Preterm birth, birth weight, and subsequent risk of female breast cancer

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We have previously found an increased risk of breast cancer among women born preterm. To confirm or refute the results, an enlarged study was conducted. The results from this study do not confirm the initial findings and suggest that preterm birth can be ruled out as a risk factor for breast cancer.

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Oestrogens are fundamental hormones in breast cancer aetiology, and it has been suggested that the surge in oestrogens during pregnancy may influence subsequent breast cancer risk in the offspring (Trichopoulos, 1990). Infants born extremely preterm, that is before 28 weeks of gestation, has been found to have high levels of oestrogens postnatally (Sedin et al., 1985). In a study of women born between 1925 and 1932, we found that birth before 32 weeks of gestation constituted a strong risk factor for adult breast cancer (Ekbom et al., 2000). The study, however, was hampered by low statistical power. We therefore conducted an enlarged study of the association between preterm birth and female breast cancer.

MATERIALS AND METHODS

We identified the study subjects by examining manually all birth records for the period 1925–1949 from Uppsala University Hospital, Sundsvalls County Hospital, and two major delivery units in Stockholm (Allmänna BB and Södra BB). Approximately 250 000 birth records were examined. Female subjects born before 35 completed weeks of gestation according to date of last menstrual period, and females born after 35 weeks with a birth weight of less than 2000 g were included. At the point of initial inclusion into the cohort, data on pregnancy and delivery, such as date of last menstrual period and birth weight, was recorded.

By use of the National Registration Number, the cohort was followed up in the Swedish Register of Population and Population Changes from 1 January 1958 to 31 December 1998 or to death or emigration prior to this time. Subjects who deceased before 1958 were excluded, since incident cancers could not be identified with any completeness prior to the initiation of the Swedish Cancer Register. We identified cases of cancer by linking our subjects to the Swedish Cancer Register. The completeness of this register is over 98% (Mattsson and Wallgren, 1984).

We used last menstrual period to estimate gestational duration. We categorised gestational duration into three groups: 32 completed weeks or less, 33–34 weeks, and 35 weeks or more. Birth weight was categorised into less than 2000 g, 2000–2999 g, and 3000 g or more. Since an unreasonably high birth weight for a short duration of gestation indicated an underestimation of gestational duration or an overestimation of birth weight, we excluded subjects born before 32 weeks of gestation with a birth weight of 3000 g or more (n = 155), and subjects with a gestational duration of 33–34 weeks with a birth weight of 3600 g or more (n = 94).

We calculated standardised incidence ratios (SIR) for all female cancer, female breast cancer, and early onset female breast cancer using age, sex, and time-specific cancer incidence rates for the Swedish population during the period 1958–1998 provided by the Cancer Registry. We analysed all cancer occurrences as well as the first cancer occurrence. As these estimates were essentially the same, we only present the results from the analysis of first cancer occurrence.

RESULTS

There were 1483 women who met the eligibility criteria and of these, 1208 were born before 35 weeks of gestation and 275 were born after week 35 with a birth weight of less than 2000 g (Table 1).

The overall risk of cancer among the women was not increased (Table 2). The risk of breast cancer was neither associated with preterm birth nor with low birth weight, but a birth weight of more than 3000 g was associated with an increased risk of breast cancer (Table 2). This increase in breast cancer risk was not dependent on age at onset.
Table 1: Study subjects by category, gestational duration, and birth weight

| Study subjects                        | No.  | Percentage (%) |
|---------------------------------------|------|----------------|
| All                                   | 1483 | 100            |
| Small-for-gestational-age<sup>a</sup> | 275  | 18.5           |
| Preterm                               |      |                |
| All                                   | 1208 | 81.5           |
| ≤ 32 weeks                            | 514  | 34.7           |
| 33–34 weeks                           | 694  | 46.8           |
| Birth weight                          |      |                |
| 0.5–1.99 kg                           | 703  | 47.4           |
| 2–2.99 kg                             | 691  | 46.6           |
| ≥ 3 kg                                | 89   | 6.0            |

| Gestational duration by birth weight  |      |                |
| ≤ 32 weeks                           |      |                |
| 0.5–1.99 kg                           | 292  | 19.7           |
| 2–2.99 kg                             | 222  | 15.0           |
| 33–34 weeks                           |      |                |
| 0.5–1.99 kg                           | 136  | 9.2            |
| 2–2.99 kg                             | 469  | 31.6           |
| ≥ 3 kg                                | 89   | 6.0            |

