An investigation of gene-environment interactions between 47 newly identified breast cancer susceptibility loci and environmental risk factors

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Abstract

A large genotyping project within the Breast Cancer Association Consortium (BCAC) recently identified 41 associations between single nucleotide polymorphisms (SNPs) and overall breast cancer (BC) risk. We investigated whether the effects of these 41 SNPs, as well as six SNPs associated with estrogen receptor (ER) negative BC risk are modified by 13 environmental risk factors for BC.

Data from 22 studies participating in BCAC were pooled, comprising up to 26,633 cases and 30,119 controls. Interactions between SNPs and environmental factors were evaluated using an empirical Bayes-type shrinkage estimator.

Six SNPs showed interactions with associated p-values ($p_{\text{int}}$) $<1.1 \times 10^{-3}$. None of the observed interactions was significant after accounting for multiple testing. The Bayesian False Discovery Probability was used to rank the findings, which indicated three interactions as being noteworthy at 1% prior probability of interaction. SNP rs6828523 was associated with increased ER-negative BC risk in women ≥170cm (OR=1.22, $p=0.017$), but inversely associated with ER-negative BC risk in women <160cm (OR=0.83, $p=0.039$, $p_{\text{int}}=1.9 \times 10^{-4}$). The inverse association between rs4808801 and overall BC risk was stronger for women who had had four or more pregnancies (OR=0.85, $p=2.0 \times 10^{-4}$), and absent in women who had had just one (OR=0.96, $p=0.19$, $p_{\text{int}}=6.1 \times 10^{-4}$). SNP rs11242675 was inversely associated with overall BC risk in never/former smokers (OR=0.93, $p=2.8 \times 10^{-5}$), but no association was observed in current smokers (OR=1.07, $p=0.14$, $p_{\text{int}}=3.4 \times 10^{-4}$).

In conclusion, recently identified breast cancer susceptibility loci are not strongly modified by established risk factors and the observed potential interactions require confirmation in independent studies.

Keywords

gene-environment interaction; breast cancer; risk factor; genetic susceptibility

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Conflict of Interest Statement

The authors declare no conflict of interest.
Introduction

Genetic and environmental factors are known to contribute to the risk of breast cancer. The biological interplay between them may lead to varying associations of the genetic factors with breast cancer risk depending on the exposure to an environmental factor. This can be assessed as departure from multiplicativity of the risk ratios of the genetic variant and the environmental factor (gene-environment (G×E) interaction). Several studies have investigated whether the relative risks associated with common genetic breast cancer susceptibility loci are modified by environmental risk factors. In the most recent investigation of 23 single nucleotide polymorphisms (SNPs) using data from the Breast Cancer Association Consortium (BCAC), we were able to replicate a previously reported interaction between rs3817198 in LSP1 and number of full-term pregnancies in parous women, and identify an interaction between rs17468277 in CASP8 and varying levels of mean lifetime alcohol consumption (>20 g/day vs. ≥20 g/day). The identification of G×E interactions may improve our understanding of breast cancer aetiology by suggesting potential biological pathways involved.

A recently conducted large genotyping project (Collaborative Oncological Gene-environment Study (COGS)) identified 41 novel genetic susceptibility loci for breast cancer, explaining an additional 5% of the familial breast cancer risk. The project also led to the identification of four loci associated with risk of estrogen receptor (ER) negative breast cancer additional to the three previously established ER-negative breast cancer susceptibility loci. G×E interactions with these newly identified variants have not been investigated so far.

Here, we evaluated G×E interactions on overall breast cancer risk between 47 single nucleotide polymorphisms (SNPs) and the following environmental factors: age at menarche, parity, age at first birth, breastfeeding, use of menopausal hormone therapy (MHT), body-mass index (BMI), adult height, smoking and alcohol consumption. The 47 SNPs represent 41 newly identified genetic susceptibility loci for overall breast cancer as well as 6 loci associated with risk for ER negative breast cancer (genotype data for the seventh ER-negative breast cancer SNP (rs2284378) was not available). We also assessed G×E interactions regarding risk for ER-positive and ER-negative breast cancer separately, as different pathways may be involved in the development of these subtypes. This investigation uses the largest dataset available at present, including genotype data on the newly identified breast cancer susceptibility loci and comprehensive data on environmental risk factors.

Materials and Methods

Study samples

We pooled data from 22 studies participating in BCAC (20 case-control studies, 2 cohort studies), which mainly recruited participants of European descent (Supplementary Table 1). Selected studies comprised at least 200 cases and 200 controls with genotype data and information on at least one of the environmental risk factors of interest.
We excluded participants from this analysis if they were male, were prevalent cases at recruitment (in MCCS and pKARMA), were not of European descent, or had a missing value for reference age (age at diagnosis/interview), the specific environmental variable of interest, or the related adjustment variables. Therefore, the number of participants available for analysis varied depending on the investigated environmental factor. The dataset with subjects included in at least one of the analyses comprised 31,850 cases and 34,816 controls. The largest sample was available for the G×E interaction analysis between SNPs and ever being parous, which included 26,633 cases and 30,119 controls and the smallest sample was available for the analysis involving lifetime average intake of alcohol, which included 3,811 cases and 4,053 controls.

All studies were approved by the relevant ethics committees and informed consent was obtained from all participants.

