

Are Black Girls Exhibiting Puberty Earlier? Examining Implications of Race-Based Guidelines

Adeiyewunmi (Ade) Osinubi, MD,^a C. Paula Lewis-de los Angeles, MD, PhD,^{a,b} Patricia Poitevien, MD, MSc,^{a,b} Lisa Swartz Topor, MD, MMSc^{a,b,c}

From 1977 to 2013, the average age of thelarche, the first sign of puberty in girls, decreased by nearly 3 months per decade.¹ This trend, along with rising rates of precocious puberty in girls, has significant implications for girls' physical and psychosocial development, especially because early puberty has been linked to future health risks.^{2,3}

In 1997, the Pediatric Research in Office Setting study introduced race into the discourse surrounding pubertal timing trends.⁴ In a study of approximately 17 000 girls, Black girls at every age had more advanced breast development compared with White girls. Breast development had begun at age 6 years for 6.4% of Black and 2.8% of White girls and, by 8 years, 37.8% of Black and 10.5% of White girls.⁴ Because of these findings, the Pediatric Endocrine Society Drugs and Therapeutic Committee recommended new race-based criteria for defining precocious puberty.⁵ Proposed recommendations stated that breast development should be considered precocious in Black girls younger than 6 years and White girls younger than 7 years.⁵ Although these guidelines were never formally recognized, race has become widely accepted

as a factor in the pubertal timing of girls and has been included in recent American Academy of Pediatrics (AAP) clinical reports⁶ and Pediatric Endocrine Society guidelines.⁷ As a result, many patients, caregivers, medical students, and physicians have been taught that Black girls experience puberty at earlier ages. In addition, in scientific and educational resources^{8,9} and the lay media,^{10,11} race continues to be associated with precocious puberty without considering the potential causes of this association, including the impact of racism.¹² Although the AAP has made it clear that race-based medicine is faulty and detrimental, its eradication from everyday practice remains a challenge.¹³ This article will examine the inclusion of Black race in discussions surrounding puberty, the implications of these race-based approaches to pubertal norms, and recommendations for reframing these notions.

POTENTIAL INFLUENCES ON PUBERTAL TIMING

The most recent AAP clinical report on precocious puberty⁶ includes that race/ethnicity, among other factors, should be taken into account when evaluating early puberty in girls.

^aWarren Alpert Medical School of Brown University, Providence, Rhode Island; and ^bDepartment of Pediatrics and ^cDivision of Pediatric Endocrinology, Hasbro Children's Hospital, Providence, Rhode Island

Drs Osinubi, Topor, Poitevien, and Lewis-de los Angeles conceptualized the paper, drafted the initial manuscript, and reviewed and revised the manuscript; and all authors approved the final manuscript as submitted and agree to be accountable for all aspects of the work.

DOI: <https://doi.org/10.1542/peds.2021-055595>

Accepted for publication Apr 13, 2022

Address correspondence to L isa Swartz Topor, MD, MMSc, Division of Pediatric Endocrinology, 593 Eddy St, Providence, RI 02903. E-mail: lisa_swartz_topor@brown.edu

PEDIATRICS (ISSN Numbers: Print, 0031-4005; Online, 1098-4275)

Copyright © 2022 by the American Academy of Pediatrics

FUNDING: No external funding.

CONFLICT OF INTEREST DISCLOSURES: The authors have indicated they have no potential conflicts of interest to disclose.

To cite: Osinubi A(A), Lewis-de los Angeles C.P, Poitevien P, et al. Are Black Girls Exhibiting Puberty Earlier? Examining Implications of Race-Based Guidelines. *Pediatrics*. 2022;150(2):e2021055595

Race is a social construct, without any biological or genetic basis,¹⁴ and it may inadvertently be used as a proxy for variables including obesity, environmental exposures, psychological stress, and, importantly, racism itself.

Food insecurity and lower socioeconomic status are factors that affect rates and trajectories of obesity in youth,¹⁵ which may be related to the higher rates of obesity observed in Black girls.¹⁶ Obesity can influence differences in pubertal timing because increased adipose tissue is associated with more estrogen.^{5,17} However, the association between obesity and early puberty remains under investigation.¹⁸ One recent study reported that total body fat has a variable influence on puberty because girls with more total body fat had slower progression of breast development, yet achieved menarche earlier than girls with lower total body fat levels.¹⁹ In addition, a recent review suggests that obesity alone does not explain early puberty, and girls with obesity should continue to have a full workup for precocious puberty despite elevated body mass index.²⁰

