Research article

Different measures of smoking exposure and mammographic density in postmenopausal Norwegian women: a cross-sectional study

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Abstract

Background Recent cohort studies have suggested an increased risk of breast cancer with long duration of smoking, and with smoking initiation before first birth. Cigarette smoking may have both carcinogenic effects and antiestrogenic effects on the breast tissue. We decided to examine the relationship between different measures of smoking exposure and mammographic density.

Methods Lifetime smoking history was collected through interview and questionnaires among 907 postmenopausal participants in the Tromsø Mammography and Breast Cancer study. The mammograms were obtained from the governmental Norwegian Breast Cancer Screening Program. Mammograms were classified according to the percentage and absolute mammographic densities using a previously validated computer-assisted method.

Results Sixty-five percent of the women reported having ever smoked cigarettes, while 34% were current smokers. After adjustment for age, age at first birth, parity, age at menopause, postmenopausal hormone therapy use, and body mass index, smoking was inversely associated with both measures of mammographic density (both trends \( P < 0.01 \)). Both current smokers and former smokers had significantly lower adjusted mean percentage mammographic density compared with never smokers (\( P = 0.003 \) and \( P = 0.006 \), respectively). An inverse dose–response relationship with mammographic density was found between both the number of cigarettes and the number of pack-years smoked among current smokers. Current smokers who smoked 11 cigarettes or more daily had a 3.7% absolute (36% relative difference) lower percentage mammographic density compared with current smokers who smoked seven cigarettes or less daily (\( P = 0.008 \)). When former smokers were stratified according to time since smoking cessation, we found that women who had stopped smoking less than 24 years ago had a significantly lower mean percentage mammographic density compared with never smokers (\( P < 0.001 \)).

Conclusion We found modest inverse dose–response associations between numbers of cigarettes and of pack-years smoked and both measures of mammographic density among current smokers. Former smokers who had stopped smoking less than 24 years ago also had a statistically significantly lower mean percentage mammographic density when compared with never smokers. These findings are consistent with an antiestrogenic effect of cigarette smoking on the breast tissue.

Introduction

Constituents in tobacco smoke may have carcinogenic effects on the breast tissue [1-3]. Tobacco smoking, however, may also have antiestrogenic effects that can reduce breast cancer risk [4,5]. These opposing effects may explain the overall inconsistent results from epidemiologic studies on the association between smoking and breast cancer risk [2,6-9]. Although most case–control studies do not find any positive associations [2,10], however, several recent cohort studies have indicated an increased breast cancer risk among women.

BMI = body mass index; HT = postmenopausal hormone therapy.
who are long-term smokers [11-14], and also among those who start to smoke before their first birth [12-17].

Mammographic density is one of the strongest independent risk factors for breast cancer [18,19], and possibly an intermediate marker for breast cancer [20]. Women with high mammographic density have a fourfold to sixfold increase in breast cancer risk compared with those with low mammographic density [18,19].

The results so far published on the association between smoking and mammographic density have also been conflicting [21-27]. One explanation for this may be that most studies have used crude (for example, ever/never) measures of smoking exposure.

The objective of the present cross-sectional study was to examine the relationship between cigarette smoking and mammographic density among postmenopausal women with a high smoking prevalence, according to different measures of smoking exposure.

**Materials and methods**

**Study population**
The Tromsø Mammography and Breast Cancer Study was conducted among postmenopausal women, aged 55–71 years, residing in the municipality of Tromsø, Norway, and attending the population-based Norwegian Breast Cancer Screening Program at the University Hospital of North Norway [28]. Women were recruited in spring 2001 and in spring 2002. After the women had undergone their screening mammograms, they were interviewed by a trained research nurse about reproductive and menstrual factors, previous history of cancer, current smoking status, and use of postmenopausal hormone therapy (HT) or other medications. The participants had their height measured to the nearest centimeter and their weight measured to the nearest half kilogram. Women had blood samples drawn, and each participant was subsequently given a questionnaire to be completed at home, eliciting information on demographics, additional menstrual and reproductive factors, lifetime smoking history, as well as lifestyle and dietary factors. All women signed an informed consent. The National Data Inspection Board and the Regional Committee for Medical Research Ethics approved the study. Altogether, 1,041 women were included in this cross-sectional study. This accounted for 70% of the women attending the Norwegian Breast Cancer Screening Program during the recruitment period.