**Data harmonization and variable definitions**

Data from the different studies were harmonized in a multi-step process according to a common data dictionary. In both case-control and cohort studies, time-dependent variables were assessed at reference date, which was defined as the date of diagnosis for cases and the date of interview for controls: for controls and cases from the two included cohort studies, data from the baseline interview were considered, or if available, follow-up information. The median time between last interview/questionnaire and diagnosis was 7.6 years in the MCCS cohort and 2.0 years in the UKBGS cohort.

Current use of any MHT was defined as use within 6 months prior to the reference date and current smoking as smoking within one year prior to the reference date. An age surrogate was used to define menopausal status. Women aged ≤54 years at reference date were considered to be premenopausal and women aged > 54 years postmenopausal. BMI was calculated based on usual adult weight or weight one year prior to the reference date (studies ABCFS, BREOGAN, CECILE, GENICA, kConFab/AOCS, KBCP, MARIE, MCBCS, OFBCR, PBCS, SASBAC) or weight in early adulthood (age around 20 years, studies ESTHER, pKARMA, SEARCH). For the two cohort studies (MCCS, UKBGS), we used the weight reported at baseline interview.

**Genetic information**

The genotyping data used in this study for all studies except BREOGAN were generated as part of the COGS project (www.nature.com/icogs). Participants from studies in BCAC were genotyped using an Illumina iSelect array (iCOGS). Approximately 61,000 of the 211,155 SNPs included on the iCOGS array were selected to follow-up on a meta-analysis of nine breast cancer genome-wide association studies (GWAS). A subsequent association study in 45,290 cases and 41,880 controls led to the identification of 41 SNPs associated with overall breast cancer risk. Similarly, three GWAS studies were meta-analysed to identify loci associated with ER-negative breast cancer risk, and 13,276 SNPs were selected to be genotyped on the iCOGS array for the replication stage, comprising 6,514 ER-negative breast cancer cases and 41,455 controls. Four new loci showing a specific association with ER-negative breast cancer were detected.

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In the current study, we use the original quality-controlled genotype data that was used for identification of the 41 SNPs and the four SNPs associated with ER-negative breast cancer risk, for all studies except for subjects in BREOGAN. Two of the three previously identified SNPs specifically associated with ER-negative breast cancer risk were also genotyped with the iCOGS array (rs10069690 in TERT on chromosome 5, rs8170 in BAMB1 on chromosome 19); genotype data for rs2284378 on chromosome 20q11 were not available.

Study participants were excluded from analyses if the overall genotyping call rate was below 95% over the whole iCOGS array or if heterozygosity deviated significantly from that expected in the general population (either lower or higher, \( p < 10^{-6} \)).

Genotyping of the 47 SNPs for BREOGAN was performed at the CeGen-ISCIII (Spanish National Genotyping Center), using Sequenom MassARRAY Genotyping system (technology iPLEX GOLD) and following the manufacturer’s instructions. DNA was dispensed in 384 well plates by a Tecan Freedom Evo robot, each plate included case and control samples, a trio of Coriell samples: Na10830, Na10831 and Na12147, and negative controls (minimum 6 per plate). We included >5% concordant duplicates. The laboratory was equipped with Life Technologies GeneAmp 9700 dual cyclers, a RS1000 Nanodispenser and a MA4 mass spectrometer. Data analysis was done using the software Typer analyzer v4.0.20. The SNPs were analyzed in 4 assays (Assay Design v4 software). The genotyping data were quality checked using the same criteria as for iCOGS.

To evaluate potential functional implications of selected SNPs and SNPs in high linkage disequilibrium (LD) with selected SNPs we used HaploReg v2.11 and the UCSC genome browser12.

**Statistical analysis**

We employed an efficient empirical Bayes procedure to calculate the interaction log odds ratio that corresponds to a weighted average of the case-only and case-control estimators. In this way, the method makes use of the greater precision of the case-only estimator by simultaneously reducing the chance of generating biased estimates due to violations of the assumption of gene-environment independence in controls13. The method is implemented in the R package “CGEN”, version 2.2, which was used within R 2.15.2.

In total, 13 variables representing the environmental risk factors of interest were used in GxE analyses (Supplementary Figure 1). The variables were: age at menarche (per 2 years), ever parous (no vs. yes), number of full-term pregnancies (among parous, 1, 2, 3, ≥4 pregnancies), ever breastfed (yes vs. no), age at first full-term pregnancy (per 5 years), adult BMI in premenopausal women (per 5 kg/m^2), adult BMI in postmenopausal women (per 5 kg/m^2), adult height (per 5 cm), current use of combined estrogen-progesterone therapy (no vs. yes), current use of estrogen-only therapy (no vs. yes), lifetime average intake of alcohol (per 10 g/day), current smoking (no vs. yes), smoking amount (per 10 pack-years). The variables age at menarche, number of full-term pregnancies, age at first full-term pregnancy, adult BMI, adult height, lifetime intake of alcohol, and smoking amount were entered into models as linear continuous variables. For SNPs, we included the number of minor alleles.
(0-1-2) as a continuous variable. Subjects with missing data for a particular SNP or an environmental factor were excluded from the respective analysis.

The models were adjusted for study, reference age and seven principal components to account for population substructure. Additionally, to account for potential differential main effects of environmental variables by study design, we included in all models an interaction term between the environmental variable of interest and an indicator variable for study design (non-population-based vs. population-based).