<sup>a</sup>Birth weight < 2000 g and gestational duration > 35 weeks.

DISCUSSION

This study is, to our knowledge, the largest cohort study hitherto conducted on long-term cancer risk among women born preterm or with low birth weight. Neither preterm birth nor low birth weight were associated with an increased risk for cancer overall or for breast cancer.

We have previously reported an increased risk for breast cancer among women born before 32 weeks of gestation (Ekbom et al., 1997; Ekbom et al., 2000). Other studies of the association between gestational duration and breast cancer have been mostly negative (Le Marchand et al., 1988; Michels et al., 1996; Sanderson et al., 1996; Sanderson et al., 1998), but these studies have categorised preterm birth either as a dichotomous variable (Le Marchand et al., 1988; Sanderson et al., 1996; Sanderson et al., 1998), or as preterm birth of more or less than 4 weeks (Michels et al., 1996). Since the majority of preterm infants are born after 32 weeks of gestation, the number of infants born before 32 weeks has been low. In a recent case–control study, however, a reduced risk for breast cancer was found among women born before 32 weeks (Innes et al., 2000), and as suggested by the present data, our previous results were either due to chance, or confined to a subset of women with specific characteristics.

Consistent with other studies (Potischman and Troisi, 1999; Innes et al., 2000; Stavola et al., 2000), we found a positive association between birth weight and breast cancer risk, and this association was independent of age at diagnosis. Some studies have reported the association between birth weight and breast cancer to be J-shaped, with an increased risk also among women with a very low birth weight (Potischman and Troisi, 1999; Innes et al., 2000). According to the present data, low birth weight is not a major risk factor for breast cancer.

When Trichopoulos (1990) put forward his hypothesis of early environmental exposures, the proposed mechanism was oestrogen exposure. Since then, a vast number of studies have supported his hypothesis, although hormones other than oestrogen may be instrumental (Potischman and Troisi, 1999). Our primary study of preterm women was originally initiated because of reports of an ovarian hyper stimulation syndrome among female infants born before 28 weeks of gestation, which lead to high levels of oestrogens perinatally (Tapanainen et al., 1981; Sedin et al., 1985). The increase in breast cancer risk was interpreted as indicating an important role of early oestrogen exposure (Ekbom et al., 2000). We conclude from the present study that there is no substantial increase in breast cancer risk among women born preterm. We cannot, however, draw conclusions about the role of hormones or ovarian hyper stimulation syndrome on basis of our data.

The use of self-reported date of last menstrual period leads to some misclassification in the calculation of gestational duration. The nature of this misclassification is both nondifferential and differential.

Table 2: Standardised incidence ratios (SIR) and 95% confidence intervals (95% CI) for all female cancer, female breast cancer and female early onset breast cancer<sup>a</sup> by category, gestational duration, and birth weight

