Maternal and Infant Anthropometric Characteristics and Breast Cancer Incidence in the Daughter

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The intrauterine and early life environments have been linked to the etiology of breast cancer in prior studies. We prospectively examined whether maternal and newborn anthropometric factors as reported by the mother are related to an increased incidence of adult breast cancer in the daughter. We used data from 35,133 mother-daughter dyads of the Nurses’ Health Study (NHS) II and the Nurses’ Mothers’ Cohort Study. In 2001, living mothers of NHS II participants who were free of cancer completed a questionnaire on their pregnancy with the nurse and their nurse daughter’s early life experience. During 403,786 years of follow-up, 865 daughters developed incident cases of invasive breast cancer. Nurses with a birthweight of ≥4000 g had a 32% greater risk for breast cancer (multivariable-adjusted hazard ratio (HR) = 1.32, 95% confidence interval (CI) = 1.02–1.71, p-trend = 0.09) compared with those with birthweights of 3000–3499 g. Higher birth length tended to increase risk of premenopausal breast cancer (p for trend = 0.05). We further noted a modest U-shaped relation between maternal weight gain during pregnancy and premenopausal breast cancer incidence in the daughter. Fetal growth may contribute to shaping later life risk for breast cancer, especially prior to menopause.

Breast cancer is the most common malignancy among women worldwide, with an estimated 1.67 million new cancer cases diagnosed in 2012¹. Known and suspected risk factors for breast cancer operating during adult life leave substantial variation in rates of this tumor unexplained. Over the past couple of decades perinatal and early life characteristics have emerged as novel breast cancer risk factors with consistency across different study populations but with some international divergence.

In prior epidemiologic studies, high birthweight has been associated with greater risk of breast cancer²–⁶, although this is not true for all studies⁷. Intrauterine exposure to sex steroid hormones, growth hormone, insulin, insulin-like growth factors (IGF)-1, and IGF-2, and epigenetic variation are potential key pathways linking anthropometric variables in early life to adult breast cancer risk⁸. Because investigations from retrospective case-control studies relying on information collected from the cases and controls themselves leave room for differential misclassification, data from prospective longitudinal analyses are warranted to overcome this bias. In previous literature, fewer data have been available on maternal factors such as pre-pregnancy body mass index (BMI), and weight gain during pregnancy in relation to the risk of breast cancer in the daughter. However, with the global epidemic of obesity maternal body mass index prior to and during pregnancy has substantially increased during the past couple of decades, with 39% of women with normal, 59% with overweight, and 56% with obese prepregnancy BMI exceeding the current U.S. Institute of Medicine (IOM) recommendations for gestational weight gain⁹. The implications of these rapidly changing intrauterine conditions for the daughters’ future breast cancer risk need to be investigated.

We therefore used data from the prospective Nurses’ Mothers’ Cohort Study to explore the relation of anthropometric variables in newborns including birthweight and birth length to their risk of developing breast cancer in adulthood; we also examined whether maternal pre-pregnancy BMI, height, and weight gain during pregnancy

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as reported by the mothers were associated with the incidence of breast cancer among daughters participating in the Nurses’ Health Study (NHS) II.

Methods

Study population. The NHS II cohort was established in 1989 when 116,680 nurses aged between 25 and 42 years from 14 U.S. states completed a mailed questionnaire on lifestyle factors and medical history. Follow-up questionnaires were mailed to participating nurses every two years updating information on lifestyle factors and health. In 2001, 35,830 living mothers of NHSII who were cancer-free participants completed a questionnaire on their pregnancy with their nurse daughter and on her early life exposures forming the Nurses’ Mothers’ Cohort Study. NHS II participants who were adopted or whose adoption status was unknown and those with missing information on the exposures of interest were excluded from the respective analyses.