Environmental racism, a phenomenon that describes the disproportionate exposure to and impact of environmental hazards on minority populations, must also be considered.²¹ Endocrine disrupting chemicals (EDCs) interfere with hormone signaling²² and have been linked to increased obesity rates and early puberty.²³ One EDC is bisphenol A,²⁴ a chemical found in plastic bottles. Studies have demonstrated that Black communities have higher exposures to bisphenol A and other EDCs.²⁵ Hair products marketed to Black communities are another environmental exposure that may influence earlier pubertal onset^{26–28} because they have been shown to

contain estrogens, human/bovine placenta, and EDCs.^{29–31} One study reported that 49.4% of Black people used products containing EDCs compared with 7% of White people.³⁰

Early childhood stress may also affect pubertal timing. The Weathering Hypothesis posits that the cumulative effects of social, economic, and political adversity fuel the early health deterioration of Black women.³² Early life stress may disproportionately affect Black girls³³; studies have shown associations between household stress³⁴ and both earlier puberty³⁵ and menarche.^{35,36}

IMPLICATIONS

Race Pathologization

The idea that racial health disparities exist because of biological differences between racial groups perpetuates a tradition of race pathologization.³⁷ Race pathologization is the practice of attributing poor health outcomes to an individual race rather than to the sociopolitical factors that influence such outcomes and perpetuating false ideas that health disparities are due to biological differences. In working toward health equity, it is essential to recognize that the burden of disease affects certain racial groups more than others and to acknowledge the roles of institutional and structural racism in fueling health inequities when identifying such disparities.

Adultification of Black Girls

The notion that Black girls achieve puberty at earlier ages may perpetuate adultification bias, a form of racial prejudice in which Black children are treated and judged as more mature than others of the same age.³⁸ This bias has important implications for Black children across social structures.

Black girls may be subject to unwanted sexual advances, harsher punishment by educators in schools, and experience greater use of force and/or harsher penalties in the criminal justice system.^{38–40}

Suboptimal Patient Care

Race-based medicine can lead to the inappropriate withholding of diagnostic and therapeutic interventions from patients. If physicians and other providers are taught that Black girls “naturally” achieve puberty at earlier times, they may fail to ask important questions, provide appropriate counseling, and unmask diagnoses. Precocious puberty can be caused by central nervous system tumors and genetic conditions such as McCune-Albright syndrome.⁴¹ Commonly held biases and expectations about pubertal timing in Black girls may prevent physicians from implementing a thorough workup on their patients, leading to missed or delayed diagnoses.

RECOMMENDATIONS

Applying Principles of Race-Conscious Medicine

In evaluating pubertal timing disparities, it is important to move away from race-based medicine and adopt principles of race-conscious medicine.⁴² Race-conscious medicine decenters race and identifies racism as a primary driver of disparities.⁴²

Specific Approaches for Girls With Early Puberty at the Bedside

Pediatricians can prioritize the judicious evaluation of all girls who present with early puberty, despite commonly taught principles of what is “natural” for Black girls. Pediatricians must also attend to Black girls’ psychosocial development and needs. An expectation of early maturity among Black girls can have long-lasting

implications on physical and emotional health.³⁸ Earlier pubertal maturation has been associated with higher rates of unwanted sexual advances and sexual harassment.^{39,43} It is essential that clinicians are aware of these risks and support their patients by asking questions, providing age-appropriate guidance, and reiterating the importance of consent.

Future Opportunities to Improve Individual and Population-Based Health Care

In the short term, pediatricians are uniquely positioned to counsel patients and their parents about environmental factors affecting early puberty. One example is helping families identify and avoid personal care products that contain EDCs, estrogens, and placental materials. Pediatricians must continue to advocate for legislation to combat childhood obesity, including neighborhood and city-wide initiatives that increase green space and eliminate food deserts.

Further research is essential in understanding recent trends in pubertal development. Relying on race as an explanation for differences in pubertal timing halts the intellectual curiosity and questioning needed to further scientific inquiry. Although many studies have identified differences in pubertal timing by race, few have focused on why these differences may exist. The effects of stress and weathering on Black girls' pubertal presentation must be further elucidated. Additionally, investigating how to best support the psychosocial needs of girls undergoing early puberty will provide a more holistic approach to patient care.

CONCLUSIONS

Transforming our approach to disparities in pubertal timing can

lead to improved patient care for Black girls. From a population health perspective, studying factors that influence pubertal timing can help researchers gain a better understanding of pubertal trends and ways to halt the drift toward earlier puberty. Individualizing care rather than categorizing care based on race allows for optimal diagnostic evaluation, age-appropriate counseling, and works toward dismantling harmful structures such as adultification bias, leading to a more equitable and accessible health care system for all.

