Effects of childhood body size on breast cancer tumour characteristics

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Abstract

Introduction: Although a role of childhood body size in postmenopausal breast cancer risk has been established, less is known about its influence on tumour characteristics.

Methods: We studied the relationships between childhood body size and tumour characteristics in a Swedish population-based case-control study consisting of 2,818 breast cancer cases and 3,111 controls. Our classification of childhood body size was derived from a nine-level somatotype. Relative risks were estimated by odds ratios with 95% confidence intervals, derived from fitting unconditional logistic regression models. Association between somatotype at age 7 and tumour characteristics were evaluated in a case-only analysis where P values for heterogeneity were obtained by performing one degree of freedom trend tests.

Results: A large somatotype at age 7 was found to be associated with decreased postmenopausal breast cancer risk. Although strongly associated with other risk factors such as age of menarche, adult body mass index and mammographic density, somatotype at age 7 remained a significant protective factor (odds ratio (OR) comparing large to lean somatotype at age 7 = 0.73, 95% confidence interval (CI) = 0.58-0.91, P trend = 0.004) after adjustment. The significant protective effect was observed within all subgroups defined by estrogen receptor (ER) and progesterone receptor (PR) status, with a stronger effect for ER-negative (0.40, 95% CI = 0.21-0.75, P trend = 0.002), than for ER-positive (0.80, 95% CI = 0.62-1.05, P trend = 0.062), tumours (P heterogeneity = 0.046). Somatotype at age 7 was not associated with tumour size, histology, grade or the presence or absence of metastatic nodes.

Conclusions: Greater body size at age 7 is associated with a decreased risk of postmenopausal breast cancer, and the associated protective effect is stronger for the ER-negative breast cancer subtype than for the ER-positive subtype.

Introduction

There is considerable evidence that childhood anthropometric measurements are associated with postmenopausal breast cancer risk. It has been consistently shown that variables that approximate body shape and size early in life are inversely associated with breast cancer risk in adulthood. For example, a study conducted in 1998 on the same data set as used in the current study [1] reported that a larger somatotype at age seven years was associated with a lower postmenopausal breast cancer risk. Likewise, Hilakivi-Clarke and colleagues [2] found that a shorter height and higher body mass in girls from age 7 to 15 years were associated with a decreased incidence of breast cancer. Berkey and colleagues [3] also found extremely lean body mass at age 10 years to be associated with elevated breast cancer risk. In another study performed in 141,393 Danish girls, a high childhood body mass index (BMI) at age 14 years was shown to be protective against breast cancer later on in life [4]. In addition, a study performed on the large Nurses’ Health Study dataset concluded that average body fatness between the ages of 5 and 10 years are inversely associated with mammographic density [5], which is generally considered to be an intermediate phenotype of breast cancer [6].

Although a role of childhood body size in adult breast cancer risk has been established, less is known about its influence on tumour characteristics. One study by Bardia and colleagues [7] looked into the risk of developing postmenopausal breast cancer stratified by estrogen receptor (ER) and progesterone receptor (PR) subtypes and reported that an increase in weight at age 12 years was
associated with a decrease in adult breast cancer risk, with the most pronounced effects exhibited by ER-positive/PR-negative tumours. No significant heterogeneity, however, was observed between the tumour subtypes studied. To our knowledge, no other study has been conducted to assess whether pre-/peri-pubertal measurements of body size can also influence tumour characteristics. We thus followed up on the work of Bar-dia and colleagues and in the present study examined the relations between childhood body size to address if the far-reaching effects of childhood body size have any influence on tumour characteristics in adult cancers.

Materials and methods

Subjects

The subjects included in the current study are drawn from a population-based case-control study of postmenopausal breast cancer in Swedish-born women aged 50 to 74 years at the time of enrolment, which was between 1 October, 1993 and 31 March, 1995. Controls were randomly selected from the Swedish registry and frequency matched to the expected age distribution of the cases. Details on data collection and subjects have been described previously [1]. The final study group included 2,818 cases and 3,111 controls. Approval of the study was given by the ethical review board at the Karolinska Institutet (Stockholm, Sweden) and six other ethical review boards in the respective regions from which the subjects were based.