Exposure assessment. On the Mothers’ questionnaire, the nurses’ mothers were asked to report their height and weight before pregnancy in open-ended questions; this information was used to calculate pre-pregnancy BMI as kg/m². Pre-pregnancy BMI was categorized following standard World Health Organization definitions of underweight (<18.5 kg/m²), normal weight (18.5–24.9 kg/m²), overweight (25–29.9 kg/m²), and obese (≥30 kg/m²). Mothers were also asked to report their gestational weight gain in pre-specified categories (<10, 10–14, 15–19, 20–29, 30–40, >40 lbs). Mothers further reported birthweight and birth length of their daughters as open-ended values. Birthweights recalled by the nurses’ mothers were highly concordant with birthweight information obtained from birth certificates (r = 0.85). Birthweight was coded using standard categories (<2500, 2500–2999, 3000–3499, 3500–3999, ≥4000 grams). Birth length was categorized in equally spaced intervals capturing the narrow distribution (<47, 47–49, 50–52, 53–55, >55 cm). Ponderal index was calculated from birth weight and length [Ponderal index = birthweight (grams) x 100/(birth length, cm)³] and categorized in quintiles.

Covariate information. Covariate information on tobacco use during pregnancy and gestational age was obtained from the Mothers’ questionnaire. Additional information on daughter’s characteristics, including race/ethnicity, family history of breast cancer, month and year of birth of the nurse, menopausal status, adult caloric intake, alcohol consumption, smoking, and BMI was available from the Mothers’ questionnaire. Additional information on daughter’s characteristics, including race/ethnicity, family history of breast cancer, month and year of birth of the nurse, menopausal status, adult caloric intake, alcohol consumption, smoking, and BMI was available from the Mothers’ questionnaire.

Ascertained breast cancer. On follow-up questionnaires administered every two years, NHS participants were asked whether they had been diagnosed with breast cancer in the previous two years. To confirm the diagnosis, participants who reported a breast cancer diagnosis were asked for permission to review their relevant medical records and pathology reports. Due to the high accuracy of self-reported breast cancer diagnosis (>98%), nurse participants who reported a diagnosis of breast cancer but for whom medical records could not be obtained were classified as breast cancer cases.

Ascertained deaths. Deaths were identified by reports from the next of kin, postal authorities, or the National Death Index.

Statistical analysis. Cox proportional hazards regression models were used to estimate the hazard ratios (HRs) and 95% confidence intervals (CIs) of the incidence of breast cancer in the daughters across categories of
Table 2. Age-standardized characteristics of NHS II participants and their mothers participating in the Nurses’ Mothers’ Cohort Study according to the mothers’ weight gain during pregnancy. BMI = body mass index; values are means (SD) or percentages and are standardized to the age distribution of the study population (except age).

| Weight gain during pregnancy (lbs) | All Women | Age-adjusted | Multivariable-adjusted model 1 | Multivariable– + adult variables adjusted model 2 |
|-----------------------------------|-----------|--------------|-------------------------------|-----------------------------------------------|
| <10                               | 53/25444  | 0.97 (0.72–1.29) | 0.92 (0.67–1.25) | 0.92 (0.67–1.25) |
| 10–14                             | 155/77004 | 0.92 (0.76–1.12) | 0.91 (0.75–1.10) | 0.91 (0.75–1.10) |
| 15–19                             | 343/16295 | 1.00          | 1.00              | 1.00              |
| 20–29                             | 208/100081| 0.99 (0.83–1.17) | 0.98 (0.83–1.17) | 0.98 (0.83–1.17) |
| 30–40                             | 72/26626  | 1.30 (1.01–1.68) | 1.32 (1.01–1.71) | 1.32 (1.02–1.71) |
| ≥40                               | 25/10407  | 0.13          | 0.10              | 0.09              |

