boston. In a review of the body of research behind these and other large, publicly funded programs, Phillips et al. (2017) concluded that there is robust evidence of short-term benefits, especially in cognitive and academic skills. However, "the available evidence about the long-term effects of state pre-k programs offers some promising potential but is not yet sufficient to support confident overall and general conclusions about long-term effects" (p. 10). For example, Lipsey et al. (2018) found that although children who participated in the Tennessee pre-K program demonstrated better cognitive skills than the control group, this advantage was lost or even reversed by second or third grade.

Still, the promising results from large-scale, publicly funded pre-K programs, in light of the impacts that Head Start and New York City's UPK program have on young children's health outcomes, suggest that ECE programs that serve significant proportions of young children can be a platform for interventions that promote health equity.

[2] For example, immunization and a valid and reliable developmental screening tool to identify students with potential developmental delays and English Language Acquisition support needs.

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Home-Based Child Care Programs

Home-based child care is regulated family child care and family, friend, and neighbor care (Porter et al., 2010b). As mentioned earlier, it is a common arrangement for many young children in the United States, particularly those from low-income families and families of color (Porter et al., 2010b). As Porter et al. (2010a) note, "parents use these arrangements for a variety of reasons, including convenience, flexibility, trust, shared language and culture, and individual attention from the caregiver" (p. 1). Home-based child care also serves as a primary nonparental care arrangement for infants and toddlers (Corcoran et al., 2019). (See Chapter 3 for statistics by race, ethnicity, and income.)

While most young children who are not yet enrolled in kindergarten participate in some kind of weekly center- or school-based early childhood program, 41 percent of them receive care weekly from a relative, and 22 percent participate in nonrelative care in a home environment (Corcoran et al., 2019). (Twelve percent of young children receive care in more than one of these settings on a weekly basis.) The number of children in center-based care increases as children get older. Furthermore, about 3.6 million of the approximately 3.7 million home-based providers are "unlisted": not registered with, licensed by, or regulated by a public agency. Together, these home-based providers serve more than 7 million young children (National Survey of Early Care and Education Project Team, 2016). Children of color are more likely to receive care from a relative, while white children are more likely to participate in nonrelative home-based care. Infants and toddlers are more likely to be cared for in a home setting—whether with a relative or nonrelative—while preschool-aged children are more likely to be enrolled in a center-/school-based program. Families with a household income of $75,000 or less are also more likely to put their young children in relative care (Corcoran et al., 2019). Thus, to the extent that regulations and policies related to safety and quality promote better care and child development, the youngest children, low-income children, and children of color could disproportionately lack access to ECE opportunities that are more equipped to support their cognitive, social-emotional, and healthy development.

The majority of evidence linking ECE to children's health, education, and well-being is primarily from center- and school-based programs. Some studies have shown a link between home-based programs and children's academic skills and socio-emotional development. For example, Iruka and Forry (2018) found that children in home-based programs that were high quality and engaged frequently in enriching literacy and numeracy activities (e.g., learning names of letters, learning the conventions of print, using manipulatives, using a measuring instrument, learning about shapes and patterns) were likely to have stronger reading and math skills compared to children in home-based programs that were low quality and engaged in fewer enriching activities. This is consistent with prior work by Forry et al. (2013) using data from a multistate study of a professional development intervention showing that the quality of home-based programs, their child-centered beliefs (e.g., progressive beliefs that children should have autonomy and be allowed to express their ideas), and their perceptions of job demands were related to children's school readiness, emotional health (e.g., initiative, self-control, and attachment), and internalizing and externalizing problem behaviors.

However, mostly correlational data indicate that children in centers compared to family child care homes had higher cognitive, language, and school readiness scores but increased likelihood of contracting communicable illnesses and otitis media (ear infection), which is likely due to the large group size (Bradley and Vandell, 2007). While the data are mixed, there is indication that children in home-based programs have stronger socio-emotional competence

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compared to children who attended center-based programs (Belsky et al., 2007). In their analyses examining multiple child care arrangement and children's academic and behavioral outcomes, Gordon et al. (2013) found that preschool children, on average, scored higher on reading and math assessments when they attended centers alone or centers in combination with home-based programs than home-based programs only or parental care. There were no differences in children's socio-emotional development between families who used or did not use multiple care arrangements. The stronger benefits for children's cognitive and school readiness skills for center-based compared to home-based programs has also been seen for Latino children (Ansari and Winsler, 2012).

These better academic outcomes for center- versus home-based programs are likely due to higher teacher education and more training opportunities (Bradley and Vandell, 2007). However, the larger group size in center-based programs may preclude sensitive individual care and attending to children's socio-emotional needs, which could exacerbate problem behaviors (Gordon et al., 2013). There is still a need for more rigorous examination of the differential impact based on program type and accounting for differences in teacher education and training, sociodemographics of children and families, and more robust health-related outcomes.

Quality Rating and Improvement Systems (QRISs)

Motivated in part by ECE research, states and localities have implemented QRISs to promote and enhance ECE program quality across various sectors and settings, including schools, community-based organizations (center- and home-based), and Head Start. State and local policy makers have used research linking high-quality early childhood education and children's outcomes in developing QRISs to ensure that children, especially disadvantaged children, are attending high-quality education programs during the early years. QRISs could serve as a unifying framework for defining quality across ECE programs and a defined pathway for achieving it. Moreover, without a strategy such as a QRIS, ECE programs could have inequitable resources for improvement, exacerbating the variance of quality among programs and leading to inequitable outcomes for children, families, and communities. When funded adequately and supported as a unifying strategy for ECE, QRISs can raise the overall quality of the ECE system and create more equity across communities.

Almost all QRISs measure staff training and education and assess the classroom or learning environment (Burchinal et al., 2015). Factors such as parent-involvement activities, business practices, child-staff ratios, and national accreditation status vary by state (Burchinal et al., 2015; Zellman and Perlman, 2008). QRISs serve multiple purposes, including providing a standardized method to rate program quality—based on a set of criteria—and to make the program rating information available to parents, as is done with restaurant ratings. The rating system is built on the primary assumption that parents often lack good information about program quality and that such information would inform their decisions on program selection (Burchinal et al., 2015). Consequently, providers who work with lower-quality programs would be incentivized to enhance the quality of their program or leave the market (Burchinal et al., 2015; Zellman and Perlman, 2008). In addition, QRISs represent a systematic approach to providing a range of technical assistance, resources, and incentives for programs to improve their quality (Burchinal et al., 2015). This could entail consultation on quality improvement, increased investments for professional development scholarships, microgrants for other targeted efforts, and increased subsidy payments for more highly rated programs (Burchinal et al., 2015). As noted by Burchinal et al. (2015), "the goal of these efforts is to foster and support providers'

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efforts to improve the quality of care they provide. Thus, [QRISs] attempt to improve quality by affecting both the demand for high-quality care and the supply of such care. Of course, the success of such efforts rests on the ability of rating systems to accurately identify and measure key aspects of quality and the willingness of providers to participate in a rating system" (Burchinal et al., 2015, p. 255; see also Zellman and Perlman, 2008).

Validation of QRISs has yielded mixed findings. The Race to the Top—Early Learning Challenge Grant resulted in a proliferation of QRIS validation studies. Prior to this, most research on QRISs was descriptive and focused on issues of implementation. A recent synthesis of the validation studies by Tout et al. (2017) from 10 states found that while these ECE rating systems were valid (i.e., independent observations indicated meaningful differences across levels), most programs were, on average, providing moderate level of quality and inconsistently associated with child outcomes, mostly for social-emotional development and EF outcomes.

There has been a limited focus on children's physical health. In their synthesis of states' QRIS validation studies, Tout et al. (2017) found that only two states focused on physical development, which included BMI and fine and gross motor skills. One state found a link between its QRIS and children's fine motor development, indicating that higher-rated programs were associated with improved fine motor development. Several states have included nutrition, physical activity, and screen time as part of their QRIS standards (Gabor and Mantinan, 2012). In their report to examine state efforts to address obesity prevention in QRISs, Gabor and Mantinan (2012) found specific standards, including a focus on nutrition (including standards), physical activity and screen time limits, professional development for staff and teachers, and sharing information about nutrition and physical activity with families.

The differences in system designs across states make it difficult to draw general conclusions from these validation studies about their links to various domains of children's development, especially health. The voluntary nature of QRISs in most states and the varying standards also make it difficult to establish a causal link between them and child outcomes. Even with these limitations, the QRIS is one *potential*, and perhaps underused, platform that could increase the use of evidence-based, health-promoting practices and standards in a mixed-delivery system by unifying leadership and governance, standards, financing, stakeholder engagement, improvement supports, accountability, and continuous quality improvement. As QRISs expand, mature, and become better funded, they could serve as the one point of entry that promotes high-quality programming across various settings (e.g., home, school, centers) and provides more children, especially children with the greatest needs, with access to beneficial ECE experiences that meet their comprehensive needs regardless of program funding.

Early Intervention for Children with Developmental Disabilities

Early Intervention[3] services support the early development of children with delays (meaning they are far behind other children) or specific health conditions that could lead to delay (e.g., genetic disorder, birth defect, hearing loss). These services and programs intend to help children catch up and increase their chances for school and life success, though most of this work falls to parents and families. Early intervention services are provided under IDEA. Eligible children are able to receive services free of charge (or reduced rate) through federal grants to states. Each state has its own definition of developmental delay and its own process for

[3] Early intervention refers to services for children ages 0–3. Early childhood special education refers to services for children ages 3–5.

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determining eligibility and identifying eligible children. Families with children under age 3 who quality for early intervention receive an Individualized Family Service Plan (IFSP) that defines goals and the types of services that will support the family and child. Children older than three who are eligible for special education services under IDEA meet with school professionals to develop an Individualized Education Program (IEP) to support their educational goals.

Some of the specialists who work with children include speech-language pathologists, who help with communication speech and language delays; physical therapists, who strengthen children's movement, gross motor skills, and physical development; occupational therapists, who improve fine motor, cognitive, sensory processing, and communication skills; nurses, who support children's health status and address feeding and growth concerns; social workers, who assess and support children's social and emotional development; and developmental therapists, who design learning activities to promote children's learning and social interaction skills.

Early Intervention (Children Under 3 Years Old)

A substantial body of research supports the effectiveness of early intervention for children's functioning (Bruder, 2010; Guralnick, 2005). However, these studies suffer from methodological limitations, such as sample heterogeneity, lack of control groups, narrowly defined outcomes, and inappropriateness of standardized measures of intelligence (Bruder et al., 2010). Nevertheless, there is a body of research indicating that children who receive early intervention services (Part C or Part B) are less likely to see a decline in their functioning over time, with effect sizes of .5 to .75 of the standard deviation (Guralnick, 1998). This is supported by a foundational meta-analysis of 31 studies examining the effect of early intervention (Shonkoff and Hauser-Cram, 1987), which found that early intervention was "effective in promoting developmental progress in infants and toddlers with biologically based disabilities" (p. 650). The mean effects of early intervention services ranged from .43 for motor development to 1.17 for language development. In particular, they found that "programs that served a heterogeneous group of children, provided a structured curriculum, and targeted […] parents and children together appeared to be the most effective" (p. 650).

In their study of community-based early intervention services for children who were in neonatal intensive care units (NICUs), Litt et al. (2018) analyzed retrospective data from the U.S. Department of Education's National Early Intervention Longitudinal Study and found that longer and more intensive services were associated with higher kindergarten skills ratings and the importance of following up after children left the NICU. These findings are consistent with those from McManus et al. (2012) in their longitudinal study of mother-infant dyads from three newborn intensive care units in southeastern Wisconsin. They matched pairs of dyads using propensity-score matching to reduce selection bias and estimate the effect of early intervention services on cognitive function trajectories. They found that service receipt was positively associated with children's cognitive functioning and trajectory and more maternal supports (e.g., mothers' report of emotional, informational, child care, financial, respite, and other support) was associated with better outcomes for families over time. Unfortunately, national data indicate that children who qualify for early intervention services are not likely to receive them, with this issue especially pronounced for black children (Boyd et al., 2018).

Most of the recent evidence about early intervention has primarily focused on children with autism spectrum disorder (ASD), which "is characterized by severe and sustained impairment in communication and social interaction and restricted patterns of ritualistic and stereotyped behaviors manifested [before] 3 years old" (APA, 2013). Children with ASD often

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qualify for early intervention services, usually because they are not developing in social, play, language, and cognitive domains at the expected pace (Landa, 2018). In her review of the efficacy of EIs for young children with or at risk for ASD, Landa (2018) found that greater intervention intensity (hours and duration in months) and fidelity of implementation were associated with greater child gains. One of the applied behavior analysis approaches for children with ASD is called Early Intensive Behavioral Intervention (EIBI), which focuses on remediation of deficient language, imitation, preacademics, self-help, and social interaction skills (Peters-Scheffer et al., 2011). In a meta-analysis examining the effectiveness of EIBI, Peters-Scheffer and colleagues (2011) found the experimental groups outperformed the control groups on IQ, nonverbal IQ, expressive and receptive language, and adaptive behavior, with differences of 4.96–15.21 points on standardized tests. Reichow (2011) offered a similar conclusion in his overview of five meta-analyses of EIBI for young children with ASD, but he also stressed the importance of more information about child characteristics, additional knowledge on the characteristics of EIBI programs used in real-world settings, and guidelines focused on the intensity, duration, level of treatment fidelity, and therapist experience and/or training necessary to achieve optimal outcomes.

There are disparities in service access for economically disadvantaged and racial and ethnic minority families who have children with ASD (Boyd et al., 2018), and these children are also "at risk for poorer outcomes in comparison to their white and higher-income counterparts, including a more severe symptom presentation (e.g., more severe language and cognitive delays)" (Boyd et al., 2018, p. 20; CDC, 2014; Cuccaro et al., 2007; Fountain et al., 2012). One posited rationale for these poorer outcomes is lower-quality or fewer services (Boyd et al., 2018) and lower likelihood of being referred for services (Delgado and Scott, 2006).

Early Intervention/Special Education (Children 3 Years Old and Older)

Special education provides children with disabilities with specialized services designed to "prepare them for further education, employment, and independent living."[4] "Practitioners are responsible for providing specific services, instructional strategies or routines, and resources that mitigate the impact of the disability on a child's learning or behavior" (Morgan et al., 2010, p. 236). "Helping the child to benefit from the school's curriculum should in turn increase subsequent educational and societal opportunities" (Morgan et al., 2010, p. 236; U.S. Department of Education, 2018).

The majority of students with disabilities performed in the "below basic" achievement level in all four areas of measurement (mathematics and reading, in fourth and eighth grade) in 2017. The gaps between students with disabilities and those without disabilities are substantial (Advocacy Institute, 2019). Youth with disabilities are also more likely to drop out of school, be delinquent, be unemployed, earn less, and be unsatisfied with their adult lives (Blackorby and Wagner, 1996; Horowitz et al., 2017; Thurlow et al., 2002). There is some evidence that at the end of the school year, youth placed in special education classrooms sometimes score lower on measures of reading, writing, and mathematics skills than they did at the start of the school year (Lane et al., 2005; Morgan et al., 2010).

Establishing rigorous evidence for special education services through randomized controlled trials (RCTs) is not possible because of the legal entitlement to these services for children meeting eligibility criteria, the small sample sizes, and the distinct categories and

[4] Individuals with Disabilities Education Improvement Act. Pub.L. 108–446.

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severities of disabilities (Hocutt, 1996). Thus, different quasi-experimental approaches (e.g., propensity matching) are used to gauge the impact of special education services on children's outcomes. In one study using propensity-score matching with data from the Early Childhood Longitudinal Study—Kindergarten Class (ECLS-K), 1998–1999, which is a large-scale, nationally representative sample of U.S. schoolchildren, Morgan et al. (2010) examined whether children receiving special education services displayed (a) greater reading or mathematics skills, (b) positive learning-related behaviors, or (c) less frequent externalizing or internalizing problem behaviors than closely matched peers not receiving such services. The results indicated that special education services had either a negative or statistically no significant impact on children's learning or behavior but a small positive effect on children's learning-related behaviors, such as their attention on task. Part of these findings may be due to the settings in which children are receiving services. For example, a review conducted by Ruijs and Peetsma (2009) examining the effects of inclusion on students with and without special education showed neutral to positive effects of inclusive education. Specifically, they found that students with special educational needs performed academically better in inclusive than noninclusive settings, possibly because they could learn from more able students or be more motivated to succeed because of the academic focus. Mixed results were found for socio-emotional development, with some positive outcomes on social and emotional ratings and negative outcomes based on peer perceptions.

Other Issues in Special Education

Disproportionality in identification. Skiba et al.'s (2005) analyses of cross-sectional state-level data indicated that black and Hispanic children were overrepresented in special education. They found this for multiple disability conditions, including intellectual disability, emotional disturbance, speech/language impairment, and learning disability. This supported prior work conducted by Oswald et al. (1999) using cross-sectional, nationally representative, and district-level data showing that minorities were overrepresented in special education, specifically in the mild mental retardation and serious emotional disturbance (SED) categories. Oswald et al. (1999) defined disproportionality as "the extent to which membership in a given ethnic group affects the probability of being placed in a specific special education disability category" (p. 198), such as mild mental retardation. Morgan et al. (2015), using ECLS-K 1998 data, found minority students were underrepresented, which contradicts previous findings. Analyses of ECLS-K data of multiyear longitudinal observations and extensive covariate adjustment for potential child-, family-, and state-level confounds showed that minority children were consistently less likely than similar white, English-speaking children to be identified as having a disability and so to receive special education services. From kindergarten entry to the end of middle school, racial and ethnic minority children were less likely to be identified as having learning disabilities, speech or language impairments, intellectual disabilities, health impairments, or emotional disturbances. Language-minority children were less likely to be identified as having learning disabilities or speech or language impairment (Morgan et al., 2015). Many reasons likely account for the mixed findings, including cross-sectional versus longitudinal data, national compared to state or local data, child- versus school-level data, special education category, and trying to equate white and ethnic minority children when the latter are more likely to experience larger risk factors for developmental delay (e.g., poverty).

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Nutrition Support Programs

For young students in elementary grades, schools are critical settings to support their health, such as in the area of nutrition. The National School Lunch Program (NSLP) was established under the National School Lunch Act (NSLA), signed by President Harry Truman in 1946, to "safeguard the health and well-being of the Nation's children and to encourage the domestic consumption of nutritious agricultural commodities and other foods" (USDA, 2017). The largest of the five school- and center-based programs, the NSLP fed about 30 million children each school day in 2014 and cost $12.7 billion (CBO, 2015). In 2014, 52 percent of school-aged children (ages 5–18) participated in the NSLP, and 23 percent participated in the School Breakfast Program (SBP) (CBO, 2015). Almost half of all lunches served are provided free to students, with an additional 10 percent at reduced prices. Ethnic minority students participate in NSLP at slightly higher levels than white students, and students from low-income households participate at higher rates than those from higher-income households (Ralston et al., 2008). 94 percent of schools, both public and private, choose to participate in the program (though they are not required to offer NSLP meals). NSLP accounts for 17 percent of the total federal expenditures for all food and nutrition assistance programs (Ralston et al., 2008). School meals are required to meet nutritional targets for calories, protein, calcium, iron, and vitamins. Recent changes in standards have made these meals healthier by reducing salt and saturated fat and increasing servings of fruits and vegetables. While such policies should theoretically lead to improvements in health and nutrition, there is emerging evidence that children may be less likely to consume healthier meals. Some schools have also stopped participating in the NSLP because of the increased costs. To be sure, these findings are from only a few studies, and more research on the overall impact on higher nutrition standards is needed (Gundersen, 2015).

Under separate legislation, this program provides free, reduced-price, and full-price breakfasts to students. Other related programs include the Summer Food Service Program (also known as the Summer Meals Program), which extends the availability of free breakfasts and lunches into the summer months in low-income areas; the Special Milk Program, which provides subsidized milk to school; and the After-School Snack Program, which "reimburses schools for healthy snacks given to students in educational after-school programs" (Ralston et al., 2008, p. 5). Food insecurity increases during the summer months, when children do not have access to the NSLP, and the Summer Food Service Program could alleviate this problem. As of 2012, the Summer Food Service Program has a budget of under $400 million and serves a fraction of the children that NSLP serves (Gundersen, 2015). Alternatively, SNAP benefits can be increased during the summer months. Based on one demonstration project, it is estimated that an increase of about $2 billion in SNAP could minimize the spike in food insecurity during the summer (Gundersen and Ziliak, 2014).

Based on the report from Ralston et al. (2008), there is mixed evidence on the impact of NSLP on obesity and nutrition. For example, some studies show that children who participate in NSLP have higher intake of key nutrients and lower intake of sweets compared to nonparticipants. Other studies find high intakes of fat and sodium, which may be due to school programs not following the guidelines about fat and sodium levels. This mixed finding may be due to selection effect. For example, in their examination of NSLP on children's behavior, health, and academic outcomes, Dunifon and Kowaleski-Jones (2003) found that NSLP was associated with an increase in children's externalizing behavior and health limitation (i.e., limitation in being able to participate in regular activities) and a decrease in their math scores. However, once family-level factors associated with the selection of children's NSLP

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participation were adjusted, the effects attenuated. Even after addressing selection effects, there are still mixed findings about the impact of NSLP, with some studies finding participants likely to be overweight (e.g., Schanzenbach, 2009) and others (Gundersen, 2015; Hofferth and Curtin, 2005) finding no effect on obesity. Recent studies to address missing counterfactuals and systematic underreporting of program participation (Gundersen et al., 2012) through causal analytical approaches found some indication of a positive link between NSLP and health outcomes. For example, Gunderson et al. (2012), using data from the 2001–2004 National Health and Nutrition Examination Survey (NHANES) study conducted by the National Center for Health Statistics, Centers for Disease Control (NCHS/CDC), found that NSLP reduced the prevalence of food insecurity by at least 3.8 percent, the rate of poor health by at least 29 percent, and the rate of obesity by at least 17 percent.

While questions remain about the impact of the NSLP on child outcomes, there is consistent evidence about the SBP. Whereas almost all schools participate in NSLP, about 75 percent are part of the SBP, serving 13.2 million children in 2013 (Gundersen and Ziliak, 2014). Frisvold (2015) used the National Assessment of Educational Progress (NAEP) and ECLS-K to determine the impact of SBP on school achievement. Using the NAEP data, he found that availability of SBP increases math by 9 percent of a standard deviation and reading achievement by 5–12 percent of a standard deviation. ECLS-K data show that SBP increases math achievement by 2.7 percent of a standard deviation, reading achievement by 2.0 percent of a standard deviation, and science achievement by 0.9 percent of a standard deviation each school year. Gleason and Dodd (2009) also found a positive effect of SBP but not NSLP on children's health. Specifically, they found that participation in the SBP is associated with lower BMI but saw no evidence for participation in the NSLP and BMI. This association is strongest among white children and not significant for Hispanic children. The mechanism of the effect between SBP and learning is likely through improvement in nutrition, such as increased milk and fruit consumption and decreased soda consumption (Frisvold, 2015).

Beyond the NSLP and SBP, there is a need to conduct in-depth examination of other food and nutrition programs. The Child and Adult Care Food Program (CACFP) reimburses child care programs (both centers and homes) and after-school programs for meals and snacks. Funded at about $3 billion and serving 3.3 million children in 2013, it is much smaller than NSLP and SBP. One study using the ECLS-B dataset found that participating in CACFP was not associated with any changes in the experience of food insecurity (Gundersen and Ziliak, 2014).