MHT was classified into estrogen-only therapy and estrogen-progesterone therapy. Models used to assess associations with current use of the type of MHT of interest were further adjusted for former use of any MHT and use of MHT preparations other than the one of interest, and the analysis was restricted to postmenopausal women. Also, when assessing G×E interactions with adult BMI in postmenopausal women, the study sample was restricted to never or former users of any MHT.

All analyses were conducted with overall breast cancer risk as the outcome, as well as with ER-negative and ER-positive breast cancer risk. Heterogeneity between risk associations for ER-negative and ER-positive breast cancer was evaluated using case-case analysis with ER-status as the dependent variable and the SNP, the environmental variable, the multiplicative interaction term, ancestry informative principal components and study as independent variables. The association between SNP and breast cancer risk in strata defined by categories of the environmental risk factor was evaluated using logistic regression. Stratified analyses were conducted using SAS 9.2.

To evaluate between-study heterogeneity in G×E interaction OR estimates, we calculated these by study and performed Cochrane’s Q-test and calculated the \( I^2 \) index, using the R package “meta” (version 2.2).

We selected G×E interactions showing \( p \)-values for interaction \(<1.1\times10^{-3} \) for overall breast cancer (all subtypes combined) or a subtype of breast cancer, but in the latter case, only if significant subtype heterogeneity (\( p \)-values for heterogeneity between ER-positive and ER-negative disease \(<0.05 \)) was also observed. The \( p \)-value threshold for selection was derived by dividing the conventional \( p \)-value threshold of 0.05 by the number of SNPs investigated.

To account for chance findings due to multiple hypothesis testing, we applied the Bayesian False Discovery Probability (BFDP)\(^{14} \) to assess noteworthiness of selected G×E interactions in terms of generating new hypotheses. We assumed a four-fold cost of a false non-discovery compared to the cost of a false discovery, considering interactions with a BFDP of less than 80% as being noteworthy, as suggested by Wakefield et al.\(^{14} \). The OR corresponding to the 97.5% point of the prior was 1.50 for positive G×E interactions and 0.66 for negative G×E interactions, i.e. we assumed that the prior probability of observing an OR for interaction larger than 1.5 or smaller than 0.66 was 5%. We calculated the BFDP for each selected interaction assuming six different prior probabilities for true interaction (20%, 10%, 5%, 1%, 0.1% and 0.01%).
Results

A brief description of the BCAC studies included in this analysis of GxE interactions is provided in Supplementary Table 1. The number of included cases and controls as well as the mean reference age for each study is shown in Table 1. Overall, the mean age was 56.7 years for cases and 55.6 years for controls. Further descriptions of the environmental variables are displayed in Supplementary Table 2.

The associations between SNPs and breast cancer risk were very similar in the study sample used for GxE interaction analysis (N = 66,666) to those reported by Michailidou et al. (sample size N = 87,170)\(^6\) (Supplementary Table 3). The largest difference was observed for rs11814448, which was previously reported to be associated with overall breast cancer with an OR of 1.26 (95% CI 1.18 – 1.35) and showed a slightly attenuated effect size in the GxE dataset (odds ratio (OR) = 1.21, 95% confidence interval (CI) 1.13 – 1.31). Supplementary Table 3 shows further information for each SNP such as the minor allele frequency and SNP location.

Forest plots for meta-analyses of the associations between the 13 environmental risk factors and breast cancer risk by study can be found in Supplementary Figure 1. The risk factor associations based on the population-based studies were consistent with previous reports. Age at menarche, ever been parous, and number of full-term pregnancies among parous women were significantly associated with a decreased breast cancer risk. Significant associations with increased breast cancer risk were observed for breastfeeding (among parous women, no vs. yes), age at first full-term pregnancy, BMI in postmenopausal women not currently using MHT, body height, current use of postmenopausal combined estrogen-progesterone therapy, and average lifetime intake of alcohol. No significant associations were found between breast cancer risk and BMI in premenopausal women, current use of estrogen-only therapy, current smoking and smoking amount (pack-years).

We identified six GxE interactions with \(p\)-values for interaction \((p_{\text{int}}) < 1.1\times10^{-3}\) (Figure 1). Estimates for each investigated GxE interaction and ORs for association between SNP and breast cancer stratified by categories of the environmental factors are presented in Supplementary Table 4. Estimates from empirical Bayes and case-control analysis (data not shown) were very similar, but \(p\)-values for interaction from empirical Bayes analysis were usually more extreme, possibly reflecting a small gain in power. Three of the six interactions were considered noteworthy according to a BFDP <80% at a 1% prior probability of interaction (Table 2). However, none of the observed interactions was noteworthy by this criterion assuming more conservative prior probabilities for interaction <1%.

The interaction with the lowest BFDP (BFDP = 36.0% at 1% prior probability of interaction) was observed regarding ER-negative breast cancer risk, between the SNP rs6828523 located in an intron of ADAM29 and adult height (ER-negative OR for interaction \((OR_{\text{int}}) = 1.14, 95\% \text{ CI } 1.06 – 1.22, p_{\text{int}} = 1.9\times10^{-4}\)). The interaction was not observed for ER-positive breast cancer risk \((OR_{\text{int}} = 1.00, 95\% \text{ CI } 0.96 – 1.03, p_{\text{int}} = 9.0\times10^{-1}, p\)-value for heterogeneity by ER status \((p_{\text{het}}) = 0.003\). SNP rs6828523 was associated with increased risk for ER-negative breast cancer in women of 170cm height or taller (ER-
negative OR = 1.22, 95% CI 1.04 – 1.44, \( p = 0.017 \)), but showed an inverse association in women shorter than 160cm (ER-negative OR = 0.83, 95% CI 0.70 – 0.99, \( p = 0.039 \)) (Figure 1A). The five additional G\times E interactions are reported below, ordered by their corresponding BFDP, as reported in Table 2.

