Onset of Breast Development in a Longitudinal Cohort

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KEY WORDS
puberty, girls, breast development, obesity

ABBREVIATIONS
BCERP—Breast Cancer and the Environment Research Program
NHANES—National Health and Nutrition Examination Survey
PROS—Pediatric Research in Office Settings

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WHAT’S KNOWN ON THIS SUBJECT: Several studies have documented earlier onset of pubertal maturation in girls, with several potential factors attributed to the earlier onset.

WHAT THIS STUDY ADDS: This study demonstrates earlier maturation in white non-Hispanic girls, with greater BMI linked as a major factor. The entire distribution of pubertal timing has shifted to a younger age, suggesting redefinition of ages for both early and late maturation.

abstract

BACKGROUND AND OBJECTIVES: There is growing evidence of pubertal maturation occurring at earlier ages, with many studies based on cross-sectional observations. This study examined age at onset of breast development (thelarche), and the impact of BMI and race/ethnicity, in the 3 puberty study sites of the Breast Cancer and the Environment Research Program, a prospective cohort of >1200 girls.

METHODS: Girls, 6 to 8 years at enrollment, were followed longitudinally at regular intervals from 2004 to 2011 in 3 geographic areas: the San Francisco Bay Area, Greater Cincinnati, and New York City. Sexual maturity assessment using Tanner staging was conducted by using standardized observation and palpation methods by trained and certified staff. Kaplan-Meier analyses were used to describe age at onset of breast maturation by covariates.

RESULTS: The age at onset of breast stage 2 varied by race/ethnicity, BMI at baseline, and site. Median age at onset of breast stage 2 was 8.8, 9.3, 9.7, and 9.7 years for African American, Hispanic, white non-Hispanic, and Asian participants, respectively. Girls with greater BMI reached breast stage 2 at younger ages. Age-specific and standardized prevalence of breast maturation was contrasted to observations in 2 large cross-sectional studies conducted 10 to 20 years earlier (Pediatric Research in Office Settings and National Health and Nutrition Examination Survey III) and found to have occurred earlier among white, non-Hispanic, but not African American girls.

CONCLUSIONS: We observed the onset of thelarche at younger ages than previously documented, with important differences associated with race/ethnicity and BMI, confirming and extending patterns seen previously. These findings are consistent with temporal changes in BMI. Pediatrics 2013;132:1019–1027
Over the past several years, multiple studies have reported an earlier age at onset of breast development. In 2007, a consensus panel reported there were sufficient data to suggest a trend toward earlier breast development in the United States over the second half of the 20th century. This trend has been noted internationally as well; for example, girls in the Copenhagen Puberty Study experienced breast development nearly a year earlier than those born 15 to 16 years previously. These findings have been linked temporally to the increase in BMI and prevalence of obesity. The relationship between higher BMI and earlier onset of puberty in girls has been noted previously; in 2 large cross-sectional studies, Pediatric Research in Office Settings (PROS) and the National Health and Nutrition Examination Survey (NHANES) III, earlier maturation occurred in those girls with greater BMI and in those with BMI ≥85th percentile. Here we report on timing of breast development from a longitudinal cohort of girls recruited at ages 6 to 8 years, to examine pubertal timing in association with higher BMI and by race/ethnicity.

**METHODS**

This project was carried out as part of the National Institute of Environmental Health Sciences/National Cancer Institute Breast Cancer and the Environment Research Program (BCERP). The Puberty Study of the BCERP is investigating environmental exposures and onset of puberty in girls, and has been described in detail elsewhere. Girls were enrolled between 2004 and 2008 at 6 through 8 years of age. They were recruited through the 3 puberty study sites of the Breast Cancer and the Environment Research Program: the San Francisco Bay Area in California (through members of Kaiser-Permanente of Northern California), the greater Cincinnati metropolitan area in Ohio and Kentucky (through local schools, and Breast Cancer Registry of Greater Cincinnati), and east and central Harlem in New York City, New York (through community centers, clinics, and local schools). Recruitment was through a combined convenience sample at each site, with the sampling frame defined as those age-eligible girls in Kaiser Permanente membership at defined sites, selected schools in greater Cincinnati, or with a clinic appointment in Harlem. Each site had a time frame for recruitment, and used printed material for recruitment, describing the study as “a project of girls growing up today.” The current report includes longitudinal data in scheduled semiannual (Cincinnati) or annual visits (Mount Sinai and Kaiser Permanente of Northern California) through March 2012, with mean follow-up of 4.3 years. The local institutional review board approved the study at each center; consent was obtained from the parent/legal guardian, and assent was obtained from participants once they reached 10 years of age.