Summary

The PPP and ABC studies were seminal because they inspired research on *why* the early years are so important to cognitive and health outcomes. They also inspired the design and implementation of large-scale, publicly-funded ECE programs that the field continues to evaluate, improve, and refine over time—many of which are discussed in this chapter. The evidence linking health equity to ECE programs and services, including Head Start, pre-K, early intervention, special education, and nutrition programs, is mixed. However, the totality of the research including and since PPP and ABC demonstrates effects that are generally positive, albeit small effect sizes and non-findings, in some instances. This indicates that ECE can play an important role in improving health outcomes that could lead to health equity and there is value in continuing to ensure that children have high-quality early learning experiences from birth, though programs may differ in standards, practices, auspices, workforce, dosage, and timing. In order to maximize the impact of ECE, the field will have to understand better how the lessons of

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PPP and ABC apply (or not) in the current context, with a different counterfactual (i.e., more children with access to some ECE), and more diverse children and programs that operate under different funding systems and auspices. The field will also have to learn from contemporary large scale, publicly-funded programs and identify essential ingredients that lead to robust health outcomes and health equity. Only then can ECE be meaningfully part of a health promotion strategy that ensures children with the greatest needs are being served, that their needs are met early and consistently (e.g., through early intervention and nutrition), and in a diversity of settings (e.g., home- or center-based care) through a unifying system that provides high-quality access for all children.

LINKAGES BETWEEN ECE AND HEALTH EQUITY THROUGH SOCIAL-EMOTIONAL DEVELOPMENT

Social-emotional/Behavioral Development

Social-emotional skills, including emotional processes, social/interpersonal skills, and cognitive regulation (Jones and Bouffard, 2012), in young children have been shown to predict later health outcomes and behaviors, such as substance use or abuse, mental health problems (e.g., depression), and teen pregnancy (Conti et al., 2010; Jones et al., 2015; Moffitt et al., 2011).[5] Conversely, externalizing behaviors are associated with later behavioral and academic problems, such as grade retention, school dropout, and lower school engagement (Schindler et al., 2015). Social-emotional skills are especially critical for children and families who experience trauma because of the ways in which such adverse events can impact brain development and affect children's cognitive and healthy development (see Chapter 2 for more details). For these children and families, those adults, professionals, and programs in both early childhood settings and public schools that provide nurturing and safe environments and bolster self-regulation and social-emotional skills can help mitigate the effects of and build resilience in the face of traumatic experiences (Bartlett et al., 2017). (See Box 7-3 for definitions of key terms.)

How effective are ECE programs in providing such environments? Some reviews of ECE programs have found adverse effects on children's social behaviors, especially externalizing behaviors (D'Onise et al., 2010). Analyses of data from the Early Childhood Longitudinal Study (Loeb et al., 2007; Magnuson et al., 2007) and the NIHCD SECCYD (Belsky et al., 2007) found that participation in ECE was associated with more reports of behavior problems, including increased externalizing behaviors and lower self-regulation, but also better cognitive outcomes, such as vocabulary and early reading and math skills. More time in ECE appears to be correlated with more pronounced behavioral issues, and the relationship sometimes persisted beyond kindergarten. See Figure 7-2 for an organizing framework for promoting social-emotional outcomes.

[5] Jones and Bouffard (2012) define these core skills further: "Emotional processes include emotional knowledge and expression, emotional and behavioral regulation, and empathy and perspective-taking. Social/interpersonal skills include understanding social cues, interpreting others' behaviors, navigating social situations, interacting positively with peers and adults, and other prosocial behavior. Cognitive regulation includes attention control, inhibiting inappropriate responses, working memory, and cognitive flexibility or set shifting" (p. 4).

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BOX 7-3
Key Terms Related to Social Emotional Development

- **Socio-emotional development** is the change over time in children's ability to react to and interact with their social environment. It is complex and includes many different areas of growth, such as temperament, attachment, social skills or social competence, and emotional regulation (The Urban Child Institute, 2019).
- **Socio-emotional learning** is the process through which children and adults acquire and effectively apply the knowledge, attitudes, and skills necessary to understand and manage emotions, set and achieve positive goals, feel and show empathy for others, establish and maintain positive relationships, and make responsible decisions (CASEL, n.d.).
- **Executive function** (as along with self-regulation skills) is made up of the mental processes that enable someone to plan, focus attention, remember instructions, and juggle multiple tasks successfully. The brain needs this skill set to filter distractions, prioritize tasks, set and achieve goals, and control impulses (Center on the Developing Child at Harvard University, n.d.).

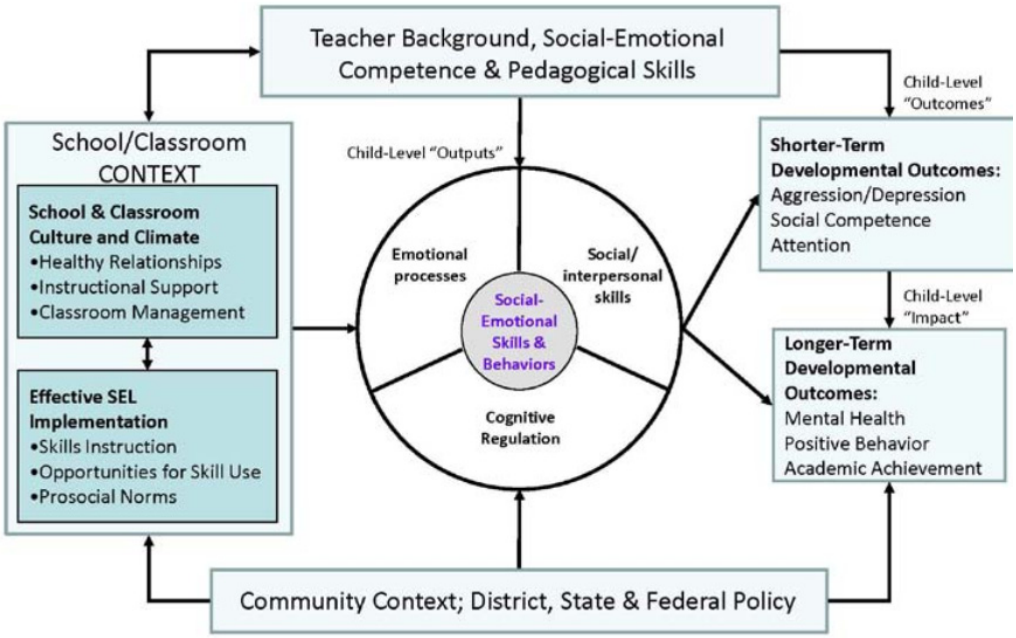

FIGURE 7-2: Organizing framework for social-emotional learning.
SOURCE: Jones and Bouffard, 2012
NOTE: Adapted from collaborative work conducted with Celene Domitrovich as part of the Preschool to Elementary School SEL Assessment Workgroup, Collaborative for Academic, Social and Emotional Learning (CASEL).

However, studies that examined different kinds of ECE or specific interventions in these programs reveal that ECE and public education can be vehicles for cultivating social-emotional skills, and reduce bullying behaviors for older children, through effective professional development, coaching, and use of evidence-based curricula. For example, an evaluation of the Chicago School Readiness Program shows that the combination of training in classroom

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management and job-embedded coaching helped Head Start teachers create more "emotionally and behaviorally supportive classroom environments" and reduced children's emotional and behavioral challenges and improved their EF skills (Raver, 2012, p. 683). Based on the broader body of research, the evaluators further suggested that results like these may lead to "biobehavioral benefits with health impact," such as lower cortisol in reaction to stress and lower risk of obesity (p. 684).

Another teacher training program implemented in Head Start programs, Incredible Years, produced "small but statistically significant improvements in children's knowledge of emotions, social problem-solving skills, and social behaviors" but no impact on children's problem behaviors or EFs (inhibition, working memory, cognitive flexibility), except among those who were exhibiting the most challenging behaviors at the beginning of the school year (Morris et al., 2014, p. 9 of executive summary). The Good Behavior Game is a behavior management program for elementary schools, and evaluations show that children who participated in it were less likely to behave disruptively. As young adults, they were also less likely to receive diagnoses of conduct or personality disorders or use mental health services than a control group (NASEM, 2016c).

Other trainings for educators offer more direct support inside the classroom. The Early Childhood Consultation Partnership provides mental health consultation to ECE programs that serve children from birth through age five in Connecticut. Consultants work with early childhood educators for 8 weeks, at 4–6 hours per week, to improve the social-emotional environment of the classroom, behavior management strategies, and interactions and support for specific children with social-behavioral challenges. An evaluation using randomized assignment showed that preschoolers whose teachers participated in the intervention were rated as exhibiting less externalizing behaviors, such as hyperactivity, restlessness, and impulsivity (Gilliam et al., 2016a).

Research on curricula designed to improve young children's social-emotional skills yields generally positive but sometimes mixed findings. For instance, a study of Head Start programs that were randomly assigned to implement Preschool PATHS showed that it had "small to moderate positive impacts on . . . children's knowledge of emotions, social problem-solving skills, and social behaviors" but no impact on their problem behaviors or EF (Morris et al., 2014, p. 13 of executive summary). Another RCT of the PATHS curriculum led to high levels of social competence, reduced aggression, and improved print knowledge but only for children who started the year with low levels of EF (Bierman et al., 2008). The researchers followed these children for 1 more year, examining 13 child outcomes at the end of kindergarten. In general, they found a sustained, small to moderate impact on social-emotional skills (as opposed to language and literacy), including enhanced learning engagement, improved social competence, and reduced aggression. The effects were especially strong among children who entered low-achieving schools (Bierman et al., 2014).

Another curriculum, Tools of the Mind, was designed to increase EF in young children—especially preschoolers and kindergarteners. The curriculum relies heavily on play-based learning, and teachers who use it with fidelity spend about 80 percent of each day promoting EF. In a study using randomized assignment, children who experienced the curriculum consistently outperformed those who did not on a variety of EF tasks (Diamond et al., 2007). Other studies showed that the curriculum has greater impact on children who may need more support, such as those who attend high-poverty schools (Blair and Raver, 2014) or have issues with hyperactivity or inattention (Solomon et al., 2017). However, Farran et al. (2015) found that children who

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participated in the Tools curriculum generally performed no better than those in the control group in EF tasks, and in some cases, the control group children performed better. Importantly, the researchers found that the teachers in the treatment group spent less than half of their class time on activities from the curriculum. In addition, while these teachers generally implemented the activities, *how* they interacted with children (e.g., time spent on content areas, listening versus talking, positive behavioral reinforcement, scaffolding) was not significantly different than their counterparts in the control group. These findings led the researchers to wonder whether a "reliance on a curriculum to affect child outcomes may be less important than changing interaction patterns in the classroom" (p. 83).

Looking at older, school-aged children, a meta-analysis of 213 school-based social-emotional learning (SEL) programs showed that "students demonstrated enhanced SEL skills, attitudes, and positive social behaviors following intervention, and also demonstrated fewer conduct problems and had lower levels of emotional distress" (Durlak et al., 2011, p. 412−413). For example, Second Step is a violence prevention curriculum aimed at a wider range of children, from age 4–14. Through short (20–50 minute) lessons, classroom management support, and parent training, the program has been found to reduce aggressive behaviors (NASEM, 2016c, p. 198).

Across the above interventions, one key ingredient for success appears to be curricula that are intentionally designed to promote targeted skills and strong training and professional development (McClelland et al., 2017; Morris et al., 2014). In addition, when they are effective, they appear to have compensatory effects for those children who live or learn in more challenging or adverse environments. As described earlier, SEL can encompass a range of skills. Effective curricula or interventions are explicit about which specific skills are targeted and have intentionally designed activities or program components that target those skills. In their meta-analysis of 31 studies that focused on externalizing behaviors, Schindler et al. (2015) found that ECE programs that implemented "enhancements" or interventions that target specific social-emotional competencies were more effective at improving children's behaviors than those that relied on a "global" curriculum that addressed children's learning and developmental domains comprehensively. Durlak et al. (2011) suggest that effective programs use curricula that are "SAFE": sequenced (activities that are designed to connect and build on each other to strengthen social-emotional skills), active (instruction based on active learning strategies), focused (curricula designed to support SEL), and explicit (targeting specific social-emotional skills rather than general development in this domain).

Beyond the design of the intervention or curriculum, the support provided to early childhood educators who implement SEL programs also appears to matter. In their study of Head Start programs, Morris et al. (Morris et al., 2014) attribute successful interventions to high-quality training followed by opportunities to practice new strategies in the classroom and ongoing support from coaches who provide feedback. Raver (2012) and Gilliam et al. (2016a) also emphasize this level of support for educators. However, Schindler et al. (2015) found that interventions that focused more on improving children's social-emotional skills directly (e.g., through specific activities or lessons in a curriculum) are more effective than training teachers to use more effective behavioral management strategies: while "the addition of a caregiver behavior management training program enhancement was not associated with significant reductions in externalizing behavior problems" (p. 253), the "addition of a child social skills training enhancement . . . resulted in half of a standard deviation reduction in externalizing behavior problems . . . which is nearly twice as large as the effect found in a previous study for high-

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quality social and emotional learning programs implemented in primary and secondary schools" (p. 257).

That said, as described above, research on early childhood mental health consultation programs may point to a promising way to improve teachers' performance in the classroom. In a qualitative analysis of six statewide or local early childhood mental health consultation programs, Duran at al. (2009) found that effective programs share three core components:

- A robust infrastructure for implementation, including strong leadership, clear program design, clear organizational structure, effective hiring, training, support and supervision of staff, strong partnerships, evaluation, and funding;
- Highly qualified consultants, defined as having a master's degree in a mental health field, demonstrating core knowledge and skills, and able to develop strong relationships with colleagues, providers, and families; and
- High-quality and comprehensive services that are child centered and targeted to both classrooms and homes, including referral to services that providers and families may need beyond consultation.

In addition, interventions at the classroom or teacher level are more likely to be effective if they are supported by the leadership and broader culture of the school or ECE program. Effective educators are more able to demonstrate their competencies if they work under supportive leadership and policies (IOM and NRC, 2015). Similarly, in its review of the research on antibullying efforts, which focuses mostly on school-aged children, the National Academies Committee on the Biological and Psychosocial Effects of Peer Victimization emphasized the importance of implementing interventions across all school contexts (not just the classroom) and involving all school staff (NASEM, 2016c). For example, the playground or the lunch room can be a "hotspot" for aggressive behaviors as well as an opportunity to promote prosocial interactions. (See Box 3-4 in Chapter 3 for this committee's findings on the effects of bullying in early childhood.)

Finally, it is unclear how critical it is for SEL interventions—whether through a curriculum or consultation—to include a parent education or engagement component. Many programs do so to ensure that the strategies implemented in the classroom are reinforced at home (Duran et al., 2009; McClelland et al., 2017). For example, Fast Track is an intervention for students in grades 1 through 10 that is designed to improve children's social, cognitive, and problem-solving skills by addressing the "interactions of influences" across the school, the home, and the individual (NASEM, 2016c, p. 201). Longitudinal RCTs of the program found that participants showed lower incidence of diagnoses of psychiatric or behavioral disorders through high school and "reduced adult psychopathology at age 25 among high-risk early-starting conduct-problem children" (NASEM, 2016c, p. 201). Unfortunately, the evaluation was not able to disaggregate the impact of the parent engagement component.

Other research had ambiguous findings. In one small, quasi-experimental study of the Incredible Years program, Williford and Shelton (2008) found that parents in the intervention group were more likely to report the use of effective parenting skills, but they did not observe a significantly lower or different level of disruptive behavior when compared with those in the control group. The two groups also did not differ in their experience of stress. The researchers posited that more robust and targeted interventions may be needed for families, as opposed to relying on a supplement to a classroom-based or teacher-focused intervention. In their qualitative review, Duran et al. (2009) also found that "engaging parents/caregivers can be difficult because

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they believe the services are unwarranted, unfamiliar, or stigmatizing, or because various factors impede their ability to actively participate in consultation activities (e.g., transportation, time constraints)" (p. 8 of executive summary).

Perhaps a better approach to strengthening social-emotional development through ECE is to consider the comprehensive array of cross-sector strategies that meet the needs of children and families. The evidence described above suggests that early childhood educators are more effective at promoting social-emotional development when they have access to effective training and consultation and evidence-based curricula. But even when those supports, ECE programs and educators may lack the capacity to fully provide what children and families need, especially recipients who have experienced trauma, chronic stress, or adverse experiences. ECE programs and teachers may need to partner with other community agencies that provide services, such as screening, referrals, and enrollment in programs outside the ECE sector (e.g., mental health, legal, child welfare) (Caringi et al., 2015). Such an approach would be aligned with the way the National Child Traumatic Stress Network conceives of a "trauma-informed child- and family-services system" (Bartlett et al., 2017, p. 8). (See Box 7-8).

An example of such a cross-sector strategy to support the multiple domains of children's development, especially the environments of school-aged children, is through implementation of full-service schools (Zigler and Finn-Stevenson, 2007). Full-service community schools (FSCS) focus on "integrat[ing] academic, health, and social supports with youth and community development strategies," which is especially critical for children and families experiencing multiple challenges. "The goal is to more efficiently use resources to bolster students' learning, strengthen families, and promote healthy communities" (Biag and Castrechini, 2016, p. 157–158; Blank et al., 2003). By coordinating services at school (by colocating or other mechanisms), community schools try address service fragmentation, and encourage more communication and collaboration among providers and educators (Biag and Castrechini, 2016). There is mixed evidence relating FSCS with student outcomes. Using longitudinal data from six high-poverty majority-Latino community schools, Biag and Castrechini (2016) examined how participation in a FSCS influenced students' educational outcomes. The results indicated that participating in FSCS that included family-engagement opportunities and extended-learning programs was associated with modest gains in students' attendance and achievement in math. In another example of FSCS, Whitehurst and Croft (2010) found that the Harlem Children's Zone (HCZ) did not produce higher academic gains than some other charter schools not identified as FSCS. These mixed findings are likely due to the variation in FSCS and their communities (Sanders, 2016), thus indicating a need for more effectiveness studies. Unfortunately, many of the evaluations on FSCS to date have focused on academic outcomes, which calls for an intentional focus on health and socio-emotional outcomes. See Box 7-4 for an example of a promising model that was designed to close the achievement gap by providing wraparound services for families.

BOX 7-4
The Northside Achievement Zone: Promising Model[a]

The Northside Achievement Zone's (NAZ) goal is to permanently close the achievement gap and end generational poverty in North Minneapolis. Together with its partner organizations, NAZ works with low-income families as they put their children on a path to college. NAZ's wraparound framework effectively supports low-income children of color so that they will graduate from high school prepared for college—the programming begins in early childhood.

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To end the achievement gap, NAZ offers family classes; early childhood education scholarships; general improvements to public, public charter, and parochial schools in the area to support academic excellence for all children; after-school and full-day summer tutoring programs; and health and wellness programs. The health and wellness focus is rooted in the knowledge that behavioral issues are often related to traumatic experiences or adverse childhood experiences (ACEs) and can result in removal or expulsion from schools and after-school programs. A licensed clinical social worker meets with families who have experienced trauma and/or children who present behavioral health issues and follows a diagnostic process to help support children and their families.

The target age group is young children (ages 0–11) and their families, with some other developing programs supporting adolescents and young adults through college. The program, which was first established in 2003, originally focused on reducing violence in North Minneapolis. In 2008, residents, community organizations, and NAZ joined forces to find solutions to "seemingly intractable" issues in their neighborhood. Together, they developed the model of wraparound services for North Minneapolis's low-income families to close the "achievement gap."

By providing wraparound services focused on education, career training and financial education, and housing stabilization for children and their families, NAZ addresses several social determinants of health (SDOH), including education, employment, the social environment, housing, income and wealth, and public safety.

The program is a collaborative effort among a broad range of nonprofit direct service providers, nonprofit advocacy organizations, school systems (parochial, public charter, and public district), universities, county government agencies, and Minneapolis's Public Housing Authority. These partners provide critical components of best-practice supports from prenatal development through college.

As of early 2019, NAZ has proven to reach the families that are furthest behind and in greatest need—73 percent of NAZ families make less than $30,000 per year; 98 percent are families of color, and 79 percent are African American. Between the 2016–2017 and 2017–2018 school years, NAZ family and scholar participation increased by almost 15 percent. NAZ data also support two or more layered strategies for closing the achievement gap—NAZ scholars with two or more layers of educational support (coach, expanded learning opportunities, anchor school) are more than twice as likely to be proficient in both reading and math, compared to just one layer.

SOURCE: Northside Achievement Zone, 2019, n.d.
NOTE: Data provided by program results summary.

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

Other school/education reform efforts found to have an effect on children's learning and behavior are the Comer School Development Program (SDP) and the 21st Century Community Learning Centers (21st CCLC). The Comer SDP was developed by Dr. James Comer and the Child Study Center at Yale University in 1968 to improve the educational experiences of low-income ethnic minority children. This model includes three mechanisms (School Planning and Management Team, Student and Staff Support Team, and Parent/Family Team); three operations (Comprehensive School Plan, Staff Development Plan, and monitoring and assessment); and three guiding principles (collaboration, consensus decision making, and no-fault problem solving) (Lunenberg, 2011). SDP is implemented in 1,150 schools across the world. Studies of SDP schools show significant student gains in achievement, attendance, behavior, and overall adjustment (Lunenburg, 2011). It is theorized that the SDP model effect manifests through improvement in school climate, indicated by improved relationships among staff and students, collaboration among staff, and central focus on students (Lunenberg, 2011). Quasi-experimental design studies have shown that students in SDP schools in comparison to students in matched

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non-SDP schools showed significant gains in achievement, behavior, and overall school adjustment (Haynes and Comer, 1990a,b). Studies have emphasized the importance of implementation of key components of the SDP to find evidence of effectiveness (Cook et al., 1999; Haynes et al., 1998).

The 21st CCLC program provides students in high-poverty communities across the nation with the opportunity to participate in academic enrichment and youth development programs designed to enhance their well-being (U.S. Department of Education, 2010). Programs provide the following activities: academic enrichment learning programs; tutoring; supplemental educational services; homework help; mentoring; recreational activities; career or job training for youth; drug and violence prevention, counseling, and character education programs; expanded library service hours; community service or service-learning programs; and activities that promote youth leadership (U.S. Department of Education, 2010). Studies of 21st CLCC have been mixed, with limited significant findings associated with gains in achievement scores and some gains in parental involvement in school and student commitment to work (Durlak and Weissberg, 2013). In their review of after-school programs beyond the 21st CLCC, Durlak and Weissberg (2013) found four evidence-based practices that formed the acronym SAFE: (S) step-by-step training approach, (A) emphasis on active forms of learning through practicing new skills, (F) focused specific time and attention on skill development, and (E) explicitness in defining the skills being promote. Students who participated in SAFE programs had more positive social behaviors (effect size of .29 versus.06 for other programs), reduction in problem behaviors (effect size of .30 versus .08 for other programs), school grades (effect size of .22 versus .05 for other programs), self-perception (effect size of .37 versus .13 for other programs), and academic achievement (effect size of .20 versus .02 for other programs). Thus, there is a need to examine and incorporate these evidence-based approaches in all aspects of learning during the school day or extended day.

Socio-emotional Learning, Trauma-informed Care, and Suspensions and Expulsions

Suspensions and expulsions in ECE or K–12 settings are often used as a deterrent for misbehavior, which could be due to many factors, including learning disabilities and social-emotional needs in response to trauma, chronic stress, and adverse experiences. “Suspension” is either out-of-school or in-school suspension, which often lasts from 1 to 10 days but varies across states and localities. “Expulsion” is the permanent removal of students from an ECE or school setting. The U.S. Department of Education, Office of Civil Rights (2016) found that students with disabilities served by IDEA (11 percent) are more than twice as likely to receive one or more out-of-school suspensions compared to students without disabilities (5 percent). More than one out of five American Indian or Alaska Native (AI/AN) (22 percent), Native Hawaiian or other Pacific Islander (23 percent), black (23 percent), and multiracial (25 percent) boys with disabilities served by IDEA received one or more out-of-school suspensions, compared to 1 out of 10 white (10 percent) boys with disabilities served by IDEA. Black students are 2.3 times as likely to receive a referral to law enforcement or be subject to a school-related arrest as white students (U.S. Department of Education, Office of Civil Rights 2016), emphasizing the need for trauma-informed approaches in schools.