Table 3. HRs (and 95% CIs) for invasive breast cancer incidence between 2001 and 2013 by categories of anthropometric characteristics at birth among NHS II participants whose mothers participated in the Nurses’ Mothers’ Cohort Study. Ponderal index = birthweight (g) x 100/(birth length, cm)^3 HR = hazard ratio, CI = confidence interval; Age-adjusted models adjusted for age of nurse (continuous). Multivariable-adjusted model 1 includes age of nurse (continuous), race (White, Non-White), family history of breast cancer (yes, no), pre-pregnancy body mass index (<18.5 kg/m^2, 18.5–24.9 kg/m^2, 25.0–30.0 kg/m^2, >30.0 kg/m^2), weight gain during pregnancy (<10 lbs, 10–14 lbs, 15–19 lbs, 20–29 lbs, 30–40 lbs, ≥40 lbs), smoking during pregnancy (no smoking, quit smoking during pregnancy, light smoker (>15 cigarettes/day), heavy smoker (>15 cigarettes/day)), and gestational age (<38 weeks, 38–42 weeks, >42 weeks). Multivariable-adjusted model 2: same as model 1 but with additional adjustment for adult caloric intake (quintiles), adult alcohol intake (quintiles), adult smoking (never, past, current <15 cigarettes/d, current ≥15 cigarettes/d), and adult body mass index (<20.9 kg/m^2, 21–22.9 kg/m^2, 23–24.9 kg/m^2, 25–29.9 kg/m^2, 30–34.9 kg/m^2, ≥35 kg/m^2).
Table 4. HRs (and 95% CIs) for pre- and postmenopausal breast cancer incidence between 2001 and 2013 by categories of anthropometric characteristics at birth among NHS II participants whose mothers participated in the Nurses’ Mothers’ Cohort Study. Ponderal index \( = \) birthweight (g) \( \times 100 \) / (birth length, cm\(^3\)). HR = hazard ratio, CI = confidence interval. Age-adjusted models adjusted for age of nurse (continuous). Multivariable-adjusted model 1 includes age of nurse (continuous), race (White, Non-White), family history of breast cancer (yes, no), pre-pregnancy body mass index (<18.5 kg/m\(^2\), 18.5–24.9 kg/m\(^2\), 25.0–30.0 kg/m\(^2\), >30 kg/m\(^2\)), weight gain during pregnancy (<10 lbs, 10–14 lbs, 15–19 lbs, 20–29 lbs, 30–40 lbs, ≥40 lbs), smoking during pregnancy (no smoking, quit smoking during pregnancy, light smoker (<15 cigarettes/day), heavy smoker (≥15 cigarettes/day), and gestational age (<38 weeks, 38–42 weeks, >42 weeks). Multivariable-adjusted model 2: same as model 1 but with additional adjustment for adult caloric intake (quintiles), adult alcohol intake (quintiles), adult smoking (never, past, current <15 cigarettes/d, current ≥15 cigarettes/d), and adult body mass index (<20.9 kg/m\(^2\), 21–22.9 kg/m\(^2\), 23–24.9 kg/m\(^2\), 25–29.9 kg/m\(^2\), 30–34.9 kg/m\(^2\), ≥35 kg/m\(^2\)).
In this study, a high birthweight was associated with a greater incidence of breast cancer later in life. Moreover, a positive trend with increasing birth length was noted for premenopausal breast cancer risk (Tables 5 and 6). Daughters whose mothers had a pre-pregnancy BMI of ≥25 kg/m² experienced a multivariable-adjusted HR of breast cancer of 0.98 (95% CI: 0.74–1.30) compared to those whose mothers were normal-weight (BMI 18.5–24.9 kg/m²; Table 5). We noted a modest U–shaped relation of maternal weight gain during pregnancy with the risk of premenopausal breast cancer in the daughters, but not for postmenopausal breast cancer (Table 6). For premenopausal breast cancer a positive trend with increasing birth length was noted for premenopausal breast cancer. We further observed a statistical trend of borderline significance for a greater risk of premenopausal breast cancer in daughters comparing low (<15 cigarettes/day) and heavy smoker (≥15 cigarettes/day), and models for pre-pregnancy BMI and weight gain during pregnancy are mutually adjusted for weight gain during pregnancy (<10 lbs, 10–14 lbs, 15–19 lbs, 20–29 lbs, 30–40 lbs, >40 lbs) and pre-pregnancy BMI (>18.5 kg/m², 18.5–24.9 kg/m², 25.0–30.0 kg/m², 30.1–34.9 kg/m², >30 kg/m²). Multivariable-adjusted model 2: same as model 1 but with additional adjustment for adult caloric intake (quintiles), adult alcohol intake (quintiles), adult smoking (never, past, current <15 cigarettes/d, current ≥15 cigarettes/d), and adult BMI (<20.9 kg/m², 21–22.9 kg/m², 23–24.9 kg/m², 25–29.9 kg/m², 30–34.9 kg/m², >35 kg/m²).