Sexual maturity was established through a standardized method based on Tanner staging. Breast development was assessed through both observation and palpation. Professional staff were trained and certified, and periodic cross-site validation was performed by a master trainer visiting all 3 sites. Onset of breast development was defined as attaining breast stage 2 or greater. We had noted previously that examiners had 87% agreement (ie, same breast stage) with a master trainer in blinded field assessments; the remaining assessments were within 1 stage.

During the examination visits, trained and certified staff members obtained standardized anthropometric measurements, including height and weight, and made 2 measurements of each parameter. If the difference exceeded a preset amount, or the amount was outside the 5th to 95th percentile values, a third measurement was taken and an average of the values was used for analysis. BMI was calculated from the mean values of height and weight measurements, as weight divided by the square of height. BMI percentile and z score were determined by using the 2000 growth charts from the Centers for Disease Control and Prevention (www.cdc.gov/nccdphp/dnpao/growthcharts/resources/sas.htm). We tested for the difference in median values for BMI percentile across sites by using the 2-sample median test, and for the difference proportions in race/ethnicity and BMI percentile groups across sites using the $\chi^2$ and Fisher's exact tests. Survival analyses were performed by using SAS PROC LIFEREG (SAS Institute, Inc, Cary, NC) first to estimate age of breast development and then SAS PROC LIFETEST to determine the effect of race, education of head of household (as a proxy for socioeconomic status), site, and BMI on age of breast development.

The first step in these analyses was to determine an estimated date of breast development (thearche). For these analyses, a “breast development interval” was defined for each participant. For girls who reached Tanner breast stage 2 or greater during follow-up (interval censored), the interval was defined as the period from the last research study visit at breast stage 1 to the first visit where the girls was noted to be at breast stage 2 or greater. For some girls, progression through breast stages was not consistent, such as apparent regression of breast stage 2 to breast stage 1 at a subsequent visit. For these girls, the breast development interval encompassed the period of inconsistency, until a consistent B2 was observed. For girls who were left-censored (breast stage 2 or greater at the first study visit), the beginning of the interval was defined as...
the race-specific 1.0 percentile value of age from the PROS study data (data supplied by M. Herman-Giddens, PhD (personal communication, 2012), and the end of the interval was the date of the initial research study visit when breast stage 2 was observed. For both interval-censored and left-censored girls, an estimated date of breast development was calculated within the interval using age at the beginning and end of the interval as covariates. The estimated date of thelarche was calculated as the date associated with the median probability over the breast development interval. These probabilities were calculated for each girl by using an SAS macro developed by us, which incorporated $\sigma$ ($\sigma$, the scale parameter of the Weibull distribution from LIFEREG) and the $\times B$ matrix for the girl, and the dates at the beginning and end of her thelarche interval.

Kaplan-Meier analyses were used to determine mean and median age at breast development, incorporating interval censored data, with the SAS application LIFETEST. The Wilcoxon test was used to test trend across strata. For hazard function estimations, the validity of the proportionality assumption was assessed by plotting the log-log of the estimated survival function against survival time; these plots were parallel, so the proportionality assumption was valid. Time ratios and hazard ratios, with 95% confidence intervals, were calculated from the accelerated failure time model using a Weibull distribution derived by Carroll. This model accounts for varying amounts of time and varying ages of the girls while under observation. For these analyses, girls who were right-censored (never at breast stage 2 or greater while under observation) contributed to observational time, but no breast development event, to the analyses. We used PROC GLM to determine the proportion of the variance attributable to each of the covariates.

We compared results of age at breast development with cross-sectional data sets from PROS published by Kaplowitz et al and by Herman-Giddens et al. Differences between our data and the mean age of attaining breast stage 2 in the PROS data set were tested with a 2-sample $t$ test, assuming unequal variance. Differences in age-specific prevalence of breast stage 2 or greater were examined by using $\chi^2$ and Fisher’s exact test.

