Race, Ethnicity, and Socioeconomic Status in Research on Child Health

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An extensive literature documents the existence of pervasive and persistent child health, development, and health care disparities by race, ethnicity, and socioeconomic status (SES). Disparities experienced during childhood can result in a wide variety of health and health care outcomes, including adult morbidity and mortality, indicating that it is crucial to examine the influence of disparities across the life course. Studies often collect data on the race, ethnicity, and SES of research participants to be used as covariates or explanatory factors. In the past, these variables have often been assumed to exert their effects through individual or genetically determined biologic mechanisms. However, it is now widely accepted that these variables have important social dimensions that influence health. SES, a multidimensional construct, interacts with and confounds analyses of race and ethnicity. Because SES, race, and ethnicity are often difficult to measure accurately, leading to the potential for misattribution of causality, thoughtful consideration should be given to appropriate measurement, analysis, and interpretation of such factors. Scientists who study child and adolescent health and development should understand the multiple measures used to assess race, ethnicity, and SES, including their validity and shortcomings and potential confounding of race and ethnicity with SES. The American Academy of Pediatrics (AAP) recommends that research on eliminating health and health care disparities related to race, ethnicity, and SES be a priority. Data on race, ethnicity, and SES should be collected in research on child health to improve their definitions and increase understanding of how these factors and their complex interrelationships affect child health. Furthermore, the AAP believes that researchers should consider both biological and social mechanisms of action of race, ethnicity, and SES as they relate to the aims and hypothesis of the specific area of investigation. It is important to measure these variables, but it is not sufficient to use these variables alone as explanatory for differences in disease, morbidity, and outcomes without attention to the social and biologic influences they have on health throughout the life course. The AAP recommends more research, both in the United States and internationally, on measures of race, ethnicity, and SES and how these complex constructs affect health care and health outcomes throughout the life course.
INTRODUCTION
In the United States, the racial and ethnic diversity and socioeconomic disadvantage of the child population are increasing dramatically. The US Census Bureau reported that in 2011, the country reached a historic tipping point, with Latino, Asian, mixed-race, and African American births constituting a majority of births. It is projected that by 2019, fewer than half of all children will be white, non-Latino. By 2050, 36% are projected to be white, non-Latino, and 36% are projected to be Latino. Furthermore, children are disproportionately affected by poverty. In 2009, children younger than 18 years of age constituted 25% of the population in the United States but represented 36% of people in poverty, and 42% lived in low-income families (<200% federal poverty level). Those most likely to be poor are African American, Latino, and Native American children, children in single-mother families, children of immigrant parents, and children younger than 5 years of age. Low parental education is also prevalent among today’s children. In 2012, less than one-third (31.4%) of children ages 6 to 18 years old lives with a parent who has a college education, and there are significant racial and ethnic disparities in parent education, with 40.4% of non-Hispanic white children living in a home with a college-educated parent, compared with 20.3% of non-Hispanic black and 13.1% of Hispanic children. Disparities in health and health care related to both race or ethnicity and socioeconomic status (SES) are well documented and have become an important focus for many health-related organizations, including the American Academy of Pediatrics (AAP), which included health equity in its strategic plan in 2008 and in 2010 published a policy statement on health equity and children’s rights. A technical report on racial and ethnic disparities in the health and health care of children was also published that year. Furthermore, the AAP cosponsored a conference in 2008 titled “Starting Early: A Life-Course Perspective on Child Health Disparities: Developing a Research Action Agenda,” which resulted in white papers and research recommendations published in a 2009 supplement to Pediatrics.

Understanding the mechanisms on how race, ethnicity, and SES create disparities is critical to alleviating them. In the report Children’s Health, the Nation’s Wealth, the National Research Council and Institute of Medicine model children’s health and its influences as the interaction over time of biology, behavior, and the social and physical environments. The model acknowledges the influence of developmental stages and indicates that all these factors operate in the larger context of services and policy (Fig 1). As children age, health is reflected in a kaleidoscope. As individual pieces of colored glass are arrayed in a fixed form, they create dynamic visual patterns of how multiple influences affect the child’s present and future health into adulthood. This model incorporates an ecologic perspective, recognizing individual, family, and community influences, and a life course perspective, acknowledging that a child’s health sets the trajectory for adult health. This model, coupled with marked increase in understanding of the physiologic and psychosocial bases of diseases affecting all age groups, provides a useful lens for understanding the roles of race, ethnicity, and SES in children’s health. The model highlights the mediating and moderating pathway (social and physical environments, behaviors, biology) through which race, ethnicity, and SES influence health across childhood.