Data collection and classification

With the exception of clinical data on tumour characteristics and mammographic density, all other covariate data were derived from the parent case-control study. Anthropometric measurements at age seven years and one year prior to enrolment were collected by means of a nine-level somatotype (Figure 1) featured in the study questionnaire, and the validity of this measurement method has been previously described [1]. These pictograms have been validated against BMI within a cohort of 100 Caucasian women from middle-class communities with an average age of 73.1 years [8]. In a population-based validation study, 111 Swedish women aged 51 to 66 years were found to have a correlation coefficient between BMI from school records and adult report of somatotype at age seven years of 0.6 [1]. The somatotypes were subsequently grouped as lean (S1 to S2), medium (S3 to S4) and large (S5 to S9) prior to analysis. Other covariate data that was collected using the self-reported study questionnaire and examined in this study include age of menarche (continuous, in years), parity (continuous, number of live births), history of benign breast disease (binary, never/ever), BMI (continuous, in kg/m²), history of hormone replacement therapy (HRT) (binary, never/ever), and family history of breast cancer (binary, no/yes). Age at menopause (continuous, in years) was also derived from information collected in the study questionnaire and the definition used in this study has been previously described [1]. It is defined as the age at the last menstrual period or the age at bilateral oophorectomy, if one year or more prior to data collection. Women who have had a hysterectomy, or who have not ceased menstruation due to HRT, or with missing information on age at menopause were considered to be postmenopausal if the age reported at time of questionnaire was equal to or above the 90th percentile of age at natural menopause of study subjects (current smokers: 54 years old; nonsmokers: 55 years old, independent of case/control status). Subjects classified as postmenopausal in this manner were assigned an age at menopause according to their current smoking status and the mean ages at natural menopause in our data. Otherwise, women were considered to be premenopausal and were excluded.

Information regarding the retrieval of tumour characteristics from the medical records of all participants from surgical and oncological units throughout Sweden have been presented in detail elsewhere [9,10]. The tumour characteristics in the present study included tumour size (categorical, groups in cm), grade (categorical, classified according to the Nottingham histological grade or Bloom-Richardson scale), as well as ER and PR status (binary, absent/present).

The process of collecting mammographic density data in this study has been described previously [11]. Film mammograms of the medio-lateral oblique view were digitised using an Array 2905HD Laser Film Digitizer (Array Corporation, Tokyo, Japan), which covers a range of 0 to 4.7 optical density. For controls, breast side was randomised. For cases, the side contralateral to the tumour was used. The density resolution was set at 12-bit spatial resolution. The Cumulus software used for the computer-assisted thresholding was developed at the University of Toronto [12]. For each image, a trained observer (LE) set the appropriate gray-scale threshold levels defining the edge of the breast and distinguishing dense from non-dense tissue. The software calculated the

Figure 1 Nine-level somatotype pictogram.
total number of pixels within the entire region of interest and within the region identified as dense. These values were used to calculate the percentage of the breast area that is dense. A random 10% of the images were included as replicates to assess the intra-observer reliability, which was high with a Spearman rank correlation coefficient of 0.95. However, as not all women attended mammographic screenings, and some mammograms were missing, such information was available for only a subset of the subjects (n = 3232, 54.5%).

**Statistical analyses**

The distribution of baseline characteristics of known breast cancer risk factors were summarised as means and standard deviations or proportions. Odds ratio (OR) estimates with corresponding 95% confidence intervals (CI) were computed by fitting unconditional logistic regression models with breast cancer risk status as the response variable, adjusting for age.