Turning to children in general education, more than 250 preschoolers are suspended or expelled daily (Malik, 2017). Estimates show that as many as 8,710 of 3- and 4- year-old children may be expelled from or pushed out of their state-funded preschool or pre-K classrooms annually—a rate nearly three times that of students in kindergarten through 12th grade. In child

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care centers, expulsion rates are 13 times what they are in kindergarten through 12th grade. These rates are particularly pernicious for black children, who are suspended and expelled at much higher rates than their peers. Black preschoolers are 3.6 times more likely to receive one or more suspensions than white preschoolers (Meek and Gilliam, 2016). Black children represent only 19 percent of preschool enrollment but are 47 percent of children receiving one or more out-of-school suspensions; in comparison, white children are 41 percent of preschool enrollment but 28 percent of children receiving one or more out-of-school suspensions (Boyd et al., 2018; Meek and Gilliam, 2016).

Various scholars, and more recently, Gilliam et al. (2016b), suggested that teachers' implicit bias may be the underlying cause for this "pushout" of black children, especially boys, from these early learning settings, as teachers view black children as older and more culpable (Goff et al., 2014). This biased views and attitudes about black children are likely to contribute to children's disengagement in school. In a meta-analytic study using 53 studies examining the link between suspension and student outcomes, Noltemeyer et al. (2015) found an inverse relationship between suspension and achievement and a positive relationship between suspension and school dropout. Effect sizes ranged from .10 for in-school suspension and .24 for out-of-school suspension to .32 for the effect of suspension on achievement tests. There was an effect size of .28 for the effect of suspension on school dropout. Noltemeyer et al. (2015) also found that the percentage of male ethnic minority students and family socioeconomic status (SES) moderated the relationship between suspension and achievement; this was also the case for school dropout. This indicates that the exclusionary practices are not beneficial for students' achievement and learning process and may heighten risk for children from ethnic and low-SES households. These practices are especially harmful when they contribute to pushing students of color into the school-to-prison pipeline (see Box 7-5 for more information).

BOX 7-5
The School-to-Prison Pipeline

The school-to-prison pipeline refers to "policies and practices that are directly and indirectly pushing students of color out of school and on a pathway to prison" (National Education Association, n.d.). These policies and practices "disproportionately [place] students of color, including those who identify as LGBTQ, have disabilities, and/or are English language learners, into the criminal justice system for minor school infractions and disciplinary matters, subjecting them to harsher punishments than their white peers for the same behaviors" (National Education Association, n.d.).

Disproportionate rates of school discipline, such as suspensions and expulsions, in ECE are one means through which children of color are driven to the criminal justice and welfare systems. Based on Department of Education (ED) national civil right data from the 2013–2014 school year, a GAO analysis (2018) found that black students, boys, and students with disabilities experienced disproportionately high rates of suspension and expulsion (GAO, 2018). For detailed information on the welfare and criminal justice systems and their effects on adolescent health and well-being through a life course perspective, see the 2019 National Academies report *The Promise of Adolescence: Realizing Opportunity for All Youth*.

Addressing the school-to-prison pipeline and its deleterious effects on the health, well-being, and development of children of color (into adolescence and across the life course) requires attention from ECE providers, as the origins of the pipeline are often established in ECE settings through disproportionate rates of discipline to young children of color, particularly boys. ECE providers can help to dismantle the pipeline through the aforementioned strategies—specifically, countering unconscious biases and cultural stereotyping and fostering greater cultural competency through trainings on antiracism, implicit bias, and mindfulness.

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In response to suspension and expulsion, there have been two general approaches for training professionals—trauma-informed care (TIC) and implicit bias training (See the section on supports for the ECE professional for an overview of both approaches to help address suspension and expulsion issues).

Social-emotional Learning and Dual Language Learners

Research suggests that children who are bilingual tend to have better self-regulation skills than their monolingual peers (Castro et al., 2013). They have enhanced ability to control their attention and greater EF skills, such as planning, working memory, and cognitive flexibility. Researchers attribute this advantage to bilingual children's experiences at paying attention to their environment and other contextual cues to understand when to use one language over another (Castro et al., 2013). A 2017 National Academies report concluded that DLLs "who have a strong base in their [home language] and acquire high levels of English proficiency will realize the cognitive, linguistic, social, and cultural benefits of becoming bilingual" (NASEM, 2017b) (p. 175). Conversely, DLLs' loss of proficiency in their home language may also compromise their social-emotional development. For instance, "children who do not develop and maintain proficiency in their home language may lose their ability to communicate with parents and family members and risk becoming estranged from their cultural and linguistic heritage" (NASEM, 2017b, p. 175).

Thus, to strengthen DLLs' social-emotional health and self-regulation, it is important for ECE and elementary schools to foster their home language skills. Research also shows that when educators implement evidence-based practices that supports bilingual development, DLLs' maintenance of their home language does not come at the cost of their proficiency in English (NASEM, 2017b).

LINKAGES BETWEEN ECE AND HEALTH EQUITY THROUGH PARENTING AND THE HOME ENVIRONMENT

Another crucial pathway through which ECE leads to health equity is the family and home environment (see Chapter 4 for the importance of family cohesion and support). As indicated in the pathway by Hahn et al. (2016), ECE programs support a positive, enriching, and stable home environment through parents' participation in educational, social, health, and job training opportunities. This pathway can lead to improvement in children's and families' cognitive, social, and health outcomes, which then impacts children's outcomes that lead to higher education attainment and health equity. This pathway is particularly supported by the National Academies report *Parenting Matters: Supporting Parents of Children, Ages 0 to 8 Years Old* (NASEM, 2016b), which emphasized the evidence about strengthening parental knowledge, attitudes, and practices to improve children's cognitive and socio-emotional well-being. Specifically, it found that parental knowledge of child development is positively associated with parent behaviors. Parents with knowledge of evidence-based parenting practices, especially those related to promoting children's physical health and safety, such as vaccination and use of seat belts, are more likely to engage in those practices.

Ample evidence exists that parental knowledge about of how to meet their children's basic physical (e.g., hunger) and emotional (e.g., wanting to be held or soothed) needs, as well as of how to read infants' cues and signals, ensures proper child growth and development (Bowlby,

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2008; Chung-Park, 2012; Regalado and Halfon, 2001; Zarnowiecki et al., 2011). Parents' attitudes about specific practices, such as breastfeeding and engagement in children's education, was related to parents' behaviors and use of services. They found strong evidence that the following parenting practices are important to children's physical health and safety and emotional and behavioral, social, and cognitive competence: (a) contingent responsiveness ("serve and return")—adult behavior that occurs immediately after a child's behavior and is related to the child's focus of attention, such as a parent smiling back at a child; (b) warmth and sensitivity; (c) routines and reduced household chaos; (d) shared book reading and talking to children; (e) practices that promote children's health and safety—in particular, prenatal care, breastfeeding, vaccination, ensuring adequate nutrition and physical activity, monitoring, and household/vehicle safety; and (e) use of appropriate (less harsh) discipline (McClelland et al., 2017).

The pathway linking ECE to health equity through supporting responsive parenting and a quality home environment is generally supported by many programs, including the three foundational early childhood programs, each of which included a significant family-engagement component—the HighScope PPP, focused on 3- and 4-year olds, the Carolina Abecedarian Study, focused on children birth to age 3, and the Chicago Child-Parent Centers (CPC) (see Box 7-6). While these studies are quite old, focused on particular populations (i.e., African Americans), and specific to local areas, in addition to being unable to disentangle the impact of teacher practices compared to family-engagement practices, they provide long-term and intergenerational evidence of the importance of engaging with families while also providing high-quality learning experiences in classrooms. Other early childhood programs, including home visiting programs that serve mothers prenatally and last until the children are 3 years old and family-engagement programs for preschool- and school-age children, provide additional support for the links between ECE and healthy equity through parenting.

BOX 7-6
Chicago Child-Parent Centers

The Chicago CPC "began in the Chicago public schools in 1967 through federal funding from the Elementary and Secondary Education Act of 1965. . . . The 24 centers provide comprehensive services under the direction of the Head Teacher and in collaboration with the elementary school principal, as well as parent resource teachers, the school-community representative, bachelor's level classroom teachers, nurses, speech therapists, and school psychologists" (Temple and Reynolds, 2007, p. 133–134). The CPC's comprehensive services included an intensive parent-involvement component, outreach, and attention to health and nutrition. Participation in the CPC preschool intervention relative to the usual enrichment program was associated with significantly higher rates of school completion by age 24 years, significantly lower rates of juvenile arrest for both violent and nonviolent offenses, lower rates of school remedial services, stable employment, health insurance coverage, and lower likelihood of depressive symptoms (Temple and Reynolds, 2007). School-age intervention was associated with lower rates of school remedial services and receipt of public aid. Extended intervention for 4–6 years was linked to significantly lower rates of remedial education, juvenile arrests for violent offense, receipt of disability assistance, and private insurance coverage.

Family-engagement Programs

The focus on family engagement has been bolstered by the recent federal funds for parent and family community engagement centers (U.S. Department of Education, Statewide Family

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Engagement Center Program Through the Consolidated Appropriations Act of 2018). As reviewed in the National Academies *Parenting Matters* (2016b) report, some ECE programs "provide full- or part-time classroom-based services (center or family child care) for children from birth to age five" that "often include parenting education and other services for families (sometimes starting prenatally)" (p. 159). To address the overall conditions of families, these programs are designed to improve parenting knowledge, attitudes, and practices with the goal of supporting children's cognitive and social-emotional development and success in school (Brooks-Gunn et al., 2000; Chase-Lansdale and Brooks-Gunn, 2014; Fantuzzo et al., 2013). These parent-focused programs have several different structures. For example, some include parent supports (and parent self-sufficiency support) as well as intensive classroom-based services for children that are multipronged. Others offer services that are primarily classroom-based with either some parenting education services or some parent self-sufficiency services (NASEM, 2016b).

In their examination of the impact of Head Start on child outcomes through parenting, Puma et al. (2012) did not find significant differences on parenting-related measures, included disciplinary practices, educational supports, parenting styles, parent participation in and communication with the school, and parent and child time together (NASEM, 2016b). With respect to Early Head Start, there was some evidence that programs that were mixed delivery (center and home) had favorable and consistent impact on parenting outcomes, including sensitive parenting and language-rich environments.

Other school-based parent engagement programs have been found to be effective for children's outcomes. For example, the Companion Curriculum uses Head Start teachers to encourage parents' participation in the classroom and provides workshops and activity spaces in the classroom that focus on training parents to engage in parent–child learning activities (NASEM, 2016b; Mendez, 2010). There were no benefits for parents, but there was significant improvement in children's vocabulary in a quasi-experimental study (Mendez, 2010). Another example is documented in the National Academies *Parenting Matters* report (p. 163):

> The Kids in Transition to School (KITS) Program is a short-term, targeted, evidence-based intervention aimed at increasing early literacy, social skills, and self-regulatory skills among children who are at high risk for school difficulties. This program provides a 24-session readiness group for children that promotes social-emotional skills and early literacy as well as a 12-session parent workshop focused on promoting parent involvement in early literacy and the use of positive parenting practices. In a pilot efficacy trial with 39 families, Pears and colleagues found that children in families who received the KITS intervention demonstrated early literacy and social skill improvements as compared with their peers who did not receive the intervention (Pears et al., 2014). In randomized controlled studies, foster children who received the intervention exhibited improvements in social competence, self-regulation skills, and early literacy skills (Pears et al., 2007, 2012, 2013).

Two-generational model programs, such as Head Start, Project Redirection, the New Chance Demonstration, Ohio's Learning and Earning Program, the Teen Parent Demonstration, and the Comprehensive Child Development Program from the 1980s and 1990s, offered mothers a wide range of services, including parenting classes, job training, mandatory schooling, and child care (Granger and Cytron, 1999; NASEM, 2016b; Polit, 1989). Many of these programs have not

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resulted in significant positive outcomes, at least based on older versions of the models (NASEM, 2016b). The new iterations of two-generation models focus "on the education benefits to children of high-quality ECE programs and higher parental levels of education and labor force motivation. Parenting knowledge, attitudes, and practices may be improved, but the improvement comes indirectly through higher parental job skills and education and reduced household stress rather than explicit programming directed at parenting skills" (NASEM, 2016b, p. 166). Examples of these new iterations that have evaluations planned or under way are the CareerAdvance Community Action Project of Tulsa, Oklahoma; the Annie E. Casey Foundation Atlanta Partnership; and the Housing Opportunity and Services Together project (see Chase-Lansdale and Brooks-Gunn, 2014 for others) (NASEM, 2016b).

There are also other family-focused interventions for school-age children. In one example, the ParentCorps program, parent groups (see Box 7-7; also documented in NASEM, 2016b)—"co-facilitated by teachers and mental health professionals with expertise in behavior management—are used to help parents establish structure and routines for children, teach positive parenting practices (e.g., positive reinforcement and consistent consequences), and provide opportunities for facilitator-observed parent–child interactions" (NASEM, 2016b, p. 171). In addition to reduced behavior problems among children in the treatment group, parents displayed more effective parenting practices than in the control group. Parents in the treatment group reported using more effective disciplinary practices and receiving higher scores on tests of knowledge of effective behavior management strategies; in addition, higher-quality parenting was observed for parent–child interactions by the research team (Brotman et al., 2011).

As documented in the National Academies *Parenting Matters* report (2016b), Getting Ready is an evidence-supported intervention that targets parents' decision-making role at school (Sheridan et al., 2010). The program includes parent-teacher conferences, monthly family socialization, and teacher home visits with parents using structured interactions. The goal is to actively engage parents in learning and behavior goal setting and decision making. "Together, teachers and parents identify learning opportunities at home and school and plan how educators and parents can complement each other's efforts to promote learning and track children's growth. Priorities include affirming parents' competence, increasing their access to information on child development, and reinforcing positive parenting practices" (NASEM, 2019b, p. 172; Knoche et al., 2012). Knoche et al., (2012) identified treatment effects for parental warmth and sensitivity, learning support, and autonomy support. Sheridan et al., (2014) found that, relative to children in the control group, children in the Getting Ready intervention had a significantly greater decline in disruptive behaviors such as difficulty standing still and tendency to run around; however, no differences were seen for other learning-related behaviors. Children's language and literacy were also improved, and one study found some evidence that the program's effects on achievement were greatest for children at highest risk for underachievement (that is, those children whose parents have less than a high school education and those who did not speak English prior to treatment) (Sheridan et al., 2011). Thus, there is some evidence that schools' partnership and engagement with parents through specific and structured approaches are related to improved parenting and child academic and behavioral outcomes; unexpectedly, these outcomes seem stronger in older age groups. However, caution is warranted because these are not national studies and, in many instances, are localized and researcher controlled, unlike the Head Start and Early Head Start studies.

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BOX 7-7
ParentCorps: Promising Model[a]

ParentCorps is a universal school-based program for all children in pre-K or early childhood settings that aims to help students develop foundational skills for learning. The program takes a family-centered approach by building on the strengths of culturally diverse families and engaging parents as partners. ParentCorps is now offered as an "evidence-based enhancement" to Pre-K for All programs throughout New York City (NYU School of Medicine, 2019). ParentCorps includes three key components that are designed to strengthen home–school connections and provide high-quality (i.e., safe, predictable, and nurturing) environments for students based on the scientific evidence that this will help students to develop strong social, emotional, and behavioral regulation skills.

1. Program for pre-K students ("Friends School"): 14- week curriculum for students on social, emotional, and behavioral regulation skills (materials offered in English and Spanish).
2. Program for parents of pre-K students: 14-week program for parents to promote the use of evidence-based strategies for enhancing social, emotional, and behavioral regulatory skills (materials offered in English and Spanish).
3. Professional development for staff: group and individual learning opportunities for pre-K and kindergarten teachers and assistants, mental health professionals, parent support staff, and school leaders to promote the use of evidence-based strategies for enhancing home–school connections and to strengthen social, emotional, and behavioral regulation skills.

ParentCorps works to buffer the effects of early adversity, such as poverty, by engaging and supporting parents and early childhood professionals in the community. The program takes a multisector approach by engaging professionals in ECE and health. The primary SDOH that it targets are education, the social environment, and health services. ParentCorps has been found to yield positive outcomes for children through age eight in urban schools in areas of concentrated poverty. These include improvements in areas of academic achievement, behavioral outcomes, and obesity outcomes (Brotman et al., 2011, 2012, 2013, 2016; Dawson-McClure et al., 2015). In addition, a cost-effectiveness analysis estimated $4,387 long-term cost savings per individual in health care, criminal justice, and productivity expenditures after factoring program costs and increased life expectancy by 0.27 quality-adjusted life years (Hajizadeh et al., 2017).

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

Home Visiting Programs

One particular early childhood program focused on supporting parents and enhancing positive parenting and reducing/preventing child abuse and maltreatment is home visiting (see Chapter 4 for additional background on home visiting, including health impacts and a committee recommendation). Home visiting (which targets families with pregnant women and children from birth through age 5—though most attend until children are 3 years old) has been found to increase parental knowledge and practices and reduce parental stress and depression while also supporting child health and reducing maltreatment. The general short-term outcomes of home visiting programs include (1) decreasing parent stress, depression, and isolation; (2) increasing family self-sufficiency; (3) enhancing parenting knowledge of child development; (4) increasing self-efficacy in the parenting role; (5) healthier parent–child relationships and interactions; (6) increasing positive guidance and decreasing harsh punishment; and (7) supporting children's learning. The long-term outcome is to enhance children's well-being, social competence, and school readiness.

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Based on strict criteria for what counted as evidence and a systematic review of that evidence (adapted in part from the U.S. Department of Education's What Works Clearinghouse),[6] the U.S. Department of Health and Human Services' (HHS) Home Visiting Evidence of Effectiveness (HomVee) identified many effective models that impact various outcomes, including positive parenting practices, family economic self-sufficiency, child health, child development and school readiness, and reductions in child maltreatment. Regarding positive parent practices, the review shows "while many individual evaluations of home visiting programs have shown impacts on parenting practices tied to positive developmental outcomes, the average impacts of home visiting on parenting practices are not large. Nor is there a strong pattern of effects on parenting practices across evaluation studies and home visiting models" (NASEM, 2016b, p. 151). None of the home visiting models were linked to reductions in juvenile delinquency, family violence, and crime. Healthy Families America, however, had one or more favorable impacts in each of the eight domains, and Nurse-Family Partnership (NFP) had favorable impacts in seven domains, followed by Early Head Start—Home Visiting with favorable impacts in five domains. See the National Academies *Parenting Matters* report for synthesis of home visiting and various models, and the Home Visiting Evidence for Effectiveness website by the HHS, Administration for Children and Family for a recent examination of all home visiting models.[7] These findings are consistent with the 2019 results from the MIHOPE and MIHOPE—Strong Start Studies of Evidence-Based Home Visiting, namely, that MIHOPE found positive effects on some family outcomes (e.g., home environment) but MIHOPE—Strong Start found little effect on birth outcomes and prenatal behaviors (Michalopoulos et al., 2019).

In summary, there is some evidence linking family engagement, including home visiting and support, to children's health and well-being. While it is challenging to disentangle family engagement and support from ECE classroom practices, there is indication that supporting family functioning and processes and providing resources and strategies will likely lead to better outcomes for parents themselves and their children, resulting in health equity. Current evidence indicates the importance of tailoring services and programs to meet the needs of individual families and ensuring that families and children with the greatest needs are engaged throughout the duration of the program/services. With new approaches to family engagement and support, such as the Two Gen 2.0 models, more actionable evidence is emerging about how best to support children and their families in the early years.

LINKAGES BETWEEN ECE AND HEALTH EQUITY THROUGH SUPPORTS FOR THE ECE PROFESSIONAL

The salutary effects that ECE can have on the physical, mental, and social-emotional health of children—whether through the inclusion of comprehensive services, evidence-based curriculum, high-quality professional development and supports, or effective family engagement—can be disrupted if the ECE professionals themselves are not trained or supported to implement best practices or navigate systems of services for children and families outside the

[6] For more information, see Institute of Education Sciences. n.d. *What Works Clearinghouse. Find What Works*, https://ies.ed.gov/ncee/wwc/FWW (accessed July 15, 2019).
[7] For more information, see U.S. Department of Health & Human Services. n.d. *Home Visiting Evidence of Effectiveness*, https://homvee.acf.hhs.gov (accessed June 14, 2019).

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ECE sector. Since the release of the National Academies report *Transforming the Workforce for Children Birth Through Age 8: A Unifying Foundation* (IOM and NRC, 2015), there has been increased interest from funders, policy makers, and advocates in overhauling the professional preparation and ongoing supports for early childhood educators so that these experiences—and the policies and funding that shape them—better reflect the science of early development.[8] For example, states and national organizations have developed statements of core professional competencies that inform the training of these educators. The report found that, in most cases, these statements need to be updated to follow recent developments in science and research. Some of these areas are especially relevant to promoting young children's health and well-being, such as knowledge and skills related to working effectively with children who have experienced chronic stress, trauma, and adversity; collaboration with professionals about of the ECE or education sector; promotion of self-regulation and related EFs, and support for DLLs. Incorporating these issues into efforts to improve policies, systems, and programs that prepare and support early childhood educators is critical to maximizing the potential of ECE programs to improve health outcomes and equity. For example, teachers, especially those who work with students who experience trauma, can be more effective if they receive more training and education on mental health issues and skills to access relevant services from that sector (Hydon et al., 2015).

Training and supports for early childhood educators are needed to help them adjust their practice to acknowledge their own biases and their understanding that children's behavioral challenges and learning difficulties are often due to the toxic stress response and undiagnosed and untreated trauma (Matthews et al., 2018; Ramirez et al., 2012). It is important to note that these trainings are critical for supporting children's learning and healthy development beyond addressing suspension and expulsion. Whether or not children are suspended or expelled, many are exposed to traumatic experiences, including abuse and neglect, family and media violence, community or school violence, loss of a parent, parents dealing with substance abuse, and mental or other health challenges. In turn, children who experience trauma can exhibit behaviors in ECE programs and schools that may indicate socio-emotional challenges.

BOX 7-8
Trauma-informed Child- and Family-service Systems

A trauma-informed child- and family-service system is one in which all parties recognize and respond to the impact of traumatic stress on those who have contact with the system, including children, caregivers, and service providers. Programs and agencies within such a system infuse and sustain trauma awareness, knowledge, and skills into their organizational cultures, practices, and policies. They act in collaboration with all those who are involved with the child, using the best available science, to facilitate and support the recovery and resiliency of the child and family.

SOURCE: The National Child Traumatic Stress Network, n.d.

Trauma-informed Care or Practices

TIC is defined as a "recognition of the pervasiveness of trauma and a commitment to identify and address it early . . . [and it] involves seeking to understand the connection between

[8] See initiatives from national organizations, such as Power to the Profession, National Governors Association, and Council of Chief State School Officers, and state-led initiatives, such as in Colorado, Minnesota, and Nebraska.

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presenting symptoms and behaviors and the individual's past trauma history" (Hodas, 2006, p. 5) (see Box 7-9 for more information). TIC can be provided in multiple settings by trained, committed professionals who understand the principles of a TIC system, including

a. trauma and how it may impact children's identity, how they will view the world, and coping mechanisms;
b. children and their contexts, including their family and community contexts, and supporting children to be active in their healing process;
c. services that are strengths based and promote children's self-control and coping skills; and
d. the service relationship, which is based on relationships and trust that is earned over time.

BOX 7-9
Trauma-Informed Care Strategies for Educators

The following TIC strategies for educators are adapted from the National Child Traumatic Stress Network (NCTSN) child welfare trauma training toolkit.

