

Fine scale mapping of the 5q11.2 breast cancer locus reveals at least three independent risk variants regulating *MAP3K1*.

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Abstract

Genome Wide Association Studies (GWAS) revealed SNP rs889312 on 5q11.2 to be associated with breast cancer risk in women of European ancestry. In an attempt to identify the biologically relevant variants, we analysed 909 genetic variants across the 5q11.2 locus in 103,991 breast cancer cases and controls from 52 studies in the Breast Cancer Association Consortium. Multiple logistic regression analyses identified three independent risk signals: The strongest associations are with 15 correlated variants (iCHAV1) where the minor allele of best candidate, rs62355902, associates with significantly increased risks of both estrogen receptor-positive (ER+: OR=1.24, 95% CI 1.21-1.27; P -trend= 5.7×10^{-44}) and estrogen receptor-negative tumors (ER-: OR=1.10, 95% CI 1.05-1.15; P -trend= 3.0×10^{-4}). After adjustment for rs62355902, we found evidence for the association of a further 173 variants (iCHAV2) containing three subsets with a range of effects, of which the strongest is SNP rs113317823 (P -cond= 1.61×10^{-5}); and five variants comprising iCHAV3 (lead rs11949391; ER+: OR=0.90, 95% CI 0.87-0.93; P -cond= 1.4×10^{-4}). Twenty six percent of the prioritized candidate variants coincide with four putative regulatory elements that interact with the *MAP3K1* promoter through chromatin looping and affect *MAP3K1* promoter activity. Functional analysis indicates the cancer risk alleles of four candidates increase *MAP3K1* transcriptional activity: rs74345699 and rs62355900 (iCHAV1); rs16886397 (iCHAV2a); and rs17432750 (iCHAV3). Chromatin immunoprecipitation analysis revealed diminished GATA3 binding to the minor (cancer protective) allele of rs17432750, indicating a mechanism for its action. We propose that the cancer risk alleles act to increase *MAP3K1* expression *in vivo* and may promote breast cancer cell survival.

Introduction

One of the first Genome Wide Association Studies (GWAS) for breast cancer [MIM 114480] susceptibility identified a SNP at 5q11.2, rs889312, associated with risk of breast cancer in women of European ancestry.¹ In the most recent analyses by the Breast Cancer Association Consortium (BCAC), the minor allele of rs889312 was associated with a per-allele Odds Ratio (OR)=1.12 (95% Confidence Intervals (CI) 1.10-1.15; P -trend= 1.8×10^{-26}).² The association was stronger for ER-positive disease (OR=1.14, 95% CI 1.11-1.17; $P=1.1 \times 10^{-26}$ in the most recent BCAC analysis) but also seen for ER-negative (OR=1.06, 95% CI 1.03-1.10; $P=0.0024$) and triple negative disease (OR=1.11, 95% CI 1.02-1.20; $P=0.016$).³ SNP rs889312 was also reported to be associated with an increased breast cancer risk in *BRCA2* [MIM 600185] mutation carriers.⁴

The GWAS SNP, rs889312, lies approximately 80kb centromeric of *MAP3K1* [MIM 600982], the gene encoding Mitogen-Activated Protein Kinase Kinase Kinase1, also known as MEK kinase 1 (MEKK1) – a stress-induced serine/threonine kinase with apparent dual functions: MEKK1 induces cell proliferation through a RAS/RAF/MEK/ERK signaling pathway⁵ but, upon caspase cleavage, generates a fragment which has a pro-apoptotic function.^{6,7} Furthermore, MEKK1 regulates transcription of key cancer related genes such as *MYC*⁸ [MIM 190080], *TP53*⁹ [MIM 191170] and *JUN*¹⁰ [MIM 165160] through its signal transduction pathway. There is already evidence of a role for *MAP3K1* in breast cancer pathogenesis: *MAP3K1* driver mutations have been observed in luminal A and B type breast tumors¹¹ and *MAP3K1* expression has been associated with specific breast tumor subtypes.¹²

In this study, we have performed genetic epidemiological analyses on all common variants at 5q11.2, together with *in silico* and *in vitro* analyses of candidate causal variants, and identified strong candidates that we propose are functionally related to breast cancer risk. Specifically, we provide evidence that these associations are mediated through *MAP3K1*.

Materials and Methods

SNP selection and genotyping

We identified all SNPs from a 305kb interval on 5q11.2 (GRCh37 positions 55,983,657–56,288,810) with a minor allele frequency (MAF) >0.02 in Europeans using the March 2010 release of the 1000 Genomes Project.¹³ SNPs with an Illumina designability score of >0.8, and $r^2 > 0.1$ with rs889312, together with a tagging set ($r^2 > 0.9$) for all other known SNPs in the interval, were selected for inclusion on the iCOGS custom array.² In total, 352 SNPs were selected, of which 300 passed post-genotyping quality control criteria.² To improve SNP coverage across the locus, 16 further SNPs, selected from the October 2010 release of the 1000 Genomes Project were genotyped in a subset of two BCAC studies (SEARCH and the combined Copenhagen studies CPGS and CCHS) using a Fluidigm array according to the manufacturer's instructions. These two datasets were used to impute all genotypes of other common variants in this interval, using IMPUTE2 and the January 2012 1000 Genomes Project release as reference.¹⁴ All participants provided written informed consent. Study characteristics and iCOGS methodology have been previously reported.²

Statistical analyses

Associations with breast cancer risk in BCAC were evaluated by comparing SNP genotype frequencies in cases and controls using unconditional logistic regression. Analyses were adjusted by study and seven principal components.² The primary analysis fitted each SNP as an allelic dose and tested for association using a 1-degree-of-freedom trend test (*P*-trend) with associated odds ratio (OR) and 95% Confidence Intervals (95% CI). To identify independent risk signals, stepwise conditional analysis was performed in R with the function "step", including any variant with $P\text{-value} < 10^{-4}$ in the single SNP analysis to calculate the most parsimonious model using a penalty value of $k=10$. The null model included study and principal components. Haplotype analysis was performed in R using the package "haplo.stats", and the analyses adjusted for study and principal components.

Cell lines

The normal breast epithelial cell line Bre-80 was cultured as described previously.¹⁵ The breast cancer cell lines MCF7, T-47D and MDA-MB231 were cultured in RPMI1640 medium supplemented with 10% fetal calf serum, antibiotics, sodium pyruvate and, in the case of MCF7 and T-47D cells, 10 $\mu\text{g/ml}$ insulin.

Chromatin conformation capture (3C)

3C libraries were generated with EcoRI from the cell lines above as described previously.¹⁶ 3C interactions were quantitated by real-time PCR (qPCR) with primers designed within the EcoRI restriction fragments spanning the 5q11.2 risk locus (**Table S1**). qPCR was performed as described previously¹⁷ using at least two independent 3C libraries from each cell line with each experiment quantified in

duplicate. Two BAC clones (RP11-378G4 and RP11-1146C6) covering the 5q11.2 region were used to create an artificial library of ligation products in order to normalize for PCR efficiency. As an internal control, interaction frequencies were normalized to that of the EcoRI fragment immediately upstream of the promoter/bait fragment.

Plasmid construction

A *MAP3K1* promoter-driven luciferase reporter construct was generated by inserting a 1928 bp fragment containing the *MAP3K1* promoter and the transcription start site (GRch37 chr5:56109070-56110997) into the MluI and HindIII sites of pGL3-Basic. To assist cloning, AgeI and SbfI sites were inserted into the BamHI and Sall sites downstream of the luciferase gene. A 1575 bp PRE-A fragment, a 1765 bp PRE-B2 fragment, a 2357 bp PRE-B3 fragment, a 2203 bp PRE-C fragment and a 1519 bp PRE-D fragment were PCR generated using primers designed with AgeI and SbfI sites and cloned into the modified pGL3-*MAP3K1* promoter construct. PRE-B was too large (~7 kb) to be cloned in its entirety so three subregions containing iCHAV1 (PRE-B1), iCHAV2b (PRE-B2) and iCHAV3 (PRE-B3) variants were cloned separately. The minor alleles of individual SNPs were introduced into promoter and PRE sequences, containing the major alleles of any other causal candidate variants, by overlap extension PCR. All constructs were sequenced to confirm variant incorporation (AGRF, Brisbane, Australia). PCR primers are listed in **Table S2**. For the PRE-B1 construct, a 2129 bp region spanning chr5:56028968-56031097 (GRCh37) was synthesized with AgeI and SbfI sites incorporated at the 5' and 3' ends (GenScript, Piscataway, NJ, USA) to assist cloning into the *MAP3K1* promoter construct. The cloned regions are highlighted in **Figure 2B**.

Reporter assays and estrogen induction

Bre-80 and MCF7 cells were transfected with equimolar amounts of luciferase reporter plasmids and 50 ng of pRLTK transfection control plasmid with Lipofectamine 2000. The total amount of transfected DNA was kept constant at 600 ng for each construct by adding pUC19 as a carrier plasmid. Luciferase activity was measured 24 h post-transfection by the Dual-Glo Luciferase Assay System. To correct for differences in transfection efficiency or cell lysate preparation, *Firefly* luciferase activity was normalized to *Renilla* luciferase. For the assays under basal conditions, the activity of each construct was calculated relative to the construct containing the *MAP3K1* promoter alone, the activity of which was defined as 1.

For estrogen induction assays, we treated cells as described.¹⁸ Briefly, 24 h after plating MCF7 cells into wells, medium was replaced with that containing 10 nM fulvestrant for 48 h, which was used to inhibit estrogen induced gene expression, thereby creating a baseline of expression for reporter assays. Cells were then incubated with fresh medium containing either 10 nM estrogen (17 β -estradiol) or DMSO (as vehicle control) and transfected with reporter plasmids. Luciferase assays were performed as above after 24 h. Statistical significance was tested by repeated-measures ANOVA, using the Greenhouse-Geisser correction for non-sphericity, followed by Dunnett's multiple comparisons test in GraphPad Prism.

ChIP assays

Chromatin immunoprecipitation (ChIP) assays were carried out as previously

described.¹⁹ GATA3 specific mouse monoclonal antibodies were obtained from Santa Cruz Biotech (sc268). Precipitate and input were used in qPCR using SYBR green master mix as described.¹⁹ Primer sequences are listed in **Table S3**. Both rs17432750 primer sets gave identical enrichment and the identity of the larger fragment was confirmed by Sanger sequencing. All values obtained were normalized to input and enrichment is given relative to the negative *CCND1* [MIM 168461] control. Allele-specific ChIP was carried out by using a TaqMan SNP genotyping assay (Applied Biosystems) on the ChIP material. In the TaqMan assay two different fluorophores are each linked to a probe detecting the two different alleles. Amplification of each allele was followed using an Applied Biosystems Real Time PCR machine (7900HT) and the data analysed using the SDS software. The SDS software converts the raw data to fluorescence intensity for each allele and then plots the results as a scatter graph of allele X versus allele Y. We tested the accuracy of this assay by genotyping known mixtures of homozygous ZR751 and T47D (C/C), and MDA-MB0-468 (A/A) cell line DNA (**Figure S1**). For the allelic discrimination three independent experiments were carried out and gave similar results - a representative example is shown.