Regarding overall breast cancer risk, an interaction between the number of full-term pregnancies and rs4808801 was observed (\( \text{OR}_{\text{int}} = 0.96, 95\% \text{ CI} 0.94 – 0.98, p_{\text{int}} = 6.1\times10^{-4}, \text{BFDP} = 51.6\% \) at 1% prior probability of interaction). The interaction did not differ by ER status (\( p_{\text{het}} = 0.40 \)) (Supplementary Table 5). The SNP is located in an intron of \( \text{ELL} \) on chromosome 19. The association between breast cancer risk and rs4808801 was stronger in women with four or more full-term pregnancies (\( \text{OR} = 0.85, 95\% \text{ CI} 0.77 – 0.93, p = 2.0\times10^{-4} \)), and weaker in women with one full-term pregnancy (\( \text{OR} = 0.96, 95\% \text{ CI} 0.90 – 1.02, p = 0.19 \)) (Figure 1B).

Another interaction on overall breast cancer risk was found between current smoking and rs11242675, located on chromosome 6 near \( \text{FOXQ1} \) (\( \text{OR}_{\text{int}} = 1.13, 95\% \text{ CI} 1.06 – 1.21, p_{\text{int}} = 3.4\times10^{-4}, \text{BFDP} = 60.5\% \) at 1% prior probability of interaction). Again, the interaction was not substantially different for ER-negative and ER-positive breast cancer (\( p_{\text{het}} = 0.82 \)) (Supplementary Table 5). As shown in Figure 1C, rs11242675 was associated with a decreased breast cancer risk in women who did not smoke at reference time (\( \text{OR} = 0.93, 95\% \text{ CI} 0.89 – 0.96, p = 2.8\times10^{-5} \)), but this association was not observed in women who smoked at reference time (\( \text{OR} = 1.07, 95\% \text{ CI} 0.98 – 1.16, p = 0.14 \)).

The three remaining interactions of the six G\times E interactions in total with \( p_{\text{int}} <1.1\times10^{-3} \) could not be considered noteworthy according to their BFDP estimated using a prior probability of interaction of 1% or lower (Table 2). One of the interactions was observed for ER-positive breast cancer risk, between rs16857609 and adult height (ER-positive \( \text{OR}_{\text{int}} = 0.95, 95\% \text{ CI} 0.93 – 0.98, p_{\text{int}} = 1.7\times10^{-4}, p_{\text{het}} = 0.018; \) Supplementary Table 5). The variant rs16857609 is located in an intron of \( \text{DIRC3} \) on chromosome 2. In the stratified analysis, rs16857609 was associated with an increased risk of estrogen receptor positive breast cancer in women shorter than 160cm (ER-positive \( \text{OR} = 1.15, 95\% \text{ CI} 1.07 – 1.23, p = 2.0\times10^{-4} \)), whereas it was not associated with breast cancer risk in women of 170cm height or taller (ER-positive \( \text{OR} = 0.97, 95\% \text{ CI} 0.90 – 1.04, p = 0.40 \)) (Figure 1D).

Two further G\times E interactions were observed specifically for ER-negative breast cancer risk, one between rs12422552 located on chromosome 12 and adult height (ER-negative \( \text{OR}_{\text{int}} = 1.09, 95\% \text{ CI} 1.04 – 1.15, p_{\text{int}} = 7.4\times10^{-4}, p_{\text{het}} = 0.006; \) Supplementary Table 5). The minor allele of rs12422552 was associated with risk for ER-negative breast cancer in women of 170cm height or taller (ER-negative \( \text{OR} = 1.18, 95\% \text{ CI} 1.04 – 1.34, p = 0.011 \)), but not in women shorter than 160cm (ER-negative \( \text{OR} = 0.92, 95\% \text{ CI} 0.81 – 1.04, p = 0.16 \)) (Figure 1E). The other interaction specific for ER-negative breast cancer risk was between rs941764 located in an intron of \( \text{CCDC88C} \) on chromosome 14 and alcohol consumption (ER-negative \( \text{OR}_{\text{int}} = 0.53, 95\% \text{ CI} 0.36 – 0.76, p_{\text{int}} = 6.8\times10^{-4}, p_{\text{het}} = 0.042; \) Figure 1F, Supplementary Table 5). As shown in Figure 4F, rs941764 was inversely associated with risk of ER-negative breast cancer risk in women having a lifetime average consumption of at least 20 g alcohol per day (ER-negative \( \text{OR} = 0.61, 95\% \text{ CI} 0.38 – 0.97, p = 0.037 \)), while...
this association was not present in women with a lower lifetime average consumption of alcohol (ER-negative OR = 0.96, 95% CI 0.84 – 1.10, \( p = 0.59 \)).

There was no significant heterogeneity between study-wise estimates for G\(\times\)E interactions: \( p \)-values from \( Q \)-test ranged from 0.36 to 0.78 (Supplementary Figure 2).