RESULTS

The baseline cohort included 1239 girls (Table 1). The percentage of BCERP girls with BMI $\geq 85$th percentile included 39% of black, 44% of Hispanic, 26% of white non-Hispanic, and 12% of Asian girls. Differences in the distribution of BMI percentile groups by race and site were examined; the only significant difference noted was that at the end of follow-up, the distribution of BMI percentile groups for blacks differed significantly between Greater Cincinnati and the San Francisco Bay Area ($P = .028$), but not New York City and the San Francisco Bay Area ($P = .174$) or New York City and Greater Cincinnati ($P = .150$). The proportion of girls who had attained breast stage 2 during the study varied by age, race/ethnicity, BMI percentile, and site (Table 1 and 2). Mean and median ages of breast stage 2 varied by race/ethnicity, and were 8.8 and 8.8 years for black, 9.2 and 9.3 for Hispanic, 9.6 and 9.7 for non-Hispanic white, and 9.9 and 9.7 for Asian participants (Table 2). Hispanic BCERP participants had maturation at significantly older ages than black participants, and younger than non-Hispanic white BCERP participants. Girls in all BMI categories $>50$th percentile were progressively more likely to have reached breast stage 2 than those $<50$th percentile ($P$ value for trend = .001), adjusting for race/ethnicity and site (Table 2); differences by race/ethnicity remained significant in adjusted models.

We examined the cumulative prevalence of breast stage 2 or greater by age, race/ethnicity, and BMI status ($<85$th percentile vs $\geq 85$th percentile). As noted in Figs 1 and 2, regardless of race (as well as ethnicity, not shown), participants with BMI $\geq 85$th percentile matured earlier than those $<85$th percentile.

Non-Hispanic white BCERP participants matured earlier than white participants in the PROS study (mean age 9.62 vs 9.96 years, $P = .0005$) (Fig 1). However, non-Hispanic white participants with BMI $<85$th percentile did not mature earlier than PROS white participants (mean age 9.84 vs 9.96, $P = .34$). Black BCERP participants matured at similar ages in both studies ($P = .36$), although those of higher BMI were younger (Fig 2).

Non-Hispanic white BCERP participants matured earlier than white PROS participants at every age between 7 and 12 years (Table 3). Of note, BMI accounted for the greatest amount of variance (14.2%) of all covariates included in the model, contrasted to race, which accounted for 4.4% of the variance. The proxy for socioeconomic status, education level of head of household, did not remain in the model.

DISCUSSION

This study examined timing of onset of breast development in an ongoing longitudinal study of girls. We observed the onset of breast development in white girls at younger ages than reported in previous publications, suggesting a continued trend to earlier ages of breast development; black girls continue to experience breast development earlier than white girls.

Higher BMI was the strongest predictor of earlier age at breast stage 2 in our study. Similar findings have been reported that noted the association between BMI and body fat with earlier timing of puberty in girls, although these data support, but do not establish, causality.
Longitudinal studies have demonstrated this relationship, where BMI,
BMI z scores, and adiposity as early as 3 years of age were related to pubertal
outcomes, including onset of breast development, onset of the pubertal
growth spurt, and age of menarche.2,14–19

The obesity epidemic appears to be a
prime driver in the decrease in age at onset of breast development in contem-
porary girls. In our study, white non-
Hispanic BCERP participants with BMI
### TABLE 2 Age at Thelarche and Hazard Ratios by BMI Percentile During Follow-Up, Race/Ethnicity and Site

<table>
<thead>
<tr>
<th>Strata</th>
<th>Unadjusted Age (in Years) at Thelarche From Kaplan-Meier Survival Analysis(^a)</th>
<th>Adjusted Likelihood of Thelarche By End of Follow-up From AFT Model(^b)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>Observed</td>
</tr>
<tr>
<td>All girls BMI percentile group during follow-up(^c)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt;95th</td>
<td>227</td>
<td>195</td>
</tr>
<tr>
<td>85th–94.9th</td>
<td>205</td>
<td>181</td>
</tr>
<tr>
<td>50th–84.9th</td>
<td>394</td>
<td>330</td>
</tr>
<tr>
<td>&lt;50th</td>
<td>411</td>
<td>333</td>
</tr>
<tr>
<td>Race/Ethnicity</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Non-Hispanic white</td>
<td>420</td>
<td>355</td>
</tr>
<tr>
<td>Asian</td>
<td>57</td>
<td>50</td>
</tr>
<tr>
<td>Hispanic</td>
<td>371</td>
<td>310</td>
</tr>
<tr>
<td>Black</td>
<td>391</td>
<td>324</td>
</tr>
<tr>
<td>Site</td>
<td></td>
<td></td>
</tr>
<tr>
<td>New York City</td>
<td>416</td>
<td>330</td>
</tr>
<tr>
<td>Cincinnati</td>
<td>379</td>
<td>321</td>
</tr>
<tr>
<td>San Francisco Bay Area</td>
<td>444</td>
<td>388</td>
</tr>
</tbody>
</table>

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\(^a\) Girls who did not reach thelarche before end of follow-up are censored.