Table 5. HRs (and 95% CIs) for invasive breast cancer incidence between 2001 and 2013 by categories of maternal height, pre-pregnancy BMI, and weight gain during pregnancy among NHIS II participants whose mothers participated in the Nurses’ Mothers’ Cohort Study. HR = hazard ratio, CI = confidence interval, BMI = body mass index. Age-adjusted models adjusted for age of nurse (continuous). Multivariable-adjusted model 1 includes age of nurse (continuous), race (White, Non-White), family history of breast cancer (yes, no), smoking during pregnancy (no smoking, quit smoking during pregnancy, light smoker (<15 cigarettes/day), and heavy smoker (≥15 cigarettes/day), and models for pre-pregnancy BMI and weight gain during pregnancy are mutually adjusted for weight gain during pregnancy (<10 lbs, 10–14 lbs, 15–19 lbs, 20–29 lbs, 30–40 lbs, >40 lbs) and pre-pregnancy BMI (>18.5 kg/m², 18.5–24.9 kg/m², 25.0–30.0 kg/m², >30 kg/m²). Multivariable-adjusted model 2: same as model 1 but with additional adjustment for adult caloric intake (quintiles), adult alcohol intake (quintiles), adult smoking (never, past, current <15 cigarettes/d, current ≥15 cigarettes/d), and adult BMI (<20.9 kg/m², 21–22.9 kg/m², 23–24.9 kg/m², 25–29.9 kg/m², 30–34.9 kg/m², >35 kg/m²).

| Maternal height | N cases/person-yrs | Age-adjusted | Multivariable-adjusted model 1 | Multivariable + adult variables adjusted model 2 |
|-----------------|--------------------|--------------|-------------------------------|-----------------------------------------------|
| <155 cm         | 15/40560           | 0.91 (0.72–1.16) | 0.92 (0.72–1.16) | 0.91 (0.72–1.16) |
| 155–164.9 cm    | 368/176952         | 0.92 (0.80–1.06) | 0.92 (0.80–1.06) | 0.92 (0.79–1.06) |
| 165–174.9 cm    | 380/170167         | 1.00          | 1.00                          | 1.00                                          |
| ≥175 cm         | 32/16109           | 0.92 (0.64–1.32) | 0.92 (0.64–1.33) | 0.92 (0.64–1.32) |

**Pre-pregnancy BMI**

| Pre-pregnancy BMI | N cases/person-yrs | Age-adjusted | Multivariable-adjusted model 1 | Multivariable + adult variables adjusted model 2 |
|------------------|--------------------|--------------|-------------------------------|-----------------------------------------------|
| <18.5 kg/m²      | 95/40651           | 1.08 (0.87–1.34) | 1.07 (0.86–1.32) | 1.06 (0.85–1.32) |
| 18.5–24.9 kg/m²  | 655/30860          | 1.00          | 1.00                          | 1.00                                          |
| ≥25 kg/m²        | 55/27229           | 0.98 (0.74–1.30) | 0.98 (0.74–1.30) | 1.02 (0.77–1.34) |

**Maternal weight gain during pregnancy**

| Weight gain | N cases/person-yrs | Age-adjusted | Multivariable-adjusted model 1 | Multivariable + adult variables adjusted model 2 |
|-------------|--------------------|--------------|-------------------------------|-----------------------------------------------|
| <15 lbs     | 118/54954          | 1.03 (0.83–1.27) | 1.02 (0.82–1.26) | 1.02 (0.83–1.27) |
| 15–19 lbs   | 154/78127          | 0.96 (0.79–1.16) | 0.96 (0.79–1.16) | 0.97 (0.80–1.17) |
| 20–29 lbs   | 320/156964         | 1.00          | 1.00                          | 1.00                                          |
| 30–40 lbs   | 152/64455          | 1.16 (0.95–1.41) | 1.16 (0.95–1.41) | 1.16 (0.96–1.41) |
| ≥40 lbs     | 47/20204           | 1.12 (0.82–1.53) | 1.11 (0.82–1.52) | 1.14 (0.83–1.55) |

**Discussion**

In this study, a high birthweight was associated with a greater incidence of breast cancer later in life. Moreover, a positive trend with increasing birth length was noted for premenopausal breast cancer. We further observed a U–shaped relation between maternal weight gain during pregnancy and premenopausal breast cancer in daughters.