To identify potential confounders of the association between somatotype at age seven years and breast cancer risk, linear/logistic regression models were fitted for either continuous (age of menarche, age of menopause, parity, BMI, and mammographic density) or binary (benign breast disease and HRT) outcomes including only controls in the analysis. Somatotype at age seven years was treated as a categorical (three-level) independent variable. Proportional odds logistic regression was used in situations where the outcome variable was ordinal (somatotypes at age seven years and one year prior to enrolment) from which cumulative OR estimates with corresponding 95% CIs were computed. Covariates were considered potential confounders if there was a priori evidence in the published literature of the factor being associated with both childhood body size and breast cancer risk, or if the factor was significantly associated at the 5% level with both somatotype at age seven years and breast cancer risk. Those covariates that, when added to the model, changed the coefficient by more than 10%, were considered confounders and adjusted for in the multivariate analysis. The final variables in the multivariate logistic regression model examining breast cancer risk overall, and stratified by ER and PR tumour subtypes, included age, age at menarche, benign breast disease, and BMI one year prior to enrolment (recent BMI). Adjustment for other variables did not influence the somatotype risk estimates. Mammographic density was also identified as a confounder. However, as mammographic density data are only available for a subset of the subjects, this variable was accounted for together with the other risk factors in a separate model. Women with and without mammographic density information were not found to differ significantly at the 5% level for the covariates included in the analysis models (data not shown).

Associations between somatotype at age seven years and tumour characteristics were evaluated in a case-only analysis, by fitting ordinal regression models treating tumour characteristics as dependent variables, with somatotype at age seven years included as a covariate. $P$ values for heterogeneity were obtained by performing one degree of freedom trend tests. As there exists prior evidence that certain tumour characteristics such as ER status are associated with age at diagnosis [13], and that somatotype at age seven years is significantly associated with age of diagnosis at the 5% level (regression coefficient for age in years of -0.91 with corresponding 95% CI of -1.32 to -0.50), every model fitted in the case-only analysis was also adjusted for age at diagnosis. All analyses were performed using the statistical software R for Windows version 2.8.0 (R Development Core Team, Vienna, Austria) [14]. The level of significance was set at 5%. All statistical tests were two-sided.

**Results**

Table S1 in Additional file 1 describes the characteristics of study subjects with respect to several breast cancer risk factors. Age of menarche was weakly but positively associated with the disease (OR per year increase in age of menarche = 0.96, 95% CI = 0.93 to 1.00, $P = 0.057$), a result consistent with the literature [4]. Family history, age at menopause, parity, age of first birth, benign breast disease, mammographic density, recent BMI and use of HRT were strongly significant for breast cancer risk with effects in a direction consistent with those estimated in other epidemiological studies. The first association analyses we performed between somatotypes at different ages and breast cancer risk were adjusted for age at enrolment only. Among the different measurements of somatotypes, only the time point at age seven years was found to affect breast cancer risk (OR per increase in somatotype class = 0.87, 95% CI = 0.8 to 0.95, $P = 0.001$). A larger proportion of cases than controls had a leaner body shape at age seven years. Despite somatotype one year prior to enrolment having a high correlation to recent BMI (Spearman correlation coefficient: 0.760, data not shown), it was not found to be significantly associated with breast cancer (OR per increase in somatotype class = 1.04, 95% CI = 0.94 to 1.15, $P = 0.160$).

To identify potential confounders of the association between somatotype at age seven years and breast cancer risk, we assessed whether other established risk factors for breast cancer are associated with somatotype at age seven years. An increase in childhood body size was found to exhibit strong inverse associations with age of menarche (OR comparing large to lean somatotype at age seven years = 0.61, 95% CI = 0.50 to 0.76, $P$ trend < 0.0001), benign breast disease (0.47, 95% CI = 0.25 to 0.89, $P$ trend = 0.006), and mammographic density (0.61,
Table 1: Associations of somatotype at age seven years with other breast cancer risk factors (controls only)