GATA3 siRNA knockdown for reporter assay

GATA3 (MIM 131320; L-003781-00) and non-targeting (D-001810-10-20) ON-TARGET^{plus} SMARTpool siRNAs for were purchased from Thermo Scientific. For knockdown, Bre-80 cells were co-transfected with the relevant luciferase reporter plasmids and 100 nM of either *GATA3* or non-targeting siRNAs with Lipofectamine 2000. Luciferase assays were performed as described above after 24 h. To validate *GATA3* knockdown, qPCR was performed as described previously (**Figure S9**).¹⁷

Expression quantitative trait locus (eQTL) analysis

eQTL analysis was undertaken in two sample sets of adjacent normal breast samples from women of European decent: the first set contained 135 samples collected for the METABRIC study²⁰; the second set was extracted from the TCGA breast cancer study²¹ data and contained 56 samples. Matched gene expression (Illumina HT-12 v3 microarray for the METABRIC data; Agilent G4502A-07-3 microarray for the TCGA data) and germline SNP data that was either genotyped (Affymetrix SNP 6.0) or imputed (1,000 Genomes Project, March 2012 data using IMPUTE version 2.0) were used. Correlations between all imputed and genotyped variants at the 5q11.2 locus and expression levels of eight (METABRIC) or four (TCGA) genes present in the fine-mapped region were assessed using a linear regression model in which an additive effect on expression level was assumed for each copy of the rare allele. Calculations on the METABRIC data were carried out using the eMap library in R, and on the TCGA data using the SNPTTEST software.²²

Results

Genotyping of case-control studies

Three hundred SNPs at the 5q11.2 locus (GRch37 positions 55,983,657–56,288,810) were successfully genotyped using the iCOGs chip, in 103,991 breast cancer cases and controls from 52 BCAC studies, of which 41 (46,451 cases and 42,599 controls) were of European, nine were of Asian (6,269 cases and 6,624 controls) and two were of African-American ancestry (1,116 cases and 932 controls). Using these data, together with data on a further 16 SNPs, genotyped in two BCAC studies (SEARCH and the combined Copenhagen studies CPGS and CCHS), we

imputed genotypes for 909 variants with $MAF > 0.02$ (out of a possible 911) and imputation $r^2 > 0.3$, using the January 2012 release of the 1000 Genomes Project as reference.

Potential breast cancer risk signals in European studies

Figure 1A shows the Manhattan plot of the 909 genotyped and imputed SNPs ($r^2 > 0.3$) for overall breast cancer risk in European studies. Genotype and association results for all 909 SNPs are presented in **Table S4**. Five hundred and forty one variants display association with overall breast cancer risk at $P\text{-trend} < 10^{-4}$ (**Table S5**). All associations are consistent with a log-additive model.

In a forward stepwise conditional analysis, using the SNPs listed in the **Table S5**, the best model included three SNPs: (1) rs62355902 – conditional P -value ($P\text{-cond}$) = 8.6×10^{-26} ; (2) rs113317823 - $P\text{-cond} = 2.8 \times 10^{-5}$ and (3) rs11949391 - $P\text{-cond} = 9.7 \times 10^{-5}$. No significant evidence for heterogeneity was observed among odds ratios (ORs) for these SNPs among studies of European ancestry (minimum observed $P\text{-het} = 0.14$ and maximum $I^2 = 19.3\%$ for SNP rs11949391 - **Figure S2**). Each SNP remaining in the conditional analysis model indicates the existence of a separate genetic risk signal (previously defined²³ as an Independent set of Correlated Highly Associated Variants – iCHAV), each of which will contain at least one directly causal variant.

The most significantly associated variant overall, is rs62355902 (OR (per minor allele) = 1.21, 95% CI 1.19-1.24; $P\text{-trend} = 9.5 \times 10^{-49}$). This is the most significant of 15 strongly correlated SNPs ($r^2 > 0.93$) lying within a 50Kb interval (GRCh37 positions

56,003,831–56,053,745; **Table 1** and marked in red in **Table S5**), which we have designated as iCHAV1. These iCHAV1 SNPs all have likelihood ratios of <100:1 relative to the best candidate SNP, rs62355902, and thus cannot be excluded from further analysis, remaining as strong candidate causal variants based on epidemiological evidence. After conditioning on iCHAV1 top SNP, rs62355902, the most strongly associated variant is SNP rs113317823: OR (per minor allele)=1.22, 95% CI 1.18-1.26; P -trend= 7.0×10^{-25} ; conditional OR=1.12, 95% CI 1.05-1.20; P -cond= 2.8×10^{-5} . One hundred and seventy two variants have likelihood ratios of <100:1 relative to rs113317823 and these constitute iCHAV2 (highlighted in shades of blue and yellow **Table S5**). SNP rs113317823 is partially correlated with iCHAV1 top candidate rs62355902 ($r^2=0.19$) and this adds complexity to the subsequent analysis – this is explored in more detail by the haplotype analysis described below.

After adjustment for the top iCHAV1 (rs62355902) and iCHAV2 (rs113317823) SNPs, the best remaining significantly associated SNP is rs11949391 (iCHAV3): OR (per minor allele)=0.91, 95% CI 0.89-0.94; P -trend= 9.4×10^{-12} ; and conditional OR=0.95, 95% CI 0.92-0.98; P -cond= 9.7×10^{-5} . Four other SNPs have likelihood ratios of <100:1 relative to rs11949391 and are highly correlated with rs11949391 ($r^2 \geq 0.95$) but not with either of the top iCHAV1 or iCHAV2 SNPs ($r^2 < 0.04$, marked in yellow in **Table S5**, listed in **Table 1**) and these are thus candidate causal variants for iCHAV3.

Effects on Estrogen Receptor-positive (ER+) and negative (ER-) tumor subtypes

Candidate causal SNPs in iCHAV1 are associated with risks of both ER+ and ER- disease (**Table 1**). However the OR is greater for ER+ (rs62355902 OR=1.24, 95% CI 1.21-1.27) than ER- disease (OR=1.10, 95% CI 1.05-1.15; P -difference= 1.5×10^{-5} ; **Table 1**). By contrast, the minor alleles of candidate causal SNPs in iCHAV3 are protective against ER+ tumor development (rs11949391 OR=0.90, 95% CI 0.87-0.93; $P=1.9 \times 10^{-10}$) but have no apparent effect on ER- tumor risk (OR=1.01, 95% CI 0.96-1.06; $P=0.66$; P -difference= 6.5×10^{-5} ; **Table 1**). The lead SNP in iCHAV2, rs113317823, remains significantly associated with ER+ tumor risk (P -cond= 9.7×10^{-5} , after adjustment for rs62355902) but not with ER- tumor risk (P -cond=0.099), but the difference in the OR by ER subtype is only borderline significant (P -difference=0.02; **Table 1**).

Effects of haplotypes on breast cancer risk

While iCHAVs 1 and 3 represent sets of highly correlated SNPs ($r^2 > 0.93$) with the lead SNP), the set of 173 SNPs, labeled iCHAV2, includes three subsets, defined according to their correlations with rs113317823 (iCHAV2) and rs62355902 (iCHAV1): these subsets are iCHAV2a - lead SNP rs113317823 - representing 90 SNPs correlated with rs113317823 ($r^2 > 0.53$) and with iCHAV1 SNP rs62355902 ($r^2 = 0.19-0.29$), marked in mid blue **Table S5**; iCHAV2b - lead SNP rs62355899 representing 66 SNPs, which are independent of rs113317823 ($r^2 \leq 0.01$) but correlated with rs62355902 ($r^2 = 0.59-0.62$) (conditional OR=0.90, 95% CI 0.86-0.95; P -cond= 3.0×10^{-5}), marked in **Table S5** in teal; and iCHAV2c - lead SNP rs7721581 - representing 17 SNPs that are modestly correlated with rs113317823 ($r^2 = 0.14-0.16$) but independent of rs62355902 ($r^2 \leq 0.01$) (conditional OR=0.96, 95% CI 0.93-0.98; P -cond= 4.8×10^{-4}) marked in **Table S5** in pale blue.

To further clarify these association signals, we performed haplotype analyses, based on the lead SNPs from iCHAVs 1, 2a, 2b, 2c and 3. These five SNPs define seven common haplotypes (**Table 2**). Two of these: **h5**, which carries the risk alleles of iCHAV1, iCHAV2a and iCHAV2b; and **h6**, which carries the risk alleles of iCHAV1 and iCHAV2b, are strongly associated with risk, with the risk being higher for haplotype **h6** ($P=1.66 \times 10^{-29}$). These results are consistent with the observation that SNPs in iCHAV1 are the most strongly associated with risk. They are also consistent with a model in which either a SNP in iCHAV2a or iCHAV2c (in combination with iCHAV1) increases risk, or a SNP in iCHAV2b reduces risk. These hypotheses are difficult to distinguish since the iCHAV1 risk allele never occurs alone – it occurs in combination with either iCHAV2a/iCHAV2c or with iCHAV2b, but not both. Some support for the iCHAV1+iCHAV2a hypothesis is provided by the fact that haplotypes **h3** and **h4**, which carry the risk alleles for iCHAV2a but not iCHAV1, though rare, are associated with an increased risk. Evidence against the iCHAV1+iCHAV2c hypothesis is provided by the fact that **h1**, which carries the risk allele for iCHAV2c alone, is not associated with an increased risk. These observations are consistent with the regression analyses, in which iCHAV2c is less likely than iCHAV2a or iCHAV2b SNPs to harbour a causal variant, (likelihood ratio ~30:1, after adjustment for iCHAV1). Haplotype **h2**, which carries the minor allele of iCHAV3 SNP rs11949391, was associated with a reduced ER+ (though not ER-) breast cancer risk, consistent with the effect of the iCHAV3 SNP in the regression analysis.

We conclude that, at least, one of the 90 SNPs in iCHAV2a (positions 56,001,002-56,270,717) or one of the 66 SNPs in iCHAV2b (positions 55,998,085-56,183,743)

are causally related to risk, together with a variant in iCHAV1 and a variant in iCHAV3.

Risk associations in Asian and African-American studies

We tested all genotyped and imputed SNPs with MAF>0.02 and imputation $r^2>0.3$ in the nine Asian studies (6,269 cases and 6,624 controls; SNPs $n=1045$) and the two African-American studies (1,116 cases and 932 controls; SNPs $n=1601$) for association with overall breast cancer risk. None reached genome-wide levels of significance ($P<5\times 10^{-8}$) but the lead SNPs of each iCHAV display compatible effects in all three ethnic groups. This is most apparent for iCHAV2a SNP rs113317823: (European–unadjusted) OR=1.22, 95% CI 1.18-1.26; P -trend= 7.0×10^{-25} ; (Asian) OR=1.19, 95% CI 1.11-1.27; P -trend= 1.4×10^{-5} ; (African-American) OR=1.04, 95% CI 0.77-1.31; P -trend=0.78, **Table S6**.

Identification of putative regulatory elements (PREs) that interact with the *MAP3K1* promoter

After the epidemiological analyses, 15 iCHAV1, 90 iCHAV2a, 66 iCHAV2b and 5 iCHAV3 variants remain as strong causal candidates (**Figure 1A** and **Table S7**), whereas the data provide weaker support for iCHAV2c being causal. As iCHAV2a and 2b were comprised of a large number of variants, we prioritized these for functional analysis using a threshold of P -cond $<1\times 10^{-4}$ and focused on the remaining 30 iCHAV2a and 10 iCHAV2b variants, in addition to the iCHAV1 and iCHAV3 candidates (**Table 1**). Next we examined whether these 62 iCHAV variants coincide with PREs that may affect gene expression.

Using publicly available ENCODE chromatin immunoprecipitation sequencing (ChIP-seq) data from MCF7 and HMEC cells, we identified regions marked by histone modifications associated with transcriptional enhancers (mono- or di-methylation of H3 lysine 4 (H3K4Me1 and H3K4Me2) and acetylation of H3 lysine 27 (H3K27Ac)) or bound by ER α , FOXA1 or GATA3, transcription factors known to play a role in breast cancer. Next, we mined RNA polymerase II ChIA-PET (chromatin-interaction analysis with paired-end tag sequencing) data, previously generated in MCF7 cells,^{24,25} and identified multiple long-range chromatin interactions between discrete regions of the iHAV loci and the promoter of *MAP3K1* (**Figure S3**). Consequently, we performed chromosome conformation capture (3C) experiments to analyze interactions between the *MAP3K1* promoter and these regions within 5q11.2. Using 3C in a normal mammary epithelial cell line, Bre-80, we found several regions that interact with the *MAP3K1* promoter (**Figure 2C**). Similar 3C profiles were observed in two ER⁺ cell lines (MCF7 and T-47D) and in ER⁻ MDA-MB231 breast cancer cells. (**Figure S4**).