1. Develop a crisis plan with resources.
2. Maximize the child's sense of safety.
3. Assist children in reducing overwhelming emotion.
4. Help children make new meaning of their trauma history and current experiences.
5. Address the impact of trauma and subsequent changes in the child's behavior, development, and relationships.
6. Coordinate services with other agencies.
7. Utilize comprehensive assessment of the child's trauma experiences and their impact on the child's development and behavior to guide services.
8. Support and promote positive and stable relationships in the life of the child.
9. Provide support and guidance to the child's family and caregivers.
10. Manage professional and personal stress.

SOURCE: Child Welfare Committee, National Child Traumatic Stress Network, 2013.

There is emerging evidence about the impact of TIC and trauma-informed practices (TIP) for children's outcomes and well-being. For example, Zakszeski et al. (2017) conducted a review to describe the implementation and evaluation of trauma-focused school practices. They found that most approaches used Cognitive Behavioral Intervention for Trauma in Schools (CBITS), followed by drama instruction, eye movement desensitization and reprocessing training, coping/social skill instruction; and comprehensive, multitiered systems. Zakszeski et al. (2017) found that most of the studies reported positive treatment outcomes, especially reduction in symptoms of trauma and internalizing problems, but effect sizes were not included to determine magnitude. The limitation of the studies in this review was that most occurred with subpopulations of children rather than across the student population. There has been a call for more rigorous study and review of TIC and TIP, beyond their implementation (Bryson et al., 2017). See Box 7-10 for an example of a promising model that employs a cognitive behavioral intervention to address trauma in schools.

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BOX 7-10
Cognitive Behavioral Intervention for Trauma in Schools: Promising Model[a]

CBITS is a mental health intervention that was designed and first implemented in 2001 to target students from 5th through 12th grade who have witnessed or experienced traumatic life events (CBITS, n.d.-a). CBITS uses cognitive behavioral techniques, implemented in a school setting. The program includes 10 group sessions, 1–3 individual therapy sessions, 2 parent psychoeducational sessions, and 1 teacher educational session.

CBITS works to address posttraumatic stress symptoms in students who are 10–18 years old. The program uses cognitive behavioral therapy techniques to address a growing issue: 20–50 percent of children in the United States are affected by violence, as victims or witnesses, and even more are exposed to natural disasters, accidents, and traumatic losses (Stein et al., 2011). These experiences, which disproportionately affect low-income and minority children, can cause posttraumatic stress symptoms, leading to behavioral problems, poorer school performance, more days of school absence, and depression and anxiety (Stein et al., 2011). CBITS addresses several SDOH (health systems and services, education, and the social environment) and engages multiple sectors (mental health providers, nonprofit community-based organizations, public schools, SAMHSA and other federal agencies, state governments, and universities).

Evidence suggests that CBITS is effective, and it has been implemented widely across the United States and abroad (Stein et al., 2011). Journey Mental Health Center in Dane County, Wisconsin, has successfully implemented CBITS in local public schools since 2004, and feedback consistently shows that it reduces symptoms, increases the potential for positive school performance, and creates a forum for open communication with students (CBITS, n.d.-b). The University of Maryland Center for Mental Health has also used CBITS in Baltimore schools since 2004. CBITS was met with such enthusiasm from participants and providers that CMH has continued provide annual trainings for its clinicians and begun conducting trainings for mental health practitioners at schools, many of whom are actively using CBITS at the request of the Maryland State Department of Education. When CBITS was implemented between 2008 and 2010 in more than 100 schools in the Los Angeles Unified School District, 81 percent of participants across grades 5–10 reported improvement in PTSD symptoms, with 63 percent falling below the clinical range.

[a] The committee used selection criteria to identify examples of promising models highlighted in this report (see Appendix A for a list of the criteria). These examples all apply developmental science and aim to advance health equity during the preconception through early childhood periods.

Implicit Bias Training

Implicit biases are unconscious and involuntary attitudes that can influence one's affect, behavior, and cognitive processes (Boysen, 2010). It has been suggested that implicit bias is one reason for the disproportionality of suspension and expulsion rates for black children and children with special needs. That is, educators may see the normative behaviors of black children in particular as dangerous and aggressive, even when there is no evidence of misbehavior (Gilliam et al., 2016b). Implicit bias training may be a potential strategy to counter these unconscious biases. Evidence is emerging on the impact of implicit bias training, mostly from the public health and nursing sectors. One core feature of implicit bias training is cultural competence, which "is the process and ability of an individual or organization to function effectively within different cultural situations" (Betancourt et al., 2003; Cross et al., 1989; Gallagher and Polanin, 2015, p. 333); it "combines a set of congruent behaviors with attitudes and knowledge that facilitate an individual or a system to work successfully in various cultural contexts other than their own culture" (Gallagher and Polanin, 2015, p. 333). See Box 7-11 for more information on the importance of cultural competence and sensitivity in ECE.

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BOX 7-11
Integrating Cultural Competence and Sensitivity into ECE

Beyond implicit bias training, it is important to broadly integrate cultural competency and sensitivity into ECE policies and practices to make progress on health equity. Chapter 4 discusses the role of culture as a contextual determinant of development. Based on this, ECE is an optimal platform to embrace and build from cultural practices of the populations being served. Integrating cultural competency can provide continuity between the home environment and the ECE setting and an opportunity to engage parents as co-teachers. Given the cultural differences that some children may experience, their understanding of specific behaviors and learning methods will inevitably differ based on their community of origin. For example, based on community values, a child may learn most effectively from child-directed interactions, observing other social actors, or observing third-party interactions in which they are not involved. An understanding of cultural norms for certain groups of children would best position ECE professionals to incorporate specific teaching practices that are consistent with what the child is experiencing at home.

One example of cultural competence training from the nursing literature is called "transcultural nursing," which "is concerned with comparing differences and similarities between cultures regarding caring values and life practices to predict the care needs of individuals and promote culturally [appropriate] care. In this approach, culture is defined as attitudes, values, beliefs, and life practices learned and shared by people in a particular social group, which are passed on down generations, affecting individuals' thinking and actions" (Allen, 2010, p. 315; Leininger and McFarland, 2002). This cultural competence training model and others similarly focus on meaningful, thoughtful, and humble interactions, and teaching individuals from different cultural backgrounds needs to be based on knowledge about specific cultural beliefs, attitudes, and practices. In a meta-analysis to examine the effect of educational interventions designed to enhance cultural competence in professional nurses and nursing students, Gallagher and Polanin (2015) found, in general, moderate to large positive effects. This is consistent with previous studies (e.g., Maina et al., 2018), including a meta-analysis conducted by Smith et al. (2006). These studies, however, were limited in their lack of an articulated definition of cultural competence, implementation, rigorous design, and self-rating measures, which calls for more rigorous, well-defined multimethod studies in this area.

SOURCES: Allen, 2010; Galindo et al., 2019; Gallagher and Polanin, 2015; Maina et al., 2018; Schneidman and Woodward, 2016; Silva et al., 2015; Smith et al., 2006; Sperry et al., 2018

Scholars caution about the focus on cultural competence without antiracist training. A literature review by Allen (2010) indicates that cultural competence training could be strengthened by antiracist training to adequately ensure that professionals are able to "challenge discrimination experienced by minority cultural groups, such as ethnocentrism, cultural biases, and overt and covert discrimination due to racial difference . . . [and cultural competence training may limit] culture to a static entity and reduces understanding of peoples' behavior to prescribed cultural norms. This promotes cultural stereotyping with the risk of discrimination and it fails to account for individual and family differences within cultural groups" (p. 315). However, there is an absence of a validated antiracist training and intervention.

Other options to address unconscious bias are mindfulness training and prejudice habit-breaking. It is theorized that mindfulness training could reduce implicit bias by (1) deactivating the "prejudice network," which involves the activation of the amygdala (or threat responses) and reduces the ventromedial prefrontal cortex (or the empathy and "humanizing" pathway) (Burgess et al., 2017); (2) meditating to increase one's ability to become aware of implicit biases, once they are activated, and to engage in self-regulatory processes to behave in less discriminatory and prejudicial ways; (3) reducing stress and internal sources of cognitive load that contribute to the activation and application of implicit biases; (4) activating empathy and compassion, which

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reduce the activation and application of implicit bias and promote willingness to engage with members of stigmatized groups; and (5) improving one's ability to communicate and focus on the other person's individual characteristics rather than their group membership. While Burgess et al. (2017) provide a strong rationale for mindfulness as a strategy to address implicit bias, there is a need for systematic implementation and examination in this area. It should also be noted that in order to maximize mindfulness, it is imperative for individuals to recognize their own implicit bias, any privilege that comes with their social status, and unintentional microaggressions (i.e., "brief and commonplace daily verbal, behavioral, or environmental indignities, whether intentional or unintentional, that communicate negative slights and insults") (Godsil et al., 2014; Sue et al., 2007).

Devine's prejudice habit-breaking framework argues that implicit biases are deeply entrenched habits developed through socialization experiences (Devine et al., 2012). "Breaking the habit" of implicit bias therefore requires learning about the contexts that activate the bias and how to replace the biased responses with responses that reflect one's nonprejudiced goals. Thus, when people who are opposed to prejudice believe they have acted with bias, they will seek out information to help them break this habit of prejudice or bias. That training includes (1) stereotype replacement (recognizing stereotypic responses within oneself and society), (2) counter-stereotypic imaging (imagining examples of outgroup members who counter popularly held stereotypes), (3) individuating (viewing others according to their personal rather than stereotypic characteristics), (4) perspective taking (adopting the perspective of a member of the stigmatized group), and (5) contact (increasing exposure to outgroup members). In an RCT, Devine et al. (2012) showed that participants in an 8-week habit-breaking training showed significant decrease in their explicit bias but also became more acutely aware of discrimination in society. This study was conducted with introductory graduate students who were likely biased in their willingness to participate. There is a need for more rigorous examination of this training approach, with educational professionals and those caring and teaching young children.

Supporting ECE Workforce Well-being

In addition, even the most competent early childhood educators will not be effective if they are not physically, mentally, and social-emotionally healthy. Educators who have strong social-emotional awareness and skills are more likely to develop close, constructive relationships with children, implement SEL programs effectively, and be more able to create a positive classroom climate. Conversely, educators who are stressed, depressed, or low in social-emotional skills are more likely to be less sensitive and warm toward children and have more conflictual relationships with them. Since children who live in high-stress communities or with adverse conditions benefit the most from warm, enriching interactions with adults, it is critically important for them to be in ECE programs in which the educators are social-emotionally healthy and strong (Becker et al., 2017; IOM and NRC, 2015; Jones and Bouffard, 2012; Roberts et al., 2019; Whitaker et al., 2015). Indeed, McClelland et al. (2017) found that effective social-emotional interventions often including building teachers' own social-emotional skills, as well as children's.

Unfortunately, it has been well documented that compared to other fields, the ECE workforce experiences a high level of stress and depression; one reason is the stress of low compensation and benefits (see IOM and NRC, 2015). Most of them (53 percent) participate in at least one of four public support or health care programs for low-income individuals, compared to 21 percent of the U.S. workforce in general (Whitebook et al., 2018). On average, child care

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teachers earn less than $11 per hour, or about $22,000 a year—barely over the federal poverty guidelines for a family of three. There is also emerging evidence (Borntrager et al., 2012; Hydon et al., 2015) that when educators work with children who experience trauma and significant adverse experiences, whether in ECE programs or public schools, they can experience "secondary traumatic stress" (STS), or "the natural consequent behaviors and emotions resulting from *knowing* about a traumatizing event experienced by a significant other—the stress resulting from helping or wanting to help a traumatized or suffering person" (Hydon et al., 2015, p. 320).

What supports or interventions can help educators manage or mitigate their general social-emotional health and their experience of stress, depression, or STS? For early childhood educators, an analysis of the National Survey of Early Care and Education found that "informal workforce supports" that help teachers feel a part of a team and respected as professionals were related to lower stress. Studies of public school teachers by Borntrager et al. (2012) and Caringi et al. (2015) similarly found that working in schools that provided or encouraged peer support was associated with lower STS. In addition, stable classroom assignments were related to lower stress, as was (unsurprisingly) higher pay. Teachers with higher household incomes (over $45,000) experienced less stress than those earning less (Madill et al., 2018). In another analysis, Roberts et al. (2019) found that early childhood educators who had no health insurance, perceived high professional demands with few work-related resources, and had multiple jobs were more likely to have depressive symptoms. Interestingly, Madill et al. (2018) found that more formal professional supports, such as small group size and access to professional development or coaching, were not related to early childhood educators' experience of stress.

There is emerging evidence that professional support that is targeted to improve the mental and social-emotional well-being of educators could be helpful. In their small, qualitative study, Caringi et al. (2015) reported that teachers' practice of "self-care" techniques helped them manage their stress. In her small study of educators and caregivers in a residential program for AIDS-affected children and youth, Lucas (Lucas, 2007) found that training on "reframing" and realistic goal setting with children and parents led to a decrease in emotional exhaustion. (Reframing has teachers identify children's and families' strengths and assets rather than their faults, which helps teachers approach problems and obstacles collaboratively with those they serve rather than treating these as issues they need to "fix" on their own.)

Another promising type of intervention is to improve "dispositional mindfulness" (Becker et al., 2017; Hydon et al., 2015). Mindfulness is related to the ability to regulate one's emotions, thoughts, and attention. People who are mindful are more aware of their thoughts, feelings, and reactions to their surroundings and better able to refrain from judgment, which enables them to react more calmly to a given situation and to better understand others' perspectives and behaviors before responding. In one study based on a survey of 1,001 Head Start teachers, those who were more mindful had closer relationships with children and less conflict with them, partly because mindfulness is also associated with fewer depressive symptoms.

> A mindful disposition may help teachers view challenging interactions with more equanimity. Alternatively, mindful teachers may have fewer challenging interactions. Their attention and focus may help them to be proactive in their guidance of young children, structuring the classroom for successful interactions and diffusing potentially difficult situations before they begin . . . Teachers who are more mindful may be better able to disengage from depressive or ruminative thoughts, and experience fewer difficult exchanges with children. Additionally, a mindful disposition may help teachers appraise

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> stressors differently, such that children's difficult behavior may not be perceived as a threat. Mindful teachers may be aware of situations that may elicit challenging behavior, and take action to engage in reappraisal to regulate their emotional response (Becker et al., 2017, p. 48).

One program, called Cultivating Awareness and Resilience in Education (CARE), has demonstrated impact on teachers' mindfulness. CARE provides 30 hours of training in group-based settings and individual sessions with trainers. The program includes "emotion skills instruction, mindfulness practices, and compassion-building activities" (Becker et al., 2017, p. 49). RCTs with elementary teachers showed improvements in their mindfulness and emotional regulation. In one of those studies, researchers also observed improvements in teachers' emotional support for students (Becker et al., 2017).

This study also indicated that for programs of this nature to be viable and sustainable, school leaders' buy-in is necessary. Strong leadership is influential in early childhood educators' professional support system (IOM and NRC, 2015). While local program administrators and leaders may not be able to improve educators' professional preparation programs, compensation levels, or benefit packages on their own, they are critical to establishing an organization and environment that supports their staff's mental health and minimizes stress—whether through opportunities for them to collaborate or appreciate each other's work, training on topics such as TIC, or self-care interventions, like mindfulness training.

ECE CONCLUSIONS AND RECOMMENDATIONS

Below, the committee provides conclusions and recommendations, based on the information reviewed in this chapter, to ensure adequate resources for ECE programs and educators, support and improve competencies for the ECE workforce, and improve access to ECE for eligible children. The goal of these recommendations is to advance the ECE system to promote health equity during the early childhood period to set the course for good health and well-being into adulthood.

Allocation of Adequate Resources to Support Health-Promoting ECE Programs and Educators

ECE programs, including systems that support and ensure standards for high quality (e.g., QRISs), can be comprehensive platforms for ensuring all children are healthy and prepared for school and life. That can only occur if these programs and systems adopt specific practices and processes that have been shown to have such impacts and set a threshold that ensures all participating programs are sufficiently high quality, as opposed to more general benchmarks related to quality (e.g., teacher education levels, class size, professional development). Further, current funding levels for child care, pre-K, and Head Start generally are not based on systematic estimates of the cost of quality, including those elements that promote better health outcomes. It is also critical to consider the cost of funding programs to ensure that they reach all eligible children. Until that has been remedied, our nation will not maximize the potential of ECE programs to promote the health and school readiness of young children, especially those who tend to fall on the wrong side of the health equity equation (e.g., low-income children and children of color).

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For ECE to be part of a system that ensures children are healthy and ready for school and life, these programs need

- Adequate resources to support the "whole child," including cognitive, emotional, social, and physical development, regardless of setting (e.g., home, center, or school). This includes adequately supporting programs' implementation of standards and content that are health promoting (e.g., high-quality nutrition and physical activity) for child, family, and staff outcomes, ensuring children's access to effective early intervention and special education services, and tailoring specific resources and supports for various settings and providers (e.g., home-based providers);
- The ability to implement the recommendation from the National Academies reports *Transforming the Workforce for Children Birth Through Age 8* (IOM and NRC, 2015) and *Transforming the Financing of Early Care and Education* (2018) addressing the level of compensation and access to basic benefits (such as health care and housing); ensuring work environments and conditions that respect the demands of the ECE profession, including adequate resources and opportunities for teamwork (e.g., planning time, substitutes); offering ongoing training to maintain educators' own emotional well-being; and developing school and program leaders' knowledge and competencies in creating an organizational culture that supports the social-emotional health and well-being of children, families, and educators;
- The capacity (funding, staffing, skill sets) to provide or make referrals (and follow up) to various agencies or integrated partnerships with professionals from various sectors or organizations (e.g., home visitors, social workers, health services) that meet the various needs of children and their families;
- An SEL component that has the following characteristics:
    - Strong training and support (e.g., job-embedded coaching, early childhood mental health consultants) for educators on specific SEL skills;
    - Curricula intentionally designed to build specific SEL skills through sequenced and active learning experiences;
    - Promotion of a supportive organizational culture and climate by engaging the involvement of the leaders, paraprofessionals, and other staff of the school or ECE program; and
    - Supports for the social-emotional health of parents and early childhood professionals and staff, including leaders.
- Educators who are well trained and well supported to develop critical professional competencies and support children's social, emotional, and physical health. Educators need to develop specialized skills that ensure children are healthy and prepared for school. In order for them to use these skills effectively, their own physical, emotional, and mental health needs to be supported; and
- Adequate resources and support to systematically identify factors that support or hinder integration at the federal, state, and local levels, as well as the impact of this effort for children, families, and communities.

Conclusion 7-1: For early care and education programs to contribute significantly to a health promotion and equity strategy, there is a need to intentionally, cohesively, and simultaneously address adequate funding that supports a comprehensive equity-promoting ECE system, well-compensated and competent workforce, connection to

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community resources and support, continuous quality improvement, and systematic examination of effectiveness at multiple levels.

Recommendation 7-1: The committee recommends that early care and education (ECE) systems and programs, including home visiting, adopt a comprehensive approach to school readiness. This approach should explicitly incorporate health promotion and health equity as core goals. Implementing this approach would require the following actions:

- **Federal, state, local, tribal, and territorial governments and other public agencies (e.g., school districts, city governments, public–private partnerships) that have decision-making power over ECE programs should establish program standards and accountability systems, such as a quality rating and improvement system, linked with better school readiness and health outcomes and provide adequate funding and resources to implement and sustain these standards effectively.**
- **The Office of Child Care and the Office of Head Start at the federal level, along with state, local, tribal, and territorial early care and other education agencies, should assess the full cost of implementing standards that promote health outcomes and equity as described above, including supporting educators' own health and well-being, and work with Congress to align funding levels of the major federal ECE programs—child care subsidy and Head Start—accordingly.**
- **Health and human service entities, the federal Early Learning Interagency Policy Board, state Early Childhood Advisory Councils, and federal, state, local, tribal, and territorial agencies that oversee home visiting and ECE programs should ensure greater programmatic coordination and policy alignment to ensure effective allocation of resources.**
- **The Office of Planning & Research, and Evaluation in the Administration for Children and Families along with the U.S. Department of Education, should examine the feasibility and seek resources to conduct (a) an implementation study to examine the design and implementation of this comprehensive ECE approach that incorporates health standards and (b) an outcomes study that examines the impact on children's school readiness and achievement, and health outcomes, with particular attention to eliminating disparities and gaps prior to school entry.**

Health-focused Competencies of the Workforce

The ECE workforce and other professionals are critical in implementing evidence-based practices in ECE settings that support children's well-being, leading to health equity. However, there is a need for systems and supports to strengthen their competencies and skills, as well as stability, in supporting health-promoting and health equity practices.

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Conclusion 7-2: Policies and systems that prepare and support early childhood educators and program leaders, including those in public schools, need to incorporate the latest evidence about how to support children's school readiness and success by fostering the health and well-being of children. This would entail providing comprehensive supports and resources to degree granting institutions and preparation programs, including the development of curricula, textbooks, practicum experiences, toolkits, and fact sheets.

Recommendation 7-2: Building off the 2015 Institute of Medicine and National Research Council report *Transforming the Workforce for Children Birth Through Age 8*, the committee recommends that degree granting institutions, professional preparation programs, and providers of ongoing professional learning opportunities develop or strengthen coursework or practicums that focus on competencies of educators, principals, and early care and education program directors that are critical to children's health, school readiness, and life success.

Specific areas include:

- strengthening professionals' understanding of and capacity to collaborate with professionals from other sectors, such as health and social work (interprofessional learnings);
- implementing practices and policies based on the understanding of the link between biology and children's learning and development and impact on children's readiness for school and school success;
- educating professionals about unconscious biases and practices that undermine the learning and socio-emotional health of diverse children and their families;
- training and coaching on effective antibias and cultural responsive practices that strengthen professionals' effectiveness in supporting the learning, social-emotional health, and well-being of diverse children and families (including dual language learners); and
- effectively implementing practices and policies informed by understanding of trauma, ACEs, toxic stress, and racism.

Access and Affordability

Children who experience comprehensive high-quality ECE early in life and for multiple years are likely to show stronger cognitive, academic, and socio-emotional outcomes over time. Unfortunately, many children who could benefit the most from ECE are less likely, for various reasons, to access these high-quality ECE environments early and for longer periods. To ensure health promotion and health equity in the early years, there is a need to ensure that these programs are available to and affordable for families as early as possible and as long as possible.

Conclusion 7-3: Maximizing the impact of ECE on positive childhood development and health and well-being at the community or population level will require increasing

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public funds for ECE programs. Currently, eligibility for ECE programs is limited, and among eligible families, access is low due to lack of funding and availability of programs and services. Therefore, even if existing publicly funded programs have the resources to provide robust supports that improve young children's health and well-being, they will not reach most children, especially those who live in low-income households or experience adverse experiences and toxic stressors.

Recommendation 7-3: Recommendation 7-3: Federal, state, local, tribal, and territorial policy makers should work with the U.S. Department of Health and Human Services (HHS), the Office of Head Start, and Office of Child Care develop and implement a plan to:

a) improve the quality of early care and education (ECE) programs by adopting the health-promoting standards discussed in Recommendation 7-1, such as building on the performance standards of Early Head Start and Head Start, and

b) within 10 years, expand access to such comprehensive, high-quality, and affordable ECE programs across multiple settings to all eligible children. Disproportionately underserved populations should be prioritized.

The Secretary of HHS should conduct a process evaluation to inform the expansion effort, and, once implemented, conduct rigorous and comparative outcomes studies to ensure that the expansion is having the intended impacts on children and families, with particular attention on what group(s) may be benefitting.