![FIGURE 1](image)

National Research Council and Institute of Medicine model of children’s health and its influences.
In the United States, data on research participants and populations often include race and ethnicity as categorical variables, with the assumption that these variables exert their effects through innate or genetically determined biologic mechanisms. There has been an explosion of information on genetic factors involved in the pathogenesis of many diseases, including those that directly affect children and those with childhood roots that manifest in adulthood. The capacity to apply newly derived information from molecular and genetic science toward child health and health care will continue to grow. In addition, much research has documented the importance of social influences on health as fundamental causes or contributors to health potential and disease. Advances in genetics, epigenetics, and gene–environment interaction provide additional insight into the complexity of biological and social influences on health and their interaction and are important areas of study. Research exploring the genetic basis for childhood diseases and the developmental origins of adult diseases will undoubtedly lead to important advances in our understanding of how race, ethnicity, and SES influence health. However, a growing body of research suggests that the sociological and psychological dimensions of these variables also have strong, and in some areas predominant, effects on health.13–15 Because research questions are often framed and some data are collected with a focus on biological mechanisms, information on relevant social mechanisms is often lacking. Likewise, some research that focuses on social science questions lacks attention to biological mechanisms. Thus, for research in which both biological and social causation is relevant, it is often difficult to disentangle the contributions of these 2 dimensions, the social and biological, from one another. Inadequate attention to either biological or social influences in research limits the scope and impact of research conclusions and can lead to erroneous attribution.

The purpose of this statement is to highlight the relationships between race, ethnicity, and SES and to stimulate appropriate definition, measurement, and analysis of the variables in any study that proposes mechanisms, be they biological or social. In addition, this statement highlights some of the advances in measurement of these constructs and understanding of the mechanisms by which racial, ethnic, and SES disparities influence health and health care starting early and through the life course.

**RACE AND ETHNICITY**

**Conceptualization**

The Institute of Medicine16 report *Unequal Treatment: Confronting Racial and Ethnic Disparities in Healthcare* has defined race: “1. As many physical anthropologists abandon racial taxonomies altogether, race can more objectively considered a sociocultural concept wherein groups of people sharing certain physical characteristics are treated differently based on stereotypical thinking, discriminatory institutions and social structures, a shared worldview, and social myths; 2. A term developed in the 1700s by European analysts to refer to what is also called a racial group (a social group that persons inside or outside the group have decided is important to single out as inferior or superior, typically on the basis of real or alleged physical characteristics subjectively selected).” Ethnicity is defined as “a concept referring to a shared culture and way of life, especially as reflected in language, folkways, religious and other institutional forms, material cultures such as clothing and food, and cultural products such as music, literature and art.”16 The term *ethnicity* grew out of the field of anthropology, in which it was used to classify human populations on the basis of shared ancestry, heritage, culture, and customs.

Although race historically has been viewed as a biological construct, it is now known to have biologic and social dimensions that change over time and vary across societies and cultures.17 In the United States, there is much discussion about race, but other societies place less emphasis on race and more on class or other characteristics. Unlike a biologic characteristic such as gender, race and ethnicity categorization has developed and changed as geographic, social, and cultural forces have shifted.

The use and misuse of self-reported race data in research have been the subject of much discussion and debate. The use of race as a proxy for genotype is highly controversial.18,19 Those supporting evidence of genetic differentiation between races point out that ancestral tree diagrams show branching relationships corresponding to the major racial groups; major genetic clusters are associated with racial categories, and self-defined ancestry correlates with these clusters; and the frequency of alleles underlying disease or normal phenotypes can vary between racial groups.20 There are higher rates of specific genetic diseases among racial and ethnic groups, such as sickle cell disease in African American individuals and certain thalassemias among Southeast Asians, and it is argued that this probabilistic information can improve clinical decision making.18 However, others argue that genetic differences between racial groups are small compared with genetic differences within groups.21,22 There is a great deal of variability within a racial category. Depending on the geographic location of a population, genetic studies of African American people have documented a range of
7% to 30% white admixture.\textsuperscript{23,24} For most diseases, racial differences are currently unexplained purely on a genetic basis.\textsuperscript{19,25,26} Although sickle cell disease is a genetic disease more common in African American people, this racial association also reflects social forces: the enslavement and transport of Africans to the United States. Other origins of the HbS gene were in the Middle East and Indian subcontinent, where malaria was prevalent.\textsuperscript{24} Although the probabilistic association of sickle cell disease with African American people can be useful, not considering the possibility of sickle cell disease in other races can be problematic. Thus, a person’s race or ethnicity should not be used to assume health-related values, beliefs, or health susceptibility in a “clinical form of stereotyping.”\textsuperscript{18} Caution is needed when applying race-related population research findings to individual patients.