PREs were defined as the loci encompassing functional elements identified from the ENCODE data within a *MAP3K1* promoter interacting region. This analysis revealed four PREs (A-D) that coincide with the iHAV candidates prioritized for functional analyses (**Figure 2**). Consistently, ENCODE ChIA-PET data demonstrates that all four PREs interact with the *MAP3K1* promoter (**Figure S3A**) but not with the promoters of other nearby genes in MCF7 cells (**Figure S3B**). It should be noted that all four PREs contain iHAV2a or 2b variants ($P\text{-cond} > 1 \times 10^{-4}$) that were not prioritized for functional analyses but could not be excluded as causal candidates after the log likelihood testing (**Figure 2A**). Furthermore, additional PREs, some

containing such iCHAV2a or 2b variants, are apparent at this locus (**Figure 2**). Twenty six percent of the 62 iCHAV variants, prioritized for functional analyses, are coincident with a PRE. This is an enrichment of the 16% of SNPs at this locus with MAF in the range of the prioritized iCHAV variants (dbSNP 138, MAF=0.04-0.18, accessed through the UCSC Genome Browser) that are located in a PRE.

The risk alleles of iCHAV1 candidates rs74345699 and rs62355900 further induce PRE-C enhancer activity after estrogen stimulation

For functional analysis of iCHAV1, we focused on seven of the 15 candidate causal variants, coincident with a PRE (PREs A-C; **Figure 2**). We first examined the effects of these PREs on *MAP3K1* promoter activity by cloning the relevant genomic regions into luciferase reporter gene constructs containing the *MAP3K1* promoter. All three of the PREs had effects on *MAP3K1* promoter activity: PRE-A acted as a silencer and reduced promoter activity by 62% ($P=0.006$) in Bre-80 cells (**Figure 3A**) with a similar but non-significant trend ($P=0.056$) in MCF7 cells (**Figure S5A**). A subregion of PRE-B, termed PRE-B1, containing two candidate variants acted as an enhancer in Bre-80 cells, increasing promoter activity by 42% ($P=0.047$; **Figure 3B**), but had no significant effect in MCF7 cells – indicating that PRE-B1 may function in specific breast cell types (**Figure S5B**). PRE-C acted as an enhancer and increased *MAP3K1* promoter activity by 90% ($P=0.034$) in MCF7 (**Figure 3C**) and 77% ($P=0.034$) in Bre-80 (**Figure S5D**) cells. Introduction of the iCHAV1 minor alleles into the PRE-A, PRE-B1 and PRE-C reference constructs did not detectably alter *MAP3K1* promoter activity (**Figure 3A-C; S5A-B and D**).

ChIP-seq data from the ENCODE project indicate that at least two different transcription factors implicated in estrogen signaling, FOXA1 and ER- α , bind within PRE-B1 and PRE-C, respectively (**Figure 2B**). We thus tested whether the iCHAV1 candidates within PRE-B1 and PRE-C confer estrogen dependent effects on *MAP3K1* promoter activity. We first confirmed that *MAP3K1* expression is up-regulated by estrogen stimulation (**Figure S6**). Then, using reporter assays, we examined the effects of estrogen induction on PRE-B1 and PRE-C by measuring the changes in *MAP3K1* promoter activity between estrogen stimulated and unstimulated cells. We showed that the PRE-C enhancer containing the protective (major) alleles (**Figure 3D**), but not PRE-B1 (**Figure S5C**), has a significant induction in activity, compared with the promoter construct (72%, $P=0.012$), after estrogen stimulation. Compared to this reference PRE-C enhancer, induction was 23% ($P=0.032$) and 15% ($P=0.011$) greater in PRE-C enhancers containing the risk (minor) alleles of iCHAV1 candidate SNPs, rs74345699 and rs62355900, respectively (**Figure 3D**). Of note, neither of these candidates significantly affected *MAP3K1* promoter activity in the absence of estrogen.

The risk allele of iCHAV2a candidate rs16886397 in PRE-D enhances *MAP3K1* promoter activity

Of the 30 candidate causal variants for iCHAV2a at $P\text{-cond}<1\times10^{-4}$, one variant, rs77371588, coincided with a PRE (**Figure 2**) and we thus prioritized this SNP for functional analysis. Using luciferase reporter assays, we demonstrated that the reference PRE-D acted as an enhancer of the *MAP3K1* promoter in Bre-80 cells, increasing *MAP3K1* promoter activity by 69% ($P=0.013$; **Figure S7A**). The PRE-D enhancer containing the risk (minor) allele of rs77371588 had 23% greater enhancer

activity than the reference PRE-D, but this effect did not reach statistical significance ($P=0.103$; **Figure S7A**). By contrast the same PRE-D reference construct did not impact *MAP3K1* promoter activity in MCF7 cells nor did the introduction of the risk allele of rs77371588 into PRE-D significantly alter its activity (**Figure 3E**). As we had generated the PRE-D construct it was straightforward to test rs16886397, an additional iCHAV2a causal candidate located in PRE-D that did not reach the threshold for the functional prioritization ($P\text{-cond}<1\times 10^{-4}$) but did pass the likelihood ratio threshold of <100:1 for defining the causal iCHAV candidates. In MCF7 cells, the construct containing the minor (risk) allele of rs16886397 had 21% ($P=0.049$) greater *MAP3K1* promoter activity than the reference PRE-D (**Figure 3E**) and, thus, rs16886397 appears to confer enhancer activity on PRE-D. In Bre-80 cells, in contrast, the minor allele of rs16886397 had no effect on PRE-D activity (**Figure S7A**).

The minor allele of iCHAV2b candidate, rs62355881, increases PRE-B2 enhancer activity

Of the ten iCHAV2b candidates at $P\text{-cond}<1\times 10^{-4}$, three variants were coincident with PRE-B and two flanked the boundaries of PRE-C (**Figure 2**). We prioritized the three variants in PRE-B as these were the most compelling functional candidates due to their central location in several functional elements (**Figure 2B**). Using reporter assays, we demonstrated that the reference PRE-B2 construct (a subregion of PRE-B containing the iCHAV2b variants) acted as enhancer and increased the activity of the *MAP3K1* promoter by 152% ($P=0.032$) and 143% ($P=0.048$) in MCF7 and Bre-80 cells, respectively (**Figure 3F**). The introduction of the minor (potentially protective) allele of the iCHAV2b candidate rs62355881 into the reference PRE-B2

construct led to a 29% ($P=0.017$) increase in the enhancer activity of PRE-B2 in MCF7 cells and the haplotype construct containing the minor alleles of all three iCHAV2b candidates demonstrated a similar effect ($P=0.030$; **Figure 3F**). However, these effects were not seen in Bre-80 cells, indicating another cell type specific effect, nor did the other iCHAV2b candidates have any effect on PRE-B2 activity in either cell line (**Figures 3F and S7B**).

The protective allele of iCHAV3 candidate, rs17432750, reduces PRE-B3 enhancer activity and GATA3 binding

Of the five iCHAV3 candidates, we focused on two variants coincident with PRE-B3 (**Figure 2B**), for functional analysis. Using reporter assays, we demonstrated that the reference PRE-B3 construct (a third subregion of PRE-B), containing the risk (major) iCHAV3 alleles, increased *MAP3K1* promoter activity by 166% ($P=0.041$; **Figure 4A**) and 110% ($P=0.035$; **Figure S8A**) in Bre-80 and MCF7 cells, respectively. Reversing the orientation of PRE-B3 in the reporter gene construct had no effect on its activity in either cell line (**Figure S8B and C**), indicating that it acts as a typical enhancer. Next, we introduced the protective (minor) alleles of the iCHAV3 candidates into the reference PRE-B3 construct. The protective A allele of rs17432750 had a repressive effect and reduced PRE-B3 enhancer activity by 43% ($P=0.024$; **Figure 4A**) in Bre-80 cells. The same allele had a similar but non-significant effect ($P=0.150$) in MCF7 cells (**Figure S8A**). By contrast the protective allele of the second iCHAV3 candidate, rs11956804, had no significant effect on enhancer activity in either cell line (**Figures 4A and S8A**). The haplotype construct containing both iCHAV3 variants also had no significant effect on enhancer activity either (**Figures 4A and**

S8A), suggesting the possibility of an interaction between the two iCHAV3 minor alleles in this construct.

We observed from MCF7 ChIP-seq data that a region containing rs17432750 binds the GATA3 transcription factor (**Figure 2B**) and that the sequence around rs17432750 shows homology to the GATA3 position weight matrix (**Figure 4B**). Using GATA3 ChIP-assays, followed by qPCR detection, we confirmed that there was a consistent 2-fold enrichment of the sequence surrounding this SNP in precipitated DNA, when compared to a *CCND1* negative control (**Figure 4C**). We also tested the allele-specificity of GATA3 binding using a Taqman genotyping assay for rs17432750 on ChIP samples from MCF7 cells. The allelic discrimination plot of these data showed an enrichment of the risk (major) C allele in the GATA3 ChIP samples (**Figure 4D**). The ratio of the two alleles in three independent ChIP experiments indicated a 3.7-fold greater GATA3 binding to the risk C allele, compared to the protective A allele, in MCF7 cells.

To determine whether differential GATA3 transcription factor binding may explain the effects of rs17432750 in the reporter assays, we knocked down *GATA3* using siRNA and found the enhancer activity of the reference PRE-B3 construct, containing the risk C allele, was reduced by 33% in Bre-80 cells ($P=0.001$; **Figure 4E**). *GATA3* knockdown had no effect on the construct containing the protective A allele of (rs17432750) or the *MAP3K1* promoter alone (**Figure 4E**). Diminished GATA3 binding to the protective A allele thus appears to be responsible for the decrease in PRE-B3 enhancer activity observed under basal conditions (**Figure 4A**).

Expression quantitative trait (eQTL) analyses

Given the findings of these functional studies, an obvious hypothesis is that the candidate causal variants in the iCHAVs would be associated with differences in expression of *MAP3K1*, and possibly other local genes in normal breast cells. We therefore explored potential eQTL associations of all locus SNPs with genes lying within ~1 Mb of the locus in 135 normal breast tissue samples from the METABRIC study and 56 further normal breast samples from TCGA. Summary results for representative SNPs from the three iCHAVs are presented in **Table S8**. None of the iCHAV-representative SNPs show detectable differences in *MAP3K1* expression in this dataset. This locus has a positive control eQTL: SNP rs832402 is the most strongly associated SNP with *SETD9* expression in both METABRIC ($P=5.93 \times 10^{-9}$) and TCGA ($P=1.96 \times 10^{-7}$), but it is not a strong candidate breast cancer risk SNP ($P\text{-cond}=1.46 \times 10^{-3}$). The positive control SNP is in iCHAV2c (correlated with lead SNP, rs7721581 at $r^2=0.74$) and SNP rs7721581 is consequently also associated, although less significantly, with *SETD9* expression (METABRIC $P=4.38 \times 10^{-8}$; TCGA $P=2.78 \times 10^{-4}$). However, as none of the other representative iCHAV SNPs are associated with *SETD9* expression, it appears unlikely that detectable *SETD9* expression differences in normal breast cells are the underlying cause of breast cancer generated by the candidate functional variants we have identified.

Discussion

In this fine scale mapping study, we have found clear evidence for at least three independent breast cancer risk variants in European women: SNPs in iCHAV1 and iCHAV2, each have the greatest effects on breast cancer in the unadjusted analysis, with minor alleles conferring increased risks of 25-30% for ER+ and ~10% for ER-

tumor-development, while the minor alleles of SNPs in iCHAV3 have a protective effect of ~10% against ER+ breast cancer, although no apparent effect on ER- tumor risk (**Table 1**). The originally detected GWAS tag SNP, rs889312, is most correlated with iCHAV1 ($r^2=0.5$) and can be excluded from causality within iCHAV1 with a likelihood ratio of $>10^{21}:1$. Within iCHAV2, we have additionally found evidence of three subsets of variants with a range of effects: iCHAVs 2a, 2b and 2c. The epidemiological analyses suggest iCHAV2c is least likely to be causally related to risk and we conclude that at least one of the variants in iCHAV2a or iCHAV2b is functional. It should be noted that the correlations between candidate causal SNPs in iCHAVs 1 and 2 have added a level of complexity to this analysis and its interpretation that we have not recognized in previous fine-scale mapping studies of breast cancer risk loci.