ABBREVIATIONS

AAP: American Academy of Pediatrics

EDC: endocrine disrupting chemical

REFERENCES

1. Eckert-Lind C, Busch AS, Petersen JH, et al. Worldwide secular trends in age at pubertal onset assessed by breast development among girls: a systematic review and meta-analysis. *JAMA Pediatr*. 2020;174(4):e195881
2. Collaborative Group on Hormonal Factors in Breast Cancer. Menarche, menopause, and breast cancer risk: individual participant meta-analysis, including 118 964 women with breast cancer from 117 epidemiological studies. *Lancet Oncol*. 2012;13(11):1141–1151
3. La Vecchia C, Franceschi S, Decarli A, Gallus G, Tognoni G. Risk factors for endometrial cancer at different ages. *J Natl Cancer Inst*. 1984;73(3):667–671
4. Herman-Giddens ME, Slora EJ, Wasserman RC, et al. Secondary sexual characteristics and menses in young girls seen in office practice: a study from the Pediatric Research in Office Settings network. *Pediatrics*. 1997;99(4):505–512
5. Kaplowitz PB, Oberfield SE; Drug and Therapeutics and Executive Committees of the Lawson Wilkins Pediatric Endocrine Society. Reexamination of

the age limit for defining when puberty is precocious in girls in the United States: implications for evaluation and treatment. *Pediatrics*. 1999; 104(4 Pt 1):936–941

6. Kaplowitz P, Bloch C; Section on Endocrinology, American Academy of Pediatrics. Evaluation and referral of children with signs of early puberty. *Pediatrics*. 2016; 137(1)
7. Bangalore Krishna K, Fuqua JS, Rogol AD, et al. Use of gonadotropin-releasing hormone analogs in children: update by an international consortium. *Horm Res Paediatr*. 2019;91(6):357–372
8. Pediatric Endocrine Society/American Academy of Pediatrics Section on Endocrinology Patient Education Committee. Pediatric Endocrinology Fact Sheet Precocious Puberty: A Guide for Families. Available at: <https://pedsendo.org/patient-resource/precocious-puberty>. Accessed February 26, 2022
9. Harrington J, Palmart MR. Definition, etiology, and evaluation of precocious puberty. Available at: <https://www.uptodate.com/contents/definition-etiology-and-evaluation-of-precocious-puberty>. Accessed February 26, 2022
10. Cleveland Clinic. Puberty. Available at: <https://my.clevelandclinic.org/health/articles/22192-puberty>. Accessed March 3, 2022
11. Mayo Clinic. Precocious puberty. Available at: <https://www.mayoclinic.org/diseases-conditions/precocious-puberty/symptoms-causes/syc-20351811>. Accessed November 21, 2021
12. Lalwani S, Reindollar RH, Davis AJ. Normal onset of puberty have definitions of onset changed? *Obstet Gynecol Clin North Am*. 2003;30(2):279–286
13. American Academy of Pediatrics Board of Directors and Executive Committee. AAP perspective: race-based medicine. *Pediatrics*. 2021;148(4):e2021053829
14. Collins FS. What we do and don't know about 'race', 'ethnicity', genetics and health at the dawn of the genome era. *Nat Genet*. 2004;36(11 Suppl):S13–S15
15. Tester JM, Xiao L, Tinajero-Deck L, Juarez L, Rosas LG. Food insecurity influences weight trajectory in children with obesity [published online ahead of print February 15, 2022]. *Child Obes*.