\(^b\) Hazard ratios and 95% confidence intervals calculated as described by Carroll.\(^8\)

\(^c\) For girls who were breast stage 2+ at enrollment, BMI% during follow-up is BMI% at enrollment. For girls in whom breast stage 2+ was first observed during the study, BMI% during follow-up is the BMI% at the last examination with data before thelarche. For girls who had not attained breast stage 2+ by the end of follow-up, BMI% during follow-up is the BMI% at their last examination.

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**FIGURE 1**
Comparing the cumulative prevalence of Breast Stage 2+ for non-Hispanic white participants between the BCERP Puberty Study and PROS.\(^9\)
<85th percentile had similar age at breast development as white girls in PROS (9.96 years).\textsuperscript{3,9} The percentage of girls 6 to 11 years of age who were obese in NHANES III (1988–1994), approximately the same time as PROS, was 9.8% for non-Hispanic white and 17.0% for non-Hispanic black girls. By the time of NHANES 2009–2010 (encompassing the age of girls in our study) it was 14.4% for non-Hispanic white and 24.0% for non-Hispanic black girls.\textsuperscript{20}

Several studies have suggested that earlier onset of breast development may occur independent of activation of the hypothalamic-pituitary-ovarian axis,\textsuperscript{2} perhaps through endocrine-disrupting chemicals,\textsuperscript{1,21,22} which we will examine in future manuscripts. In a previous report, Parent et al\textsuperscript{23} questioned whether the earliest onset had shifted downward, or if the entire distribution shifted downward. Our data suggest that white, non-Hispanic girls throughout the entire distribution of relative timing of puberty (early, on-time, and late-maturing girls) are maturing at younger ages than previously reported.

The impact of earlier maturation in girls has important clinical implications. Clinicians may need to examine additional contemporary studies to decide whether to lower the age for late maturation in girls, and possibly age of precocious puberty. Both the PROS study and our study excluded girls with pathologic conditions known to modify age of pubertal maturation. However, both studies likely included girls with undiagnosed conditions, such as ovarian cysts, which may explain some of the very early ages at breast development and some of the apparent breast development regression that we observed. Previous authors have commented on the impact of timing of pubertal maturation, with several psychosocial and biologic outcomes, perhaps due to a discrepancy between biological and psychological transitions.\textsuperscript{24} Girls with earlier maturation are at risk for lower self-esteem\textsuperscript{25} and higher rates of depression.\textsuperscript{26} They are more likely to be influenced by older peers and more deviant peers,\textsuperscript{27} and initiate intercourse, substance use, and other norm-breaking behaviors at younger ages.\textsuperscript{28–30} Although the greatest impact on these psychosocial outcomes appears during the adolescent years, the impact on adult women who matured early includes greater rates of depression,\textsuperscript{26,29} lower levels of academic achievement,\textsuperscript{30} and greater number of sexual partners.\textsuperscript{26} The longer-term impact of early maturation on

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**FIGURE 2**

Comparing cumulative prevalence of Breast Stage 2+ for non-Hispanic black participants between the BCERP Puberty Study and PROS.\textsuperscript{9}
psychosocial functioning may represent a complex interaction between factors associated with earlier maturation, such as family stress, with developmental outcomes during adolescence, such as depression. Several authors have described that girls maturing earlier are perceived to be aging more rapidly or have an accelerated life course, described as "weathering." It is unclear if these adverse psychosocial outcomes associated with early maturation will be sustained when many girls mature at a younger age.

The study does have several potential limitations. The participants in this study are not nationally representative, so the findings may not be generalizable. The BMI distribution of our cohort is similar to those published recently regarding the NHANES data.9 We did not observe exact age of transition; however, girls in this study with later breast development have lower BMI, as several girls were noted to have breast development at time of intake and we did not observe exact age of transition. Additionally, the accelerated failure time model, which both allows determination of event rates as well as extension in survival time, was used in this study. The determination of age of breast development used a somewhat novel application of a well-established approach, an accelerated failure time model (specifically the Weibull model).