We have separately identified at least four PREs within the 5q11.2 locus that contain iCHAV candidate variants and interact with the *MAP3K1* promoter in normal and cancer mammary epithelial cells. 3C analysis indicates that there are long range chromatin interactions between these PREs and the *MAP3K1* promoter, while interactions between the PREs and the promoters of other nearby genes are not evident from available ChIA-PET studies.^{24,25} Although we cannot rule out interactions between the iCHAVs and the promoters of other genes in the region such as *SETD9* or *MIER3*, a proposed candidate gene for this risk locus,²⁶ we propose that *MAP3K1* is the likely target gene of the 5q11.2 breast cancer susceptibility locus. Consistent with this proposition and our analyses, Corradin *et al.* identified an enhancer (chr5:56052477-56053943) coincident with the element we

have termed PRE-C that is predicted to regulate *MAP3K1*, based on correlation of cell-type specific H3K4me1 modification and *MAP3K1* expression data.²⁷

Our reporter assays indicate that PREs B, C and D act as enhancers while PRE-A is a silencer of the *MAP3K1* promoter. Having identified these regulatory elements we have investigated whether the iCHAV candidate causal variants, within these PREs, detectably modify their regulatory activity. We found that the risk alleles of two iCHAV1 candidates, rs74345699 and rs62355900, in the PRE-C enhancer, act to further induce *MAP3K1* promoter activity in breast cancer cells under estrogen stimulation; the risk allele of iCHAV2a candidate, rs16886397, in PRE-D confers enhancer activity on this PRE for *MAP3K1* promoter activity in breast cancer cells; the potentially protective allele of iCHAV2b candidate, rs62355881, in the PRE-B2 enhancer, increases *MAP3K1* promoter activity in breast cancer cells; while the protective allele of iCHAV3 SNP rs17432750 diminishes the enhancer activity of PRE-B3 for the *MAP3K1* promoter in these cells.

Due to experimental constraints, we have only been able to examine the functions of a minority of the iCHAV causal candidates and, thus, we cannot exclude the presence of more functional variants across the recognizable iCHAVs. Of the candidates we have examined in reporter gene assays, four support a hypothesis that alleles, which increase *MAP3K1* expression, also increase breast cancer risk. Currently, the allelic effects of a fifth candidate, iCHAV2b SNP rs62355881, may be inconsistent with this hypothesis as the conditional analysis suggests iCHAV2b has a protective effect. However, we do not have clear epidemiological evidence that iCHAV2b has an individual effect on risk: iCHAV2b is only observed in the presence

of iCHAV1, which in combination on the same haplotype act to increase breast cancer risk, and we could not test such iCHAV haplotype effects in reporter gene assays, given the number of candidates and the size of the region they encompass.

Consistent with our hypothesis that candidate causal risk alleles act by increasing transcriptional activation of *MAP3K1*, Godde et al. have recently demonstrated that up-regulation of MAP kinase activity in mouse mammary basal progenitor cells is associated with ductal hyperplasia and accelerated tumor progression.²⁸ This hypothesis is also supported by one known function of *MAP3K1* protein (MEKK1): knockdown of *MAP3K1* in human breast cancer cells reduces tumor invasiveness and progression in a mouse model.²⁹ Furthermore, studies have shown that MEKK1 has an anti-apoptotic effect and enhances cancer cell survival,^{30,31} although, upon stress stimulus, caspase cleavage of the protein generates a fragment that plays a pro-apoptotic role.^{6,7} These dual functions of MEKK1 suggest it has a complex role in cell fate decisions. In this context, it is intriguing that somatic *MAP3K1* driver mutations, found in tumor sequencing studies, are mostly truncating and are predicted to disrupt MEKK1 signaling,¹¹ since inactivation of the kinase domain at the protein terminus reduces apoptotic responses in cells exposed to stress.⁷ Thus, it appears that germline cancer risk alleles act by increasing *MAP3K1* expression, but once a tumor has developed, somatic mutations drive cancer progression by disrupting MEKK1 signaling within the tumor. Dysregulation of *MAP3K1* expression or MEKK1 function may thus promote tumorigenesis by perturbing a balance between cell apoptosis and survival. Similar conflicting effects of germline risk variants and somatic mutations have been observed at other breast cancer risk loci. For example, *CCND1* is frequently amplified in breast tumors and here

overexpression appears to play an important role in breast cancer pathogenesis,³² even though germline breast cancer risk alleles at 11q13 reduce *CCND1* transcriptional activity.¹⁷ Amplification of *TERT* [MIM 187270] is also common within breast and other tumors³³ but, again, germline breast cancer risk alleles reduce *TERT* transcriptional activity.³⁴ These observations therefore challenge the notion that variants at loci such as 5q11.2 act in the same manner as somatic tumor driver mutations to confer germline risk of tumor development.

iCHAV1 spans multiple PREs (PREs 1, 2 and 3) and is consistent with a recent proposal that genetic susceptibility to common diseases can be explained by multiple enhancer variants in linkage disequilibrium, each with modest effects on gene expression, which cooperatively act to alter gene expression.²⁷ There is also some suggestion from the haplotype data that iCHAV2a, which is in linkage disequilibrium ($r^2=0.19-0.29$) with iCHAV1, may in combination with iCHAV1 have a cooperative effect on risk. The modest effect sizes observed in the reporter assays for iCHAV1 and 2a variants may be a consequence of the fact that, due to size limitations of reporter gene constructs, we could not examine these variants in combination.

Some SNP effects in our reporter assays were suggestive of cell-line and stimulus-dependence, highlighting the importance of cellular and environmental context when assessing SNP functionality.²³ Similarly, we defined our PREs on the basis of chromatin modification and conformation states, both of which can change during development or in response to stimulus. Additionally, the effects of some SNPs may not have been observed because plasmid reporter gene constructs do not reflect the native genomic context nor the chromatin or methylation state of genomic DNA. The

chromatin state of transiently transfected DNA, for example, is considered to be more open and disorganized than the corresponding chromatin in the native genomic context and may not have the repressive chromatin structures, found in genomic DNA, that inhibit binding of ubiquitous transcription factors.³⁵ A disorganized chromatin state may explain the inconsistent effect of rs17432750 in the PRE-B3 constructs (**Figure 4A**). The abrogation of the effect of rs17432750 in the haplotype construct suggests some interaction or cooperative effect of the minor alleles of rs17432750 and rs11956804 to enhance promoter activity. This effect may be possible due to the more open and permissive chromatin structures associated with transfected plasmid DNA.

It is noteworthy that, in available normal breast tissue, the top candidate causal variants show no association with *MAP3K1* expression, although iCHAV2c variants are associated with significant differences in *SETD9* mRNA levels. Similarly, normal breast tissue eQTL studies with strong candidate causal variants at the 11q13,¹⁷ 10q26,¹⁵ 2q35³⁶ and 5p15.33³⁴ breast cancer loci have indicated that available normal breast samples may be inappropriate for these studies. Sample sizes are large enough to detect significant eQTLs at these loci but those detected do not appear to drive breast cancer risk. It is possible that tissue heterogeneity, developmental stage or stimulus-dependent effects prevent the detection of risk driving eQTLs in currently available normal breast samples. Indeed, the finding that up-regulation of MAP kinase activity in mammary progenitor cells is associated with mammary tumorigenesis²⁸ suggests that increased *MAP3K1* expression in specific breast cell populations, possibly at a specific point in time, could drive breast cancer risk.

Transcription factor binding studies indicate that the C (risk) allele of rs17432750 preferentially binds GATA3 over the A (protective) allele. Increased binding of the C allele by GATA3 appears to explain the activity of the PRE-B3 enhancer in which it is located (**Figure 4E**). These findings suggest that SNP rs17432750 is a strong causal candidate for the protective effect of iCHAV3. The transcription factor GATA3 has multiple regulatory roles and can affect histone modifications associated with enhancers, and the binding of other transcription factors related to breast cancer such as ER- α and FOXA1.³⁷ We have previously identified GATA3 to be a mediator of breast cancer risk across multiple loci³⁸ and specifically at the 11q13 locus.¹⁷

In conclusion, we have found evidence for the existence of at least three breast cancer risk iCHAVs that partially coincide with four *MAP3K1* gene regulatory elements at 5q11.2. Genetic epidemiological studies within BCAC reduced the catalogue of potentially causal variants from 909 to 193 candidates within five iCHAVs, of which at least three must be functional. Functional studies on candidates that lie within the identified regulatory elements have shown that the effects of strong candidate cancer risk alleles, in iCHAVs 1, 2a and 3, are compatible with the hypothesis that they act via increased expression of *MAP3K1*. Moreover, the function of *MAP3K1* protein (MEKK1) suggests that increased expression may alter the balance between apoptosis and cell survival in breast cancer cells, thus explaining the risks conferred by the candidate alleles.

Supplemental Data

Supplemental Data include Acknowledgements, nine figures and eight tables.

Web Resources

1,000 Genomes Project: <http://www.1000genomes.org/>

R: <http://www.bios.unc.edu/~weisun/software/eMap>

UCSC Genome Browser: <http://genome.ucsc.edu>

Online Mendelian Inheritance in Man: <http://www.omim.org>

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Figure Titles and Legends

Figure 1. Genetic mapping and chromatin state of the 5q11.2 locus. (A) Manhattan plot of overall breast cancer risk in Europeans at the 5q11.2 locus. SNPs are plotted based on their chromosomal position on the x axis and P -values (\log_{10} values) for association. The span of the iCHAVs in terms of chromosomal location and P -value is displayed using shading and candidate causal variants from the iCHAV are colored black. The lead SNPs from each iCHAV, the original GWAS tag SNP (rs889312) and the three genes present in the region are shown. The dotted line intersects the y-axis at $P=10^{-8}$ and indicates conventional genome-wide significance. (B) The chromatin state of the 5q11.2 locus in human mammary epithelial cells (HMECs) is shown using ENCODE ChIP-seq data from H3K4me3, H3K4me2, H3K4me1 and DNaseI studies accessed from the UCSC Genome Browser (<http://genome.ucsc.edu>). Transcription factor (TF) binding from ENCODE ChIP-seq studies of 161 TFs in 91 cell lines is also displayed.

Figure 2. Candidate causal variants are located in PREs that interact with the *MAP3K1* promoter. (A) The candidate causal variants associated with breast cancer risk from iCHAVs 1, 2a, 2b and 3 were mapped to PREs at the 5q11.2 locus. (B) PREs (highlighted) were identified using ChIP-seq data (H3K4Me1 studies in 7 ENCODE cell lines (GM12878, H1-hESC, HSMM, HUVEC, K562, NHEK and NHLF); H3K4Me1 and H3K4Me2 in MCF7 and HMEC cells; transcription factors ER- α , FOXA1 and GATA3 in MCF7 cells) accessed from the UCSC Genome Browser. Regions cloned into reporter gene constructs are also shown. (C) 3C analysis of

interactions between EcoRI fragments at the 5q11.2 locus, encompassing the PREs coincident with candidate causal variants, and the *MAP3K1* promoter in Bre-80 cells (error bars represent standard deviation and a representative graph is shown).

Figure 3. Risk (minor) alleles of iCHAV1 and iCHAV2a SNPs enhance *MAP3K1* promoter activity in luciferase reporter assays. PRE-A, PRE-B, PRE-C and PRE-D regions containing the major allelic variants of iCHAV1, iCHAV2a and 2b SNPs were cloned downstream of a *MAP3K1* promoter-driven luciferase construct (Promoter) to create reference (Ref) PRE constructs. Minor allelic variants of the iCHAV1, iCHAV2a and 2b SNPs were engineered into the constructs and are designated by the rs ID of the corresponding SNP. Constructs containing minor allelic haplotypes (Haplotype) were also generated. Cells were transiently transfected with each of these constructs and assayed for luciferase activity after 24 h. Panels (A-B) show results from assays of PRE-A and PRE-B1, respectively, in Bre-80 cells. Panels (C, E and F) show results from assays of PRE-C, PRE-D and PRE-B2 constructs, respectively, in MCF7 cells under basal conditions. Panel (D) shows results after estrogen induction of MCF7 cells. For each reporter construct in this assay, the luciferase activity of estrogen treated cells was normalized to the activity of the corresponding vehicle treated cells. Error bars denote standard error of the mean (SEM) from three experiments performed with triplicates. *P*-values were determined by repeated-measures ANOVA followed by Dunnett's multiple comparisons test (**P*<0.05 and ***P*<0.01).