16. Anderson PM, Butcher KF, Schanzenbach DW. Understanding recent trends in childhood obesity in the United States. *Econ Hum Biol.* 2019;34:16–25
17. Biro FM, Greenspan LC, Galvez MP, et al. Onset of breast development in a longitudinal cohort. *Pediatrics.* 2013; 132(6):1019–1027
18. Maione L, Bouvattier C, Kaiser UB. Central precocious puberty: recent advances in understanding the aetiology and in the clinical approach. *Clin Endocrinol (Oxf).* 2021;95(4):542–555
19. Ortega MT, McGrath JA, Carlson L, et al. Longitudinal investigation of pubertal milestones and hormones as a function of body fat in girls. *J Clin Endocrinol Metab.* 2021;106(6):1668–1683
20. Tenedero CB, Oei K, Palmert MR. An approach to the evaluation and management of the obese child with early puberty. *J Endocr Soc.* 2021;6(1):bvab173
21. Bullard RD. Environmental racism and invisible communities. *West VA Law Rev.* 1994;96(4)
22. Kiess W, Haeusler G. Endocrine-disrupting chemicals. *Best Pract Res Clin Endocrinol Metab.* 2021;35(5):101566
23. Lopez-Rodríguez D, Franssen D, Heger S, Parent A-S. Endocrine-disrupting chemicals and their effects on puberty. *Best Pract Res Clin Endocrinol Metab.* 2021;35(5):101579
24. Leonardi A, Cofini M, Rigante D, et al. The effect of bisphenol a on puberty: a critical review of the medical literature. *Int J Environ Res Public Health.* 2017; 14(9):E1044
25. Ruiz D, Becerra M, Jagai JS, Ard K, Sargis RM. Disparities in environmental exposures to endocrine-disrupting chemicals and diabetes risk in vulnerable populations. *Diabetes Care.* 2018;41(1):193–205
26. James-Todd T, Terry MB, Rich-Edwards J, Deierlein A, Senie R. Childhood hair product use and earlier age at menarche in a racially diverse study population: a pilot study. *Ann Epidemiol.* 2011;21(6):461–465
27. McDonald JA, Tehranifar P, Flom JD, Terry MB, James-Todd T. Hair product use, age at menarche and mammographic breast density in multiethnic urban women. *Environ Health.* 2018; 17(1):1
28. McDonald JA, Llanos AAM, Morton T, Zota AR. The environmental injustice of beauty products: toward clean and equitable beauty. *Am J Public Health.* 2021;112(1):50-53
29. Tiwary CM. Premature sexual development in children following the use of estrogen- or placenta-containing hair products. *Clin Pediatr (Phila).* 1998; 37(12):733–739
30. James-Todd T, Senie R, Terry MB. Racial/ethnic differences in hormonally-active hair product use: a plausible risk factor for health disparities. *J Immigr Minor Health.* 2012;14(3):506–511
31. James-Todd T, Connolly L, Preston EV, et al. Hormonal activity in commonly used Black hair care products: evaluating hormone disruption as a plausible contribution to health disparities. *J Expo Sci Environ Epidemiol.* 2021; 31(3):476–486
32. Geronimus AT. The weathering hypothesis and the health of African-American women and infants: evidence and speculations. *Ethn Dis.* 1992;2(3):207–221
33. Warner J. The unequal toll of toxic stress: how the mental burdens of bias, trauma, and family hardship impact girls and women. Available at: <https://americanprogress.org/article/unequal-toll-toxic-stress/>. Accessed November 4, 2021
34. Sun Y, Mensah FK, Azzopardi P, Patton GC, Wake M. Childhood social disadvantage and pubertal timing: a national birth cohort from Australia. *Pediatrics.* 2017;139(6):e20164099
35. Aghaee S, Deardorff J, Greenspan LC, Quesenberry CP Jr, Kushi LH, Kubo A. Early life household intactness and timing of pubertal onset in girls: a prospective cohort study. *BMC Pediatr.* 2020;20(1):464
36. Ruttle PL, Shirtcliff EA, Armstrong JM, Klein MH, Essex MJ. Neuroendocrine coupling across adolescence and the longitudinal influence of early life stress. *Dev Psychobiol.* 2015; 57(6):688–704
37. Amutah C, Greenidge K, Mante A, et al. Misrepresenting race - the role of medical schools in propagating physician bias. *N Engl J Med.* 2021;384(9):872–878
38. Epstein R, Blake JJ, Gonzalez T. Girlhood interrupted: the erasure of black girls' childhood. Available at: <https://www.law.georgetown.edu/poverty-inequality-center/wp-content/uploads/sites/14/2017/08/girlhood-interrupted.pdf>. Accessed April 29, 2022
39. Moore SR, Harden KP, Mendle J. Pubertal timing and adolescent sexual behavior in girls. *Dev Psychol.* 2014;50(6):1734–1745
40. Nanda J. Blind discretion: girls of color & delinquency in the juvenile justice system. Available at: <https://www.uclalawreview.org/blind-discretion-girls-of-color-delinquency-in-the-juvenile-justice-system/>. Accessed April 29, 2022
41. Grandone A, Capristo C, Cirillo G, et al. Molecular screening of MKRN3, DLK1, and KCN9 genes in girls with idiopathic central precocious puberty. *Horm Res Paediatr.* 2017;88(3-4): 194–200
42. Gerdeña JP, Plaisime MV, Tsai J. From race-based to race-conscious medicine: how anti-racist uprisings call us to act. *Lancet.* 2020;396(10257): 1125–1128
43. Skoog T, Özdemir SB. Explaining why early-maturing girls are more exposed to sexual harassment in early adolescence. *J Early Adolescence.* 2015;36(4):490-509