| Risk factor (dependent variable) | Somatotype (independent variable) | n   | OR   | 95% CI       | P trend* |
|---------------------------------|-----------------------------------|-----|------|--------------|---------|
| Age of menarche (years)         | Lean                              | 1456| 1.00 | reference    | <0.0001 |
|                                 | Medium                            | 669 | 0.72 | 0.64 0.82    |         |
|                                 | Large                             | 187 | 0.61 | 0.50 0.76    |         |
| Age of menopause (years)        | Lean                              | 1572| 1.00 | reference    | 0.697   |
|                                 | Medium                            | 736 | 1.19 | 0.85 1.68    |         |
|                                 | Large                             | 204 | 0.93 | 0.53 1.65    |         |
| Parity (Number of live births)  | Lean                              | 1578| 1.00 | reference    | 0.217   |
|                                 | Medium                            | 745 | 0.93 | 0.83 1.05    |         |
|                                 | Large                             | 207 | 0.93 | 0.76 1.13    |         |
| Benign breast disease           | Lean                              | 1578| 1.00 | reference    | 0.006   |
|                                 | Medium                            | 745 | 0.76 | 0.56 1.03    |         |
|                                 | Large                             | 207 | 0.47 | 0.25 0.89    |         |
| Somatotype one year prior to enrolment | Lean                     | 1571| 1.00 | reference    | <0.0001 |
|                                 | Medium                            | 739 | 1.72 | 1.44 2.05    |         |
|                                 | Large                             | 206 | 2.33 | 1.70 3.18    |         |
| BMI (kg/m²)                     | Lean                              | 1562| 1.00 | reference    | <0.0001 |
|                                 | Medium                            | 742 | 1.85 | 1.30 2.65    |         |
|                                 | Large                             | 205 | 2.66 | 1.47 4.83    |         |
| Percent mammographic density (%)†| Lean                             | 862 | 1.00 | reference    | 0.001   |
|                                 | Medium                            | 428 | 0.72 | 0.58 0.91    |         |
|                                 | Large                             | 108 | 0.61 | 0.41 0.90    |         |
| HRT                             | Lean                              | 1569| 1.00 | reference    | 0.868   |
|                                 | Medium                            | 739 | 0.99 | 0.83 1.18    |         |
|                                 | Large                             | 206 | 0.98 | 0.73 1.32    |         |

Other independent variables

| Birthweight (g) on somatotype at age 7 | ≤2500 | 2500-3000 | 3000-3500 | 3500-4000 | >4000 |
|----------------------------------------|------|----------|----------|----------|-------|
|                                        | 49   | 229      | 470      | 397      | 135   |
| Birthweight (g) on somatotype at age 7 | 1.00 | 1.18     | 1.29     | 1.44     | 1.89  |
|                                        | reference | 0.61   | 0.68     | 0.76     | 0.95  |
|                                        | 2.29 | 2.43     | 2.73     | 3.76     |       |

| Family history on somatotype at age 7 | No   | Yes      |          |          |       |
|---------------------------------------|------|---------|----------|---------|
|                                       | 2258 | 227     | 1.00     | 1.10    | 1.44  |
| Family history on somatotype at age 7 | reference | 0.64   | 0.84     |         |       |

* Based on Wald tests for regression coefficients in continuous, ordinal or logistic regression models (see statistical analyses section). All regression models were adjusted for age at enrolment. † Subset with phenotypic data. BMI, body mass index; CI, confidence interval; HRT, hormone replacement therapy; OR, odds ratio.

95% CI = 0.41 to 0.90, P trend = 0.001; Table 1). Associations in the opposite direction were found for proxy measures of physique at other time points, such as birthweight (OR comparing birthweight >4000 g to ≤2500 g = 1.89, 95% CI = 0.95 to 3.76, P trend = 0.014), somatotype one year prior to enrolment (OR comparing large to lean somatotype at age seven years = 2.33, 95% CI = 1.70 to 3.18, P trend < 0.0001) and recent BMI (2.66, 95% CI = 1.47 to 4.83, P trend < 0.0001). No evidence of association was found between age of menopause and somatotype at age seven years.
age seven years or between family history and somatotype at age seven years. Parity and HRT were found to be independent of somatotype at age seven years (0.93, 95% CI = 0.76 to 1.13, P trend = 0.217 and 0.98, 95% CI = 0.73 to 1.32, P trend = 0.868, respectively).