The strategic plan should be modeled after and build upon the relevant performance standards of Early Head Start and Head Start, which emphasize mixed settings, the whole child, family and community engagement, transition between home and school, and CQI. It should also strengthen the program components discussed in this chapter that lead to stronger school readiness and health outcomes, including mitigating the impact of adverse experiences and toxic stress for children, families, teachers, and staff. Critical components include a comprehensive social-emotional strategy that encompasses both the classroom (curriculum, teacher training, and support) and program/school (leadership, culture, and climate) levels and educators who have the competencies described in Recommendation 7-2. The plan should identify strategies to bolster capacity and resources of new and existing programs to implement these more ambitious standards, including by incentivizing collaboration among Head Start, pre-K, and child care programs. Implementation of this plan will likely require funding from Congress.

A national evaluation led by the Office of Head Start, Office of Child Care, Office of The Assistant Secretary for Planning and Evaluation (ASPE), and Office of Planning, Research and Evaluation, in the Administration of Children and Families, is needed to examine and inform the design, implementation, and effect of this expanded access and strengthening of Early Head Start and Head Start to enhance children's school readiness and achievement and health outcomes.

Heeding the findings from the Head Start Impact Study that program implementation varies widely from site to site and workforce challenges, which leads to variable outcomes (Phillips et al., 2017), the improvement and scale-up strategy described in this recommendation

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should include mechanisms that help new providers incorporate program features and conditions that are associated with stronger outcomes. For example, an evaluation can provide rapid-cycle feedback (see Chapter 8 for more on this) to ensure faithful implementation of the "upgraded" program model and fidelity and to inform ongoing midcourse corrections as needed to reach targeted health outcomes. This study will help to identify factors that supported or hindered expansion and access at the federal, state, and local levels, as well as the impact of this effort for children, families, and communities.

State policy makers (e.g., governors, legislators, agency leaders, pre-K administrators) should also consider how pre-K funding and policies can support the program and workforce characteristics discussed in this chapter that are associated with health outcomes. State pre-K programs are highly variable, and some will be better positioned to serve as a platform for promoting health equity than others. In the final analysis, the committee believes that among the major publicly funded ECE programs, Head Start's history, program design, quality standards, targeted populations, and evidence base make a useful platform from which to build expanded access to comprehensive high-quality ECE programs. Health promotion (including social-emotional health) and family engagement are already important goals of the program which can be strengthened based on evidence from this report. The program's historical focus on children furthest from opportunity can help address inequities among different racial and income groups. With an intentional effort to bolster its health-promoting strategies, provide adequate resources for educators and leaders to implement them effectively, and expand access to all eligible children, Head Start can be a critical element of our nation's cross-sector approach to improving child health and reducing inequities.

CONCLUSION

The ECE system is a critical setting to provide young children with a strong foundation for skill building and positive learning, as well as shaping socio-emotional, cognitive, and physical health. This chapter delves into the evidence on ECE programs and childhood outcomes with respect to the many different service settings (e.g., home-, school-, and center-based care). In addition, the committee highlights salient issues and populations related to health equity throughout the chapter, such as early intervention for children with developmental disabilities, DLLs, and implicit bias training for educators. Based on its review of the evidence and committee expertise, the committee applies the evidence to provide recommendations in the areas of allocating adequate resources to support ECE programs and educators, supporting and training the workforce, and improving access to quality ECE for eligible children. By targeting these key areas that are instrumental to an effective and equitable system, the committee identifies a comprehensive approach to leverage and enhance the current ECE system to promote health equity. The following chapter integrates the crosscutting themes from the report and applies them to inform a systems approach to promote equitable prenatal and childhood development.

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8

A Systems Approach to Advance Early Development and Health Equity

INTRODUCTION

Advancing health equity in the preconception through early childhood periods cannot be achieved by any one sector alone. It will take action, collaboration, and alignment across all sectors that frequently interact with children and families and the professionals who serve them. Additionally, better alignment among systems will need to be accompanied by an increase in overall investment in early life: many of the systems best positioned to address early life drivers of health inequities are chronically underresourced, and improved collaboration may only be possible if resources are available to adopt, scale, and spread best practices within a redesigned and better aligned cross-sector ecosystem. There is likely no single, sweeping change that will create a new and better system of care that can address the variety of needs and challenges identified in this report; rather, steady progress integrating and connecting the efforts of the systems already in place, along with improved investments in those systems, will lead incrementally but steadily to improvements in health equity over time. Systems change is not an easy strategy, it seldom yields speedy returns, and it may not be sufficient without an investment of resources designed to take advantage of new and better aligned approaches. However, given that disparities are systematically generated, it is likely a necessary precursor to real and widespread advances in health equity. (See Box 1-1 for a brief overview of this chapter.)

BOX 8-1
Chapter in Brief: A Systems Approach

There are many opportunities to overcome key barriers to strengthen a systems approach to advance health equity in the preconception through early childhood periods. The crucial stakeholders who need to be involved and the alignment, measures, and research that are needed for systems change are discussed in this chapter, based on the committee's assessment of the literature in Chapters 2–7 in this report.

Recommendations in brief:

- Develop cross-sector initiatives that align strategies to address barriers to data sharing and integration, cross-sector financing, and other challenges to cross-sector collaboration.
- Enhance detection of early-life adversity and improve response systems.
- Develop adversity and trauma-informed systems.
- Build a diverse, culturally informed workforce in all relevant systems.

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- Improve access to programs and policies across systems that provide parental or caregiver supports and help build or promote family attachments and functioning. For families with intensive support needs, develop programs or initiatives designed to provide comprehensive wraparound services.
- Integrate care and services across the health continuum, including the adoption of models that provide comprehensive support for the whole person by leveraging and connecting existing community resources.
- Support payment reform to allow for upstream investment.
- Support research that advances the state of the science in several critical ways to advance health equity, including exploring alternative methods to address complex causality and expanding research into individual differences in response to adversity and treatment.

As defined in Chapter 1, systems are a collection of interacting, interdependent parts that function as a whole. For the purposes of this report, most systems are social constructs and organized around a key functional area (e.g., education, health care, criminal justice). Systems have existing patterns and structures that define how people tend to move through them. This chapter summarizes the opportunities to overcome key barriers to strengthen a systems approach, the crucial stakeholders who need to be involved, and the necessary alignment, measures, and research based on the committee's assessment of the literature in Chapters 2–7 in this report. The committee identified eight crosscutting recommendation areas where multiple sectors need to take action.

Systems Characteristics That Impede Advancing Health Equity

There are many systems characteristics that act as barriers to spreading and scaling up evidence-based and promising programs and approaches to reduce health disparities and advance health equity. Sometimes barriers are simply financial—scaling programs requires significant and sustained investments in a world with limited resources and many competing priorities. However, the barriers often stem from policy or structural arrangements that could be altered with sufficient political will or from knowledge gaps that could be addressed with appropriate research, well-targeted dissemination, and thoughtfully considered implementation assistance. Some of the key system barriers that prevent moving forward with what is known to work include the following.

Systems are Designed to React, Not to Prevent

Many systems are designed in response to a challenge or crisis—people are sick or lack jobs—and operate with the primary goal of mitigating the negative impacts of those challenges. Few systems are set up to think ahead to the root causes of the problem and to address those causes, and as a result, resources are applied downstream. In fact, many systems are explicitly not allowed to spend resources upstream (on prevention, for example), because the upstream factors are seen as a different system's challenge to address, and that other system has its own budget and goals. Current programs and policies are operating in an interconnected world, where one system's cause is another system's effect, and the carefully partitioned systems each address only the portion of the problem that falls explicitly within their purview. From the perspective of a life course approach—where failing to invest early means missing critical windows to set positive health trajectories, and the later an intervention comes, the more difficult it is to change negative trajectories—this fractured approach is an impediment to reaching health equity in the preconception through early childhood periods and beyond.

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Systems are Structured to Take the Short-Term View

Currently, systems are poorly structured to incentivize long-term thinking and planning. The pressures of annual budget and performance cycles make investment in long-term gains challenging, and savings are often dis-incentivized by penalizing systems that reduce costs with a subsequent reduction in payment rates or budget allocations. Often, payments or budgets for systems are based on prior experiences that do not reflect transformed realities. For example, early efforts to integrate behavioral health into primary care faced formidable barriers because payment rates are based on prior experience that does not include many mental or behavioral health services, making transformation extraordinarily difficult to catalyze. From an economic and political standpoint, the benefits of early care and education (ECE) are generally not realized until after children enter school, and sometimes even later in life. Additionally, when the root causes of poor outcomes are interconnected across systems but the financial stakes are not, misaligned incentives emerge whereby each partner fears they may put in significant work and expense to change an outcome, only to see the savings realized primarily by someone else. As Teutsch and Berger noted, "If everyone is focused only on his own task, no one is responsible for ensuring that our nation's investments are well utilized, let alone best utilized, to improve health" (Teutsch and Berger, 2005, p. 486).

Systems Typically have a Singular Focus

A multisystem approach to health equity is difficult when involved systems are primarily built to specialize on one aspect of the health continuum (and whose main goal might not even be health). Which system responds to risks that might generate poor health outcomes, how that work is organized, which system is allowed to address it, and the way funds flow to support a response are all built within highly specialized silos with distinct rules, regulations, strategies, and normative practices. There are relatively few system architectures ready made to advance cross-sector work and few easy pathways to conceptualize, implement, or pay for it. In addition, those who work within systems might feel a strong sense of special expertise and ownership over their area of focus, which may cause a reluctance to invite others to share hard-earned professional "turf." Overcoming both the structural and cultural barriers to integrating systems will be no easy task, but structures that align the interests of different sectors—for example, by creating shared savings or other models that reward systems or actors within systems for contributing to one another's positive outcomes—may help overcome some of these inherent challenges.

Systems Take a Narrow View of the Biological and Social Context

Scientific and institutional systems tend to segment the biological and social; in the context of early life, they are often poorly set up to address the symbiotic relationship between biological development and social context. Even within settings that address health and development explicitly, such as pediatric clinics, there tends to be a segmentation of biological risk (assessed by clinicians or pharmacists) from social risk (assessed by social workers or therapists), lacking clear processes for developing a plan for each that is informed by the other.

Systems Undervalue Community Expertise

Current systems tend to value formal knowledge attained through traditional, credentialed academic channels but often undervalue knowledge of culture, community, populations, or other

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context gained through lived experiences. Those making decisions about how to serve some communities, or even those providing the service, often do not reflect or live in the communities being served. When the experiences of those most impacted by inequities are not represented in strategies to address them, there is a risk of developing solutions that do not connect with, will not be used by, or do not meet the actual needs of the population for whom they are intended, which may ultimately exacerbate the very disparities that are intended to be addressed.

OPPORTUNITIES TO STRENGTHEN A SYSTEMS APPROACH

Throughout this report, the committee has explored key factors that help set the odds for long-term health and health equity outcomes and made recommendations suitable for actors within key systems and institutions to help improve those odds. While the actions taken by each system identified in this report are important components of improving health equity, one of the committee's important findings—that the forces driving health inequities are systemic and profoundly interconnected across those systems—suggests that a larger strategy is also needed to advance health equity in the long term. When outcomes are driven by forces that cut across multiple systems, even doing everything perfectly within one system is not enough. Multisector causality requires a multisector response.

Based on the committee's assessment of the needed changes in Chapters 2–7, the corresponding recommendations contained in each chapter, and its collective expertise, the committee identified eight crosscutting recommendations that need to be adopted by all systems that frequently touch the lives of children and families and the professionals who serve them. These recommendations represent part of a comprehensive strategy for improving the overall system's ability to address the early-life drivers of health inequity. These strategies draw upon the evidence presented in this report and reflect the key insights from the core principles presented in Chapter 1. These recommendations can optimize the impact of the strategies in this report and create a framework where what is known to work can be made to work for more people, under more circumstances, and with a greater overall impact on health equity.

Support Cross-sector Initiatives Across the Health Continuum

Achieving health equity is a systems challenge. It will not be improved solely by developing and deploying programs aimed at individuals experiencing poor outcomes—until the root causes are addressed, negative health outcomes will persist. Young people experience adverse and positive exposures that cumulatively help shape their odds for good health over the life course, but within systems, those exposures occur at systematically different rates for different groups of people. They also intersect profoundly—the early-life exposures and experiences that shape health are multidimensional and fall under the purview of multiple social and cultural systems (see Chapters 2 and 3 for more information). These different exposures and experiences result in different cumulative odds for good health over the life course, odds that are ultimately expressed in the form of disparate outcomes between groups.

Early efforts to adopt cross-sector approaches were often impeded by the challenges of data sharing. The regulatory structure that governs data use and data privacy offers important protections for individual consumers, but it was also largely developed without considering the limitations it might place on cross-sector interventions by communities seeking to collectively address complex problems with roots in multiple systems. The distinct regulations for different

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sectors (e.g., HIPAA[1] in health care and FERPA[2] in education) have different criteria for data use, and while some allow data to be combined for the purposes of research or evaluation, there are few legal pathways available to make *operational* use of cross-sector data in ways that would help comanage common populations or address common root causes with coordinated, cross-sector approaches. By providing a regulatory architecture to support data sharing between the key systems identified in this report, policy makers and systems leaders could dramatically improve communities' ability to adopt, scale, and spread cross-sector interventions.

A second key challenge of cross-sector work stems from how public dollars are organized. Resources tend to be allocated in a very siloed manner: this money is for health care, that money is for education, and so on. In general, most systems have strict administrative rules on how their funding can be spent and limited ability to direct it at work that occurs outside of their system, even if that work is critical to that same system to achieve its desired outcomes. However, in reality, these issues are not confined to the convenient boundaries within which finances are organized, and if the problem that is creating poor outcomes for a system does not happen to fall within its boundaries, that system is left with only a limited ability to address it. Creating pathways for funds within a system to flow to activities outside the system, especially when there is strong evidence that those activities will result in improved outcomes, is critical to building effective cross-sector initiatives. A recent example of this is an effort underway by health care provider Kaiser Permanente—it plans to spend $200 million on fighting homelessness and building more low-cost housing in eight states, plus DC.[3]

In addition to allowing for investments across sectors, it is important to consider how any *savings* generated by those investments might be shared. In the absence of shared savings models, a "wrong pocket" problem is often created, whereby system partners fear they may invest in the work only to see the outcomes or savings accrue to others. There are currently few validated methodologies for measuring shared savings—especially given the data sharing challenges that have already been highlighted—or mechanisms for capturing those savings and distributing them among the participating partners. Making headway on shared savings models could help create powerful aligned incentives that help cross-sector initiatives create, sustain, and spread their work. One strategy to address this problem is to develop and test shared savings models that allow sectors whose work helps the outcomes in another sector "share the savings" generated by that work. This would help create aligned incentives that acknowledge intersectional causes and reward intersectional work.

Finally, licensure and certification requirements vary dramatically across fields; in many emerging fields, they are virtually nonexistent. However, services that impact health are sometimes also delivered by paraprofessionals, peers, or other workforces that either lack formal professional licensure standards or use standards that are not recognized by larger, more established systems that would pay for such services (IOM, 2001). Payment rules for many systems often require more traditional certification, so this barrier hinders using nontraditional workforces, which may be closer to the communities being served and could have similar lived experiences, to deliver services that help create health within the context of collaborative cross-sector work.

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[1] Health Insurance Portability and Accountability Act of 1996.
[2] The Family Educational Rights and Privacy Act of 1974.
[3] See https://about.kaiserpermanente.org/community-health/news/kaiser-permanente-announces-three-initiatives-to-improve-communi for more information (accessed April 16, 2019).

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While there has been significant work around the key components necessary to establish successful cross-sector initiatives (see examples in Chapters 5, 6, and 7), a number of key barriers remain, including cultural, ideological, or normative barriers around perceived ownership of specific content areas that may limit cross-sector communication and cooperation that policy makers and leaders can help address.

> **Recommendation 8-1: Policy makers and leaders in the health care, public health, social service, criminal justice, early care and education/education, and other sectors should support and invest in cross-sector initiatives that align strategies and operate community programs and interventions that work across sectors to address the root causes of poor health outcomes. This includes addressing structural and policy barriers to data integration and cross-sector financing, and other challenges to cross-sector collaboration.**

These initiatives could include collective impact strategies, Accountable Communities of Health (ACH),[4] Health in All Policies[5] initiatives, or other models of aligned, cross-sector community action in service to shared health and health equity goals. (See Chapter 8 of the 2017 National Academies report *Communities in Action: Pathways to Health Equity* for more information on Health in All Policies and other collective impact strategies to foster multisector collaboration.)

Enhance Detection of Early-life Adversity and Improve Response Systems

As discussed throughout this report, adversities in early life—including Adverse Childhood Experiences (ACEs) and other adverse experiences or exposure to key health-related social determinants, such as housing instability or food insecurity—help set the odds for poor health outcomes later in life. Early-life adversity is especially important for a number of reasons: first, it occurs at points of high plasticity for key biological systems that shape health over the long term; second, it occurs during a period of critical development in the social, emotional, and cognitive domains of self; third, it increases the likelihood of additional exposures to adversity later in the life course; and fourth, it may impact future resilience to additional exposures (see Chapter 2 for more information). Early adversities make a negative health trajectory more likely; left unaltered, such trajectories will play out across decades and result in differences in health outcomes within and across generations. Early detection and rapid response are essential to help mitigate the long-term effects of exposure to adversity by adjusting the trajectories back toward positive health as early as possible.

Early detection and rapid response would require a number of steps across systems, such as developing and promoting screening tools and procedures that can be adopted, implemented, and scaled to improve monitoring and fast reaction. Screening approaches that can be adopted within

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[4] "Accountable health initiatives, most commonly referred to as accountable communities for health (ACHs), have been implemented nationwide in response to or as a result of contributions from state innovation model grants and community transformation grants, through collaborations with state Medicaid programs, or through other policy and financial incentives" (Mongeon et al., 2017). They bring together partners from health, social service, and other sectors to improve population health and clinical-community linkages within a geographic area (Spencer, and Bianca, 2016).

[5] "Health in All Policies is a collaborative approach to improving the health of all people by incorporating health considerations into decision-making across sectors and policy areas. The goal of Health in All Policies is to ensure that all decision-makers are informed about the health consequences of various policy options during the policy development process" (Rudolph et al., 2013, p. 6).

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and connected across settings, such as the health care and early learning systems, and that make frequent contact with young families during the first years of a child's life are especially critical because they represent the best opportunities to detect early and act decisively in response.

Professionals who frequently interact with children and caregivers—clinicians, teachers, or personnel in social service or other agencies, for example—need training to understand the effects of early-life adversity and the associated toxic stress response on the physiological and psychosocial development of young people. These trainings need to move beyond awareness training to include how to respond effectively within the context of their field and how to refer across sectors when needed.[6] Dedicated staff are also needed to help professionals refer children and families to services outside of that system (for example, social workers embedded within the relevant systems). These professionals need to know how their approach to treatment or service provision will vary based on the results of a child's adversity screen. (See below for more on transdisciplinary needs.)

It is also necessary to develop rapid response or referral systems that can bring a range of community resources to bear when early-life trauma and adversity is detected. These systems should include information and service pathways and the ability to "close the loop" back to the referring agency or partner so that responses are coordinated across the continuum of health. Accomplishing this will require changes to regulatory limitations on how data can be shared across sectors or between professionals with different qualifications and licensure systems.

> **Recommendation 8-2: Policy makers and leaders in the health care, public health, social service, criminal justice, early care and education/education, and other sectors should adopt and implement screening for trauma and adversities early in life to increase the likelihood of early detection. This should include creating rapid response and referral systems that can quickly bring protective resources to bear when early-life adversities are detected, through the coordination of cross-sector expertise, as covered in Recommendation 8-1.**

Rapid response and referral systems will require implementing and scaling screening across key settings, enhancing trauma response training, and assuring support for these systems.

Develop Adversity and Trauma-informed Systems

Early-life trauma and adversity are key factors that help set the odds for poor health outcomes across the life course, but *how systems respond* to that trauma is nearly as important. Once adversity or trauma occurs, it cannot be erased. However, its effects also are not destiny: effective services can help mitigate the impact of adversity on health across the life course. Most systems are designed to capture discrete data elements about their service domain, but exposure to social adversity or trauma are often contextual, captured via narrative interactions with patients or clients. These are also sensitive personal data with additional layers of privacy protections. Because these data are not easily shared within or across systems, however, people seeking help at multiple points of service often have to tell the story of their trauma over and over again. Mechanisms that allow service providers to have the needed data and context about

[6] For example, the Alberta Family Wellness Initiative in Canada has developed a "Brain Story Certification Course" that teaches the foundational science of brain development to help professionals in all fields who interact with families with children. See https://www.albertafamilywellness.org/training for more information.

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their clients at the point of care could ensure that their service options are appropriate to clients' financial, social, cultural, and personal situations. For example, when service options are presented that require resources a client cannot possibly access (e.g., financial cost-sharing, transportation), the client's status as an "outsider" is reinforced. When systems have mechanisms to meet clients where they are, clients will be more supported and understood and be more likely to adhere to recommendations, remain engaged with the system, and receive the help they need.

Many clients feel distress not just from past life events but from interactions with the system itself: they feel unwelcome or stigmatized when they seek services due to their race, poverty, sexual or gender identity, or other factors. It is critical to train service providers on discrimination and stigma to ensure all persons feel welcome receiving services in any setting, so they will engage in and benefit from those important services. As noted in Chapter 7, trauma-informed care (TIC) and implicit bias training reflects the understanding that a child's or family's behavioral or health challenges are often due to experiences of toxic stress and undiagnosed and untreated trauma and with implicit bias among those they interact with in the education or other systems. Further, systems delivering services to people with traumatic histories are also at risk of retraumatizing clients if they do not act in a trauma-informed manner. There are strategies for TIC and that can be used to prevent and mitigate the impact of implicit biases (see Chapter 7 for more strategies and Recommendation 8-8 for research needs on this topic). Further, harmonizing eligibility criteria across programs and systems is needed so that when children or families experiencing trauma or adversity enter a system, professionals can take a holistic perspective and refer them to cross-sector services; children and families would also be more likely to be eligible for needed services in other systems.

Recommendation 8-3: Policy makers and leaders in the health care, public health, social service, criminal justice, early care and education/education, and other sectors should adopt best practices and implement training for trauma-informed care and service delivery. They should implement trauma-informed systems that are structured to minimize implicit bias and stigma and prevent retraumatization. Standards for trauma-informed practice exist in a variety of service sectors, including health care and social services; those standards should be replicated and implemented across systems.

Build a Diverse, Culturally Informed Workforce

Building a high-quality health, early learning, or social services system will not improve outcomes if people do not engage with the system(s), and the populations where health inequities are most strongly expressed are often the least likely to engage in systems designed to serve them, because they have not historically felt welcomed in those systems. Most systems are built to specifications that are responsive to dominant culture norms and practices, but there is no one perfect system that works for everyone. Systems need to move beyond *what works* and address the question of *what works for whom, and under what circumstances*? Some communities may prefer to receive care and services via alternative modes, or in a manner of their choosing, or from providers who look and speak like them or understand their unique cultural or community identity. Offering services to caregivers and children in culturally and linguistically appropriate ways is critical to ensuring not only that good systems exist but that the people who have historically had the poorest outcomes engage in these systems as partners in generating health.

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System actions to offer culturally and linguistically appropriate services include training service providers to be attuned to and respect cultural and other identity differences and ensuring that systems have a range of appropriate linguistic services available for providers. Furthermore, systems that ensure that signage, forms, and data processes that clients use are available in multiple modes and languages facilitate participation in these systems. Another system action relates to the integration of workforces across sectors in the early-life period. Complementary disciplines can be drawn from to create teams that promote transdisciplinary service delivery. This might include traditional service providers and the expanded use of paraprofessionals, community health workers, peer support specialists, parent advisors, or others who bring expertise in, and lived experiences that are relevant to, the communities or populations being served. Bringing transdisciplinary providers together is just the first step; standards and workflows that allow these teams to collaborate effectively may need to be developed so that each functions at the top of its license to provide comprehensive wraparound support for families that need it.