|                    | Obs | Exp | SIR  | 95% CI         | Obs | Exp | SIR  | 95% CI         | Obs | Exp | SIR  | 95% CI         |
|--------------------|-----|-----|------|----------------|-----|-----|------|----------------|-----|-----|------|----------------|
| **Small-for-gestational-age<sup>b</sup>** |     |     |      |                |     |     |      |                |     |     |      |                |
| All                | 20  | 26.7| 0.75 | 0.46–1.16      | 10  | 9.2 | 1.09 | 0.52–2.01      | 4   | 3.8 | 1.06 | 0.29–2.71      |
| Preterm            |     |     |      |                |     |     |      |                |     |     |      |                |
| All                | 127 | 112.2| 1.13 | 0.94–1.35      | 39  | 39.4| 0.99 | 0.70–1.35      | 19  | 16.8| 1.13 | 0.68–1.77      |
| ≤ 32 weeks         | 55  | 47.8| 1.15 | 0.87–1.50      | 18  | 16.7| 1.08 | 0.64–1.70      | 10  | 7.1 | 1.40 | 0.67–2.57      |
| 33–34 weeks        | 72  | 64.4| 1.12 | 0.88–1.41      | 21  | 22.7| 0.92 | 0.57–1.41      | 9   | 9.7 | 0.93 | 0.43–1.77      |
| Birth weight<sup>c</sup> |     |     |      |                |     |     |      |                |     |     |      |                |
| 0.5–1.99 kg        | 45  | 39.9| 1.13 | 0.82–1.51      | 16  | 14.0| 1.14 | 0.65–1.85      | 9   | 5.9 | 1.53 | 0.70–2.91      |
| 2–2.99 kg          | 63  | 64.7| 0.97 | 0.75–1.25      | 16  | 22.7| 0.71 | 0.40–1.15      | 7   | 7.9 | 0.72 | 0.29–1.49      |
| ≥ 3 kg             | 19  | 7.6 | 2.51 | 1.51–3.91      | 7   | 2.7 | 2.55 | 1.03–5.25      | 3   | 1.2 | 2.46 | 0.51–7.19      |
| Gestational duration by birth weight |     |     |      |                |     |     |      |                |     |     |      |                |
| ≤ 32 weeks         |     |     |      |                |     |     |      |                |     |     |      |                |
| 0.5–1.99 kg        | 32  | 26.9| 1.19 | 0.81–1.68      | 13  | 9.4 | 1.38 | 0.73–2.36      | 8   | 4.0 | 1.99 | 0.86–3.92      |
| 2–2.99 kg          | 23  | 20.9| 1.10 | 0.70–1.65      | 5   | 7.3 | 0.69 | 0.22–1.60      | 2   | 3.1 | 0.64 | 0.08–2.31      |
| 33–34 weeks        | 13  | 13.1| 1.00 | 0.53–1.70      | 3   | 4.6 | 0.65 | 0.13–1.91      | 1   | 1.8 | 0.54 | 0.01–3.01      |
| 0.5–1.99 kg        | 40  | 43.7| 0.91 | 0.65–1.25      | 11  | 15.5| 0.72 | 0.36–1.28      | 5   | 6.6 | 0.76 | 0.25–1.77      |
| ≥ 3 kg             | 19  | 7.6 | 2.51 | 1.51–3.91      | 7   | 2.7 | 2.55 | 1.03–5.25      | 3   | 1.2 | 2.46 | 0.51–7.19      |

<sup>a</sup>Onset before 50 years of age. <sup>b</sup>Birth weight < 2000 g and gestational duration > 35 weeks. <sup>c</sup>Among women born before 35 weeks of gestation.

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systematic, the latter with a tendency to overestimate gestational duration (Henriksen et al., 1995; Savitz et al., 2002). By exclusion of subjects whose birth weights indicated an underestimation of gestational duration, however, misclassification was reduced, and a systematic overestimation of gestational duration can only lead to a more conservative inclusion of subjects in the cohort. The main limitation of the study is that study subjects were born between 1925 and 1949, but the follow-up did not start until 1958. The cohort will therefore consist of women born more than 50 years ago who survived beyond childhood. Given the increasing proportion of survivors among infants born preterm or small-for-gestational-age, other disease patterns may occur among infants born preterm or small-for-gestational-age today.

In conclusion, we found that neither preterm birth, nor low birth weight, are associated with an increased risk for breast cancer.

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