We excluded 22 women because of a previously diagnosed ($n=16$) or a newly diagnosed ($n=6$) breast cancer, and one woman because of an ongoing chemotherapy treatment. Among the remaining 1,018 women, we were unable to retrieve mammograms on 11 women. We therefore obtained mammographic density readings on 1,007 women. More details are described elsewhere [28]. We further excluded three women because they were equivocal for menopausal status and excluded 97 women because of a missing smoking history, leaving 907 women for the analyses.

**Mammographic classifications**
The women’s left cranio-caudal mammogram was digitized using a Cobrascan CX-812 scanner (Radiographic Digital Imaging, Torrance, CA, USA) at a resolution of 150 pixels per inch. Percentage and absolute mammographic densities were determined by an experienced reader (GU) using the University of Southern California Madena computer-based threshold method, which has been described in detail and validated elsewhere [29]. Briefly, the method works as follows. The digitized mammographic image is viewed on a computer screen. A reader defines the total breast area using a special outlining tool. Next, the region of interest – excluding the pectoralis muscle, prominent veins, and fibrous strands – is defined. The computer software program assigns a pixel value of 0 to the darkest (black) shade in the image and a value of 255 to the lightest (white) shade, with shades of grey assigned to intermediate values. The reader then uses a tinting tool to apply a yellow tint to dense pixels with grey levels at or above some threshold $X$ and a pixel value $\leq 255$. The reader searches for the best threshold where all pixels $\geq X$ within the region of interest are considered to represent mammographic densities. The software estimates the total number of pixels and the number of tinted pixels within the region of interest. The absolute density represents the count of the tinted pixels within the region of interest. The percentage density, or the fraction (%) of the breast with densities, is the ratio of the absolute density to the total breast area multiplied by 100. The absolute density (measured in centimeters squared) was calculated as the number of tinted pixels within the region of interest divided by the number of pixels per centimeter squared.

The reader of the mammograms was blinded to characteristics of the study participants. We have previously shown a high intra-rater agreement for the reader in our study (Pearson correlation coefficient $= 0.86$) [28].

**Smoking assessments**
The women were interviewed about their current smoking status. The self-administered questionnaire elicited additional information on the participant’s lifetime smoking history. Women reporting never having smoked or never having been exposed to passive smoking were categorized as ‘never active smokers’. We further categorized women who had never actively smoked but had been exposed to passive smoking at home or at the workplace as ‘passive smokers’. Never active smokers and passive smokers were also grouped together as ‘never smokers’. This group serves as the reference group in all analyses, if not specified otherwise. Current smokers and former smokers were grouped together as ‘ever-smokers’. Pack-years were calculated as the number of cigarettes
smoked daily divided by 20 and multiplied by the number of years smoked.

We categorized former smokers according to the time since smoking cessation (tertiles). We categorized current smokers according to their age at smoking initiation (tertiles), to the average number of cigarettes smoked per day (tertiles), to the number of years smoked (≤ 25 years, 26–40 years, 41+ years), to the number of pack-years smoked (tertiles), and parous women according to smoking initiation before or after first birth.

Statistical analyses
Mammographic density was not normally distributed. Both the percentage and absolute mammographic densities were log-transformed to obtain approximate normal distributions. We used analysis of variance for an unbalanced design to study the associations between cigarette smoking and mammographic densities (Proc GLM; SAS Institute Inc., Cary, NC, USA).

Each of the following factors was evaluated as a potential confounder of the association between smoking and mammographic density: age at screening (continuous), age at menarche (continuous), age at menopause (continuous), number of children (continuous), age at first birth (continuous), years of education (continuous), family history of breast cancer in first-degree relatives (yes, no), alcohol intake (g/day, continuous), HT use (never used, past use, current use), and body mass index (BMI) (weight in kilogram divided by height in meters squared; continuous).