Another system action is expanding the workforce in early-life serving sectors to ensure that service providers reflect the diversity of their communities and that people of diverse racial, ethnic, cultural, sexual orientation, gender expression, or other identities have access to more service providers who look like them and reflect their experiences. This could be accomplished through mentoring programs, outreach into culturally specific communities when recruiting or hiring, or scholarship programs to "plant the seeds" to introduce more diverse communities into fields where some communities are underrepresented.

Recommendation 8-4: Policy makers and leaders in the health care, public health, social service, criminal justice, early care and education/education, and other sectors should each develop a transdisciplinary and diverse workforce to implement culturally competent service delivery models. The workforce should reflect the diversity of populations who will engage in sector services.

Align Across Systems to Enhance Early-life Supports for Caregivers

Parental and caregiver supports are critical to promoting prosocial attachment, nurturing, and healthy family relationships that foster the healthy development of youth (see Chapter 4). Supporting caregivers should be an essential goal of multiple systems—for example, caregivers who do not have to worry about access to health care have a better opportunity to remain healthy and maintain positive relationships with their children, caregivers with adequate housing may experience reduced strain and be better able to bond with their children, and caregivers whose children are receiving effective early-life education and developmental supports may feel better equipped as effective and engaged parents.

Better support of caregivers will increase the likelihood of positive nurturing relationships between caregivers and children, and improvements in the multiple systems that are touch points for children and families should explicitly address caregiver support. Such changes may include encouraging the development of programs and policies that provide support for the whole family unit, including parents, children, and other important members of the family's extended caregiving system. Strategies include moving away from segmented policies, such as those that consider eligibility for supports or services separately for parents and children (e.g., child-only

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health insurance versus a family plan), and instead consider programs that incorporate or leverage the extended family unit or other close social networks.

Another strategy is policies and programs that provide assistance to families without requiring a separation between caregivers and children that might negatively impact attachments and family functioning, especially in early life. For example, programs could offer to families with young children certain financial or other assistance that does not include extensive requirements to leave the home in order to work or perform other tasks, which reduces the opportunities for developing, maintaining, and supporting healthy family attachments and functioning. (See Chapter 6 for examples.)

Efforts could be promoted to create wraparound services for families with greater needs, offering multiple types of assistance designed to support caregivers along a variety of dimensions within a single program or setting, even or especially if those services come from traditionally segmented or siloed systems in the community. Bundled services require less time for caregivers to navigate the systems, allowing more time for them to focus on effective caregiving. (See Chapters 4, 5, and 7 for examples of integrated and wraparound care.)

> **Recommendation 8-5: Policy makers and leaders in the health care, public health, social service, criminal justice, early care and education/education, and other sectors should improve access to programs or policies that explicitly provide parental or caregiver supports and help build or promote family attachments and functioning by engaging with the families as a cohesive unit. For families with intensive support needs, develop programs or initiatives designed to provide comprehensive wraparound supports along a number of dimensions, such as health care, education, and social services designed to address needs related to the social determinants of health that are integrated and community based.**

Support Integration of Care and Services Across All Dimensions of Health and Community

Integration refers to establishing standards by which services are delivered in ways that break down traditional silos and are informed by and responsive to the intersection of health and the key drivers of health. This might include social domains (such as social support, cultural identity, or community cohesion; see Chapter 4), clinical domains of health (such as physical, mental, behavioral, or dental health; see Chapter 5), economic domains (such as income, housing stability, or food security; see Chapter 6); educational domains (such as access to high-quality ECE programs or others; see Chapter 7). Integrated service models cohesively connect and align along the health continuum, address health holistically, and often include both primary prevention designed to forestall health crises and screening and response systems designed to act quickly when needs are identified.

A whole-family approach calls for integrating services at the point of care or intervention. Thus, the committee's vision for programs and services is that families have access to an array of clinical, early educational, family developmental, and psychosocial support and economic help in their communities. Rather than fragmented programs, the vision calls for easy and coordinated access across the breadth of needed services for households—viewed from a life course perspective and one that assures equity in access and use. Achieving this integration takes substantial work and community leadership, with programs having only limited incentives to collaborate, share accountability, and pool resources across sectors. The recent National

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Academies report, *Communities in Action*, provides principles and examples of integration at the community level (2017).

Integration and whole-family clinical care models are one example. This could include integrating the delivery of clinical care to include physical, mental, behavioral, and dental health services and connecting clinical care with other services for families (nutrition, early childhood programs, housing) either through a "one-stop" colocation strategy or a seamless and easy referral process with strong information sharing, "warm" handoffs between providers, and a "no wrong door" policy that helps people get whatever help they need easily through any entry point.

New models that integrate services to address the social determinants of health (SDOH) should follow this model. These include service delivery models that integrate supports across a range of SDOH, especially housing, transportation, food security, and social support, with a particular focus on family social supports and programs that integrate informal social networks into the health care, ECE, and services ecosystem (see Chapter 5 for recommendations on this for the clinical care system and its connections to other sectors). Another strategy is integration across settings and sectors, such as establishing community centers that include health care, nutrition, parent education, and resources to identify social needs and adversity and to help families find resources, or supporting church- or community-based programs, such as health care/community partnerships (see Chapter 5 of *Communities in Action*[7] for in-depth examples of these types of partnerships) to extend the reach of health or social service programs into culturally specific or otherwise historically underserved communities.

Recommendation 8-6: Policy makers and leaders in the health care, public health, social service, criminal justice, early care and education/education, and other sectors should integrate care and services across the health continuum, including the adoption of models that provide comprehensive support for the whole person in a contextually informed manner, leveraging and connecting existing community resources wherever possible, with a focus on prevention.

Support Payment Reform to Allow for Upstream Investment

Payment structures have a profound impact on how resources are invested and the ability to address complex, interrelated causes of poor health using multisector approaches. In many sectors, payment remains tied to the delivery of a service rather than its success in achieving an intended outcome, creating incentives to focus on the processes of providing services and maximize the associated billing of those processes rather than on creating better health outcomes. Similarly, the regulatory structure that governs how funds flow within and across systems is a major impediment to both cross-sector work and the spread and scale of interventions or programs that are known to succeed. As noted earlier, nearly every major system or sector has its own regulatory and/or funding structure, which is primarily designed to ensure accountability to spending within that system; there are usually rules that ensure the money within a sector stays within that sector. Accountability policies can also discourage upstream investment. For example, in the education system, schools and districts are not held accountable for school outcomes until third grade at the earliest, which creates disincentives for school leaders to invest in the early grades or before kindergarten.

[7] NASEM. 2017. *Communities in action: pathways to health equity*. National Academies Press: Washington, DC.

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Another strategy is to move to payment models that attach payment to the value of a provided service or its desired outcome, rather than to the simple delivery of the service. Value-based structures promote efficiency and impact over quantity of services and encourage upstream investment to address root causes rather than downstream work to deliver services. Payment models that emphasize upstream investment have substantial focus on prevention—in health care, ECE, and community services. In many cases, as documented in earlier chapters, the downstream payoff may be much later, sometimes years later.

Finally, there is a need to rethink budgeting and contracting to address the "success penalty" problem. For systems that contract with government, budget and contracting policies often set reimbursement rates or budgets by examining expenditures versus costs in previous years. When systems invest upstream and reduce costs, they risk being penalized by having their budgets reduced or rates cut the following year, disincentivizing success. Models that allow successful systems to reinvest some portion of saved dollars into scaling and spreading the approaches that helped them achieve that success will incentivize change.

Recommendation 8-7: Policy makers and leaders in the health care, public health, social service, criminal justice, early care and education/education, and other sectors should invest in programs that improve population health and in upstream programs that decrease long-term risk and poor health outcomes. These changes should be accompanied by accountability metrics to ensure that the spending is tangibly and demonstrably in service to the goals behind the original funding, but offer more flexibility in how those goals are achieved.

Support Transdisciplinary Research on the Complex Pathways of Health Equity

As described in this report, a tremendous amount is known about what works to advance health equity in early development (and the lifelong benefits of doing so), but there are still many unknowns in the area of implementing and scaling up interventions. Many interventions have shown promising results at small scale but have not been fully tested across multiple settings or in diverse communities and populations. Others have promising preliminary data but little high-quality evidence. The evidence around systems and policy changes—the work needed to address inequities with a multisector and systems-based approach—remains less certain than programmatic evidence in many cases precisely because it is complex and set in shifting environments that make it challenging to confidently attribute effects. There is also a relative dearth of research on heath equity produced through genuinely participatory methods that authentically engage the communities and populations most impacted by health inequities to help formulate, conduct, interpret, and disseminate results to community members, advocates, policy makers, and other decision makers.

The committee has identified important research needs in this report relevant to the chapter topics (see Recommendations 2-2, 4-1, 4-2, and 4-3); however; here, the committee recommends strategies focused on how to conduct research differently to help translate science to action across sectors, including needed data to inform subgroup analysis and to elucidate the complex causality related to health inequities to better target interventions across sectors. Recommendation 8-8 also identifies research needs that would support strategies identified throughout this report and relate to all systems that frequently interact with children and their families (for example, research on addressing discrimination, structural racism, and implicit bias

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training), and calls for an increase in participatory methods that engage communities, especially historically marginalized or excluded communities, as partners in research on health equity.

An important caution, however: although more targeted research is needed, enough is already known to act now to advance health equity in the prenatal and early childhood periods—this has been made abundantly clear in the preceding chapters of this report. The research recommended below is important to continually improve efforts and increase impact, but this should not impede action at the federal, state, tribal, territorial, local, and community levels. Here, the committee provides guidance on charting the course for future research to better meet the needs of the nation's children in the future and, specifically, to advance health equity.

Recommendation 8-8: The National Institutes of Health, Agency for Healthcare Research and Quality, Centers for Disease Control and Prevention, Health Resources and Services Administration, Centers for Medicare & Medicaid, U.S. Department of Education, philanthropies, and other funders should support research that advances the state of the science in several critical ways to advance health equity. Specific actions and research to support include:

Research Methods

- **Explore alternative methods to address complex causality.** Randomized controlled trials (RCTs) are often considered the gold standard for scientific evidence, and they remain a valuable tool (see Chapter 1 for more on their strengths and limitations). However, RCTs are not designed to assess the complex, interconnected causality that lies at the heart of health equity—if anything, they excel at reducing or controlling for complexity to isolate a single cause. It is necessary to embrace new approaches, including data science/data mining, multilevel modeling, integrated mixed methods, rapid-cycle analysis,[8] and other methods designed to employ a wide range of data and evidence to uncover the most effective approaches. Additionally, research needs to move beyond assessing cause and effect to explore the mediators and moderators of differential effectiveness—to identify not only what works, but what works for whom, and under what circumstances, to pursue a deeper understanding of causality that facilitates the ability to adapt promising and evidence-based models to optimally fit the needs and priorities of diverse populations.

- **Expand research into individual differences (heterogeneity) in response to adversity and treatment.** Many programs have shown some impact on important health outcomes; yet, in many cases, the effect sizes have been small to moderate. In part, this reflects the differential vulnerability and response to adversity and the differences in response to interventions. Studying heterogeneity in context and how it shapes responses to interventions and promising programs is a critical component of actionable research, but

[8] For example, the Frontiers of Innovation platform at the Harvard Center on the Developing Child employs a "structured but flexible framework that facilitates idea generation, development, implementation, testing, evaluation, and rapid-cycle iteration" (see https://developingchild.harvard.edu/innovation-application/frontiers-of-innovation for more information).

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current knowledge to identify these differences is limited. Exploring and understanding these differences will enable more targeted and tailored interventions.

- **Promote scientific research that includes individuals and families from underrepresented communities.** Even programs with a good evidence base often rely on studies that were conducted in dominant culture settings or with communities that are not representative of the full range of diversity (see Chapter 4) of characteristics such as race and ethnicity, socioeconomic or immigrant status, or sexual minority parent status. Studies are needed that seek to not only understand what works but to move beyond top-line findings to explore variation in outcomes across populations, settings, and subgroups (see Chapter 4 for a discussion on subgroup variation)—what works for whom, and under what circumstances. The results can be used to target efforts to address inequities more precisely in contextually informed ways. This scientific work should be informed by appropriate theoretical frameworks that take into consideration the multifactorial nature of early childhood development in diverse populations. Members of these populations or scientists with extensive research and or clinical experience working with these populations should be part of the investigative teams.

Research content:

- **Promote research that explicitly seeks to understand the interconnected mechanisms of health inequities**. Health equity has complex, interconnected root causes—factors that unfold across the distinct systems highlighted in this report—and also complex, interconnected mechanisms by which those factors shape health outcomes differentially across the life course. These mechanisms include biological development, social-psychological development, and differential opportunity structures and choice architectures that life presents people based on their circumstances. In addition, the biological and psychosocial responses to these mechanisms vary across the life course, with some developmental periods of high plasticity offering critical windows for establishing long-term trajectories for health outcomes. Understanding the *root causes* of inequities allows actions to prevent them; understanding the *mechanisms* by which root causes shape inequities helps to intervene more effectively when prevention is less than perfect, and understanding *variation in responsiveness* to those mechanisms across the life course will help ensure interventions are optimally targeted for maximum impact.

- **Support research that addresses discrimination and structural racism.** There is an urgent need for research on the structural roots of racism; how to stem the development of negative societal stereotypes, attitudes, and implicit biases; how to change those biases once they are formed; and how to develop applications of that knowledge that can help reduce discrimination. The impact of these forces—both the negative belief systems themselves and their structural and historical roots—on health and health outcomes are keenly felt, but there are few proven tools that allow for effective response.

- **Support research for trauma-informed care and implicit bias training:** As discussed in Recommendation 8-3, TIC and implicit bias training are critical tools for advancing health equity in the preconception through early childhood periods. However, that

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research base needs to be expanded further. Regarding research on structural racism and implicit bias, the 2017 report *Communities in Action: Pathways to Health Equity* provided the following recommendation on this topic, and the committee endorses it:

The committee recommends that research funders support research on (a) health disparities that examines the multiple effects of structural racism (e.g., segregation) and implicit and explicit bias across different categories of marginalized status on health and health care delivery; and (b) effective strategies to reduce and mitigate the effects of explicit and implicit bias.

There have been promising developments in the search for interventions to address implicit bias, but more research is needed, and engaging community members in this and other aspects of research on health disparities is important for ethical and practical reasons. . . . In the context of implicit bias in [schools], workplaces and business settings, including individuals with relevant expertise in informing and conducting the research could also be helpful. Therefore, teams could be composed of such nontraditional participants as community members and local business leaders, in addition to academic researchers (NASEM, 2017, p. 115).

- **Support systematic dissemination and implementation research.** In many sectors and increasingly across sectors, there are numerous well-tested examples of what works to help young families and improve outcomes. Enhancing and improving access to these programs will benefit from an extensive program of dissemination and implementation research to bring them to scale. Further, a mechanism to capture what has been tested but does not work is needed.

Measuring Success

Disparities have been measured for a long time and show the *outcomes* of inequity. What is lacking are good tools for measuring the various systemic and personal factors that influence and interact in complex ways to shape health outcomes over the life course. In the absence of such measures, designing the right kinds of system change remains a challenge.

The committee has identified a number of measures and indicators that can currently be measured and are important for tracking progress within each of the systems that act as key leverage points for early childhood development. For example, measures for primary caregivers include maternal depression and stress, parental feelings of rejection or hostility to the child, and support for mothers/primary caregivers. For children, measures include infants born at low or very low birth weight (LBW), breastfeeding at 6 months, blood lead levels (BLLs), social-emotional learning, meeting expectations in language development (e.g., measures of vocabulary), and kindergarten readiness. Measures for families include poverty (using the Supplemental Poverty Measure [SPM]), food insecurity, homelessness, health care insurance coverage, and exposure to toxicants through the home or early care environments. Taken together, improvements in these key metrics would represent systems that are moving in the right direction to address early-life drivers of inequities. However, other measures will be needed that are not yet available; the following section outlines these.

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The committee's conceptual model in Chapter 1 identifies two important dimensions to consider when exploring the early-life drivers of health inequities: the key systems that play the largest role in helping "set the odds" for healthy development and the interrelated mechanisms or pathways by which the influence of those systems are expressed into health outcomes over the life course (see Figure 8-1).

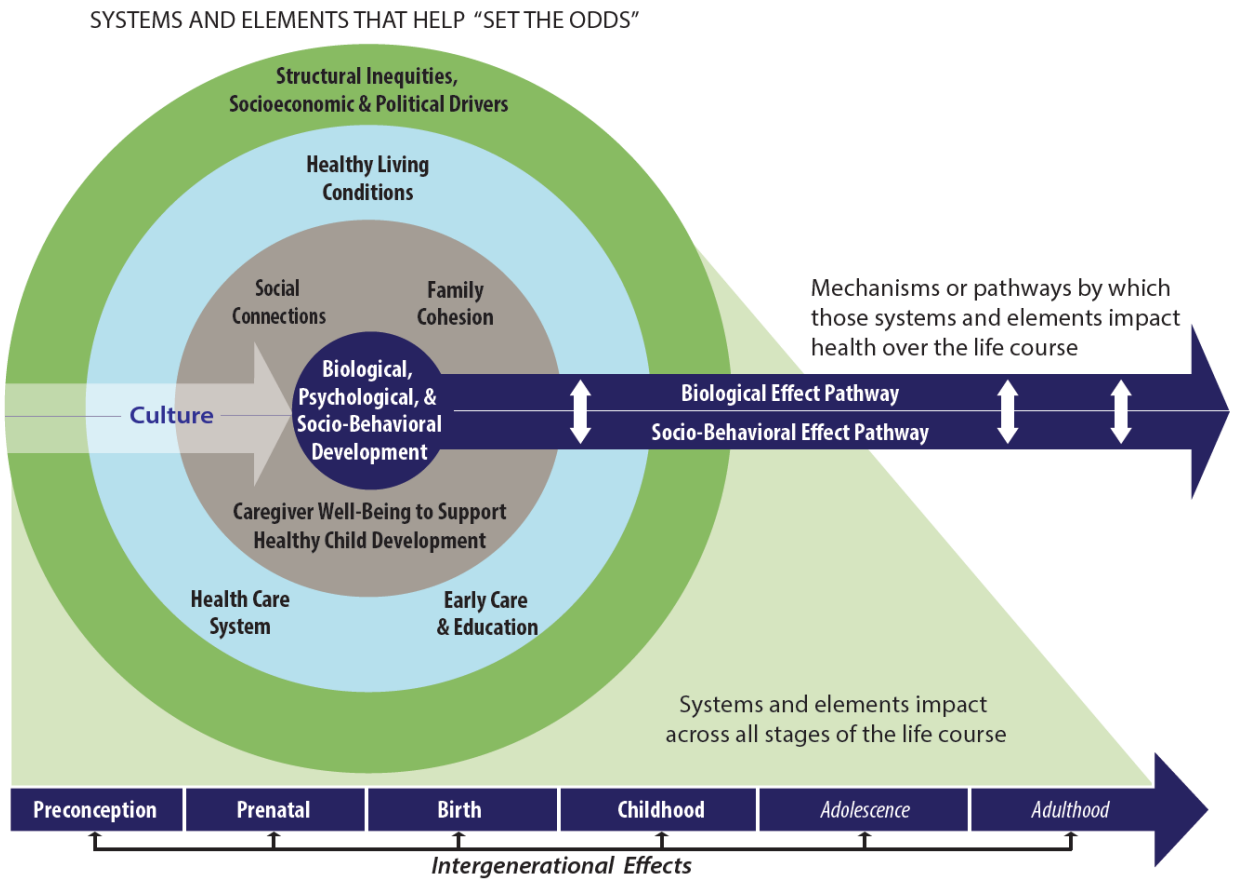

Figure 8-1: Leveraging early opportunities to achieve health equity across the life course: A conceptual framework
NOTE: The elements and systems included in the nested circles impact every stage of the life course.

The model leads naturally to several important insights about measurement. First, the systems that set the trajectory for health are explicitly connected and nested in the model—a child's biological, psychological, and socio-behavioral development (innermost ring) is nested within family and caregiver social systems, but those systems exist within and are shaped by key institutional systems like health care or ECE, which in turn exist within and are shaped by structural inequities and other historical, political, or macroeconomic forces. This means it is not enough to just have a good measure of caregiver attachment or of access to health care—it is necessary to have good measures of how those constructs *interact* to collectively set the odds of healthy early-life development and why systems present some populations with a greater or lesser probability of exposure to those constructs.

The mechanistic pathways by which those constructs act to shape health across the life course are also complex and interconnected. They might act via a biological pathway, whereby

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events or exposures that occur in childhood result in changes to key neurological or other biological systems that act over the years to enable or impede good health. They might also act via a socio-behavioral or psychological pathway, whereby events or exposures that occur in childhood change the likelihood that other events or exposures will occur later in life, alter the way people formulate their own identity or move through and react to key systems, or result in differences in the kinds of opportunity structures or choice architectures that life ultimately presents to someone. Crucially, these pathways are far from mutually exclusive: things that happen in one may profoundly impact another in positive and/or negative ways. This makes subgroup analyses based on the biological dynamics of the SDOH difficult.

None of this is deterministic. Rather, the committee conceptualizes health equity as a probabilistic challenge, with each element in the model contributing some adjustment to the odds of experiencing healthy development in early life and continued good health across the life course. Each factor contributes a probability that someone experiencing that factor will have good health outcomes; a person's overall odds of experiencing good health are a cumulative function of all of those probabilities. When populations experience different rates of exposure within systems for any reason, their cumulative odds will systematically differ. Even though the odds are not individually predictive—a given person may or may not "beat the odds" and experience any particular outcome—when applied to populations and expressed over time, different odds play out as systematically disparate outcomes between groups of people.

There is also a need to learn from and continuously collect data on both intervention successes and failures to better understand which program elements do or do not work and for which subgroup populations. In light of this approach to understanding health equity, more robust measures are needed to understand:

Understanding and measuring cumulative exposure. In this report, the committee identified a number of key factors that impact early-life development, ranging from influences in the microsocial or family environment, such as attachment, nurturing, and maternal well-being, to institutional levers, such as access to prenatal care or effective responses to trauma exposure, to macrosocial forces, such as racism and poverty. There are good tools available to measure exposure to some of these factors, but there are few methods for empirically understanding how exposures to risks or protective factors accumulate and combine over time to establish a cumulative overall risk profile. In the absence of a means to measure cumulative exposure, there is a lack of "math" for how to most effectively intervene when exposures occur.

Understanding the interaction among developmental pathways. Significant gains have been made in understanding how biological processes react to some contextual exposures—for instance, in the areas of science of trauma and toxic stress. However, there are few frameworks for understanding the multidirectional relationship between the biological, social-behavioral, and psychological development of young children. In particular, it is critical to understand how that interaction may vary across the life course in response to changing plasticity of biological systems, different stages of personal and cognitive development, and different life conditions and accumulated experiences in order to build a health equity strategy that puts the right responses in the right places at the right points of optimal potential impact.

Measuring interactions between systems. There are good methods for measuring how distinct elements of systems or policies impact health outcomes—for example, assessing whether

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systems with a given feature tend to produce better outcomes than systems without it. But understanding the dynamic interplay between systems—how a design decision in health care might interact with an economic policy or early learning curriculum to cumulatively shape the odds of good health—is not as developed. Models that can estimate "integrated risk" by combining key data from across the sectors where people live their lives are needed. Similarly, measures that examine results from cross-sector collaboration can help in documentation and accountability. As an example, school readiness at age five reflects both a child's health status and the family's access to basic income and housing, prevention of early adversity through support of maternal well-being, and community early education systems. Other measures, including variability in high school completion, third-grade reading readiness, unemployment, or arrest rates, may hold similar multisector significance as a lens on equity, though more work is needed to understand how measures like these ripple through the health continuum to impact disparities measures within other connected systems.