Racial and ethnic differences in health and disease may be related to SES, culture, bias, differential access to care, and environmental and genetic influences. Understanding the relative contribution of these factors is an important area of study, although measuring all influences with the same degree of precision and disentangling their contributions are daunting challenges, and findings should be interpreted in light of these challenges. Burchard et al\textsuperscript{20} suggest that “the evaluation of whether genetic (as well as nongenetic) differences underlie racial disparities is appropriate in cases in which important racial and ethnic differences persist after socioeconomic status and access to care are properly taken into account.” Advances in genetic admixture mapping to identify ancestral contributions may assist in the investigation of putative genetic factors related to race and ethnicity.\textsuperscript{27}

**Measurement**

Both race and ethnicity are subjective and context-specific characteristics that vary both across countries and within individuals. Although it is standard practice to describe participants and populations in terms of “race” or “ethnicity,” recommendations on how to measure these constructs have changed. For example, the decennial census classified respondents according to the 1977 Office of Management and Budget Directive 15, which includes 4 racial categories (American Indian or Alaska Native, Asian or Pacific Islander, black, and white) and 2 ethnic categories (Hispanic and not Hispanic). The 1997 revision of this directive\textsuperscript{26} expanded these categories to 5 by separating Asian from Pacific Islander and expanding the latter to Native Hawaiian or other Pacific Islander. The Revised Directive 15 rejected the use of a “multiracial” category but recommended that the 2000 census allow respondents to check more than 1 category. Use of the Hispanic ethnicity question followed by the 5-category self-reported race question has become the norm for health researchers using survey data.

In 2009, the Institute of Medicine Subcommittee on Standardized Collection of Race/Ethnicity Data for Health-Care Quality Improvement\textsuperscript{29} concluded that “there is strong evidence that the quality of health care varies by race, ethnicity, and language. Having quality metrics stratified by race, Hispanic ethnicity, granular ethnicity and language need can assist in improving overall quality and promoting equity.” They supported collection of the existing race and Hispanic ethnicity categories (Table 1) and more fine-grained categories of ethnicity (referred to as granular ethnicity and based on one’s ancestry) and language need (a rating of spoken English language proficiency and preferred language for health-related encounters). Granular ethnicity and languages relevant to the local area could be chosen from national standardized categories, with opportunity offered to individuals who want to self-identify their ethnicity and languages.\textsuperscript{29} Most recently, the Office of Minority Health published final data collection standards for race, ethnicity, primary language, gender, and disability status required by Section 4302 of the Affordable Care Act (http://minorityhealth.hhs.gov/templates/content.aspx?ID=9227&lvl=2&lvlID=208).\textsuperscript{30} Clearly, there is heterogeneity within racial and ethnic groups and a growing mixed-race or mixed-ethnicity population that may be better elucidated with the new measures. Furthermore, depending on the potential mechanisms for producing health differences, other measures may be considered, including socially assigned race (“How do other people usually classify you in this country?”)\textsuperscript{31,32} or skin color.\textsuperscript{33} For example, some researchers studying effects of vitamin D on biomarkers of disease risk will assess skin color in addition to race.