The variables of age at screening, number of children, HT use, and BMI have previously been found associated with mammographic density in this study population [28,30], and were always adjusted for as confounders in the multivariate analyses. We further identified those of the above listed variables that were associated with cigarette smoking in univariate analyses, and that also were significantly associated with mammographic density in the multivariate analyses. This procedure left the following factors in the final model: age at screening, age at first birth, number of children, age at menopause, HT use and BMI. Women with missing values were excluded from the analyses (<10%).

Reported trend-test P values correspond to analyses where the categories of cigarette smoking were treated as ordered variables. We tested for possible effect modification by analyzing the association between smoking and mammographic density stratified by the confounders, and by adding multiplicative interaction terms to the analysis of variance procedure.

The crude and adjusted mean mammographic density results were back-transformed, and are presented with 95% confidence intervals. Results were considered statistically significant if the two-sided P value was < 0.05. We conducted all statistical analyses using SAS® 9.1 for Windows (SAS Institute Inc.).

Results
Altogether, 65% of the women were ever smokers. Among the 310 (34%) women who reported current smoking, 82% smoked daily. Altogether, 318 women reported to have never been smokers, among whom 73% reported to have been exposed to passive smoking at home or at the workplace.

Table 1 presents the distribution of selected characteristics according to smoking status. Current smokers were younger at screening, were younger at time of first birth, had less formal education, reached menopause at an earlier age, were leaner, and were more likely to have ever used oral contraceptives, when compared with never smokers (all P ≤ 0.02).

Former smokers also differed from never smokers, their data values being between current smokers and never smokers, with regard to age at first birth and ever oral contraceptive use (both P ≤ 0.007). Former smokers had a significantly higher BMI when compared with never smokers (P = 0.04). Passive smokers were more likely to have ever used HT (P < 0.05), but were otherwise similar to never smokers in the other measured characteristics (results not shown).

Table 2 presents the median, the crude mean and the adjusted mean mammographic densities across smoking status. Both current smokers and former smokers had lower crude mean mammographic density compared with never smokers, according to both the percentage and the absolute mammographic densities. When the associations were adjusted for age at screening, for age at first birth, for number of children, for age at menopause, for HT use and for BMI, both current smokers and former smokers had a significantly lower adjusted mean percentage mammographic density compared with never smokers (P = 0.003 and P = 0.006, respectively). The magnitude of the difference in the mean percentage mammographic density between current smokers and never smokers was 2.2% absolute (23% relative) difference. A similar association was found between smoking status and absolute mammographic density (Table 2). These associations did not change materially when we excluded passive smokers from never smokers (results not shown), or when we excluded occasional smokers from current smokers (results not shown).

Current smokers on average smoked 10 cigarettes per day. The majority of current smokers had initiated smoking by the age of 20, and had been smoking for 39 years or more. Among those reporting to be former smokers, 71% had stopped smoking 10 years or more ago. Table 3 presents the association between different measures of smoking exposure and the percentage mammographic density among current smokers, and according to time since smoking cessation among former
smokers. We found an inverse dose–response relationship between the number of cigarettes smoked daily and the percentage mammographic density among current smokers \( (P = 0.008) \), and this remained when the analysis was further restricted to current smokers who had smoked for 25 years or more \( (P = 0.01) \). Current smokers who smoked 11 cigarettes or more daily had a 3.7% absolute (36% relative difference) lower mean percentage mammographic density compared with current smokers who smoked seven cigarettes or less daily \( (P = 0.008) \). Furthermore, an inverse dose–response relationship was found between the number of pack-years and the percentage mammographic density among current smokers \( (P = 0.09) \). When former smokers were stratified according to time since smoking cessation, we found that women who had stopped smoking less than 24 years ago had a significantly lower mean percentage mammographic density compared with never smokers \( (P < 0.001) \). These associations were similar when absolute mammographic density was used as the outcome variable (results not shown).