Improving methods to assess complex causality. As outlined in Recommendation 8-8, perhaps the biggest challenge facing health equity research is that of complex causality. As noted, many of the preferred tools of science, such as RCTs, are designed to control for and isolate single causes rather than embrace complex, interrelated causality that may include multilevel, multidirectional, and nested effects. For example, there needs to be greater exploration of effective community-based intervention approaches that use existing resources (e.g., as in "natural experiments").

CONCLUSION

These measurement needs represent a significant barrier to advancing the understanding of the biological and social pathways by which early life experiences are translated into health inequities, and the committee calls for improved measurement and research methodologies that can advance the state of the science and better inform effective societal responses. However, as noted earlier, there is no reason to wait for the science to solve all of these challenges before taking action. There are systemic disparities in health outcomes between populations, and the groundwork is laid for those disparities in early life. There are many solutions available to start now (see Chapters 4–7 and the system-level recommendations in this chapter). Measuring progress and refining approaches needs to continue, but there is no reason not to deploy the tools that are already available. Chapter 9 summarizes these actions and the key principles discussed in this report that provide a roadmap to advance health equity in preconception through early childhood.

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9

A Roadmap for Applying the Science of Early Development

This chapter highlights the main findings and concepts discussed throughout this report and summarizes the report's recommendations to lay out a roadmap for applying and advancing the science of early development. First, the chapter highlights the key big-picture findings gleaned from this report, followed by briefly reviewing the specific actions to be taken within each sector or area based on the report content.

OPPORTUNITIES FOR ACTION

Key Report Findings

The persistence of disparities between groups of people shows that there is more involved in producing them than individual behavioral choices (see Chapter 1 and NASEM, 2017[1]). The factors that drive health disparities from the preconception period through early childhood are complex, interconnected, and systemic; they result from exposures and experiences that children and families encounter throughout their lives, as well as choices they make. These exposures add up over the life course to exert a cumulative effect on health that is probabilistic, not deterministic. That is, the odds of good health are never fixed; individual exposures, experiences, and choices help set and adjust them over time. While groups of people have varying rates of exposure to key adversities and experiences, their distinct contexts shape their choices and opportunities, and thus they have different odds of experiencing good or poor health outcomes over time. For any given child, these odds may or may not lead to a particular health outcome. Applied to groups or populations over time, however, they manifest as health disparities.

To understand how certain subsets of U.S. children (e.g., racial/ethnic minority and low-income children) have heightened risk of exposure to the key drivers of poor health outcomes, the systems in which lives are lived and organized need to be examined. If those systems present different opportunity structures and choices to some groups of children and families, intentionally or not, they may generate health disparities upstream (early in life) even as those same systems might be working to address disparities downstream. Reducing health disparities by addressing their systemic roots is foundational to advance health equity; although it is a substantial challenge, it cannot be ignored.

As discussed in Chapter 8, health equity is a systems challenge. Implementing programs aimed at people in crisis alone will not advance health equity—those programs are important for

[1] NASEM (National Academies of Sciences, Engineering, and Medicine). 2017. *Communities in action: Pathways to health equity*. Washington, DC: The National Academies Press.

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addressing immediate needs, but until the systemic roots of inequity are addressed, there will likely be another set of crises around the corner.

In assessing the state of the evidence on early-life factors that shape health inequities, the committee developed a conceptual model (see Chapter 1, Figure 1-9) that summarizes the complex considerations that need to be addressed to break this cycle. Inherent in the model are three insights that are foundational to an effective health equity strategy targeted at the early-life period (preconception to early childhood):

1. The Importance of Intervening Early

The preconception through early-life periods are foundational for healthy development across the life course. Biologically, a number of critical systems are developing, and humans have high plasticity during these lifestages. Social-psychological and cognitive development are also important leverage points in the early years of life. Biological and social-psychological developmental pathways interact over the entire life course to set the trajectory for positive health outcomes, but the initial trajectories are established in early life. With the opportunity to play out across an entire life course, even small changes in initial health trajectories can result in large differences in long-term outcomes. Additionally, what happens early in life may impact not only initial health trajectories but how responsive those trajectories are to subsequent efforts to change them later in life—"resiliency" in the face of challenging circumstances. It is generally easier to change health trajectories for the better early on.

2. The Importance of Addressing Systemic and Structural Factors

This is critical because individual experiences, exposures, and choices are nested within and informed by those factors. The likelihood that a child or family will experience positive or adverse exposures is impacted by the systems they move through in their lives, including microsocial systems, such as families or immediate social networks, larger institutional systems, such as health care or education, and the cultural and historical forces that shape those institutional systems and experiences within them. In addition to affecting exposures to health-positive or health-negative factors, systems also impact how individuals *respond* to exposures or to interventions designed to mitigate the effects of negative exposures. Individual experiences within systems vary dramatically based on racial, cultural, or other personal characteristics. The effects of these systemic factors are by no means individually deterministic, but they do help set the odds, and when different odds play out over time and across groups of people, they generate systematically different health outcomes. A health equity approach requires systems to change in ways that improve opportunities for good experiences and reduce the odds of adverse exposures for populations that are currently experiencing disparate health outcomes.

3) The Interconnected Nature of Health Disparities

The systems that influence developmental and health trajectories are profoundly interconnected. The microsocial environment of children's daily lives—families and immediate social networks—is nested within and impacted by the key institutional systems that form the framework of society, and those institutional systems are nested within and impacted by cultural and historical forces that have shaped their essential character. The impact of any one of these

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systems will inevitably ripple across and shape what happens in others. In terms of the early-life drivers of health inequities, this means that there is no one-sector solution: the root causes are crosscutting, and improving outcomes in one sector is often interdependent on what happens in another. Poor outcomes in the early learning system may have roots in the family environment, which may in turn be shaped by the family's economic circumstances, which may also mean that the family is unable to access needed services in health care or other systems. Complex and interconnected root causes call for comprehensive and crosscutting solutions; one-dimensional strategies are not enough.

What Works? The Key Elements of a Successful Strategy

The above insights are not meant to suggest that the work being done to improve health outcomes within individual settings is unimportant or somehow reflects inadequate thinking. In fact, a tremendous amount has been learned about what works and what actions need to be taken—enough to identify the key building blocks of a successful strategy for addressing health equity. In this report, the committee examined the evidence to identify "what works" in a number of different critical contexts:

What Works in the Family Environment (Chapter 4)

Supporting the well-being of children starts with supporting the well-being of their c*aregivers,* and strategies designed to reduce children's potential exposure to maltreatment are particularly important. Strategies built to harness existing social resources within a family's environment or community are both cost effective and able to improve the continuity of support systems for caregivers. Context is important: culturally specific or experience-specific processes need to be considered to best respond to families' unique needs in ways that are most likely to engage them, and tiered approaches that recognize the different degrees of risk families face and offer appropriate resources based on that assessment are more likely to succeed than a one-size-fits-all approach.

What Works in the Health Care System (Chapter 5)

Health care practice, especially in the preconception, prenatal, and pediatric arenas, has not yet caught up to the current science to encompass key advances, such as the life course perspective, the role of adversity, trauma, and the toxic stress response, and the integration of social determinants of health (SDOH), to transform health care for young families into a system designed to facilitate health production. However, strengthening the content of health care alone will not advance health equity—access to health-enhancing services for all populations will be critical, by expanding public coverage and addressing nonfinancial barriers to participation. Further, holding health systems accountable for this shift through changes in equity-focused quality measures and aligned incentives that more comprehensively address health and the risks of poor health will be required. Other necessary actions include transforming the organization, payment, and delivery of health care services to allow for the adoption of integrated, whole-person care models that emphasize a life course approach and address upstream causes of poor health; facilitating the spread of multidisciplinary team-based care; developing trauma-informed systems that can respond effectively when a child is exposed to early-life adversity; and

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supporting cross-sector partnerships that intentionally connect health care services to the work of partners outside of the health system that are taking the lead to address the SDOH.

What Works in the Early Living Environment (Chapter 6).

It is critically important to provide predictability and security in the lives of children and their families by reducing childhood poverty, ensuring economic stability, and establishing a healthy and safe living environment. Money matters—having resources available to meet basic needs can improve health and reduce health and developmental disparities in early childhood, and public programs that provide resources to those families represent critical investments that "pay off" in the form of better outcomes for children as they enter school healthy and ready to learn and later move into adulthood. However, hinging these benefits on employment or earned income requirements that take caregivers from the home may be counterproductive for families if those requirements increase caregiver stress or hinder opportunities for developing healthy family relations, attachment between mother and child, or breastfeeding. Finally, the critical role of safe, stable, and affordable housing and food security are key determinants of childhood health, as are efforts to prevent and mitigate the impact of exposure to environmental toxicants during the preconception and early childhood periods.

What Works in Early Childhood Education (Chapter 7)

Early childhood education programs play a critical role in ensuring that children are healthy and ready for school life and beyond. It is necessary to incorporate health outcomes and health equity into a comprehensive approach to school readiness, including integrating it into the preparation and training of teachers. Educators—who are critically important caregivers to the children they work with—need adequate compensation and supports for their own health and well-being to ensure their effectiveness in the classroom or learning environment. Family support programs based in the home environment can provide valuable support when aligned with existing early childhood education systems. Last, there is a critical need to address access and affordability of promising models in the early childhood education world, especially for populations who have historically experienced health inequities.

Roadmap to Advance Health Equity Across the Life Course

In this report, the committee identified knowledge gaps that can be closed with greater investment in biological, behavioral, psychological, intervention, and implementation research. Each of these areas is equally important for addressing the gaps, and the committee recommends multidisciplinary research efforts to bring fresh, new ideas and practical approaches (such as innovative measurements and research methodologies) to advance efforts in tackling, head on, the serious challenges of health disparities to achieve health equity. The committee emphasized, however, that the great advances in knowledge since the release of *From Neurons to Neighborhoods* (NRC and IOM, 2000) make it very clear that policy makers, health providers, business leaders, and others in the public and private sectors do not need to wait any longer to take action. As the report brings to the fore, there are important opportunities to harness the many promising and evidence-based approaches that have come from advances in the understanding of how to address the neurobiological and socio-behavioral determinants of health

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disparities. In brief, the roadmap the committee has put forth includes the following key strategies (see Table 9-1):

- **Intervene early:** In most cases, early intervention programs are easier to implement, more effective, and less costly.
- **Support caregivers**: This includes both primary caregivers and caregivers in systems who frequently interact with children and their families.
- **Reform health care system services to promote healthy development**: Redesign the content of preconception, prenatal, postpartum, and pediatric care while assuring ongoing access, quality, and coordination.
- **Create supportive and stable early living conditions:**
    - **Reduce child poverty and address economic and food security,**
    - **Provide stable and safe housing, and**
    - **Eliminate exposure to environmental toxicants**
- **Maximize the potential of early care and education to promote health outcomes**
- **Implement initiatives across systems to support children, families, other caregivers, and communities**: Ensure trauma-informed systems, build a diverse and supported workforce, and align strategies that work across sectors.
- **Integrate and coordinate resources across the education, social services, criminal justice, and health care systems, and make them available to translate science to action.**

It is the committee's hope that this roadmap will catalyze the steps that need to be taken across systems to close the health equity gap and improve the lives of the nation's children.

See next page for Table 9-1

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TABLE 9-1 Roadmap to Apply the Science of Early Development[2]

Roadmap	Specific Action	Who
Intervene early	Implement programs that ensure families have access to high-quality, cost-effective community programs, including interventions to foster strong attachments and group-based supports in communities (Recommendation 4-4)	Policy makers at the federal, state, tribal, territorial, and local levels; philanthropic organizations
	Routinely track levels of risk among mothers and children over time using periodic assessments (Recommendation 4-5)	Health care providers
Support caregivers	Strengthen and expand evidence-based home visiting programs (Recommendation 4-3)	Federal policy makers; HRSA; ACF; federal, state, territorial, tribal, and local agencies overseeing program implementation
	Implement paid parental leave (Recommendation 6-1)	Federal, state, tribal, territorial, and local policy makers
Reform health care system services to promote healthy development	Increase access to preconception, prenatal, postpartum, and pediatric health care (Recommendation 5-1)	HHS; Medicaid agencies; public and private payers; federal, state, local, tribal, and territorial policy makers
	Expand accountability and improve quality of preconception, prenatal, postpartum, and pediatric care (Recommendation 5-2)	Public and private payers; HRSA, CDC, CMS, perinatal and pediatric quality collaboratives, and health-care-related workforce development entities
	Adopt policies and practices that improve the organization and integration of care systems from preconception through pediatric care and that focus on the caregiver and child together as the unit of care (Recommendation 5-3)	HHS; state Medicaid agencies; health systems leaders; federal, state, tribal, and territorial policy makers

[2] Some of these actions fit in more than one category but are only listed once; this table does not include recommendations from this report that are solely research based.

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Roadmap	Specific Action	Who
	Transform preconception, prenatal, postpartum, and pediatric health care to address the root causes of poor health and well-being (Recommendation 5-4)	HHS; public and private payers; medical accreditation bodies; WPSI, Bright Futures, ACOG, AAP, AAFP, and others
Create supportive and stable early living conditions • **Address economic, food, and housing security** • **Eliminate exposure to environmental toxicants**	Reduce barriers to participation to WIC and SNAP benefits; do not tie these benefits to parent employment for families with young children or for pregnant women (Recommendation 6-2)	Federal, state, local, territorial, and tribal policy makers
	Increase the supply of high-quality affordable housing that is available to families (Recommendation 6-3)	Federal, state, tribal, territorial, and local agencies
	Develop a comprehensive plan to ensure access to stable, affordable, and safe housing in the prenatal through early childhood periods (Recommendation 6-4)	Secretary of the HHS in collaboration with HUD and other relevant agencies, s
	Test new Medicaid payment models that engage providers and other community organizations in addressing housing safety concerns, especially focused on young children (Recommendation 6-5)	Center for Medicare and Medicaid Innovation
	Address the critical gaps between family resources and family needs through a combination of benefits that have the best evidence of advancing health equity, such as SNAP benefits, increased housing assistance, and a basic allowance for young children (Recommendation 6-6)	Federal, state, tribal, and territorial policy makers
	Support and enforce efforts to prevent and mitigate the impact of environmental toxicants during the preconception through early childhood periods (Recommendation 6-7, 6-8, and 6-9)	Federal, state, territorial, tribal, and local governments; CDC, EPA, FDA, the U.S. Consumer Product Safety Commission; health care providers

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Roadmap	Specific Action	Who
Maximize the potential of ECE to promote health outcomes	Develop a comprehensive approach to school readiness that explicitly incorporates health outcomes and leverages ECE systems and programs, including home visiting (Recommendation 7-1)	Federal, state, local, tribal, and territorial governments and other public agencies (e.g., school districts, city governments, public–private partnerships); Office of Child Care and Office of Head Start; Health and human service entities, the federal Early Learning Interagency Policy Board, state Early Childhood Advisory Councils, and federal, state, and local, tribal, and territorial agencies; HHS; OPRE; DOE
	Develop and strengthen coursework or practicums that focus on competencies of educators, principals, and ECE program directors that are critical to children's health, school readiness, and life success (Recommendation 7-2)	Degree granting institutions, professional preparation programs, providers of ongoing professional learning opportunities
	Develop and implement a strategic plan to (1) improve the quality of ECE programs by adopting the health-promoting standards and (2) expand access to comprehensive high-quality and affordable ECE programs across multiple settings (Recommendation 7-3)	Federal, state, tribal, and territorial policy makers in coordination with HHS, the Office of Head Start, and Office of Child Care
Implement initiatives across systems to support children, families, and other caregivers	Develop cross-sector initiatives that align strategies to address barriers to data sharing and integration, cross-sector financing, and other challenges to cross-sector collaboration (Recommendation 8-1)	For all actions in this section: policy makers and leaders in the health care, public health, social service, criminal justice, ECE/education, and other sectors who frequently interact with children and their families
	Enhance detection of early-life adversity and improve response systems (Recommendation 8-2)	
	Develop trauma-informed systems (Recommendation 8-3)	

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Roadmap	Specific Action	Who
	Build a diverse, culturally informed workforce in all relevant systems (Recommendation 8-4) Improve access to programs and policies across systems that provide parental or caregiver supports and help build or promote family attachments and functioning. For families with intensive support needs, develop programs or initiatives designed to provide comprehensive wraparound services (Recommendation 8-5) Integrate care and services across the health continuum, including the adoption of models that provide comprehensive support for the whole person by leveraging and connecting existing community resources (Recommendation 8-6)	
Resources need to be integrated and coordinated to translate science to action	Support payment reform to allow for upstream investment (Recommendation 8-7)	Policy makers and leaders in the health care, public health, social service, criminal justice, ECE/education, and other sectors who frequently interact with children and their families

TABLE NOTE: AAP = American Academy of Pediatrics; AAFP = American Academy of Family Physicians; ACF = Administration for Children and Families; ACOG = American College of Obstetricians and Gynecologists; CDC = Centers for Disease Control and Prevention; CMS = Centers for Medicare and Medicaid Services; DOE = Department of Education; ECE = Early Care and Education; EPA = Environmental Protection Agency; FDA = Food and Drug Administration; HHS = Department of Health and Human Services; HUD = Department of Housing and Urban Development; HRSA = Health Resources and Services Administration; OPRE = Office of Planning, Research and Evaluation; SNAP = Supplemental Nutrition Assistance Program; WIC = Women, Infants, and Children Program; WPSI = Women's Preventative Services Initiative.

REFERENCES

NASEM (National Academies of Sciences, Engineering, and Medicine). 2017. *Communities in action: Pathways to health equity*. Washington, DC: The National Academies Press.

NRC and IOM (National Research Council and Institute of Medicine). 2000. *From neurons to neighborhoods: The science of early childhood development.* The National Academies Press: Washington, DC.

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Appendix A

Criteria for Selecting Promising Models

The below criteria are adapted from the 2017 National Academies of Sciences, Engineering, and Medicine report *Communities in Action: Pathways to Health Equity*[1] that this report is building from, per the committee statement of task. The criteria were used to identify promising models from the prenatal through early childhood phases to highlight in this report. The committee did not evaluate the overall effectiveness of these efforts; rather, it used these promising models as examples throughout the report to highlight bright spots that have been able to use what is known from the science to advance health equity in the preconception through early childhood periods. Further, "promising" does not imply that the model is new but rather that it is a program or intervention that met the committee's core criteria, and each promising model has a unique approach and is at a different phase of development: some have been around for more than 30 years and have changed based on evaluations or input from users, while others have emerged in the past few years. These examples are not blueprints, and exact replicas might not work with all populations or locations, but the lessons learned and approaches used are valuable to those working to create positive change toward health equity during the preconception through early childhood periods. See Chapter 1 for more details.

This report applied three sets of criteria:

1. **Core criteria:** These function like inclusion criteria (i.e., to be included for consideration, the examples need to meet each of the six core criteria).
2. **"Aspirational" criteria:** The examples need to meet at least one, and preferably more, of the aspirational criteria.
3. **Contextual criteria:** These criteria are applied to the examples that meet the six core criteria and a number of the aspirational criteria to ensure that the examples are diverse in terms of communities/populations, approaches to solutions, and other characteristics.

[1] NASEM (National Academies of Sciences, Engineering, and Medicine). 2017. *Communities is action: Pathways to health equity*. Washington, DC: The National Academies Press.

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Set 1: Core Criteria

1. It focuses on preconception/prenatal and/or early childhood (whether intervention is focused on children or caregivers in their lives).

2. It is informed by findings from the neurobiological, closed-behavioral, and/or biological sciences.

3. It addresses at least one, preferably more, of the nine social determinants of health identified in the 2017 National Academies of Sciences, Engineering, and Medicine report (health systems and services, education, employment, the physical environment, the social environment, housing, income and wealth, public safety, and transportation).

4. It is designed to or has evidence of having an impact on a group or population that experiences health inequities.

5. It is multisectoral (i.e., at least two sectors engaged).

6. It includes an assessment of evidence, including data or best available information, to
    1. identify a problem and
    2. develop a solution that has a measurable outcome that there are plans to measure.

Set 2: Aspirational Criteria

1. It includes nontraditional partners and/or nonhealth domains.

Note: This is meant to be inclusive of nontraditional partners for communities to engage that may not necessarily be sectors (i.e., community organizers, PTA groups).

2. It is interdisciplinary.
    a. The solution draws on multiple sources, including practice-based experience and research from multiple disciplines.

3. It is multilevel—the intervention has multiple levels of influence, such as individual, family, organizational/institutional, or governmental.

Note: This does not mean that a solution must target each of these levels

4. It has a strong evaluation plan in place, including relevant measures to track the impact of the intervention.

5. It documents what it is trying to achieve, why that is important, how it plans to achieve the desired outcome (i.e., a theory of change), and/or the mechanisms being targeted based on scientific evidence.

6. It includes a plan for sustainability, including consideration of

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1. Long-term strategy and structure,
2. Funding, operating costs, resources, etc.,
3. Efficient use of resources,
4. Potential cost savings realized or return on investment,
5. Increased community capacity to shape outcomes,
6. Building the next generation of leaders, and
7. Clear policy solutions/changes at the local, state, or federal levels to support or scale a promising intervention or strategy.

7. It has transferable key elements[2] that could practically be applied or adapted to similar contexts in order to scale impact.

8. It incorporates the evidence required of proposed intervention(s):
    1. It addresses a significant health disparity (or disparities), based on data of a documented need or problem and data showing impact on at least one proximal or distal measure of a health disparity.

    2. The actual or projected health benefits are substantial/meaningful to the population(s) and community as a whole (not just statistically significant).

    3. There is ongoing data collection of processes and outcomes (flexibility in terms of what type of data is generated and applied).

    4. There is one or more high- or moderate-quality impact study of the approach.

Note: This includes health outcomes in a broad sense, related to social determinants (e.g., third-grade reading level rates) that are strongly linked to health outcomes.

9. The implementation process is well documented, including
    1. The key elements and subtleties of how the solution is contributing to success (not referring to legal documents/individual health data),
    2. Performance measurement,
    3. Particular practice (training, supervisory),
    4. Funding,
    5. Regulatory context, and
    6. Political context.

10. It is community-driven or informed: engagement with the community is evident preintervention and incorporated in the solution or the solution is initiated by the community/a community group/local government.

11. The solution is freely available to the community and not a proprietary resource.

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[2] Key elements are the functions or principles and activities of the solution that are necessary to achieve similar outcomes.

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Set 3: Contextual Criteria

As a whole, the set of examples selected will

1. address a range of the nine determinants of health identified in the 2017 National Academies of Sciences, Engineering, and Medicine report (health systems and services, education, employment, the physical environment, the social environment, housing, income and wealth, public safety, and transportation),

2. reflect rural, suburban, and urban contexts,

3. reflect diversity in several of the following population characteristics:
    - Race,
    - Ethnicity,
    - Age,
    - Gender identity,
    - Sexual orientation status,
    - Socioeconomic status,
    - Disability status, or
    - Other statuses (e.g., documentation status),

4. include solutions that require changes in the systems or policies within which the solution was implemented AND those that did not require changes in systems or policies to be effective, and

5. reflect various levels of political engagement.