**Discussion**

The present study identified six G\(\times\)E interactions with \( p_{\text{int}} < 1.1 \times 10^{-3} \), two regarding risk for overall breast cancer, one regarding risk for ER-positive breast cancer and three regarding risk for ER-negative breast cancer. After calculating the BFDP, none of the six interactions could be considered as being noteworthy at prior probabilities for interaction smaller than one percent although three G\(\times\)E interactions were considered noteworthy at 1% prior probability of interaction. Our results do not suggest that the relative risks associated with 47 recently identified breast cancer susceptibility loci are strongly modified by environmental risk factors for breast cancer.

For some the effect modifications assessed, our findings are based on the largest available dataset at present. The number of studies with available data was relatively small for other environmental risk factors such as alcohol consumption and use of MHT. Power was also likely diminished due to the fact that we studied mostly tag-SNPs, rather the true genetic variants affecting breast cancer risk. The power was even further reduced when looking at subtype specific associations, especially for ER-negative breast cancer risk. Although the environmental data of the contributing studies were harmonized in a standardized fashion, we still observed heterogeneity in marginal effect associations with breast cancer risk (Supplementary Figure 1). Associations were less heterogeneous between population-based studies, and we included an interaction term between study design and the environmental variable in the models to account for potentially biased estimates from non-population-based studies. The assessment of associations between environmental factors and breast cancer risk was restricted to population-based studies and the estimates were comparable to those reported in the literature\(^{15–22} \). The association was not significant for BMI in premenopausal women, which may in part be attributed to the small sample available when using only population-based studies, however the direction of association was as expected. Another limitation of our study was that the sample consisted primarily of case-control studies and comprised only two cohort studies. While case-control studies have the advantage of being able to assess exposure close to the reference date, for example, for current MHT use, the retrospective assessment of exposure is prone to recall bias. However, we did not observe any heterogeneity between study-wise estimates for G\(\times\)E interactions. Also, G\(\times\)E interaction estimates derived from the whole study sample and from a sensitivity analysis restricted to population-based studies were similar (Supplementary Table 6). The robustness of our findings is also supported by the fact that, given reasonable assumptions, selection bias is unlikely to influence the assessment of multiplicative G\(\times\)E interactions\(^{23} \). Also, both non-differential and differential misclassifications of environmental risk factors would lead to a reduction in power rather than increasing the probability of a spurious finding of an interaction\(^{24} \). The magnitude of the interactions for which strongest evidence was observed was comparable to those previously reported between breast cancer risk SNPs and

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environmental factors\(^1\). When taking into account the number of tests performed, the identified G×E interactions were not statistically significant and further evidence is needed for confirmation. However, not all of the performed tests can be considered independent as we looked at different variables that are highly correlated (e.g. parity and number of full term pregnancies) and also tests for interaction concerning all cases and subgroups of cases defined by ER status are related. We therefore calculated the BFDP to be able to rate the noteworthiness of the observed G×E interactions.

It should be noted, that the investigated susceptibility loci have been identified in a sample of European descent, and that this investigation of gene-environment interaction was also restricted to subjects with European ancestry. The potential gene-environment interactions detected here do not necessarily have to be present in study populations of different ancestry due to the varying genetic structure and possible different prevalence of risk factors.

The interaction with the lowest BFDP was found between rs6828523 and adult height on ER-negative breast cancer risk. The SNP rs6828523 itself was associated with a decreased risk of ER-positive breast cancer, but not for ER-negative breast cancer, showing significant heterogeneity by ER status in the analysis identifying the variant \((p_{\text{het}} = 1.2 \times 10^{-7})\), and also in sample analysed here \((p_{\text{het}} = 9.5 \times 10^{-6})\). SNP rs6828523 showed a positive association with ER-negative breast cancer risk in women taller than 164 cm (the median height in the study sample) (ER-negative OR = 1.13, 95% CI 1.01 – 1.26, \(p = 0.036\)). Current evidence suggests that adult body height is a risk factor for both ER-positive and ER-negative breast cancer, although the estimates for ER-negative breast cancer are not entirely consistent across studies\(^{25–29}\). The variant rs6828523 is located in an intron of ADAM29. The potentially functional implications of rs6828523 or SNPs highly correlated with rs6828523 \((r^2>0.6)\) are unclear as they are not located within any strong regulatory elements (Supplementary Figure 3). Also, a more comprehensive investigation of the functional effects of the 41 SNPs associated with overall breast cancer risk did not identify a SNP in LD with rs6828523 coinciding with a regulatory genomic feature\(^{30}\). ADAM29 encodes a disintegrin-metalloproteinase. Metalloproteinases are involved in the modification of the extracellular matrix and growth factor bioavailability, and changes in expression of metalloproteinases have been linked to breast cancer progression\(^{31}\). It is unclear however, how factors involved in growth and adult height might interplay with variants in ADAM29 to influence risk of ER-negative breast cancer.