### Table 1

**Characteristics of the study population \( (n = 907) \) by smoking status**

| Characteristic                  | Smoking status | \( P \) valuea |
|-------------------------------|----------------|---------------|
|                              | Current smokers \( (n = 310) \) | Former smokers \( (n = 279) \) | Never smokers \( (n = 318) \) |
| Age at screening (years)      | 60.7 (4.4)     | 61.6 (4.5)    | 61.9 (4.6)    | <0.001 |
| Age at menarche (years)       | 13.3 (1.4)     | 13.3 (1.4)    | 13.3 (1.4)    | 0.47   |
| Age at first birthb (years)   | 22.3 (3.6)     | 22.7 (3.3)    | 23.5 (3.9)    | <0.001 |
| Number of childrenb           | 2.9 (1.2)      | 2.8 (1.1)     | 2.9 (1.4)     | 0.82   |
| Education (years)             | 9.4 (3.2)      | 9.7 (3.4)     | 10.2 (3.6)    | 0.02   |
| Age at menopause (years)      | 47.5 (5.5)     | 48.8 (4.8)    | 49.1 (4.8)    | <0.001 |
| Body mass index (kg/m²)       | 26.1 (4.4)     | 28.4 (5.1)    | 27.6 (4.7)    | <0.001 |
| Alcohol consumptionc (g/day)  | 3.8 (3.8)      | 4.0 (3.9)     | 3.4 (4.0)     | 0.36   |
| Ever oral contraceptive use (%) | 54.8           | 49.5          | 37.4          | <0.001 |
| Parous (%)                    | 93.6           | 93.6          | 90.3          | 0.13   |
| Ever postmenopausal hormone therapy use (%) | 43.2           | 45.2          | 41.8          | 0.72   |
| Breast cancer in first-degree relative (%) | 6.8            | 9.3           | 9.8           | 0.18   |

Data presented as the mean (standard deviation) or as the frequency. aCurrent smokers versus never smokers. bAmong parous women only. cAmong alcohol drinkers only.

### Table 2

**Median, crude mean and adjusted mean percentage and absolute mammographic density by smoking status**

| Smoking status | \( n \) | Percentage mammographic density (%) | Absolute mammographic density (cm²) |
|----------------|--------|-------------------------------------|-------------------------------------|
|                |        | Median (range) | Crude mean (95% confidence interval) | Adjusted mean (95% confidence interval) | Median (range) | Crude mean (95% confidence interval) | Adjusted mean (95% confidence interval) |
| Current smokers | 310    | 9.4 (0–59.4) | 8.0 (7.0–9.1) | 7.3 (6.5–8.3) | 13.7 (0–110.1) | 10.7 (9.4–12.2) | 10.4 (9.0–12.0) |
| Former smokers  | 279    | 7.7 (0–51.1) | 6.7 (5.8–7.7) | 7.5 (6.6–8.5) | 11.9 (0–155.2) | 9.7 (8.4–11.4) | 10.6 (9.2–12.2) |
| Never smokers   | 318    | 11.9 (0–69.2) | 9.7 (8.6–11.0) | 9.5 (8.4–10.7) | 17.1 (0–152.3) | 14.0 (12.3–15.9) | 13.7 (11.9–15.7) |
| \( P \) valuea |        | 0.03     | 0.003       | 0.004        | 0.005        | <0.001       | 0.01        |
| \( P \) valuemb |        | <0.001   | 0.006       | <0.001       | 0.005        |

Analyses are adjusted for age at screening, age at first birth, number of children, age at menopause, postmenopausal hormone therapy use, and body mass index. Reported means are back-transformed from log-transformed estimated means. aCurrent smokers versus never smokers. bFormer smokers versus never smokers.
Discussion

This population-based cross-sectional study found an inverse association between smoking and percentage mammographic density among postmenopausal women, after adjustment for potential confounders. We also observed an inverse dose–response relationship among current smokers between both the numbers of cigarettes and pack-years smoked and the percentage mammographic density. Women who had stopped smoking less than 24 years ago had a significantly lower mean percentage mammographic density compared with never smokers. These associations were similar when absolute mammographic density was used as the outcome variable.