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Appendix B

Public Meeting Agendas

MEETING 1

Thursday, May 31, 2018
Keck Center of the National Academies, 500 Fifth Street NW, Washington, DC 20001

11:35 am – 12:15 pm	**Presentation of the Statement of Task, Background, and Discussion** Dwayne Proctor *Senior Adviser to the President* *Robert Wood Johnson Foundation*
12:15 pm	**ADJOURN**

MEETING 2

Monday August 6, 2018
Keck Center of the National Academies, 500 Fifth Street NW, Washington, DC 20001

8:30 am	**Attendee check-in outside Room 100**
9:00 – 9:15 am	**Welcome and Opening Remarks** Jennifer E. DeVoe, M.D., D.Phil. (*Chair*), *Oregon Health & Science University* Victor J. Dzau, M.D., *President, National Academy of Medicine* (via video)
9:15 – 9:55 am	**Opening Presentation—Early Childhood Seen Through a Health Equity Lens** Paula Braveman, M.D., M.P.H., *University of California, San Francisco* **Discussion** Moderator: Myra Parker, J.D., Ph.D., *Committee member*

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9:55 – 10:55 am

Panel 1: Translating Early Development Science into Interventions and Policies

Gene–Environment Interactions: Role in Susceptibility and Resilience
Fernando Martinez, M.D., *University of Arizona, College of Medicine*

Translating Scientific Knowledge into Action on Early Childhood Development
Phil Fisher, Ph.D., *University of Oregon*

Discussion

Moderator: Pat Levitt, Ph.D., *Committee member*

10:55 – 11:10 am

BREAK

11:10 am – 12:45 pm

Panel 2: Approaches to Promote Healthy Development During the Prenatal and Early Childhood Phases

Early Childhood Innovation Network: Moving from Science to Action
Sarah Barclay Hoffman, M.P.P., *Early Childhood Innovation Network*

All Children Thrive: A Learning Network to Promote Child Health Equity in Cincinnati
Robert Kahn, M.D., M.P.H., *Cincinnati Children's Hospital*

The First 1,000 Days on Medicaid Initiative: A Medicaid-driven, Cross-sector Approach to Improving Child Outcomes
Suzanne C. Brundage, M.S., *Children's Health Initiative*, *United Hospital Fund*

The MOMS® Partnership: Partnering with Communities to Use Neurobiological and Socio-behavioral Sciences to Address Maternal Depression
Megan Smith, Dr.P.H., M.P.H., *MOMS Partnership*, *Yale School of Medicine*

Discussion

Moderator: Iheoma Iruka, Ph.D., *Committee member*

Discussant: Lee Beers, M.D., Children's National Health System

12:45 – 1:45 pm

LUNCH ***(lunch is not provided but can be purchased in the cafeteria located on the third floor)***

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1:45 – 2:55 pm

Panel 3: Policy and Systems Changes for Prenatal–Early Childhood Development

Leveraging the Science of Early Development: Creating Systems to Help Children Thrive
Neal Halfon, M.D., M.P.H., *UCLA Fielding, School of Public Health*

Social Determinants of Health Interventions, Fatherhood, and Reproductive Health
Milton Kotelchuck, Ph.D., *Harvard University Medical School*

Discussion
Moderator: Albert Wat, M.A., *Committee member*

2:55 – 3:35 pm

Closing Presentation—The Next Step in Evidence-based Policy: Implementing and Evaluating Universal Programs
Ron Haskins, Ph.D., *Brookings Institution*

Discussion
Moderator: Cynthia García Coll, Ph.D., *Committee member*

Public Comment

3:35 – 4:00 pm

Please add your name to the public comment sign-in sheet at the registration desk if you are interested in providing brief remarks to the committee.

4:00 pm

ADJOURN

MEETING 3

Monday, October 1, 2018
Beckman Center of the National Academies, 100 Academy Way, Irvine, CA 92617

8:00 – 8:05 am

Welcome and Opening Remarks
Jennifer E. DeVoe, M.D., D.Phil. (*Chair*), *Oregon Health & Science University*

8:05 – 9:05 am

Panel 1: Translating Early Development Science into Interventions
Greg Miller, Ph.D., M.A., *Northwestern University*
Greg Duncan, Ph.D., *University of California, Irvine*

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	Discussion
9:05 – 10:20 am	**Panel 2: Policy Perspectives on Prenatal and Early Childhood Development** Representative Ruth Kagi, *Washington House of Representatives* Senator Elizabeth Steiner Hayward, *Oregon Senate* Bobby Cagle, M.S.W., *L.A. County Department of Children & Family Services* Senator David Wilson*, Alaska Senate* **Discussion**
10:20 – 10:35 am	**BREAK**
10:35am – 12:00 pm	**Panel 3: Approaches to Promote Healthy Development During the Prenatal and Early Childhood Phases** Jessica Pizarek, M.A., *PolicyLink* Helena Sabala, *Chula Vista Promise Neighborhood* Anne Mauricio, Ph.D., M.A., *Family Check-up, Arizona State University* Elisa Nicholas, M.D., M.S.P.H., *The Children's Clinic, Serving Children and Their Families*
12:00 – 1:00 pm	**Panel 4: Caregiver Perspectives** Abraham Gomez Shalice Gosey Lori Hernandez Ana De Jesus Yesenia Manzo-Meda Maria Rodgers **Discussion** *Discussants:* Patricia McKenna, *SHIELDS for Families* Reggie Van Appelen, *SHIELDS for Families* Jennifer Eich, *Western Youth Services* Alexa Bach, *Network Anaheim*
1:00 – 1:15 pm	**Public Comment**
1:15 pm PT	**ADJOURN**

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Appendix C

Committee Biographical Sketches

Jennifer E. DeVoe, M.D., D.Phil. (Chair), is Chair of and Saultz Endowed Professor in the Department of Family Medicine at Oregon Health and Science University (OHSU). As a practicing family physician and doctorally trained health-services researcher, Dr. DeVoe studies access to health care, disparities in care, and the impact of practice and policy interventions on vulnerable populations. Her research portfolio spans both OHSU Family Medicine and OCHIN, Inc., a national community health information network based in Portland, OR. Dr. DeVoe leads a multidisciplinary research team with expertise in informatics, sociology, epidemiology, biostatistics, economics, primary care, mental health, health-services research, clinical medicine, health care disparities, and anthropology. Dr. DeVoe is the senior research advisor at OCHIN, where she previously served as chief research officer and executive director of its practice-based research network of community health centers from 2010–2016. Dr. DeVoe is a Principal Investigator (PI) or co-Investigator on numerous research studies funded by the Patient-Centered Outcomes Research Institute (PCORI), the Agency for Health Care Research and Quality, the National Cancer Institute, and the National Heart, Lung, and Blood Institute, with nearly $20 million in active grant funding. She also serves as co-PI of the ADVANCE Clinical Data Research Network, part of PCORnet, which is "horizontally" integrating electronic health record data, creating a unique community laboratory to include disadvantaged and vulnerable patients across the country. She holds joint appointments in the OHSU Department of Medical Informatics and Clinical Epidemiology and the Kaiser Permanente Northwest Center for Health Research. She also serves on the National Core Team for Family Medicine for America's Health Board of Directors and is past president of the North American Primary Care Research Group. She was elected to the National Academy of Medicine in 2014. Dr. DeVoe served as a NAM Puffer/ABFM Anniversary fellow from 2012–2014, and she was on the National Academies Committee on Accessible and Affordable Hearing Health Care for Adults from 2015–2016. Dr. DeVoe earned her M.D. from Harvard Medical School in 1999. Selected as a Rhodes Scholar in 1996, she also earned an M.Phil. and D.Phil. from Oxford University in 1998 and 2001, respectively. She completed her family medicine residency at OHSU in 2004 and earned an M.C.R. from OHSU in 2010.

Cynthia García Coll, Ph.D., is currently an adjunct Professor in the Pediatrics Department at the University of Puerto Rico Medical School and the Charles Pitts Robinson and John Palmer Barstow Professor Emerita at Brown University. Previously, Dr. García Coll was a professor in the clinical Ph.D. program and Associate Director of the Institutional Center for Scientific Research at Albizu University in San Juan, Puerto Rico. Prior to moving back to Puerto Rico,

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she spent 30 years at Brown. Her research focuses on the interplay of sociocultural and biological influences on child development, with particular emphasis on at-risk and minority populations. She received her Ph.D. in Personality and Developmental Psychology from Harvard University. Dr. García Coll has served on the editorial boards of many leading academic journals, including as the senior editor of *Child Development* and *Developmental Psychology*. She is a fellow of the American Psychological Association (APA) and the Association for Psychological Science. She has received awards from Tufts and Brown University, the Erikson Institute, the Society for Developmental and Behavioral Pediatrics, the Society for Research in Child Development (SRCD), and Progreso Latino. She has been on the governing boards of the United Way of Rhode Island, Rhode Island Community Foundation, SRCD, Society for the Study of Human Development, and Foundation of Child Development. She also served as member and chair of the Young Scholars Program at the WT Grant Foundation for 11 years. Her research has been funded by NIH, the McArthur Foundation, the WT Grant Foundation, and the Spencer Foundation.

Elizabeth E. Davis, Ph.D., is a professor of Applied Economics at the University of Minnesota and recently served as a member of the National Academies of Sciences, Engineering, and Medicine Committee on Financing Early Care and Education with a Highly Qualified Workforce. Dr. Davis conducts research in economics and public policy related to low-income families, child care and early education, and low-wage and rural labor markets in the United States. Her recent research has focused on disparities in access to high-quality child care, including development of new measures of access that are family centered and take cost, proximity, and quality into account. Other studies on early childhood topics have examined the role of child care subsidies in families' decisions about employment and the type, quality and stability of child care arrangements. In related work, she has examined the dynamics of participation in the child care subsidy programs in Maryland, Minnesota, and Oregon and advised state and federal agencies on child care subsidy policy. Her other research has examined the impact of local competition on wages and job turnover in the retail food industry, income equality, and the relationship between local labor market conditions and employment outcomes for disadvantaged workers. Dr. Davis earned her Ph.D. and M.A. in Economics from the University of Michigan—Ann Arbor.

Nadine Burke Harris, M.D., M.P.H., FAAP, is the first Surgeon General of California (appointed February 2019). Before her appointment, Dr. Burke Harris was the CEO of the Center for Youth Wellness. She is a pioneer in the field of medicine, dedicated to changing the way society responds to one of the most serious, expensive, and widespread public health crises of our time: childhood trauma. As founder and CEO of the Center, Dr. Burke Harris has brought this critical work to stages at the Mayo Clinic, American Academy of Pediatrics, Aspen Institute, and Partnership for a Healthier America. Her TED Talk, "How childhood trauma affects health across a lifetime," has been viewed more than 3.8 million times. Her work has been profiled in best-selling books, including *How Children Succeed* by Paul Tough and *Hillbilly Elegy* by J.D. Vance, as well as in Jamie Redford's feature film *Resilience*. It has also been featured on CNN, NPR, and Fox News and in *USA Today* and the *New York Times*. Dr. Burke Harris wrote a book on the issue of childhood adversity and health, called *The Deepest Well: Healing the Long-Term Effects of Childhood Adversity*, which was released in January 2018. Dr. Burke Harris received the Arnold P. Gold Foundation Humanism in Medicine Award, presented by the American Academy of Pediatrics, and the Heinz Award for the Human Condition. Additionally, she serves

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as an expert advisor to the Too Small to Fail initiative and as a member of the American Academy of Pediatrics National Advisory Board for Screening.

Iheoma U. Iruka, Ph.D., is the chief research innovation officer and director of the Center for Early Education Research and Evaluation at HighScope Educational Research Foundation. Prior to joining HighScope, she was at the Buffet Early Childhood Institute at the University of Nebraska and the Frank Porter Graham Child Development Institute at the University of North Carolina—Chapel Hill. Dr. Iruka's research focuses on determining how early experiences impact the learning and development of low-income and ethnic minority children and the role of the family and education environments and systems. She is engaged in projects and initiatives focused on how evidence-informed policies, systems, and practices in early education can support the optimal development and experiences of low-income, ethnic minority, and immigrant children, such as through family engagement and support, quality rating and improvement systems, and early care and education systems and programs. She is co-PI for the IES-funded Early Learning Network, Nebraska Site, a large-scale and far-reaching study aimed at identifying malleable factors that support early learning in preschool through Grade 3 that may be effective at closing the achievement gap for disadvantaged students. In particular, she has been engaged in addressing how best to ensure excellence for young diverse learners, especially black children, such as through development of a classroom observation measure, public policies, and publications geared toward early education practitioners and policymakers. She has served on numerous national boards and committees, including the National Academies of Sciences, Engineering, and Medicine's Committee on Supporting Parents of Young Children and the National Research Conference on Early Childhood. Dr. Iruka has a B.A. in psychology from Temple University, an M.A. in psychology from Boston University, and a Ph.D. in applied developmental psychology from the University of Miami—Florida.

Pat R. Levitt, Ph.D., is the chief scientific officer, vice president, and director of the Saban Research Institute. He is also a professor of Pediatrics, the Simms/Mann Chair in Developmental Neurogenetics at Children's Hospital Los Angeles, and the W.M. Keck Provost Professor in Neurogenetics at the Keck School of Medicine of the University of Southern California. He is the chief scientific officer for Children's Hospital Los Angeles. Dr. Levitt has held leadership positions at the University of Pittsburgh, Vanderbilt University, and the University of Southern California. In 2013, Dr. Levitt was elected to the National Academy of Medicine. Named a McKnight Foundation Scholar in 2002, Dr. Levitt also was a MERIT awardee from the National Institute of Mental Health and served as a member of its National Advisory Mental Health Council. He is an elected Fellow of the American Association for the Advancement of Science (AAAS), serving as the Neuroscience Section Chair in 2015–2015, and an elected member of the Dana Alliance for Brain Initiatives. He is a senior fellow at the Center on the Developing Child at Harvard University and serves as co-scientific director of the National Scientific Council on the Developing Child, a policy council that brings the best research from child development and neuroscience to assist policymakers and business leaders in making wise program investment decisions. He is a member of scientific advisory boards for several foundations and university programs and currently serves as editor in chief of *Mind, Brain, and Education* and on several editorial boards. Dr. Levitt's research program includes basic and clinical studies to identify the genetic and environmental factors that assure healthy development of the brain architecture that controls learning and emotional and social behavior. His clinical research studies address: how

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toxic stress responses in infants and toddlers may be detected as early as possible to promote resilience and better prevention; and children with autism who also have co-occurring medical conditions, such as gastrointestinal disorders. Dr. Levitt has published 295 scientific papers and made hundreds of academic and public presentations. He received his B.A. in Biological Sciences from the University of Chicago and a Ph.D. in Neuroscience at the University of California—San Diego. He completed a postdoctoral fellowship in Neuroscience at Yale University.

Michael C. Lu, M.D., M.S., M.P.H., is the Dean of University of California at Berkeley's School of Public Health (term began in July 2019). Before his appointment, Dr. Lu was a professor and the Senior Associate Dean for Academic, Student, and Faculty Affairs at the George Washington (GW) University Milken Institute School of Public Health. Prior to joining GW, Dr. Lu was the director of the Maternal and Child Health Bureau for the U.S. Department of Health and Human Services from 2012 to 2017. During his tenure, Dr. Lu transformed key federal programs in maternal and child health, launched major initiatives to reduce maternal, infant, and child mortality in the United States, and received the prestigious Herbert H. Humphrey Award for Service to America. Dr. Lu joined the federal government from UCLA Schools of Medicine and Public Health, where he held a joint faculty appointment in obstetrics-gynecology and community health sciences for nearly 15 years. He was best known for his research on racial-ethnic disparities in birth outcomes and his leadership in developing, testing, and translating a theory on the origins of maternal and child health disparities based on the life course perspective. Dr. Lu has served on two Institute of Medicine: Committee to Reexamine IOM Pregnancy Weight Guidelines and the Committee on Understanding Premature Birth and Assuring Healthy Outcomes. Dr. Lu received his B.A. in political science and human biology from Stanford University, an M.A. in health and medical sciences and public health from UC Berkeley, a medical degree from UC San Francisco, and his residency training in obstetrics and gynecology from UC Irvine.

Suniya S. Luthar, Ph.D., is a Foundation Professor of Psychology at Arizona State University and Professor Emerita at Columbia University's Teachers College. Dr. Luthar's research involves vulnerability and resilience among various populations, including youth in poverty, families affected by mental illness, mothers under stress, and teens in high-achieving, affluent communities (who reflect high rates of symptoms relative to national norms). Previously, she served on the faculty of the Department of Psychiatry and the Child Study Center at Yale University and then at Columbia University's Teachers College. Dr. Luthar is a Fellow of the American Association for Psychological Science and APA Divisions 7 and 37. She received the Boyd McCandless Young Scientist Award from APA, a Research Scientist Development (K) Award from the National Institutes of Health, an American Mensa Foundation Award for Excellence in Research on Intelligence, and an award for Integrity and Mentorship from the Society for Research in Child Development's Asian Caucus. Dr. Luthar's work is frequently cited in major news outlets in the United States, including the *New York Times, The Washington Post, The Wall Street Journal, The Atlantic,* NPR, PBS, and CNN, as well as overseas. Dr. Luthar has served as chair of a grant review study section at NIH and a member of the Governing Council of the Society for Research on Child Development. At the APA, she has served on the Committee on Socioeconomic Status and its Council of Representatives, and she is currently president of Division 7 (Developmental). Dr. Luthar received her B.S. and M.S. from Delhi

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University in 1978 and 1980, respectively, and her Ph.D. (Distinction) in Developmental/Clinical Psychology from Yale University in 1990.

Amy Rohling McGee, M.S.W., has served as the president of the Health Policy Institute of Ohio, a nonpartisan, independent nonprofit organization that provides information and analysis to state policymakers, since 2010. Her prior public sector experience includes work in the executive branch of state government, focused on policy related to issues such as health insurance, health system improvement, health information technology, and Medicaid, and she served in the state legislature as a Legislative Service Commission intern in the mid-1990s. Her private sector experience includes 5 years as the executive director of the Ohio Association of Free Clinics, representing health clinics that served the uninsured, primarily through volunteers, and several years in a management position at FIRSTLINK (now HandsOn Central Ohio). Ms. McGee earned her B.A. and M.A. from the Ohio State University. She has received the Business First "Forty under Forty" award and the Ohio State University Alumni Association William Oxley Thompson award.

Myra Parker, Ph.D., J.D., (Mandan-Hidatsa-Cree) is an assistant professor in the Center for the Studies of Health and Risk Behavior, Department of Psychiatry and Behavioral Sciences, in the University of Washington School of Medicine. She also works at the Indigenous Wellness Research Institute at the University of Washington School of Social Work. Dr. Parker has worked for over 10 years on tribal public health program implementation and coordination with tribal communities in Arizona, Idaho, and Washington and tribal colleges and universities across the United States. She has over 5 years of experience in tribal public health research. Prior to embarking on a career in research, Dr. Parker worked for 5 years in the policy arena within Arizona state government, in tribal governments, and with tribal working groups at the state and national levels. Her research experience in public health involves community-based participatory research, cultural adaptation of evidence-based interventions, and disparities research. She received an RWJF New Connections Junior Investigator grant in 2011, 1 year into her postdoctoral fellowship. Dr. Parker's research in this project focused on alcohol-related fatalities and tribal cross-jurisdictional agreements with local non-Native communities. She has provided trainings to tribal health department staff, tribal research teams, and urban Indian service delivery teams. She has also provided indigenous health research training to University of Washington students, from undergraduates through Ph.D. students. As an enrolled member of the Mandan and Hidatsa tribes, she is aware of the historical health practices and misconduct perpetuated on tribes and other minority and disenfranchised populations in the United States. Her background in law and policy has informed a broader understanding of the principles of ethics and honed her skills in identifying methods to address the disparities in research control and access through the use of formalized agreements. She has experience in working with tribes in their ongoing efforts to balance the collective rights of communities and individuals. Dr. Parker received her B.A. in human biology from Stanford University. She received a J.D. from the James E. Rogers College of Law at the University of Arizona in 2001, with an emphasis in federal Indian law. She received her M.A. in public health from the Mel and Enid Zuckerman School of Public Health at the University of Arizona in 2002. Dr. Parker graduated with a Ph.D. in health services from the University of Washington School of Public Health in 2010.

James M. Perrin, M.D., is professor of pediatrics at Harvard Medical School, former director of the Division of General Pediatrics at the Mass General Hospital for Children, and associate chair

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of pediatrics for research at MGH. His research has examined asthma, middle ear disease, children's hospitalization, health insurance, and childhood chronic illness and disabilities, with recent emphases on the epidemiology of childhood chronic illness and organization of services for the care of children and adolescents with chronic health conditions. Dr. Perrin holds the John C. Robinson Distinguished Chair in Pediatrics, founded the MGH Center for Child and Adolescent Health Policy (a multidisciplinary research and training center with an active fellowship program in general pediatrics), and directed the center for over 15 years. He is former president of the American Academy of Pediatrics, former chair of its Committee on Children with Disabilities, past president of the Ambulatory (Academic) Pediatric Association, and founding editor in chief of its journal, *Academic Pediatrics*. He also directed the Evidence Working Group reporting to the Maternal and Child Health Bureau for the Secretary's Advisory Committee on Heritable Disorders and Genetic Diseases in Newborns and Children. Dr. Perrin was elected to the National Academy of Medicine in 2016. He currently serves on the Board on Children, Youth, and Families and has previously served on the National Academies Committee on Improving Health Outcomes for Children with Disabilities and the Committee to Evaluate the Supplemental Security Income Disability Program for Children with Mental Disorders, as well as earlier committees on long-term care quality, disability in the United States, and the evaluation of federal health care quality activities. Dr. Perrin earned his A.B. from Harvard College and his M.D. from Case Western Reserve University School of Medicine. He had his residency and fellowship training at the University of Rochester.

Natalie Slopen, Sc.D., M.A., is an assistant professor in the Department of Epidemiology and Biostatistics at the University of Maryland, College Park School of Public Health. Dr. Slopen's research focuses on social influences on health, health disparities, and psychological and biological mechanisms through which childhood experiences are embedded to increase risk for later chronic diseases. The overarching goal of her research is to identify processes and conditions that can be targeted by interventions in order to reduce health disparities and promote health over the life course. Dr. Slopen completed her Master of Arts in Social Sciences at the University of Chicago, her Doctorate of Science in Social Epidemiology at the Harvard T.H. Chan School of Public Health, and her postdoctoral fellowship training at the Center on the Developing Child at Harvard University.

Albert Wat, M.A., is a senior policy director at the Alliance for Early Success, where he supports the organization's strategy and goals for early education, including increasing access to high-quality prekindergarten, improving the early learning workforce, and enhancing alignment with K–12 policies. Before joining the Alliance, Mr. Wat was a senior policy analyst in the Education Division of the National Governors Association Center for Best Practices, where he helped governors' staff and advisors improve their early care and education policies, from early childhood through third grade. Prior to NGA, Mr. Wat was the research manager at Pre-K Now, an advocacy campaign at the Pew Center on the States, where he authored a number of policy reports, managed research activities for the initiative, and provided analysis and information about the latest pre-K and early education research and policy developments to Pre-K Now staff and its network of state partners. In 2014, Mr. Wat served on the Institute of Medicine and National Research Council's committee on the Science of Children Birth to Age 8: Deepening and Broadening the Foundation for Success, which released the report *Transforming the Workforce for Children Birth Through Age 8: A Unifying Foundation* in April 2015. He also serves on the board of the Council for Professional Recognition. Mr. Wat has worked with

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schools, school reform nonprofits, and community-based organizations in the San Francisco Bay Area, southeastern Michigan, and Washington, DC. He holds a B.A. in psychology, an M.A. in education from Stanford University, and an M.A. in education policy from the George Washington University.

Bill J. Wright, Ph.D., is director of the Center for Outcomes Research and Education (CORE), an organization devoted to conducting innovative health policy and health-services research in support of health care transformation, with an emphasis on the social determinants of health. A sociologist whose primary emphasis in in longitudinal survey research in vulnerable or underserved populations, Dr. Wright has led the design and implementation of a numerous panel studies assessing the impacts of health systems and policy changes on historically underserved or excluded communities. Dr. Wright was a PI on the Oregon Health Insurance Experiment, the first randomized trial assessing the impacts of health insurance expansion, and currently oversees a portfolio of research on the impact of social needs and adversity on health and health care outcomes in vulnerable populations. He received his Ph.D. in sociology from South Dakota State University.

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