The association of rs4808801 located on chromosome 19 in an intron of ELL with overall breast cancer risk appeared to vary according to the number of full-term pregnancies in parous women. Risk of breast cancer associated with the SNP decreased with an increasing number of pregnancies. Several SNPs in LD with rs4808801 \((r^2 > 0.6)\) are located in regulatory regions (enhancer elements, DNase hypersensitive sites, transcription factor binding sites) in the proximity of ELL and two closely-located genes, SSBP4 and ISYNA1 (Supplementary Figure 3). Three SNPs in LD with rs4808801 \((r^2 \geq 0.9)\) are located in exons of SSBP4 (rs10405636) and ISYNA1 (rs2303697, rs4595905), and all result in synonymous codon changes. ELL encodes the eleven-nineteen lysine-rich leukaemia protein, which was first identified as part of a fusion gene MLL-ELL in acute myeloid leukaemia cells, caused by a t(11;19)(q23;p13.1) translocation\(^{32}\). ELL is part of the super elongation complex, an
important regulator of transcriptional elongation. Furthermore, ELL has been found to be essential for the transcription of rapidly induced genes, and therefore plays a key role in quick responses to environmental changes. Rhie et al. identified another SNP (rs2303696) in LD with rs4808801 ($r^2 = 0.79$) located in the promoter region of ISYNA1, and likely to affect ISYNA1 expression. ISYNA1 encodes an inositol-3-phosphate synthase enzyme that catalyses the synthesis of inositol 1-phosphate from glucose 6-phosphate. ISYNA1 expression has been found to be reduced in breast cancer. Inositol containing compounds are involved in many biological processes and act as essential second messenger molecules in signalling pathways, as components of cellular membranes and regulators of chromatin remodelling. Less is known about the role of SSBP4. SSBP4 is a putative tumour suppressor, as chromosomal regions containing members of the SSBP gene family are often found to be deleted in solid tumours. How biological changes associated with multiple pregnancies in women potentially interplay with rs4808801 to influence its association with breast cancer risk is unknown.

We also observed that current smoking may modify the risk associated with rs11242675, located in close proximity to FOXQ1 on chromosome 6. Two SNPs in LD with rs11242675 are located in enhancer regions. FOXQ1 is a transcription factor, which has been found to be involved in the epithelial-mesenchymal transition of tumour cells, a process initiating metastasis. Overexpression of FOXQ1 was observed in colorectal cancer and metastatic breast cancer cell lines and a subsequent study suggested that FOXQ1 overexpression is caused by aberrant Wnt signalling. A potential biological implication of the interaction between current smoking and rs11242675 is suggested by the observation that cigarette smoking deregulates nitric oxide synthesis, which in turn may decrease the expression of the Wnt/β-catenin regulator Dickkopf-1 (DKK1) and release Wnt signalling.

This is the first evaluation of multiplicative GxE interactions between these 47 newly identified breast cancer susceptibility loci and environmental risk factors. For most of the investigated pairs of SNPs and environmental factors, there was no indication of multiplicative GxE interaction. However, despite the overall very large study sample, we cannot exclude the existence of real GxE interactions of smaller magnitude with some environmental risk factors, for which power in this study was still limited. The six potential interactions identified are largely hypothesis generating and have to be confirmed in independent studies of sufficient size. Overall, our study does not suggest that the associations between recently identified breast cancer susceptibility loci and breast cancer risk are strongly modified by environmental risk factors.

**Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

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Laboratory Medicine Program, University Health Network, Toronto, Ontario, Canada
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Peter MacCallum Cancer Center, Melbourne, Australia
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The GENICA Network: Dr. Margarete Fischer-Bosch-Institute of Clinical Pharmacology, Stuttgart, and University of Tübingen, Germany; Department of Internal Medicine, Evangelische Kliniken Bonn gGmbH, Johanniter Krankenhaus, Bonn, Germany, Institute of Pathology, University of Bonn, Germany, Molecular Genetics of Breast Cancer, Deutsches Krebsforschungszentrum (DKFZ), Heidelberg, Germany, and Institute for Prevention and Occupational Medicine of the German Social Accident Insurance, Institute of the Ruhr University Bochum (IPA), Bochum, Germany; Institute of Occupational Medicine and Maritime Medicine, University Medical Center Hamburg-Eppendorf, Germany
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Department of Molecular Genetics, University of Toronto, Toronto, Ontario, Canada
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Breakthrough Breast Cancer Research Centre, Division of Breast Cancer Research, The Institute of Cancer Research, London, UK

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**Abbreviations**

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>BCAC</td>
<td>Breast Cancer Association Consortium</td>
</tr>
<tr>
<td>BFDP</td>
<td>Bayesian False Discovery Probability</td>
</tr>
<tr>
<td>BMI</td>
<td>body-mass index</td>
</tr>
<tr>
<td>CI</td>
<td>confidence interval</td>
</tr>
<tr>
<td>ER</td>
<td>estrogen receptor</td>
</tr>
<tr>
<td>G×E</td>
<td>gene-environment</td>
</tr>
<tr>
<td>GWAS</td>
<td>genome-wide association studies</td>
</tr>
<tr>
<td>MHT</td>
<td>menopausal hormone therapy</td>
</tr>
<tr>
<td>OR</td>
<td>odds ratio</td>
</tr>
<tr>
<td>pint</td>
<td>p-value for interaction</td>
</tr>
<tr>
<td>phet</td>
<td>p-value for heterogeneity</td>
</tr>
<tr>
<td>SNP</td>
<td>single nucleotide polymorphism</td>
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**References**


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Novelty and Impact Statement

Relative risks associated with 47 recently identified susceptibility loci for overall or estrogen receptor negative breast cancer may vary depending on exposure levels of environmental (non-genetic) risk factors. In this study, gene-environment interactions between these 47 single nucleotide polymorphisms and 13 established environmental risk factors were investigated. Relative risks of breast cancer associated with the susceptibility loci were not strongly modified by environmental risk factors. This finding may have important implications for risk prediction.
Figure 1.
Odds ratios and 95% confidence intervals for association between SNP and overall breast cancer (B, C), estrogen receptor positive breast cancer (D), and estrogen receptor negative breast cancer (A, E, F) stratified by categories of environmental factors.
### Table 1