After adjustment of known breast cancer predictors and other associated risk factors, the inverse association of somatotype at age seven years with breast cancer remained highly significant (Table 2; OR comparing large to lean somatotype at age seven years = 0.73, 95% CI = 0.58 to 0.91, P trend = 0.004). The protective effect of a larger somatotype was found to be significant (P trend < 0.05) for ER-negative, PR-positive and PR-negative subtypes and marginally significant (P trend = 0.062) for the ER-positive subtype. Within the group consisting of large somatypes, the most prominent effects were shown in ER-negative (OR comparing large to lean somatotype at age seven years = 0.40, 95% CI = 0.21 to 0.75, P trend = 0.002) and PR-negative (0.63, 95% CI = 0.40 to 0.99, P trend = 0.028) tumours. The point estimates changed very little before and after additional adjustment for mammographic density as a continuous variable [see Table S2 in Additional file 2], using a subset of the data with this information available (n = 3232).

We next assessed the effects of childhood body size on tumour characteristics (ER status, PR status, tumour size, grade, histology, and absence/presence of metastatic nodes) by fitting binary/ordinal logistic regression models, adjusting for age at diagnosis in years as a confounder. We established that the protective effect of somatotype at age seven years was significantly stronger for ER-negative disease than for ER-positive disease (P heterogeneity = 0.046; Table 3). When comparing between two extreme groups, women with a larger body size at age seven years were 1.71 times (95% CI = 0.96 to 3.06) more likely to get ER-positive than ER-negative disease after menopause. Although the estimated trend suggests that women with the same physique are more likely to get the PR-positive disease in adulthood, the difference between the two tumour subtypes was not significant (P heterogeneity = 0.283). The point estimates for tumour size, histology, grade, or the presence or absence of metastatic nodes did not vary much before and after adjustment for age of diagnosis as a continuous variable.

Discussion
Our first main finding was that a large somatotype at age seven years was associated with a decreased risk of post-menopausal breast cancer. Although strongly associated with other risk factors such as age of menarche, adult BMI and mammographic density, somatotype at age seven years remained a significant protective factor (OR comparing large to lean somatotype at age seven years = 0.73, 95% CI = 0.58 to 0.91, P trend = 0.004) after adjust-
Table 2: Multivariate-adjusted OR estimates and corresponding 95% CIs of postmenopausal breast cancer for somatotype at age seven years, overall and stratified by breast cancer tumour subtype based on ER and PR status

| Type of breast cancer | Somatotype  | All subjects |
|-----------------------|------------|--------------|
|                       | Cases      | OR  | 95% CI | P trend* |
| All data              | Lean       | 1784 | 1.00  | reference | 0.004 |
|                       | Medium     | 757  | 0.90  | 0.79     | 1.02  |
|                       | Large      | 173  | 0.73  | 0.58     | 0.91  |
| ER positive           | Lean       | 963  | 1.00  | reference | 0.062 |
|                       | Medium     | 408  | 0.91  | 0.78     | 1.06  |
|                       | Large      | 98   | 0.80  | 0.62     | 1.05  |
| ER negative           | Lean       | 219  | 1.00  | reference | 0.002 |
|                       | Medium     | 81   | 0.77  | 0.58     | 1.03  |
|                       | Large      | 14   | 0.40  | 0.21     | 0.75  |
| PR positive           | Lean       | 841  | 1.00  | reference | 0.027 |
|                       | Medium     | 354  | 0.89  | 0.75     | 1.04  |
|                       | Large      | 83   | 0.76  | 0.57     | 1.00  |
| PR negative           | Lean       | 320  | 1.00  | reference | 0.028 |
|                       | Medium     | 126  | 0.86  | 0.68     | 1.08  |
|                       | Large      | 25   | 0.63  | 0.40     | 0.99  |

* Logistic regression models were used, accounting for age, age at menarche, benign breast disease and recent body mass index. CI, confidence interval; ER, estrogen receptor; OR, odds ratio; PR, progesterone receptor.

leagues [29] showed that high childhood BMI was associated with a lower Wolfe grade, and Samimi and colleagues [5] found that a rounder pre-pubertal body shape was predictive of lower mammographic density later in life.