**Racial and Other Forms of Discrimination**

Discrimination has been defined as “actions carried out by members of dominant groups, or their representatives, that have a differential and harmful impact on subordinate racial or ethnic groups,” which may result from biases, prejudices, stereotyping, and uncertainty in communication and decision making.\textsuperscript{16} Discrimination can be based on race, ethnicity, or SES, with racial discrimination being the dominant form of discrimination that has received attention in the research community and with most literature on African Americans. A recent review of racism and child health identified 40 articles since 1950, most reporting on racism and behavioral and mental health outcomes.\textsuperscript{34} Racial prejudice may influence access to and the quality of health services through patient perception of discriminatory treatment, implicit clinician biases, or...
other mechanisms. Racism also contributes to health inequities through multiple mechanisms. Evidence is growing that racial discrimination (both individual and institutional) as a social stress on groups of children and families can influence psychology, physiology, and health behaviors. In response to this growing body of evidence, Geronimus proposed the “weathering” hypothesis, which postulates that health deterioration among African American people is a consequence of cumulative economic or social adversity and political marginalization. For instance, research has shown that clinically relevant stress-related biomarkers are higher in black than in white people, and these racial differences are not explained by SES.

For children and adolescents, discrimination related to race, ethnicity, or SES may affect the child directly. Timing of exposure to discrimination may alter its influence, with adolescents having the greatest likelihood of understanding the nature and meaning of discriminatory remarks and actions of others. Furthermore, discrimination experienced by parents may influence parenting, either through parental attitudes, beliefs, and behaviors or parental mental health, thereby influencing children’s health. Biological and behavioral responses place individual children at greater risk of both short- and long-term poor health outcomes and disease. For example, the gap between black and white infant mortality rates in the United States has been persistent over time and across socioeconomic strata. Researchers have found that lifelong accumulated experiences of racial discrimination are an independent risk factor for preterm delivery.

For researchers attempting to assess discrimination, self-report has been the traditional approach. Recent reviews have documented the strengths and limitations of several measures of perceptions of race- or ethnicity-based discrimination. Measures specific for children, for a wide variety of racial and ethnic groups, and for use in health care settings are limited. Measures of SES and ethnicity-related discrimination are needed and represent an important area for future work. Recognizing that self-report measures are affected by what people are able and willing to say, a newer approach has been to measure implicit bias. The Implicit Association Test is an indirect measure of implicit social cognition or prejudice, including among clinicians. It is a computer-based reaction measure of the relative strength between positive and negative associations toward one social group compared with another. Clinician implicit bias has been associated with physician treatment recommendations and patient-rated quality of medical visit communication and care. Such a tool may be useful to researchers studying mechanisms through which discrimination and stress affect health throughout the life span.

### TABLE 1

| Institute of Medicine Recommended Variables for Standardized Collection of Race and Hispanic Ethnicity |
|---|---|
| **Construct** | **Sample Measure** |
| Hispanic ethnicity | Are you Hispanic or Latino? yes/no |
| Race | What is your race? You may give more than one answer |
| a. White |  |
| b. Black or African American |  |
| c. American Indian or Native American |  |
| d. Asian |  |
| e. Native Hawaiian or Other Pacific Islander |  |
| f. Some other race |  |

**Acculturation**

Acculturation has important health influences. Although related to race and ethnicity, it is a distinct concept that has its roots in anthropology but has been used and defined in many other disciplines. Early definitions conceptualized a process of accommodation with eventual (and irreversible) assimilation into the dominant cultural group. A more recent framework delineates 2 separate processes: maintenance of the original culture and development of relationships with the new culture. There has been much research on acculturation, and some studies have found it to be associated with worse health outcomes, behaviors, or perceptions, but others have found positive effects on other health outcomes, health care use, and self-perception of health. Others argue that the associations between acculturation and health disparities are tentative and that its mechanism of influence is uncertain. Reviews of acculturation research have found wide variation in the definition and measurement of acculturation, and many measures are, at best, proxy variables that do not fully capture the construct of acculturation.

Measures are categorized as unidimensional, bidimensional, and multidimensional. Unidimensional instruments describe a linear process related to assimilation into a new culture, usually focusing on generation, years in the United States, English proficiency, and self-reported ethnic identity. Bidimensional instruments individually assess acculturative change in the “old” and “new” culture. Multidimensional instruments assess multiple aspects of the acculturative process, including attitudes, values, and ethnic interaction. Although there has been much research on measures, they have often been developed for a specific racial or ethnic group, and their applicability to other groups is
not clear. Research reviews state that “it is of fundamental importance that public health researchers provide a clear statement of the interpretation and use of acculturation within their studies and interventions” if acculturation measures are to be meaningful to the study of health.54 With the growing population of immigrant children in this country, the need to understand the mechanisms and impact of acculturation on child health is urgent.