List of participating studies and number of Caucasian subjects included in at least one GxE analysis

<table>
<thead>
<tr>
<th>Study acronym</th>
<th>Study Name</th>
<th>Country</th>
<th>Design category</th>
<th>Cases</th>
<th>ER+ cases</th>
<th>ER− cases</th>
<th>Controls</th>
<th>Mean age (sd) cases</th>
<th>Mean age (sd) controls</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABCFS</td>
<td>Australian Breast Cancer Family Study</td>
<td>Australia</td>
<td>Population-based</td>
<td>790</td>
<td>456</td>
<td>261</td>
<td>551</td>
<td>40.0 (6.7)</td>
<td>42.4 (9.3)</td>
</tr>
<tr>
<td>ABCS</td>
<td>Amsterdam Breast Cancer Study</td>
<td>Netherlands</td>
<td>Mixed</td>
<td>1143</td>
<td>420</td>
<td>152</td>
<td>1177</td>
<td>43.4 (8.7)</td>
<td>48.1 (12.2)</td>
</tr>
<tr>
<td>BBCC</td>
<td>Bavarian Breast Cancer Cases and Controls</td>
<td>Germany</td>
<td>Mixed</td>
<td>554</td>
<td>456</td>
<td>82</td>
<td>458</td>
<td>61.2 (12.1)</td>
<td>57.6 (10.9)</td>
</tr>
<tr>
<td>BREOGAN</td>
<td>Breast Oncology Galicia Network</td>
<td>Spain</td>
<td>Mixed</td>
<td>1216</td>
<td>819</td>
<td>194</td>
<td>1806</td>
<td>56.9 (12.4)</td>
<td>43.3 (14.7)</td>
</tr>
<tr>
<td>CECILE</td>
<td>CECILE Breast cancer study</td>
<td>France</td>
<td>Population-based</td>
<td>900</td>
<td>743</td>
<td>130</td>
<td>999</td>
<td>54.8 (10.8)</td>
<td>55.3 (11)</td>
</tr>
<tr>
<td>CGPS</td>
<td>Copenhagen General Population Study</td>
<td>Denmark</td>
<td>Mixed</td>
<td>2811</td>
<td>1919</td>
<td>357</td>
<td>4626</td>
<td>62.3 (12.4)</td>
<td>58.4 (15.5)</td>
</tr>
<tr>
<td>CNIO-BCS</td>
<td>Spanish National Cancer Centre Breast Cancer Study</td>
<td>Spain</td>
<td>Mixed</td>
<td>704</td>
<td>213</td>
<td>76</td>
<td>834</td>
<td>54.7 (11.8)</td>
<td>50.5 (11.1)</td>
</tr>
<tr>
<td>ESTHER</td>
<td>ESTHER Breast Cancer Study</td>
<td>Germany</td>
<td>Population-based</td>
<td>471</td>
<td>302</td>
<td>98</td>
<td>502</td>
<td>61.0 (8.9)</td>
<td>62.5 (7.2)</td>
</tr>
<tr>
<td>GENICA</td>
<td>Gene Environment Interaction &amp; Breast Cancer in Germany</td>
<td>Germany</td>
<td>Population-based</td>
<td>465</td>
<td>328</td>
<td>119</td>
<td>427</td>
<td>57.5 (10.9)</td>
<td>57.8 (11.8)</td>
</tr>
<tr>
<td>KBCP</td>
<td>Kuopio Breast Cancer Project</td>
<td>Finland</td>
<td>Population-based</td>
<td>410</td>
<td>288</td>
<td>89</td>
<td>250</td>
<td>59.4 (14.5)</td>
<td>52.8 (11.6)</td>
</tr>
<tr>
<td>kConFab/AOCS</td>
<td>Kathleen Cuningham Foundation Consortium for Research into Familial Breast Cancer/Australian Ovarian Cancer Study</td>
<td>Australia</td>
<td>Mixed</td>
<td>410</td>
<td>135</td>
<td>50</td>
<td>897</td>
<td>46.2 (9.5)</td>
<td>58.3 (11.2)</td>
</tr>
<tr>
<td>LMBC</td>
<td>Leuven Multidisciplinary Breast Centre</td>
<td>Belgium</td>
<td>Mixed</td>
<td>2522</td>
<td>2069</td>
<td>378</td>
<td>1386</td>
<td>57.1 (12.4)</td>
<td>57.1 (9.1)</td>
</tr>
<tr>
<td>MARIE</td>
<td>Mammary Carcinoma Risk Factor Investigation</td>
<td>Germany</td>
<td>Population-based</td>
<td>1656</td>
<td>1279</td>
<td>370</td>
<td>1777</td>
<td>62.8 (6.3)</td>
<td>62.3 (6.1)</td>
</tr>
<tr>
<td>MCBCS</td>
<td>Mayo Clinic Breast Cancer Study</td>
<td>USA</td>
<td>Mixed</td>
<td>1546</td>
<td>1271</td>
<td>250</td>
<td>1931</td>
<td>57.5 (12.6)</td>
<td>57.1 (14)</td>
</tr>
<tr>
<td>MCCS</td>
<td>Melbourne Collaborative Cohort Study</td>
<td>Australia</td>
<td>Population-based</td>
<td>454</td>
<td>330</td>
<td>110</td>
<td>511</td>
<td>64.3 (8.7)</td>
<td>56.8 (8.3)</td>
</tr>
<tr>
<td>OFBCR</td>
<td>Ontario Familial Breast Cancer Registry</td>
<td>Canada</td>
<td>Mixed</td>
<td>1157</td>
<td>629</td>
<td>267</td>
<td>511</td>
<td>53.5 (10.2)</td>
<td>52.4 (9.2)</td>
</tr>
<tr>
<td>PBCS</td>
<td>NCI Polish Breast Cancer Study</td>
<td>Poland</td>
<td>Population-based</td>
<td>519</td>
<td>519</td>
<td>0</td>
<td>424</td>
<td>56.8 (10)</td>
<td>56.9 (9.8)</td>
</tr>
<tr>
<td>pKARMA</td>
<td>Karolinska Mammography Project for Risk Prediction of Breast Cancer - prevalent cases</td>
<td>Sweden</td>
<td>Mixed</td>
<td>2700</td>
<td>2238</td>
<td>387</td>
<td>5529</td>
<td>59.3 (10.5)</td>
<td>53.9 (9.5)</td>
</tr>
<tr>
<td>SASBAC</td>
<td>Singapore and Sweden Breast Cancer Study</td>
<td>Sweden</td>
<td>Population-based</td>
<td>1163</td>
<td>663</td>
<td>144</td>
<td>1378</td>
<td>63.6 (6.5)</td>
<td>63.8 (6.4)</td>
</tr>
<tr>
<td>SBBCS</td>
<td>Sheffield Breast Cancer Study</td>
<td>UK</td>
<td>Mixed</td>
<td>751</td>
<td>358</td>
<td>104</td>
<td>848</td>
<td>60.0 (12.4)</td>
<td>58.0 (5.8)</td>
</tr>
<tr>
<td>SEARCH</td>
<td>Study of Epidemiology and Risk factors in Cancer Heredity</td>
<td>UK</td>
<td>Mixed</td>
<td>9095</td>
<td>5130</td>
<td>1170</td>
<td>8064</td>
<td>54.9 (9.2)</td>
<td>58.2 (8.6)</td>
</tr>
<tr>
<td>UKBGS</td>
<td>UK Breakthrough Generations Study</td>
<td>UK</td>
<td>Population-based</td>
<td>413</td>
<td>88</td>
<td>18</td>
<td>470</td>
<td>56.8 (10.2)</td>
<td>54.7 (10)</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td></td>
<td></td>
<td></td>
<td>31850</td>
<td>20653</td>
<td>4806</td>
<td>34816</td>
<td>56.7 (11.5)</td>
<td>55.6 (12)</td>
</tr>
</tbody>
</table>