A positive association between high birthweight studies (from measurements made at birth by medical doctors, birth records/medical registers, maternal interviews, or adult self-report) and an increased risk of breast cancer in adulthood has been observed in most prior studies and confirms our earlier findings from the same
Table 6. HRs (and 95% CIs) for pre- and postmenopausal breast cancer incidence between 2001 and 2013 by categories of maternal height, pre-pregnancy BMI, and weight gain during pregnancy among NHS II participants whose mothers participated in the Nurses’ Mothers’ Cohort Study. HR = hazard ratio, CI = confidence interval, BMI = body mass index. Age-adjusted models adjusted for age of nurse (continuous). Multivariable-adjusted model 1 includes age of nurse (continuous), race (White, Non-White), family history of breast cancer (yes, no), smoking during pregnancy (no smoking, quit smoking during pregnancy, light smoker (<15 cigarettes/day), and heavy smoker (≥15 cigarettes/day), and models for pre-pregnancy BMI and weight gain during pregnancy are mutually adjusted for weight gain during pregnancy (<10 lbs, 10–14 lbs, 15–19 lbs, 20–29 lbs, 30–40 lbs, ≥40 lbs) and pre-pregnancy BMI (<18.5 kg/m², 18.5–24.9 kg/m², 25.0–30.0 kg/m², ≥30 kg/m²). Multivariable-adjusted model 2: same as model 1 but with additional adjustment for adult caloric intake (quintiles), adult alcohol intake (quintiles), adult smoking (never, past, current <15 cigarettes/day, current ≥15 cigarettes/day), and adult BMI (<20.9 kg/m², 21–22.9 kg/m², 23–24.9 kg/m², 25–29.9 kg/m², 30–34.9 kg/m², ≥35 kg/m²).

| Maternal height | Premenopausal Women | Postmenopausal Women |
|----------------|----------------------|-----------------------|
|                 | N cases/person-yrs  | Age-adjusted          | Multivariable-adjusted model 1 | Multivariable + adult variables adjusted model 2 | N cases/person-yrs | Age-adjusted | Multivariable-adjusted model 1 | Multivariable + adult variables adjusted model 2 |
| <155 cm         | 33/17643             | 0.91 (0.62–1.34)      | 0.92 (0.63–1.35) | 0.90 (0.61–1.32) | 40/18739 | 0.87 (0.62–1.23) | 0.87 (0.62–1.24) | 0.87 (0.61–1.23) |
| 155–164.9 cm    | 142/80230            | 0.90 (0.71–1.13)      | 0.91 (0.72–1.14) | 0.90 (0.71–1.13) | 189/79574 | 0.96 (0.78–1.19) | 0.97 (0.78–1.19) | 0.96 (0.78–1.18) |
| 165–174.9 cm    | 156/79486            | 1.00                   | 1.00                   | 1.00                   | 178/73777 | 1.00                   | 1.00                   | 1.00                   |
| ≥175 cm         | 14/7883              | 0.86 (0.49–1.53)      | 0.86 (0.49–1.52) | 0.87 (0.49–1.53) | 14/6628 | 0.92 (0.53–1.58) | 0.90 (0.52–1.56) | 0.91 (0.52–1.57) |
|                 |                      | P trend                | 0.53                   | 0.56                   | 0.47                   | 0.53                   | 0.55                   | 0.52                   |