Language

Like acculturation, language is also closely tied to race, ethnicity, and SES. Language preference may be a proxy for cultural differences in the perception of illness or access to care, and limited English proficiency may affect how well symptoms are communicated between the patient and the clinician. For example, 1 study found that language preference had a large effect on whether a child with symptoms consistent with asthma received a diagnosis.57 Language preference is often a part of acculturation scales. In research, language preference is typically accommodated through translation of questionnaires and supportive materials. Questionnaires may have differences in validity and reliability in different languages. Therefore, when the questionnaire is not currently available in the language preference group and is translated for the study, or when the questionnaire was custom designed and then translated for the study, it should be revalidated.

Socioeconomic Status

Conceptualization

Numerous terms describe and measure socioeconomic conditions. Such terms as social class, social stratification, and SES are often used interchangeably, although they have different theoretical foundations and interpretations.58 In this statement, SES is used to describe a complex, multidimensional concept that can be difficult to operationalize and accurately assess. Marmot and Wilkinson59 and others60 have provided ample evidence that SES is related to health status. Krieger et al61 condensed the complexity of SES into 2 aspects, both of which may exert influences on health directly or through associated experiences and behaviors. One aspect includes resources, such as education, income, and wealth. The other includes status or rank, a function of relative positions in a hierarchy. As described by Braveman et al62 in a review on SES in research, SES can affect health at different levels (eg, individual, household, neighborhood), through different causal pathways (eg, by influencing exposures, vulnerability, or direct physiologic effects), and at different times in the life course. They recommend that SES measurement involve considering plausible explanatory pathways and mechanisms, measuring as much relevant socioeconomic information as possible, specifying the particular socioeconomic factors measured rather than SES overall, and systematically considering how potentially important unmeasured socioeconomic factors may affect conclusions.

The effect of SES on current and future health is a particularly active area of research, shedding light on the complexity of mechanisms whereby this multidimensional variable influences health.58 Numerous studies have documented the direct relationship between SES and health throughout the life course. SES-associated gaps in early child development and school readiness are associated with later shortcomings in academic achievement and attainment, with implications for long-term productivity.63 Despite advances in quality and access to health care services, it is noteworthy that the discrepancy in health status between social status groups has persisted over time, even though the specific diseases that produced morbidity and mortality have changed.64 Furthermore, standard measures of health correlate with the extent of income discrepancy between rich and poor, and the extent of income inequality appears to explain more of the variation in health than is explained by other socioeconomic factors, even the absolute level of income.64–66 Across industrialized countries, the greater the discrepancy in income distributions, the worse the health status of the entire population.64,68 Data across individual states within the United States demonstrate a similar relationship.65,66 Longitudinal cohort studies have now clearly demonstrated the relationship between socioeconomic conditions during childhood and adolescence and future adult health.23,68 Two reviews provide evidence that SES during childhood is a powerful predictor of adult physical health independent of adult SES.69,70 The greater risk of low SES during childhood correlating with poorer adult outcomes has been found for overall mortality as well as morbidity and mortality from specific causes. For example, in Finland, the childhood SES of adult men correlated more closely with ischemic heart disease during middle age than did their adult SES.71 Three broad conceptual models, the timing, accumulation, and change models, have been hypothesized as potential pathways.68 The timing model suggests that the detrimental effects of low SES on adult health are greatest if low SES is experienced during specific sensitive periods of development. The accumulation model posits that the intensity and duration of SES disadvantage affect adult health, whereas the change model suggests that the directionality of SES mobility influences later health outcomes. Additional research is needed to
clarify the environmental, behavioral, and physiologic pathways and the timing, level, and duration of exposure critical to explaining how the SES of children affects both their current and future health status. Recognizing the link between early SES and later health questions the ability of SES data collected at 1 point in time to contribute to our understanding of the effects of SES factors on health. Furthermore, the depth, persistence, and intergenerational transmission of poverty have been shown to affect children’s health and health care.