1Population-based design was defined as recruiting a random sample of all cases occurring in a geographically defined population during a specified period of time, and recruiting controls that were a random sample of the same source population as cases during the same period of time. Mixed design was defined as not strictly population-based or hospital-based.
## Table 2

Bayesian False Discovery Probability of G×E interactions showing p-value for interaction <1.1×10^-3

<table>
<thead>
<tr>
<th>Breast cancer subtype</th>
<th>Environmental Factor × SNP (locus)</th>
<th>ORint (95% CI)</th>
<th>Assumed 95% probability range interaction OR</th>
<th>BFDP Prior of βGE ≠ 0</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.2</td>
</tr>
<tr>
<td>ER-negative</td>
<td>Adult height × rs6828523 (ADAM29)</td>
<td>1.14 (1.06 – 1.22)</td>
<td>0.66–1.50</td>
<td>0.022</td>
</tr>
<tr>
<td>Overall</td>
<td>Number of full-term pregnancies × rs4808801 (ELL)</td>
<td>0.96 (0.94 – 0.98)</td>
<td>0.66–1.50</td>
<td>0.041</td>
</tr>
<tr>
<td>Overall</td>
<td>Current smoking × rs11242675 (4kb 3’ of FOXQ1)</td>
<td>1.13 (1.06 – 1.21)</td>
<td>0.66–1.50</td>
<td>0.058</td>
</tr>
<tr>
<td>ER-negative</td>
<td>Alcohol intake × rs941764 (CCDC88C)</td>
<td>0.53 (0.36 – 0.76)</td>
<td>0.66–1.50</td>
<td>0.177</td>
</tr>
<tr>
<td>ER-negative</td>
<td>Adult height × rs12422552 (105kb 5’ of ATF7IP)</td>
<td>1.09 (1.04 – 1.15)</td>
<td>0.66–1.50</td>
<td>0.188</td>
</tr>
<tr>
<td>ER-positive</td>
<td>Adult height × rs16857609 (DIRC3)</td>
<td>0.95 (0.93 – 0.98)</td>
<td>0.66–1.50</td>
<td>0.224</td>
</tr>
</tbody>
</table>

1 Adjusted for reference age, study, principal components to adjust for population stratification and an interaction term between environmental factor and study design (population-based vs. non-population-based). Model used to assess association with current smoking have been adjusted for former smoking.

2 To calculate the BFDP it was assumed that with probability 0.95, the interaction OR lies within the given range.

ER: estrogen receptor; OR\textsubscript{int}: odds ratio for interaction; CI: confidence interval; BFDP: Bayesian False Discovery Probability