Figure 4. The protective (minor) allele of iCHAV3 SNP rs17432750 demonstrates diminished PRE-B3 enhancer activity in luciferase reporter assays and reduced GATA3 binding in ChIP analysis. (A) The PRE-B3 region containing the major allelic variants of iCHAV3 SNPs was cloned downstream of a *MAP3K1* promoter-driven luciferase construct (Promoter) to create reference (Ref) PRE constructs. Minor allelic variants of iCHAV3 SNPs were engineered into the constructs which are designated by the rs ID of the corresponding SNP. A construct containing the minor allelic haplotypes (Haplotype) was also generated. Bre-80 cells were transiently transfected with each of these constructs and assayed for luciferase activity after 24 h. Error bars denote SEM from three experiments performed in triplicate. *P*-values were determined by repeated-measures ANOVA followed by Dunnett's multiple comparisons test (**P*<0.05). (B) Position Weight Matrix of GATA3 is shown relative to the negative strand of the sequences surrounding rs17432750. (C) GATA3 ChIP assays demonstrate enrichment of rs17432750 relative to the *CCND1* negative control. A GATA3 site from the ER- α enhancer was included as positive control. Results from two biological repeats are shown and error bars denote SD of three technical repeats. (D) Genotyping of rs17432750 in MCF7 genomic DNA versus MCF7 GATA3-ChIP DNA. Homozygous cell lines ZR751 (C/C), T47D (C/C), MDA-MB-468 (A/A) and no template controls (NTC) were included as reference for the assay. The risk (major) C allele is preferentially precipitated in the ChIP experiment. (E) Luciferase assay in Bre-80 cells showing the effect of GATA3 siRNA silencing on the activity of the *MAP3K1* promoter alone (Promoter) and with PRE-B3 constructs containing the C allele (ref PRE-B3) and protective A (minor) allele rs17432750 (rs17432750). Error bars denote SEM from three experiments performed in triplicate. *P*-values were determined by two-way repeated-measures

ANOVA followed either by Sidak's multiple comparisons test (to analyze the effect of *GATA3* knockdown within constructs) or Dunnett's multiple comparison test (to analyze differences in activity between constructs). (* $P < 0.05$, ** $P < 0.01$ and *** $P < 0.001$). The level of *GATA3* knockdown is shown in **Figure S9**.

Tables

iCHAV	SNP	Chr Position (GRCh37)	Alleles (major/ minor)	MAF	r^2 with lead iCHAV1 SNP	Imp r^2	Overall breast cancer risk				ER-positive breast cancer risk				ER-negative breast cancer risk				ER- positive /negative
							OR	(95%CI)	P-trend	P-cond	OR	(95%CI)	P-trend	P-cond	OR	(95%CI)	P-trend	P-cond	P-diff
1	rs10461612	56003831	G/C	0.18	0.96	0.97	1.21	(1.18-1.23)	1.08E-47		1.23	(1.20-1.26)	1.71E-43		1.09	(1.04-1.14)	4.13E-04		1.47E-05
1	rs7709971	56007339	G/A	0.18	0.96	0.97	1.21	(1.18-1.23)	1.05E-47		1.24	(1.21-1.27)	1.33E-43		1.09	(1.04-1.14)	4.86E-04		1.17E-05
1	rs7714232	56011357	A/T	0.18	0.94	0.99	1.21	(1.18-1.23)	4.76E-48		1.23	(1.20-1.26)	5.09E-44		1.09	(1.04-1.14)	5.64E-04		8.03E-06
1	rs12653202	56016918	A/C	0.18	0.94	1	1.21	(1.18-1.23)	2.17E-47		1.23	(1.20-1.26)	7.65E-42		1.09	(1.04-1.14)	5.04E-04		2.68E-05
1	rs59270457	56017887	T/G	0.18	0.96	1	1.21	(1.18-1.23)	1.22E-47		1.23	(1.20-1.26)	1.02E-42		1.09	(1.04-1.14)	4.48E-04		1.76E-05
1	rs61055995	56019064	A/T	0.18	0.96	1	1.21	(1.18-1.23)	1.16E-47		1.23	(1.20-1.26)	7.36-E43		1.09	(1.04-1.14)	4.70E-04		1.62E-05
1	rs16886165	56023083	T/G	0.18	0.96	1	1.21	(1.18-1.23)	1.14E-47		1.23	(1.20-1.26)	6.22E-43		1.09	(1.04-1.14)	4.56E-04		1.55E-05
1	rs16886181	56029243	T/C	0.18	0.96	1	1.21	(1.18-1.23)	4.80E-48		1.23	(1.20-1.26)	2.87E-43		1.10	(1.05-1.15)	2.69E-04		2.26E-05
1	chr5:56030827:I	56030827	T/TA	0.18	0.97	0.99	1.20	(1.18-1.23)	2.33E-47		1.23	(1.20-1.26)	1.27E-42		1.09	(1.04-1.14)	5.82E-04		1.24E-05
1	rs66893416	56051596	G/A	0.18	0.99	0.97	1.21	(1.18-1.23)	1.19E-47		1.23	(1.20-1.26)	4.60E-43		1.09	(1.04-1.14)	4.42E-04		1.05E-05
1	rs62355900	56052695	T/C	0.18	0.98	1	1.20	(1.18-1.23)	4.26E-47		1.23	(1.20-1.26)	8.93E-43		1.09	(1.04-1.14)	5.84E-04		1.21E-05
1	rs74345699	56053479	C/T	0.18	0.99	0.99	1.20	(1.18-1.23)	5.08E-47		1.23	(1.20-1.26)	1.57E-42		1.09	(1.04-1.14)	7.01E-04		1.05E-05
1	rs62355901	56053535	T/C	0.18	0.98	0.93	1.21	(1.18-1.24)	3.90E-47		1.24	(1.20-1.27)	5.41E-42		1.10	(1.04-1.15)	5.20E-04		1.49E-05
1	rs62355902 ^a	56053723	A/T	0.18	-	0.95	1.21	(1.19-1.24)	9.50E-49		1.24	(1.21-1.27)	5.71E-44		1.10	(1.05-1.15)	3.02E-04		1.47E-05
1	rs149188233	56053745	A/AAAA C	0.18	0.99	0.98	1.20	(1.18-1.23)	3.92E-47		1.23	(1.20-1.26)	1.00E-42		1.09	(1.04-1.14)	6.48E-04		1.13E-05
2a	rs77961606	56040643	T/C	0.05	0.28	0.98	1.27	(1.22-1.31)	1.03E-28	9.21E-05	1.29	(1.25-1.34)	3.42E-25	4.55E-04	1.09	(1.01-1.18)	3.59E-02	7.58E-01	1.04E-04
2a	rs16886272	56067434	A/G	0.07	0.19	0.95	1.21	(1.17-1.25)	8.89E-24	4.14E-05	1.25	(1.20-1.29)	3.41E-23	1.01E-05	1.07	(0.99-1.14)	8.47E-02	8.08E-01	5.42E-05
2a	rs113317823 ^a	56087883	T/C	0.08	0.19	0.72	1.22	(1.18-1.26)	7.00E-25	1.61E-05	1.24	(1.20-1.29)	2.02E-21	9.74E-05	1.12	(1.05-1.20)	2.56E-03	9.90E-02	1.93E-02
2a	rs77371588	56134560	G/T	0.05	0.26	0.98	1.27	(1.22-1.31)	1.36E-27	9.03E-05	1.29	(1.25-1.34)	2.01E-24	2.68E-04	1.11	(1.02-1.19)	2.11E-02	5.37E-01	3.32E-04
2a	rs60590641	56141155	G/A	0.05	0.26	0.98	1.27	(1.22-1.31)	1.16E-27	7.93E-05	1.30	(1.25-1.35)	1.08E-24	2.01E-04	1.11	(1.02-1.19)	2.10E-02	5.33E-01	3.05E-04
2a	rs61154548	56224720	C/T	0.05	0.24	0.97	1.27	(1.23-1.32)	4.32E-27	2.76E-05	1.30	(1.25-1.36)	4.03E-24	9.06E-05	1.10	(1.01-1.19)	3.65E-02	6.19E-01	3.37E-04
2a	chr5:56237882:D	56237882	CAGTTA AGTTTA/ C	0.04	0.23	0.93	1.28	(1.24-1.33)	1.82E-25	8.94E-05	1.32	(1.26-1.37)	3.49E-23	1.57E-04	1.10	(1.01-1.20)	3.80E-02	6.02E-01	5.14E-04
2a	rs79565352	56251753	G/A	0.06	0.20	1	1.23	(1.19-1.27)	7.45E-24	6.46E-05	1.25	(1.20-1.29)	8.52E-20	7.69E-04	1.11	(1.03-1.19)	1.42E-02	3.04E-01	2.20E-03
2a	rs74571895	56254772	A/G	0.05	0.20	0.99	1.24	(1.20-1.28)	1.71E-24	3.62E-05	1.26	(1.21-1.30)	1.54E-20	3.99E-04	1.11	(1.03-1.19)	1.16E-02	2.70E-01	2.10E-03
2a	rs78743305	56256217	G/A	0.05	0.20	0.99	1.24	(1.20-1.28)	1.41E-24	3.28E-05	1.26	(1.21-1.30)	1.20E-20	3.54E-04	1.11	(1.03-1.19)	1.11E-02	2.64E-01	2.03E-03
2a	rs7726354	56256483	C/T	0.06	0.20	1	1.23	(1.19-1.27)	4.51E-24	5.13E-05	1.25	(1.20-1.30)	4.61E-20	5.90E-04	1.11	(1.03-1.19)	1.30E-02	2.88E-01	2.11E-03