**Measurement**

Multiple measures have been used to assess SES, including income, wealth, education, occupation, poverty level, neighborhood socioeconomic characteristics, past socioeconomic experiences (life course SES), and subjective social status (Table 2). Each construct contributes to the multidimensional concept of SES, and there are strengths and weaknesses in their measurement. Early work tended to use composite measures that consolidated multiple constructs into a single SES score. The Hollingshead Four Factor Index, based on occupation, education, gender, and marital status, is a classic example (A.B. Hollingshead, unpublished working paper, 1975). As research in the field of social epidemiology has progressed, it has become clear that these constructs each work through their own distinct (sometimes interactive) pathways to influence health and development. Thus, use of composite indices is not currently recommended. The choice of which construct to assess to represent social status should be based on the hypothesized pathways by which social inequalities in health accrue. Income and education remain the most widely used constructs to measure SES. Because their correlation is usually less than 0.50 and they probably represent different pathways to health; income and education should not be used as proxies for each other. Also, income and occupation are not proxies for wealth, which represents accumulated economic resources and may vary across individuals or groups with similar incomes or occupations. Failure to measure wealth may underestimate the contribution of SES to health.

It is increasingly recognized that beyond individual SES, neighborhood or contextual SES can influence health. Neighborhood SES has sometimes been used as a proxy for individual SES as addresses are linked to geocoded census tracts and census variables. Some argue that this may be a practical and population-based approach for monitoring disparities and allocating resources to address disparities. Additional research is needed to elucidate individual, family, and neighborhood contributions to SES and effects on health.

Perceived SES or subjective social status assesses how individuals perceive their relative position in the social hierarchy. Some suggest that perception of one’s social status may more fully capture the influence of SES on health by taking into account previous life experiences, context, and perceptions of the future. In addition, it is argued that perception guides behavior, and subjective social status is a new type of identity that influences health. Recent research has found that subjective social status in adolescents and adults is independently associated with a number of behaviors, outcomes, and psychological variables.

Although SES is a complex concept, understanding the mechanisms of how socioeconomic conditions influence health (eg, stress, discrimination, social capital) are critical to guide solutions. Researchers must consider the mechanisms by which SES might influence their outcomes in deciding which measures to use.

**INTERACTIONS BETWEEN RACE, ETHNICITY, SES, AND OTHER CONFOUNDERS**

Attributing causal effects to any one of these factors can be complicated by the relationships between constructs and the heterogeneity within and across the constructs. They interact with and are confounded by each other and potentially other social influences, including culture, bias, access to care, as well as geographic, environmental, and genetic influences. LaVeist et al found that racial segregation creates different exposures to economic opportunity, environmental conditions, and other resources that improve health, resulting in racial disparities confounded by disparities based on geographic location. For example, environmental pollution may be more intense in impoverished areas, and hazardous waste sites may even be intentionally located in poor and minority neighborhoods because of familial SES or discrimination based on race and ethnicity. Consequently, it is difficult to disentangle the adverse consequences of that pollution from the effects of discrimination.

Although most studies of such confounding or interaction of race, ethnicity, and SES have focused on adults, the need for inquiries into such factors affecting child health is equally strong. Of note, the AAP technical report on racial and ethnic disparities in the health and health care of children documented that 22% of studies did not perform multivariable or stratified analyses to ensure that racial and ethnic disparities persisted after adjustment for SES and other potential confounders. Little is known about the way that the relationships between these social factors influence the health of children or their effects on the trajectory of the development of adult health or disease. Nevertheless, multiple studies document racial and ethnic disparities.
### TABLE 2  SES Measurement Constructs