The strengths of our study are the large sample size and that it was a part of a population-based screening program with a high attendance rate [31]. The reader of the mammograms

| Smoking exposure                  | n    | Current smokers | Former smokers |
|-----------------------------------|------|-----------------|----------------|
| Number of cigarettes smoked per day |      |                 |                |
| ≤ 7 cigarettes                    | 87   | 10.3 (8.2–12.8) |                |
| 8–10 cigarettes                   | 79   | 9.0 (7.0–11.5)  |                |
| ≥ 11 cigarettes                   | 77   | 6.6 (5.2–84)    |                |
| P trend                           |      | 0.008           |                |
| Number of years smoked            |      |                 |                |
| ≤ 25 years                        | 56   | 8.9 (6.8–11.8)  |                |
| 26–40 years                       | 92   | 8.8 (6.9–11.3)  |                |
| ≥ 41 years                        | 95   | 8.1 (6.5–10.1)  |                |
| P trend                           |      | 0.57            |                |
| Number of pack-years smoked       |      |                 |                |
| ≤ 11 pack-years                   | 82   | 10.0 (7.9–12.6) |                |
| 12–20 pack-years                  | 85   | 8.5 (6.7–10.8)  |                |
| ≥ 21 pack-years                   | 75   | 7.5 (5.9–9.6)   |                |
| P trend                           |      | 0.09            |                |
| Age at smoking initiation         |      |                 |                |
| ≥ 21 years                        | 94   | 8.9 (7.1–11.1)  |                |
| 18–20 years                       | 114  | 8.4 (6.8–10.3)  |                |
| 13–17 years                       | 78   | 7.3 (5.8–9.3)   |                |
| P trend                           |      | 0.26            |                |
| Smoking initiation before first birtha |  |                |                |
| No                                | 79   | 9.2 (7.2–11.8)  |                |
| Yes                               | 186  | 7.8 (6.7–9.1)   |                |
| P value                           |      | 0.24            |                |
| Time since smoking cessation       |      |                 |                |
| ≥ 24 years                        | 89   | 8.5 (6.7–10.7)  |                |
| 12–23 years                       | 89   | 5.2 (4.0–6.7)   |                |
| ≤ 11 years                        | 88   | 6.5 (5.1–8.2)   |                |
| P trend                           |      | 0.12            |                |

Analyses are adjusted for age at screening, age at first birth, number of children, age at menopause, postmenopausal hormone therapy use, and body mass index. Reported means are back-transformed from log-transformed estimated means. aAmong parous women only.
was experienced and blinded to the characteristics of the women. We have previously shown a high intra-rater agreement for the reader in our study [28]. Further, we have assessed both relative and absolute measures of mammographic density as outcome variables, since they may be influenced differently by residual confounding by adiposity [32], and we found similar associations between smoking and mammographic density when we performed the analyses using the absolute mammographic density as the outcome variable. Another strength is that we have a large proportion of current smokers (34%) among the women in our study compared with previous studies among postmenopausal women, which all had less than 14% current smokers [21-24].

One limitation of our study is the possible misclassification of smoking exposure. Any misclassification of smoking would presumably be nondifferential with respect to mammographic density, and would therefore be expected to bias the results toward the null association. Another limitation is that the median mammographic density in our study is low and that mammographic density has a narrow range. In another study with a different reader, however, women from our study had significantly lower percentage mammographic density compared with Caucasians from Hawaii and Arizona [33].

In a study from the United States, Modugno and colleagues also found that current smokers had a significantly lower percentage mammographic density compared with noncurrent smokers. Their study, comprising 239 women aged 70 years or older, only had information about current smoking habits and therefore could not analyze the association in more detail [21].

Three other studies found no association between smoking and mammographic density among postmenopausal women [22-24]. In a British study with 406 women from Norfolk, Sala and colleagues found no significant association between smoking and high-risk Wolfe’s parenchymal patterns among the 313 postmenopausal women in their study, even though current smokers had a significantly reduced odds ratio of having high-risk Wolfe’s parenchymal patterns compared with never smokers when the analyses also included premenopausal women [22]. In an American study including 191 Hispanic women, Gapstur and colleagues found no association between smoking and mammographic density [23]. In another American study, Vachon and colleagues did not find any association between smoking duration and intensity and the percent mammographic density among 1,554 postmenopausal women, but an inverse association was found among the 346 premenopausal women in their study [24].