2a	rs79760198	56262639	T/C	0.05	0.23	0.98	1.27	(1.23-1.31)	1.39E-26	2.71E-05	1.30	(1.25-1.35)	3.04E-23	1.34E-04	1.11	(1.03-1.20)	1.74E-02	3.91E-01	1.23E-03
2a	rs79041328	56264375	A/G	0.05	0.23	0.99	1.27	(1.23-1.31)	1.48E-26	2.70E-05	1.30	(1.24-1.35)	3.42E-23	1.38E-04	1.11	(1.03-1.20)	1.60E-02	3.71E-01	1.42E-03
2a	rs6893174	56265025	C/T	0.05	0.22	1	1.26	(1.22-1.30)	8.59E-26	4.75E-05	1.28	(1.23-1.33)	2.83E-22	2.77E-04	1.11	(1.02-1.19)	1.88E-02	4.02E-01	1.56E-03
2a	rs113325879	56266142	G/A	0.05	0.23	0.99	1.27	(1.23-1.31)	1.30E-26	2.53E-05	1.30	(1.25-1.35)	3.73E-23	1.45E-04	1.11	(1.03-1.20)	1.52E-02	3.59E-01	1.47E-03
2a	rs142258027	56266278	G/A	0.05	0.22	0.96	1.27	(1.22-1.31)	8.33E-26	4.57E-05	1.29	(1.24-1.34)	1.83E-22	2.34E-04	1.11	(1.02-1.20)	1.97E-02	4.12E-01	1.46E-03
2a	rs113173541	56266689	C/A	0.05	0.23	0.98	1.27	(1.23-1.31)	1.34E-26	2.57E-05	1.30	(1.25-1.35)	4.05E-23	1.51E-04	1.11	(1.03-1.20)	1.52E-02	3.60E-01	1.49E-03
2a	rs78925509	56267141	A/G	0.05	0.23	0.98	1.27	(1.23-1.31)	1.41E-26	2.63E-05	1.30	(1.25-1.35)	4.38E-23	1.56E-04	1.11	(1.03-1.20)	1.55E-02	3.63E-01	1.49E-03
2a	rs73122135	56267308	G/A	0.05	0.23	0.96	1.27	(1.22-1.31)	1.93E-26	3.14E-05	1.29	(1.24-1.35)	4.03E-23	1.63E-04	1.11	(1.02-1.19)	1.98E-02	4.30E-01	1.02E-03
2a	rs80089016	56267444	G/A	0.05	0.23	0.98	1.27	(1.23-1.32)	1.43E-26	2.64E-05	1.30	(1.25-1.35)	4.61E-23	1.59E-04	1.11	(1.03-1.20)	1.54E-02	3.62E-01	1.52E-03
2a	rs73122138	56267949	G/A	0.05	0.23	0.96	1.26	(1.22-1.31)	6.61E-26	5.47E-05	1.29	(1.24-1.34)	1.54E-22	3.19E-04	1.11	(1.02-1.19)	2.27E-02	4.66E-01	1.17E-03
2a	rs80310238	56267984	C/A	0.05	0.23	0.98	1.27	(1.23-1.32)	1.45E-26	2.67E-05	1.30	(1.25-1.35)	5.00E-23	1.66E-04	1.11	(1.03-1.20)	1.55E-02	3.64E-01	1.53E-03
2a	rs77706078	56268686	G/A	0.05	0.23	0.98	1.27	(1.23-1.32)	1.40E-26	2.63E-05	1.30	(1.25-1.35)	5.05E-23	1.67E-04	1.11	(1.03-1.20)	1.52E-02	3.59E-01	1.57E-03
2a	chr5:56268884:I	56268884	C/CTGG GAGGT	0.04	0.22	0.96	1.28	(1.24-1.33)	8.44E-27	1.76E-05	1.31	(1.25-1.36)	3.05E-23	1.16E-04	1.12	(1.03-1.21)	1.15E-02	3.00E-01	2.05E-03
2a	rs2408651	56268991	T/C	0.05	0.23	0.95	1.26	(1.22-1.31)	1.08E-25	6.53E-05	1.29	(1.24-1.34)	2.10E-22	3.50E-04	1.11	(1.02-1.19)	2.19E-02	4.57E-01	1.19E-03
2a	rs79459889	56269286	G/A	0.05	0.23	0.98	1.27	(1.23-1.31)	2.09E-26	3.08E-05	1.29	(1.24-1.35)	8.65E-23	2.06E-04	1.12	(1.03-1.20)	1.43E-02	3.49E-01	1.74E-03
2a	rs111773762	56269336	T/C	0.05	0.23	0.98	1.27	(1.23-1.31)	2.53E-26	3.33E-05	1.29	(1.24-1.35)	7.04E-23	1.87E-04	1.11	(1.03-1.20)	1.50E-02	3.58E-01	1.57E-03
2a	rs80097053	56269492	C/T	0.05	0.23	0.98	1.27	(1.23-1.31)	2.50E-26	3.31E-05	1.29	(1.24-1.35)	7.03E-23	1.87E-04	1.12	(1.03-1.20)	1.49E-02	3.56E-01	1.58E-03
2a	rs59536253	56269512	A/G	0.05	0.23	0.96	1.27	(1.22-1.31)	2.84E-26	3.73E-05	1.29	(1.24-1.34)	8.40E-23	2.26E-04	1.11	(1.03-1.20)	1.64E-02	3.88E-01	1.40E-03
2a	rs112032073	56270717	G/T	0.05	0.23	0.98	1.27	(1.23-1.31)	2.40E-26	3.27E-05	1.30	(1.24-1.35)	7.36E-23	1.92E-04	1.12	(1.03-1.20)	1.41E-02	3.44E-01	1.69E-03
2b	rs12659430	56032619	A/G	0.13	0.62	1.00	1.15	(1.12-1.18)	1.73E-19	4.42E-05	1.17	(1.13-1.20)	3.95E-18	1.81E-03	1.08	(1.02-1.14)	8.95E-03	6.26E-01	1.96E-02
2b	rs62355881	56033093	T/C	0.13	0.62	1.00	1.15	(1.12-1.18)	1.61E-19	4.68E-05	1.17	(1.13-1.20)	3.62E-18	1.92E-03	1.08	(1.02-1.14)	8.95E-03	6.26E-01	1.93E-02
2b	rs21113084	56033671	T/C	0.13	0.62	1.00	1.15	(1.12-1.18)	1.97E-19	4.02E-05	1.17	(1.13-1.20)	4.80E-18	1.62E-03	1.08	(1.02-1.14)	8.92E-03	6.27E-01	2.02E-02
2b	rs62355882	56045110	C/T	0.13	0.60	1.00	1.15	(1.12-1.18)	3.69E-19	7.81E-05	1.16	(1.13-1.20)	2.64E-17	1.06E-03	1.08	(1.02-1.14)	1.22E-02	5.60E-01	2.42E-02
2b	rs62355899 ^a	56050465	A/G	0.13	0.62	1.00	1.15	(1.12-1.18)	2.98E-19	3.04E-05	1.17	(1.13-1.20)	5.52E-18	2.04E-03	1.08	(1.02-1.14)	1.11E-02	5.42E-01	1.86E-02
2b	rs10513090	56054065	T/C	0.13	0.62	1.00	1.15	(1.12-1.18)	1.91E-19	3.67E-05	1.17	(1.13-1.20)	2.92E-18	2.79E-03	1.08	(1.02-1.13)	1.43E-02	4.46E-01	1.32E-02
2b	rs6895844	56057940	G/A	0.13	0.62	1.00	1.15	(1.12-1.18)	7.44E-20	7.50E-05	1.17	(1.13-1.20)	1.17E-18	4.66E-03	1.07	(1.02-1.13)	1.63E-02	3.99E-01	1.08E-02
2b	rs6882255	56058171	T/A	0.13	0.63	0.97	1.15	(1.12-1.18)	3.03E-20	9.13E-05	1.17	(1.14-1.21)	8.02E-19	5.26E-03	1.08	(1.02-1.13)	1.47E-02	4.15E-01	9.86E-03
2b	rs12654584	56058415	T/G	0.13	0.62	1.00	1.15	(1.12-1.18)	5.90E-20	8.57E-05	1.17	(1.13-1.20)	8.79E-19	5.31E-03	1.07	(1.02-1.13)	1.61E-02	4.01E-01	1.01E-02
2b	rs12654176	56058487	A/G	0.13	0.62	1.00	1.15	(1.12-1.18)	4.29E-20	9.99E-05	1.17	(1.14-1.20)	8.04E-19	5.66E-03	1.08	(1.02-1.13)	1.33E-02	4.64E-01	1.19E-02
3	rs17432750	56031822	C/A	0.16	0.04	0.98	0.92	(0.89-0.94)	4.55E-11	1.61E-04	0.90	(0.87-0.93)	9.03E-11	1.48E-04	1.01	(0.96-1.06)	6.53E-01	2.44E-01	5.13E-05

3	rs199945768	56033233	CG/C	0.16	0.04	0.92	0.91	(0.89-0.94)	6.35E-11	2.66E-04	0.90	(0.87-0.93)	1.90E-10	3.09E-04	1.01	(0.96-1.06)	6.61E-01	2.49E-01	6.53E-05
3	rs11956804	56033233	C/A	0.16	0.04	0.94	0.91	(0.89-0.94)	7.39E-11	1.86E-04	0.90	(0.87-0.93)	1.90E-10	2.05E-04	1.01	(0.96-1.06)	6.61E-01	2.22E-01	7.44E-05
3	rs11949391 ^a	56045081	T/C	0.16	0.04	1	0.91	(0.89-0.94)	9.36E-12	5.57E-05	0.90	(0.87-0.93)	1.00E-10	1.44E-04	1.01	(0.96-1.06)	8.14E-01	3.50E-01	1.27E-04
3	rs6884514	56062614	C/T	0.16	0.04	1	0.92	(0.89-0.95)	6.04E-10	5.80E-04	0.91	(0.88-0.94)	1.90E-09	6.74E-04	1.01	(0.96-1.06)	6.15E-01	2.30E-01	1.13E-04

Table 1. iCHAVs with the strongest effects on breast cancer risk in European studies

Single SNP risk estimates for the top candidates in each iCHAV with overall breast cancer risk and subtypes by estrogen receptor status. Results are given as ORs with 95% CI (using the minor variant alleles as the reference), per-allele *P*-trend and *P*-cond, (*P*-cond being conditional on the iCHAV1 lead SNP rs62355902). The ER-positive/negative *P*-diff is from a case-only analysis, comparing the effect sizes in the subtypes by estrogen receptor status. For iCHAVs 2a and 2b, only SNPs with $P\text{-cond} < 1 \times 10^{-4}$ for overall breast cancer risk are shown here - full results are listed in **Table S5** and all other iCHAV2a and 2b candidate variants are listed in **Table S7**.

^aThe best candidate SNP in the iCHAV.

Table 2. Breast cancer risk in Europeans by haplotypes of five iCHAV representative SNPs

Haplotype	iCHAV1 ^a	iCHAV2a ^b	iCHAV2b ^c	iCHAV2c ^d	iCHAV3 ^e	Frequency	Overall Breast cancer			ER+ Breast cancer			ER- Breast cancer		
							OR	(95%CI)	P-value	OR	(95%CI)	P-value	OR	(95%CI)	P-value
h ₀ ^f	1	1	1	1	1	0.52	-	-	-	-	-	-	-	-	-
h1	1	1	1	2	1	0.12	1.01	(0.99-1.05)	4.50E-01	1.03	(0.99-1.07)	1.12E-01	1.04	(0.98-1.11)	2.08E-01
h2	1	1	1	1	2	0.15	0.95	(0.92-0.97)	2.08E-04	0.95	(0.92-0.99)	5.23E-03	1.04	(0.98-1.10)	1.67E-01
h3	1	2	1	1	1	0.02	1.08	(0.99-1.17)	7.57E-02	1.13	(1.02-1.24)	1.68E-02	1.08	(0.92-1.27)	3.45E-01
h4	1	2	1	2	1	0.01	1.05	(0.99-1.17)	3.81E-01	0.99	(0.87-1.13)	8.75E-01	1.17	(0.96-1.42)	1.12E-01
h5	2	1	2	1	1	0.11	1.16	(1.12-1.20)	2.00E-20	1.19	(1.14-1.23)	9.37E-20	1.10	(1.04-1.18)	1.82E-03
h6	2	2	1	2	1	0.05	1.31	(1.25-1.37)	1.66E-29	1.34	(1.27-1.42)	3.74E-27	1.15	(1.05-1.25)	3.64E-03

^a rs62355902, representing 15 SNPs^b rs113317823, representing 90 SNPs which have $r^2 \geq 0.53$ with rs113317823 and r^2 in [0.19-0.29] with rs62355902^c rs62355899, representing 66 SNPs which have $r^2 \leq 0.01$ with rs113317823 and r^2 in [0.59-0.62] with rs62355902^d rs7721581, representing 17 SNPs which have r^2 in [0.14-0.16] with rs113317823 and $r^2 \leq 0.01$ with rs62355902^e rs11949391, representing 5 SNPs^f h₁-h₆ are compared to h₀ (the reference haplotype carrying the major alleles of all 5 SNPs).