<table>
<thead>
<tr>
<th>Construct</th>
<th>Sample Measure</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Parent derived</strong></td>
<td></td>
</tr>
<tr>
<td>Household income</td>
<td>Which of these categories best describes your total combined family income for the past 12 mo? This should include income (before taxes) from all sources, wages, veteran's benefits, help from relatives, rent from properties, and so on.</td>
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<tr>
<td></td>
<td>&lt;$5000</td>
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<td></td>
<td>$5000–$11999</td>
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<tr>
<td></td>
<td>$12,000–$15,999</td>
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<td></td>
<td>$16,000–$24,999</td>
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<td></td>
<td>$25,000–$34,999</td>
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<td></td>
<td>$35,000–$49,999</td>
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<td></td>
<td>$50,000–$74,999</td>
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<td></td>
<td>$75,000–$99,999</td>
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<tr>
<td></td>
<td>$100,000</td>
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<tr>
<td></td>
<td>Don't know</td>
</tr>
<tr>
<td></td>
<td>No response</td>
</tr>
<tr>
<td>Education</td>
<td>What is the highest degree you earned?</td>
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<tr>
<td></td>
<td>High school diploma or equivalency (GED)</td>
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<tr>
<td></td>
<td>Associate degree (junior college)</td>
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<td></td>
<td>Bachelor's degree</td>
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<td></td>
<td>Master's degree</td>
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<tr>
<td></td>
<td>Doctorate</td>
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<td></td>
<td>Professional (eg, MD, JD, DDS)</td>
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<tr>
<td></td>
<td>Other (specify)</td>
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<tr>
<td></td>
<td>None of the above (less than high school)</td>
</tr>
<tr>
<td>Occupation</td>
<td>a. In what kind of business or industry do (did) you work?</td>
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<td></td>
<td>(For example: hospital, newspaper publishing, mail order house, auto engine manufacturing, breakfast cereal manufacturing)</td>
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<tr>
<td></td>
<td>b. What kind of work do (did) you do? (job title)</td>
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<tr>
<td></td>
<td>(For example: registered nurse, personnel manager, supervisor of order department, gasoline engine assembler, grinder operator)</td>
</tr>
<tr>
<td>Assets and wealth</td>
<td>Is the home where you live:</td>
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<td></td>
<td>Owned or being bought by you (or someone in the household)?</td>
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<tr>
<td></td>
<td>Rented for money?</td>
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<td></td>
<td>Occupied without payment of money or rent?</td>
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<tr>
<td></td>
<td>Other (specify)</td>
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<tr>
<td>Life course SES</td>
<td>What is the highest degree your parent(s) earned?</td>
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<tr>
<td></td>
<td>High school diploma or equivalency (GED)</td>
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<td></td>
<td>Associate degree (junior college)</td>
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<td>Doctorate</td>
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<td>Professional (eg, MD, JD, DDS)</td>
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<td></td>
<td>Other (specify)</td>
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<td></td>
<td>None of the above (less than high school)</td>
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<tr>
<td>Perceived social status</td>
<td>Think of this ladder as representing where people stand in the United States.</td>
</tr>
<tr>
<td></td>
<td>♦ At the top of the ladder are the people who are the best off: those who have the most money, the most education, and the most respectable jobs.</td>
</tr>
<tr>
<td></td>
<td>♦ At the bottom are people who are the worst off: those who have the least money, least education, and the least respected jobs or no job.</td>
</tr>
<tr>
<td></td>
<td>Where would you place yourself on this ladder? Fill in the circle that best represents where you think you stand, relative to other people in the United States.</td>
</tr>
<tr>
<td>Child (teen) derived</td>
<td>What is the highest grade (or year) of regular school you have completed? (Check one.)</td>
</tr>
<tr>
<td>Schooling</td>
<td>Elementary School</td>
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<tr>
<td></td>
<td>01 ☐</td>
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<td>02 ☐</td>
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<td>03 ☐</td>
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<td>06 ☐</td>
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<td>07 ☐</td>
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<td>08 ☐</td>
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disparities that persist even after adjustment for SES and the SES disparities that persist after adjustment for race or ethnicity.\textsuperscript{7,81} Disentangling the contributions and mechanisms by which race, ethnicity, and SES influence health and health care is necessary to effectively address disparities. Recognizing variability within populations, it is necessary to study those with poor health outcomes as well as those with similar exposures who are resilient to poor outcomes.

**SYSTEMATIC COLLECTION AND REPORTING OF DATA**

Children are dependent on their parents. Measurement of race, ethnicity, and SES and the related constructs of discrimination, acculturation, and language are key, but these characteristics are not fixed. Many children are of mixed race and ethnicity, which may necessitate assessment of both parents. How individuals conceptualize these factors can differ between parents and between parents and children, particularly as children age into adolescence and grow in their ability to understand and conceptualize these constructs. Selection of child or parent measurement must be based on the hypothesized pathway by which these factors affect health outcomes and the research question under study.