### Table 3: Comparing the Proportion with Breast Stage 2+ by Age Between BCERP Puberty Study Cross-Sectional Equivalent Data and PROS9

<table>
<thead>
<tr>
<th>Age Group</th>
<th>Black</th>
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</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Total No. of Girls</td>
<td>Percent at Breast Stage 2+</td>
<td>Total No. of Girls</td>
<td>Percent at Breast Stage 2+</td>
<td>χ²</td>
<td>P Value</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>6.0–6.9</td>
<td>126</td>
<td>100</td>
<td>8</td>
<td>7.43</td>
<td>.008</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<td></td>
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<tr>
<td>7.0–7.9</td>
<td>136</td>
<td>237</td>
<td>22</td>
<td>2.28</td>
<td>.131</td>
<td></td>
<td></td>
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<td>8.0–8.9</td>
<td>145</td>
<td>294</td>
<td>38</td>
<td>0.02</td>
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<td>9.0–9.9</td>
<td>115</td>
<td>232</td>
<td>62</td>
<td>0.00</td>
<td>.971</td>
<td></td>
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<td></td>
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<tr>
<td>10.0–10.9</td>
<td>136</td>
<td>139</td>
<td>98</td>
<td>0.02</td>
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<tr>
<td>11.0–11.9</td>
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<td>190</td>
<td>96</td>
<td>0.00</td>
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<td></td>
<td></td>
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<td>12.0–12.9</td>
<td>126</td>
<td>133</td>
<td>99</td>
<td>0.00</td>
<td>.999</td>
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</tbody>
</table>

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a Every B visit in the BCERP Puberty Study was treated as if it were a separate girl; missing visits or visits with refused Tanner staging not included.

b Fisher’s Exact Test used where cell frequency is ≤5.

c PROS did not separate Hispanic and Non-Hispanic white subjects; the BCERP Puberty Study did.
direct comparison of our participants, from a longitudinal study, with participants from 2 cross-sectional studies that used different BMI standards from this study, might appear to be of limited relevance; when we excluded our participants with elevated BMI, the BMI distribution is similar to the results reported by Kaplowitz et al.3 and by Rosenfeld et al.4 and the remaining participants had similar age of breast maturation. Unlike the NHANES III and most (61%) of PROS participants, we used breast palpation for all of our participants, but this should have increased the age of breast development, rather than decreased it. Although BCERP participants appear to have a younger median age at breast stage 2 than NHANES III participants, because of the differences in study design methods we were unable to conduct statistical analyses to determine the difference in age between the 2 groups. Multiple studies have cautioned against overclassification of breast tissue in girls with greater BMI, particularly after onset of the obesity epidemic. Our study procedure used palpation as well as inspection, which limits misclassification of fat tissue deposited in the chest area. Additionally, as previously reported, we validated our maturation assessment procedures with dual breast-development assessment, which demonstrated substantial agreement between examiners.3 Study site has a significant effect on the age of breast development, with effects for the Cincinnati (hazard ratio 1.58) and the San Francisco Bay Area (hazard ratio 0.77) sites being statistically different than the New York City site, which enrolled only black and Hispanic girls. The earlier age at maturation in Cincinnati may be attributed, in part, to performing maturation assessments every 6 months rather than once each year, which could account for an earlier age by 3 to 4 months (data not shown). The later age at maturation in the San Francisco Bay Area could represent other factors not included, such as country of origin differences not captured in our categorization of race and ethnicity, exposures, or lifestyle factors. In addition, our models may not have adequately accounted for differences in race/ethnicity and BMI at the 3 sites. Last, younger age of breast development may not lead to earlier age of menarche, this relationship will be examined when the sample achieves later pubertal milestones.

CONCLUSIONS

We noted earlier onset of breast stage 2 in non-Hispanic white girls, contrasted with 2 previous studies (PROS and NHANES III); this is likely due to greater obesity in the white non-Hispanic girls in our study compared with earlier reports. Girls with BMI >85th percentile matured earlier than those with lower BMI, and BMI explained much of the difference between studies. Black girls experienced breast development at a similar age to blacks from the 2 previous studies, and continued to mature at ages younger than whites.

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