Mammographic density is an independent risk factor for breast cancer, and lower mammographic density could suggest lower risk for breast cancer [18,19]. One of the suggested mechanisms for how smoking might increases the risk of breast cancer is carcinogenic effects of constituents in tobacco smoke on the breast tissue [1-3]. It is possible that this suggested mechanism does not affect mammographic density, and thus would not influence mammographic density as measured in our study.

Conversely, the other suggested effect of smoking is believed to be an antiestrogenic effect [4,5,34]. Smoking has been shown to enhance the metabolism of estradiol to metabolites believed to have minimal peripheral estrogen activity, to increase estrogen binding by sex-hormone binding globulin and, because smokers tend to be leaner than nonsmokers, to lower the amount of estrogens derived from adipose tissue [4,5]. It has been shown that mammographic density may be intentionally manipulated by hormonal treatments such as tamoxifen or a gonadotropin-releasing hormone agonist [35,36]. A change in serum estrogen levels has also been shown to influence mammographic density [37]. The findings in the present study are consistent with an antiestrogenic effect of cigarette smoking on breast tissue, reflected in lower mammographic density among smokers.

We do not, however, believe our finding that current and former smokers have lower mean mammographic density when compared with never smokers will be reflected in a lower risk of breast cancer among smokers. The magnitudes of the difference in the mean percentage mammographic density between current smokers and never smokers were modest and may be due to residual confounding. We have previously shown that, in this study population, women using a continuous estrogen and progestin combination HT for 5 years or longer had a significantly higher percentage mammographic density compared with never HT users. The magnitude of that difference was substantially higher (7% absolute difference, 100% relative difference) than what we observed in the current study [30], suggesting that the effect of cigarette smoking may not be clinically relevant. Our finding that former smokers who had stopped smoking less than 24 years ago had statistically significantly lower mammographic density when compared with never smokers may partly be due to misclassification of current smokers as former smokers, as well as due to residual confounding. Boyd and colleagues, however, have proposed that the cumulative exposure to dense mammographic breast tissue could be represented by the area under the curve of the Pike and colleagues model for breast tissue ageing [38]. If there is an antiestrogenic effect of smoking, therefore, former smoking may have increased the age-related decrease in mammographic density.

**Conclusion**

We found inverse associations between cigarette smoking and both the percentage and absolute mammographic densities among postmenopausal women in our cross-sectional study. We also observed a modest inverse dose–response association among current smokers between the amount
smoked and the mammographic density. Also, former smokers who had stopped smoking less than 24 years ago had a statistically significantly lower mean percentage mammographic density when compared with never smokers. Our results are consistent with an antiestrogenic effect of smoking on the breast tissue, as measured by mammographic density.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
YB had full access to all data in the study, and takes responsibility for the integrity of the data and the accuracy of the data analyses. ITG conceived of the study, and had full access to all data. The classification of mammograms was performed by GU. YB, GU, NB, and ITG performed the analysis and interpretation of data. YB and ITG drafted the manuscript. Critical revision of the manuscript for important intellectual content was performed by YB, ITG, GU, and NB. All authors read and approved the final manuscript.