'1' represents major alleles and '2' minor alleles in each SNP

Figure 1
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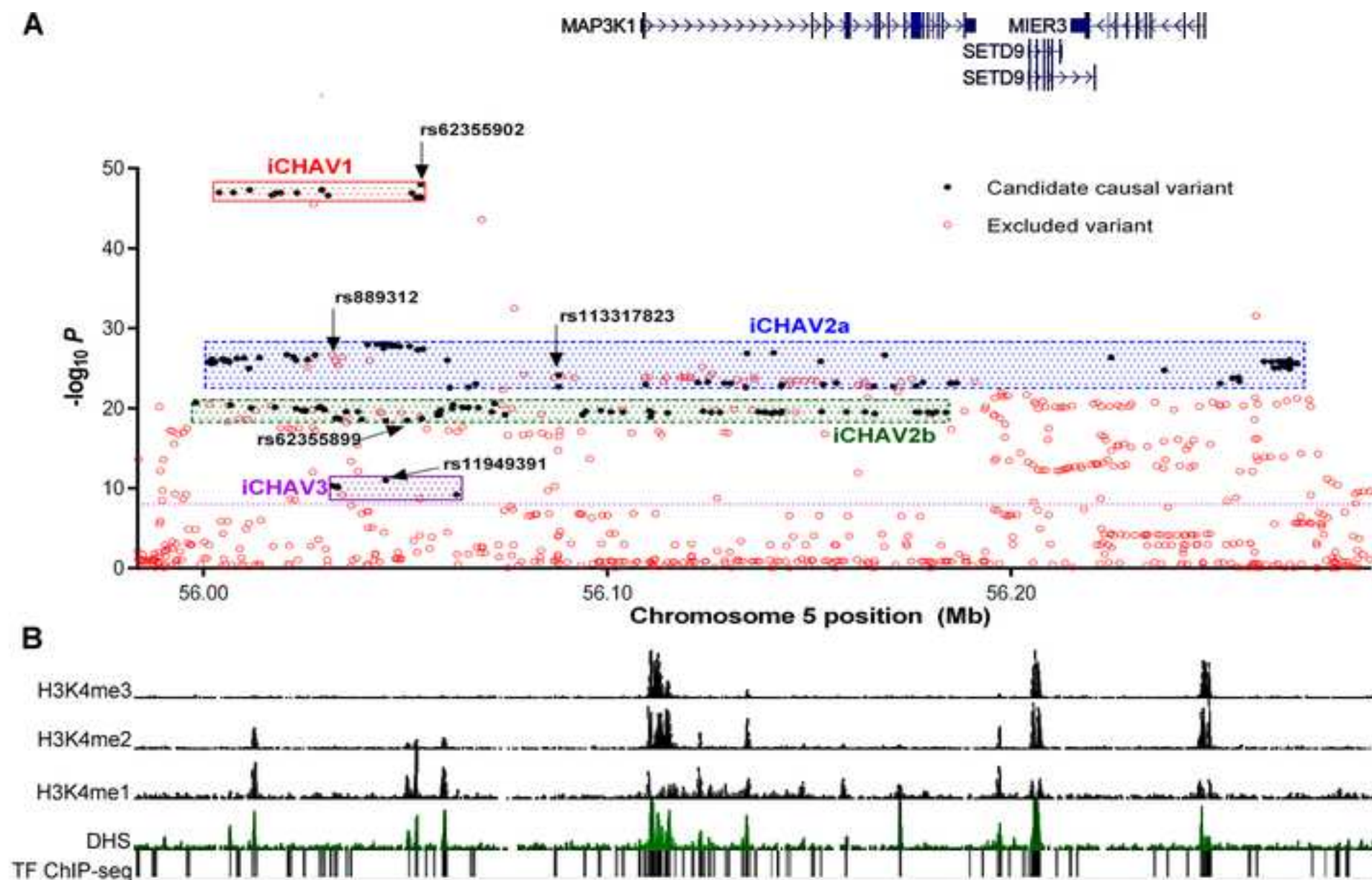


Figure 2
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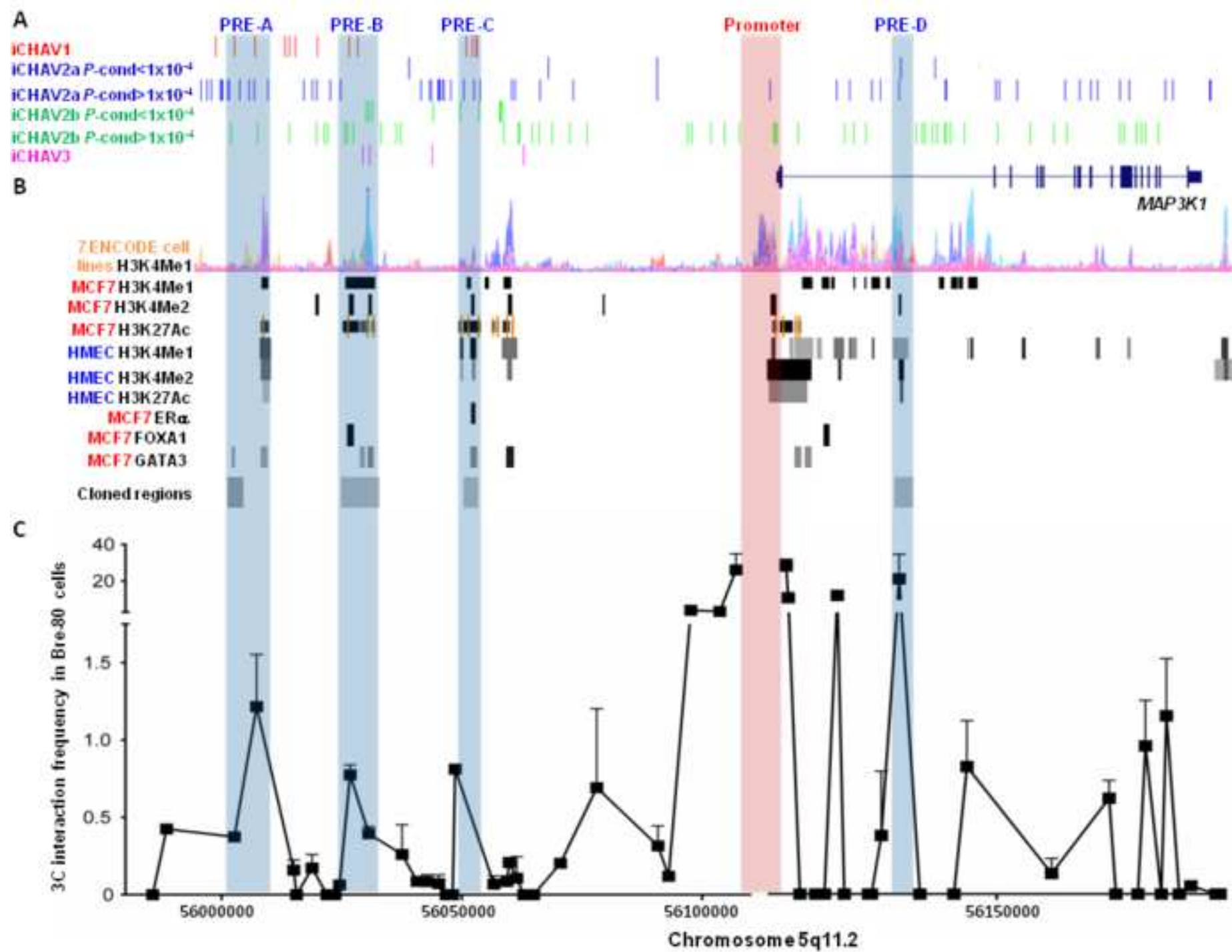


Figure 3
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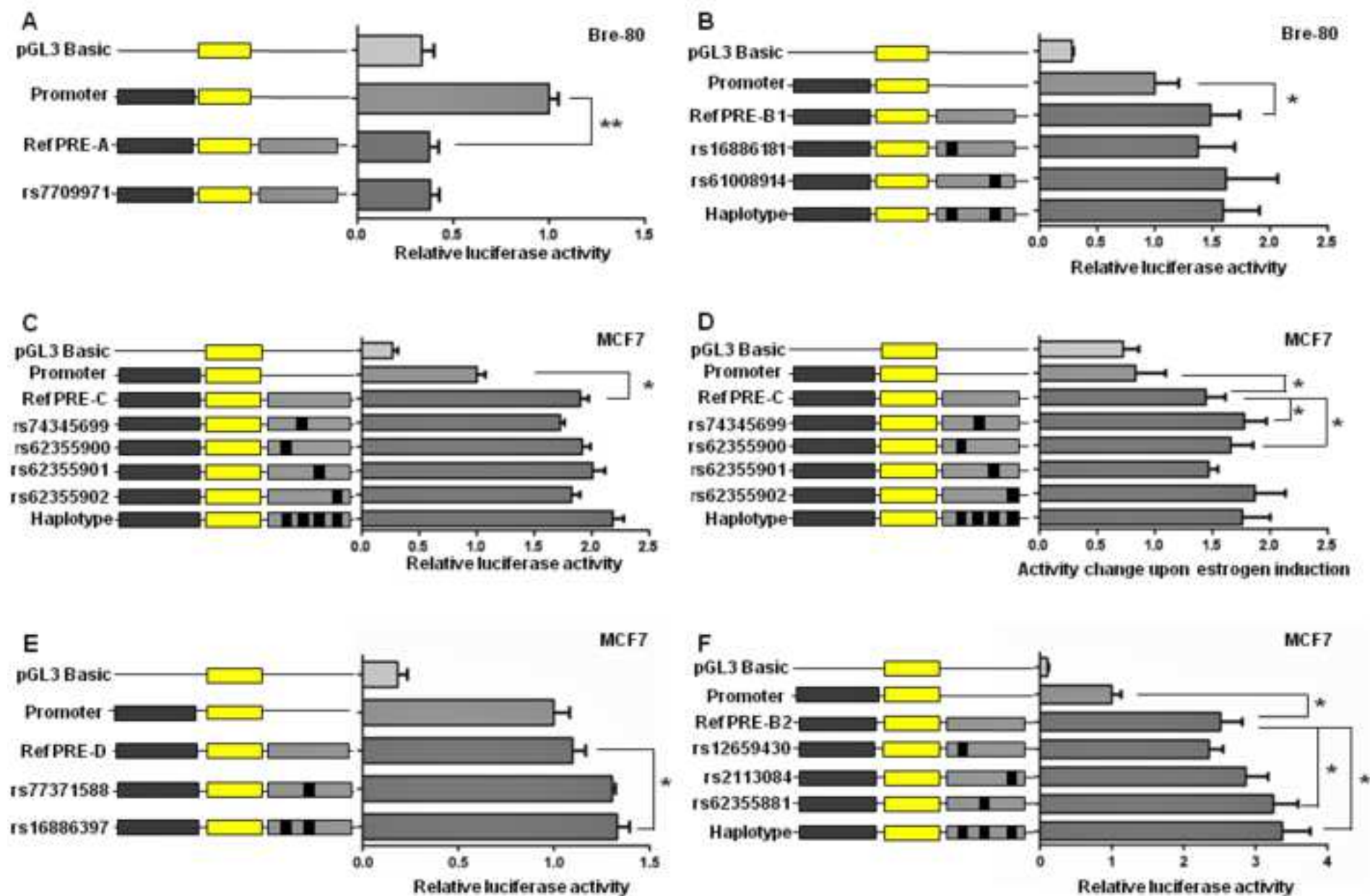
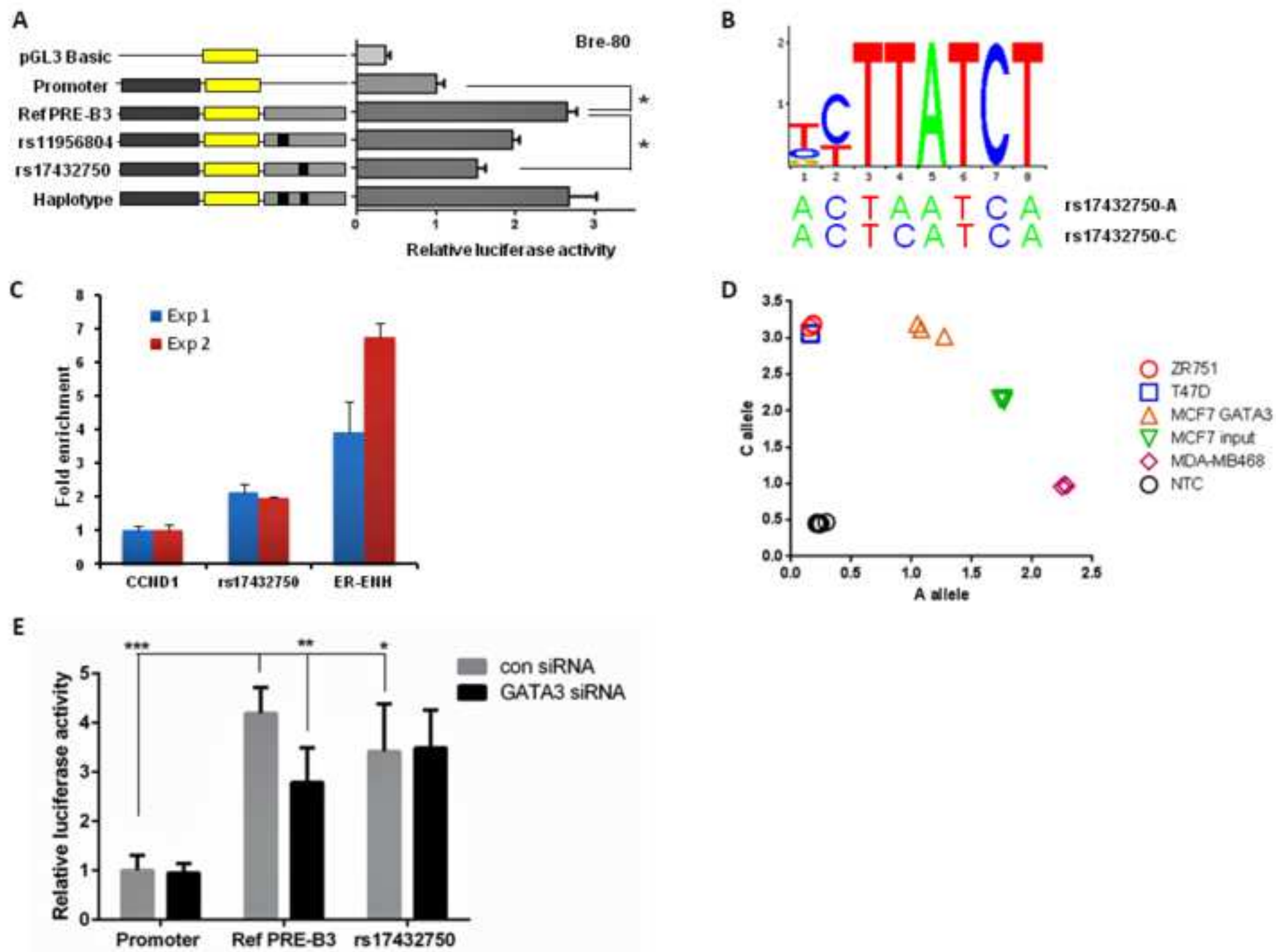