The age-adjusted case-only comparison of our data reflected a significant difference in the effects of childhood body size on the two ER subtypes (P trend = 0.046), but not the PR subtypes. However, in lieu of the fact that PR is an estrogen-induced target gene, and that its presence could serve to indicate ER functional capacity and tumour differentiation state [30], we also conducted stratified analyses on PR subtypes. We found that the protective trend conferred by a larger childhood somatotype on postmenopausal breast cancer applies to all ER and PR tumour subtypes. Overall our results were consistent with Bardia and colleagues [7], although in that study the effects were only significant for ER-positive (0.80, 95% CI = 0.67 to 0.96) and PR-negative (0.62, 95% CI = 0.43 to 0.89) tumours (comparing women with above average weight at age 12 years to women with average weight at age 12 years). Although Bardia and colleagues observed a stronger protective effect in ER-negative tumours than in their ER-positive counterparts (in agreement with our finding) when comparing women with above average weight at age 12 years to women with average weight at age 12 years, the association they observed in this subgroup was not statistically significant (0.77, 95% CI = 0.5 to 1.19).

Hormonal exposure and mammographic density are established risk factors of breast cancer that have been suggested to be independent, operating through different pathways [31]. Adjustment for these factors and other traditional risk factors did not attenuate the negative association of childhood body size on breast cancer risk (OR comparing large to lean somatotype at age seven years = 0.73, 95% CI = 0.58 to 0.91, P trend = 0.004, for association, after adjustment), thus suggesting an independent underlying mechanism. We speculate that a possible mechanism driving the negative association with breast cancer risk could be epigenetic changes that occur during mammary development. Hilakivi-Clarke [32] summarised in a review several perspectives on special windows of mammary development. Mammary tissue is postulated to undergo epigenetic extensive modelling or re-modelling during different stages in life such as fetal development, puberty or pregnancy. Such epigenetic modification can persist into adulthood if taken place in mammary stem cells, uncommitted mammary myoepithelial or luminal progenitor cells and inherited by subse-
Table 3: Relation of somatotype at age seven years to tumour-defined characteristics of breast cancer

| Tumour characteristics | Categories | Somatotype at age seven years | P heterogeneity§ |
|------------------------|------------|------------------------------|-----------------|
|                        |            | S1-S2 | S3-S4 | S5-S9 |                |
| Tumour size (cm)*      | <1         | 300   | 138   | 39    |                |
|                        | 1-2        | 752   | 299   | 70    |                |
|                        | 2-3        | 366   | 152   | 31    |                |
|                        | 3-4        | 116   | 66    | 14    |                |
|                        | 4-5        | 52    | 16    | 3     |                |
|                        | >=5        | 65    | 26    | 4     |                |
| Cumulative OR (95% CI) | 1.00 (ref.) | 1.00 (0.85-1.17) | 0.78 (0.58-1.05) | 0.255 |
| Cumulative OR (95% CI) § | 1.00 (ref.) | 1.00 (0.85-1.18) | 0.78 (0.58-1.06) | 0.266 |
| Grade*                 | Low        | 159   | 69    | 20    |                |
|                        | Medium     | 479   | 186   | 46    |                |
|                        | High       | 463   | 222   | 51    |                |
| Cumulative OR (95% CI) | 1.00 (ref.) | 1.15 (0.94-1.41) | 0.99 (0.69-1.43) | 0.443 |
| Cumulative OR (95% CI) § | 1.00 (ref.) | 1.15 (0.93-1.41) | 0.99 (0.69-1.42) | 0.463 |
| Histology*             | Ductal     | 1350  | 570   | 137   |                |
|                        | Lobular    | 206   | 77    | 16    |                |
|                        | All other  | 92    | 37    | 7     |                |
| Cumulative OR (95% CI) | 1.00 (ref.) | 0.91 (0.72-1.15) | 0.76 (0.48-1.20) | 0.192 |
| Cumulative OR (95% CI) § | 1.00 (ref.) | 0.92 (0.73-1.17) | 0.79 (0.50-1.25) | 0.265 |
| Metastatic nodes†      | Absent     | 1159  | 473   | 116   |                |
|                        | Present    | 513   | 227   | 46    |                |
| OR (95% CI)            | 1.00 (ref.) | 1.08 (0.90-1.31) | 0.90 (0.63-1.28) | 0.923 |
| OR (95% CI) §          | 1.00 (ref.) | 1.07 (0.88-1.29) | 0.86 (0.60-1.23) | 0.878 |
| ER status†             | Negative   | 219   | 81    | 14    |                |
|                        | Positive   | 963   | 408   | 98    |                |
| OR (95% CI)            | 1.00 (ref.) | 1.15 (0.87-1.52) | 1.59 (0.89-2.84) | 0.089 |
| OR (95% CI) §          | 1.00 (ref.) | 1.18 (0.89-1.56) | 1.71 (0.96-3.06) | **0.046** |
| PR status†             | Negative   | 320   | 126   | 25    |                |
|                        | Positive   | 841   | 354   | 83    |                |
| OR (95% CI)            | 1.00 (ref.) | 1.07 (0.84-1.36) | 1.26 (0.79-2.01) | 0.307 |
| OR (95% CI) §          | 1.00 (ref.) | 1.07 (0.84-1.37) | 1.28 (0.80-2.03) | 0.283 |