Pre-pregnancy BMI

| Weight gain <15 lbs | 51/24064              | 1.23 (0.88–1.70)      | 1.23 (0.88–1.71) | 1.24 (0.89–1.72) | 50/25453 | 0.80 (0.58–1.10) | 0.78 (0.57–1.08) | 0.78 (0.57–1.08) |
| Weight gain 15–19 lbs | 56/35940             | 0.94 (0.69–1.29)      | 0.94 (0.69–1.29) | 0.95 (0.69–1.30) | 77/34477 | 0.92 (0.70–1.21) | 0.92 (0.70–1.21) | 0.93 (0.70–1.22) |
| Weight gain 20–29 lbs | 132/75463            | 1.00                   | 1.00                   | 1.00                   | 159/66229 | 1.00                   | 1.00                   | 1.00                   |
| Weight gain 30–40 lbs | 66/29900             | 1.25 (0.92–1.69)      | 1.26 (0.93–1.70) | 1.28 (0.94–1.73) | 70/27878 | 1.05 (0.79–1.40) | 1.04 (0.78–1.39) | 1.04 (0.78–1.38) |
| Weight gain ≥40 lbs | 18/8872              | 1.21 (0.73–2.01)      | 1.17 (0.70–1.95) | 1.17 (0.70–1.96) | 24/9251 | 1.05 (0.68–1.62) | 1.04 (0.67–1.61) | 1.03 (0.67–1.60) |
| P trend            | 0.46                 | 0.49                   | 0.47                   | 0.13                   | 0.12      | 0.14                   |

The mechanisms underlying the association between a high birthweight and the diagnosis of breast cancer risk in later life is not entirely clear, but may be orchestrated by intrauterine exposures to growth hormones and epigenetic programming. Birthweight is likely a marker for intrauterine levels of sex estrogen and progesterone, growth hormone, IGF-1, IGF-2, and insulin itself which may increase the number of susceptible stem cells in the mammary gland or enhance cell proliferation, thereby contributing to tumor development through accumulation of DNA mutations. Since the epigenetic profile is established in utero, intrauterine stressors may affect the epigenetic pattern and thereby susceptibility to cancer. The exposure to high levels of hormones and growth factors in utero may change the response of breast tissue to hormonal inputs later.

Our study revealed a statistically significant dose-response relation between birth length and premenopausal breast cancer. A previous meta-analysis suggested a positive association between birth length and later breast cancer in studies based on birth records, but not in studies based on parental recall or self-reports, likely reflecting the difficulty in precisely capturing birth length. Unlike in our study, the birth length – breast cancer association was not modified by menopausal status in that meta-analysis. The mechanisms underlying any association between maternal pre-pregnancy BMI and breast cancer are likely similar to those connecting birthweight and breast cancer.
did not observe a link between maternal pre-pregnancy BMI and breast cancer risk in the daughters. However, both our study and that by Sanderson and colleagues were based on pregnancies several decades in the past, when in particular pre-pregnancy BMI, but also gestational weight gain recommendations were considerably lower than currently. Hence, these associations need to be further evaluated in more recent studies including a larger proportion of mothers with high pre-pregnancy BMI and maternal weight gain, reflecting current distributions.

Strengths of our study include its prospective design, the large number of study participants, and adjustment for a number of important potential perinatal confounding variables. Moreover, our study was based on a two-generation design providing the unique opportunity to use data collected directly from the mothers and combine them with data provided by the nurses.

A limitation of our study is that gestational and newborn characteristics had to be recalled by the mothers likely introducing random misclassification. However, the validity of birthweight reported by the mothers appears to be high when compared with birthweight recorded on state birth records (r = 0.85)12. Similarly, other studies have reported good agreement between pregnancy-related factors recorded during pregnancy and maternal recall18–21. The nurses’ adult life variables did not appreciably affect associations observed. However, we cannot rule out unmeasured confounding or residual confounding through covariates measured with error. Finally, with mothers being born between 1921 and 1964, we had limited power to examine substantial pregnancy weight gains and high values of pre-pregnancy BMI which have become more prevalent in recent years.

In summary, a high birthweight and possibly a high birth length may predict an increased risk of breast cancer in later life, particularly of premenopausal breast cancer. Whether maternal factors during pregnancy and other infant factors are related to adult breast cancer risk requires further clarification in other prospective studies with good exposure validity.

Data availability
All relevant data are provided within the manuscript or are available from the corresponding author upon request.

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**Author contributions**

D.S. and K.M. designed the study; K.M. and W.W. collected the data; D.S. and M.D. analyzed the data; D.S. wrote the initial draft; all authors reviewed and revised the draft.

**Competing interests**

The authors declare no competing interests.

**Additional information**

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