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References
1. Band PR, Le ND, Fang R, Deschamps M: Carcinogenic and endocrine disrupting effects of cigarette smoke and risk of breast cancer. Lancet 2002, 360:1044-1049.
2. Terry PD, Rohan TE: Cigarette smoking and the risk of breast cancer in women: a review of the literature. Cancer Epidemiol Biomarkers Prev 2002, 11:953-971.
3. Terry PD, Goodman M: Is the association between cigarette smoking and breast cancer modified by genotype? A review of epidemiologic studies and meta-analysis. Cancer Epidemiol Biomarkers Prev 2006, 15:602-611.
4. Baron JA, La VC, Levi P: The antiestrogenic effect of cigarette smoking in women. Am J Obstet Gynecol 1990, 162:502-514.
5. Tanke LB, Christiansen C: An update on the antiestrogenic effect of smoking: a literature review with implications for researchers and practitioners. Menopause 2004, 11:104-109.
6. Hamajima N, Hirose K, Tajima K, Rohan T, Calle EE, Heath CW Jr, Coates RJ, Lifit JM, Talamini R, Chantarakul N, et al.: Alcohol, tobacco and breast cancer – collaborative reanalysis of individual data from 53 epidemiological studies, including 58,515 women with breast cancer and 95,067 women without the disease. Br J Cancer 2002, 87:1234-1245.
7. Morabia A: Smoking (active and passive) and breast cancer: epidemiologic evidence up to June 2001. Environ Mol Mutagen 2002, 39:89-95.
8. Johnson KC: Accumulating evidence on passive and active smoking and breast cancer risk. Int J Cancer 2005, 117:619-628.
9. IARC: IARC Monographs on the Evaluation of Carcinogenic Risks to Humans. Tobacco Smoke and Involuntary Smoking Volume 83. Lyon: IARC Press; 2004.
10. Prescott J, Ma H, Bernstein L, Ursin G: Cigarette smoking is not associated with breast cancer risk in young women. Cancer Epidemiol Biomarkers Prev 2007, 16:620-622.
11. Terry PD, Miller AB, Rohan TE: Cigarette smoking and breast cancer risk: a long latency period? Int J Cancer 2002, 100:729-728.
12. Gram IT, Braaten T, Terry PD, Sasco AJ, Adami HO, Lund E, Weiderpass E: Breast cancer risk among women who start smoking as teenagers. Cancer Epidemiol Biomarkers Prev 2005, 14:81-86.
13. Reynolds P, Hurley S, Goldberg DE, Anton-Culver H, Bernstein L, Deapen D, Horn-Ross PL, Peel D, Pinder R, Ross RK, et al.: Active smoking, household passive smoking, and breast cancer: evidence from the California Teachers Study. J Natl Cancer Inst 2004, 96:29-97.
14. Al-Delamy WK, Cho E, Chen WY, Colditz G, Willet WC: A prospective study of smoking and risk of breast cancer in young adult women. Cancer Epidemiol Biomarkers Prev 2004, 13:398-404.
15. Egan KM, Stampfer MJ, Hunter D, Hankinson S, Rosner BA, Holmes M, Willett WC, Colditz GA: Active and passive smoking in breast cancer: prospective results from the Nurses’ Health Study. Epidemiology 2002, 13:138-145.
16. Olson JE, Vachon CM, Vierkant RA, Sweeney C, Limburg PJ, Cerhan JR, Sellers TA: Breast cancer attributable to cigarette smoking and subsequent risk of postmenopausal breast cancer. Mayo Clin Proc 2005, 80:1423-1428.
17. Ha M, Mabuchi K, Sigurdson AJ, Freedman DM, Linet MS, Doody MM, Hauptmann M: Smoking cigarettes before first childbirth and risk of breast cancer. Am J Epidemiol 2007, 166:55-66.
18. Boyd NF, Guo H, Martin LJ, Sun L, Stone J, Fishell E, Jong RA, Hislop G, Chiarelli A, Minkin S, et al.: Mammographic density and the risk and detection of breast cancer. N Engl J Med 2007, 356:227-236.
19. McCormack VA, dos Santos Silva : Breast density and parenchymal patterns as markers of breast cancer risk: a meta-analysis. Cancer Epidemiol Biomarkers Prev 2006, 15:1159-1169.
20. Yaffe M, Hendrix S, Pike M, Santen R, Eden J, Genazzani A: Is mammographic density, as currently measured, a robust surrogate marker for breast cancer? Gynecol Endocrinol 2005, 21(Suppl 1):17-21.
21. Modugno F, Ngo DL, Allen GO, Kuller LH, Ness RB, Vogel VG, Costantino JP, Cauley JA: Breast cancer risk factors and mammographic breast density in women age over 70. Breast Cancer Res Treat 2006, 97:157-166.
22. Sala E, Warren R, McCann J, Duffy S, Luben R, Day N: Smoking and high-risk mammographic parenchymal patterns: a case–control study. Breast Cancer Res 2000, 2:59-63.
23. Calangur SM, Lopez P, Colangelo LA, Wolman J, Van HL, Hendrick RE: Associations of breast cancer risk factors with breast density in Hispanic women. Cancer Epidemiol Biomarkers Prev 2003, 12:1074-1080.
24. Vachon CM, Kuni CC, Anderson K, Anderson VE, Sellers TA: Association of mammographically defined parenchymal breast density with epidemiologic risk factors for breast cancer (United States). Cancer Causes Control 2000, 11:653-662.
25. Roubidoux MA, Kaur JS, Griffith KA, Stillwater B, Novotny P, Sloan J: Relationship of mammographic parenchymal patterns to breast cancer risk factors and smoking in Alaska Native women. Cancer Epidemiol Biomarkers Prev 2003, 12:1081-1086.
26. Warwick J, Pinney E, Warren RM, Duffy SW, Howell A, Wilson M, Cuzick J: Breast density and breast cancer risk factors in a high-risk population. Breast Cancer Research 2004, 6:1041-1046.
27. Jeffreys M, Warren R, Gunnell D, McCarron P, Smith GD: Life course breast cancer risk factors and adult breast density (United Kingdom). Cancer Causes Control 2004, 15:947-955.
28. Gram IT, Bremnes Y, Ursin G, Maskarinec G, Bjurstam N, Lund E: Predicted density, Wolfe’s and Tabár’s mammographic patterns – agreement and association with breast cancer risk factors. Breast Cancer Res 2005, 7:R862-R870.
29. Ursin G, Astrahan MA, Salane M, Panisky YR, Pearce JG, Daniels JR, Pike MC, Spicer DV: The detection of changes in mammographic density. Cancer Epidemiol Biomarkers Prev 1998, 7:43-47.
30. Bremnes Y, Ursin G, Bjurstam N, Lund E, Gram IT: Different types of postmenopausal hormone therapy and mammographic density. In Norwegian women. Int J Cancer 2007, 128:880-884.
31. Hofvind S, Moller B, Thoresen S, Ursin G: Use of hormone therapy and risk of breast cancer detected at screening and