Disparities in children’s health and health care cannot be identified, monitored, addressed, or eliminated without consistent collection of race, ethnicity, language, and SES data. SES data are rarely collected systematically, although race and ethnicity have had more attention. Still, the most recent data available from US health plans reveal that 33% of health plan enrollees are covered by plans that do not collect race or ethnicity data.\textsuperscript{82} In 1 survey of patients, 80% thought health care providers should collect information about race and ethnicity, but 28%, especially patients of minority backgrounds, were uncomfortable giving the information.\textsuperscript{83} It is likely that socioeconomically disadvantaged people would feel similarly about the collection of SES data. Use of computerized systems to collect race and ethnicity data from patients has been shown to be feasible.\textsuperscript{84} The Patient Protection and Affordable Care Act of 2010\textsuperscript{85} requires that “any federally conducted or supported health care or public health program, activity or survey . . . collects and reports, to the extent practicable” data on race and ethnicity, in addition to a number of other factors. Interestingly, SES was not included in the Affordable Care Act mandate but could be easily added to computer algorithms. The systematic collection of data provides a tremendous opportunity for researchers to explore how race, ethnicity, and SES affect children’s health and health care. Monitoring of its standardized collection and appropriate use is critical. Guidelines on the use and reporting of race or ethnicity data in research have been recommended and include describing the reason for use of these variables and how they were measured, distinguishing between the variables as risk factors or risk markers, and adjusting for and interpreting differences in the context of all conceptually relevant factors, including SES and other factors.\textsuperscript{21} Furthermore, clinical or community-based research that addresses race, ethnicity, and SES should be guided by the AAP policy statement “Ethical Considerations in Research With Socially Identifiable Populations,” which emphasizes community involvement in the research process.\textsuperscript{86}

**CONCLUSIONS**

The racial and ethnic diversity of US children continues to increase dramatically, and the proportion of children who live in poverty is unacceptably high. The AAP acknowledges that race, ethnicity, and SES strongly influence health through social, physical, behavioral, and biological mechanisms (Fig 1) as fundamental causes, mediators, and moderators of child health and predictors of adult health status. Their influences are evident in the extensive and persistent racial, ethnic, and SES disparities in children’s health documented in the literature. Furthermore, these health disparities may persist through adulthood, leading to high health care expenses, increased work absenteeism (with reduced productivity), disability, and unemployment later in life. Despite these well-documented disparities and the importance of these factors, their measurement is challenging, both operationally and procedurally. The AAP strongly recommends the following:

- Recognizing that early life experiences can shape health across the life course and across generations, research to understand and address disparities related to race,
ethnicty, and SES must be prioritized.

- Child health studies including quality improvement research should measure race, ethnicity, and SES to improve their definitions and increase understanding of how these factors and their complex interrelationships affect child health. As guidelines on use and reporting of race and ethnicity data have recommended, researchers should be thoughtful and clear on the reason for use of these variables and how they were measured, distinguish between the variables as risk factors or risk markers, and adjust for and interpret differences in the context of all conceptually relevant factors, including SES.71

- Researchers should consider both biological and social mechanisms of action in relation to race, ethnicity, and SES as they relate to the aims and hypotheses of the specific area of investigation. It is important to measure these variables, but it is not sufficient to use these categories alone as explanatory for differences in disease, morbidity, and outcomes without attention to both the biological and social influences they have on health throughout the life course. If data relevant to the underlying social or biological mechanisms have not been collected or are unavailable, researchers should discuss their absence as a limitation of the presented research.

- Scientists who study child and adolescent health and development should understand the multiple measures used to assess race, ethnicity, and SES, including their validity and shortcomings. They must apply and, if need be, create research methods that will result in careful definitions of these complex constructs and their influences on child and adolescent health, analysis of interactions between them, and, ultimately, elucidation of the mechanisms of their effects on health throughout the life course.

- More research and funding on how race, ethnicity, and SES affect health and health care over the life course in the United States and internationally are needed. Potential areas for investigation include elucidation of the life course effects of race, ethnicity, and SES from prenatal through adulthood, the mechanisms underlying these effects, and ways to ameliorate negative outcomes. With this knowledge, effective intervention strategies can be developed and disseminated to improve the health of children and the adults they will become.

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Race, Ethnicity, and Socioeconomic Status in Research on Child Health
Tina L. Cheng, Elizabeth Goodman and THE COMMITTEE ON PEDIATRIC RESEARCH

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