*Proportional odds logistic regression models were used. † Logistic regression models were used. ‡ Derived from one degree of freedom trend tests. § Adjusted for age at diagnosis. CI, confidence interval; ER, estrogen receptor; OR, odds ratio; PR, progesterone receptor.
quent daughter cells [33]. Prepubertal exposure to estrogen has been shown to upregulate the expression of BRCA1, a well-known DNA repair gene [28]. Liu and colleagues [34] also demonstrated that BRCA1 is responsible for differentiating ER-negative stem/progenitor cells into ER-positive luminal cells. They also proposed that loss of expression of the DNA repair gene (BRCA1) may result in an accumulation of ER-negative stem cells with multiple genetic defects. Incidentally, loss of BRCA1 is frequently associated with ER-negative breast cancers [35]. The evidence for altered gene expression possibly caused by childhood body size helps to explain the general reduction in breast cancer risk overall. The apparent differential protection conferred to the ER-negative subtype could possibly be driven by the same underlying mechanism that operates through epigenetic modifications.

The strengths of our study include being a population-based study, its large sample size and detailed information on many variables: anthropometric measures at different time points throughout life, mammographic density, reproductive and hormonal risk factors, and tumour characteristics. To our knowledge, this is the first study to consider the effects of somatotype at age seven years on adult breast cancer with the consideration of mammographic density, and also the first to examine its effects on tumour characteristics other than ER status.

A limitation of our study is that risk factor data were self-reported, and could thus be measured with error. Although two studies have demonstrated the validity of using the nine-level somatotype diagram for the long-term recall of childhood body size via high correlations with BMI at the same ages [8,36], it is noteworthy that in those studies no woman recalled their figure as larger than level seven in these studies, and that women with large body size were more likely to misreport their childhood somatotypes than women who were lean. However, any such measurement error is most likely to attenuate any association between childhood body size and breast cancer risk [37]. In addition, as the questionnaire study was conducted post-diagnosis of breast cancer, recall bias could have been introduced. Although the nine-level somatotype measure has not been validated specifically for breast cancer, our findings may have important implications. The effects of childhood body size on the different breast cancer subtypes are independent of other breast cancer risk factors, such as mammographic density and estrogen exposure. Given the strength of the associations, and the ease of retrieval of information on childhood somatotypes retrospectively from pictures early in life, childhood body size is potentially useful for building breast cancer risk or prognosis prediction models.

Additional material

Additional file 1: Table S1. Descriptive characteristics of post-menopausal women.

Additional file 2: Table S2. Multivariate-adjusted odds ratio (OR) estimates and corresponding 95% confidence intervals (CIs) of postmenopausal breast cancer for somatotype at age seven years on a subset of women with mammographic density data; overall and stratified by breast cancer tumour subtype based on estrogen receptor (ER) and progesterone receptor (PR) status.

Abbreviations

BMI: body mass index; CI: confidence interval; ER: estrogen receptor; HRT: hormone replacement therapy; OR: odds ratio; PR: progesterone receptor.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

J Li participated in the study design, carried out the analyses and drafted the manuscript. LE digitised and obtained readings for the mammograms. KH, KC, J Liu and PH participated in study design and coordination and helped to draft the manuscript. All authors read and approved the final manuscript.

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