Available online http://breast-cancer-research.com/content/9/5/R73
between mammographic screens. *Int J Cancer* 2006, 118:3112-3117.

32. Kaaks R: Insulin-like growth factor-I and mammographic breast density. *Cancer Epidemiol Biomarkers Prev* 2005, 14:3019. (Letter to Editor)

33. Maskarinec G, Pagano I, Chen Z, Nagata C, Gram IT: Ethnic and geographic differences in mammographic density and their association with breast cancer incidence. *Breast Cancer Res Treat* 2007, 104:47-56.

34. Kendall A, Folkard EJ, Dowsett M: Influences on circulating oestrogens in postmenopausal women: relationship with breast cancer. *J Steroid Biochem Mol Biol* 2007, 103:99-109.

35. Cuzick J, Warwick J, Pinney E, Warren RM, Duffy SW: Tamoxifen and breast density in women at increased risk of breast cancer. *J Natl Cancer Inst* 2004, 96:621-628.

36. Gram IT, Ursin G, Spicer DV, Pike MC: Reversal of gonadotropin-releasing hormone agonist induced reductions in mammographic densities on stopping treatment. *Cancer Epidemiol Biomarkers Prev* 2001, 10:1117-1120.

37. Ursin G, Palla SL, Reboussin BA, Slone S, Wasilauskas C, Pike MC, Greendale GA: Post-treatment change in serum estrone predicts mammographic percent density changes in women who received combination estrogen and progestin in the Postmenopausal Estrogen/Progestin Interventions (PEPI) Trial. *J Clin Oncol* 2004, 22:2842-2848.

38. Boyd NF, Lockwood GA, Martin LJ, Byng JW, Yaffe MJ, Tritchler DL: Mammographic density as a marker of susceptibility to breast cancer: a hypothesis. *IARC Sci Publ* 2001, 154:163-169.