Figure 4
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Supplementary Data

Supplemental Acknowledgements

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BCAC investigating groups:

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Supplemental Figures

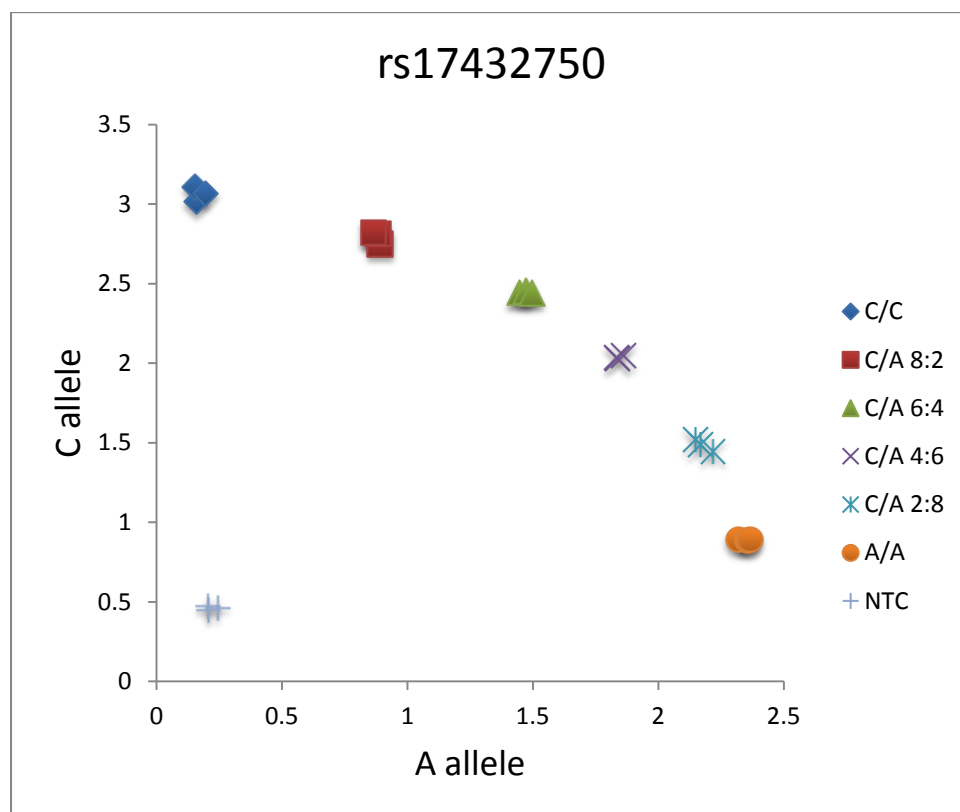
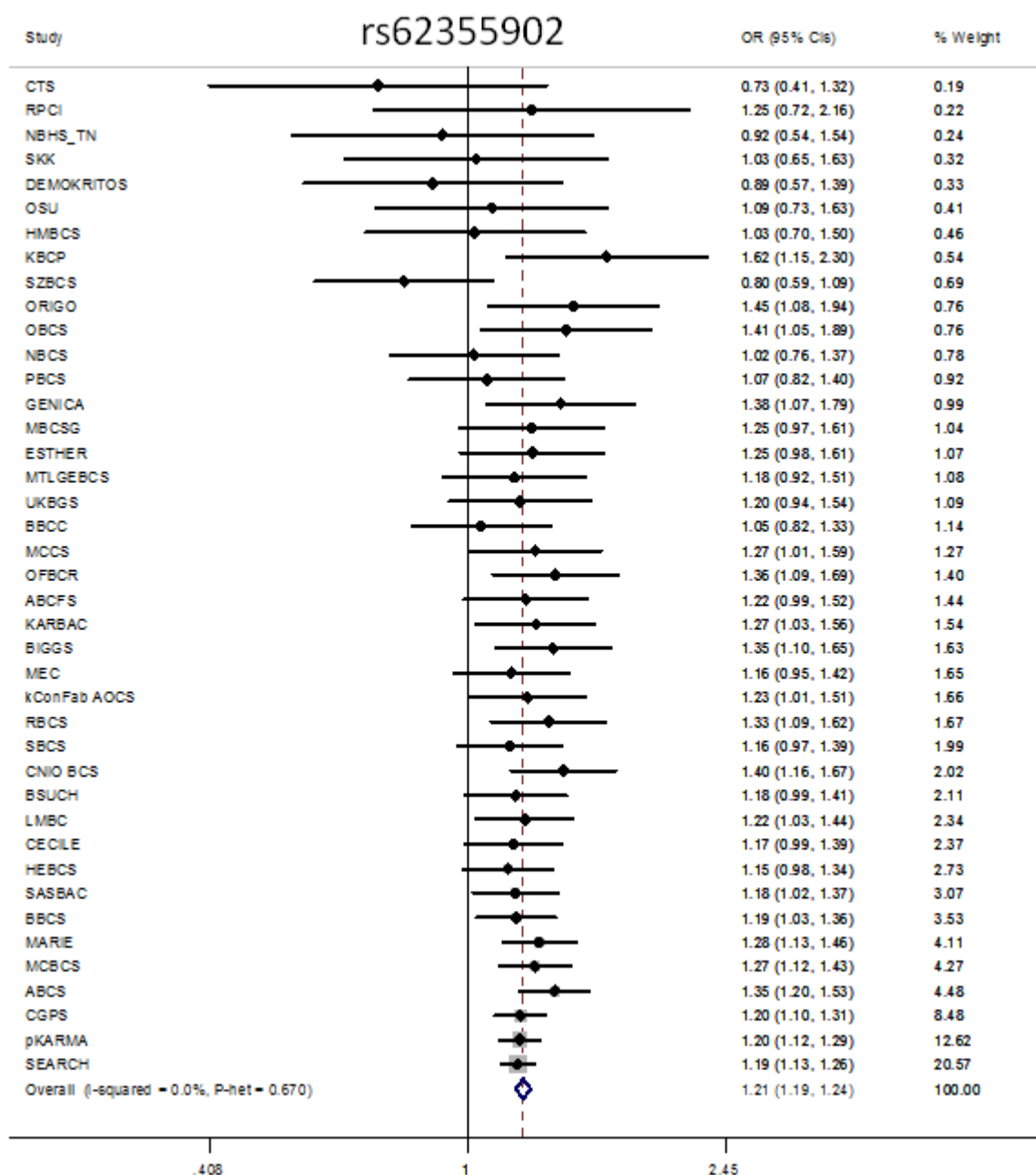
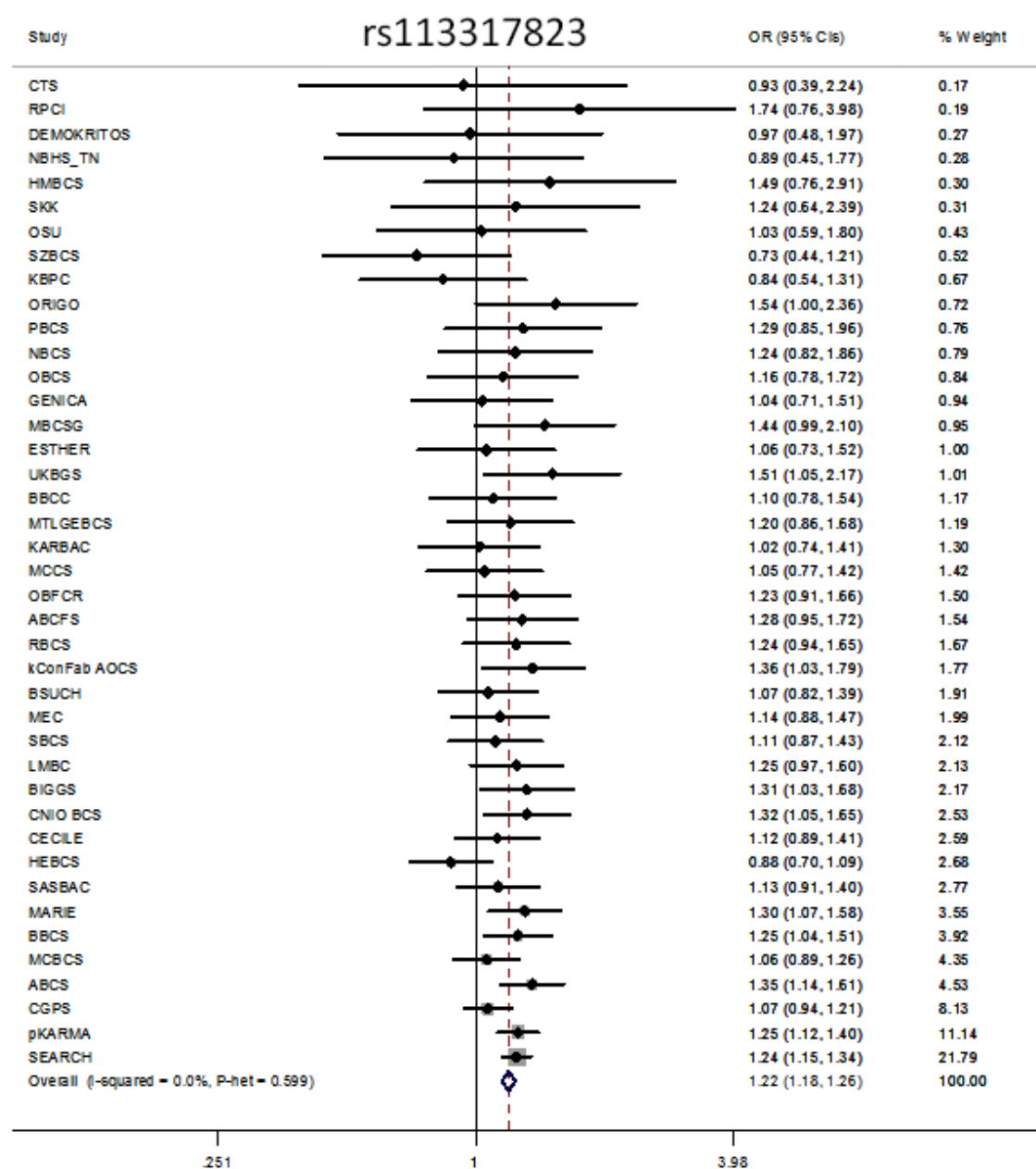


Figure S1. Allelic Discrimination plot of known ratios of genomic DNA from breast cell lines carrying the C/C (ZR751) or A/A (MDA-MB-468) at rs17432750.





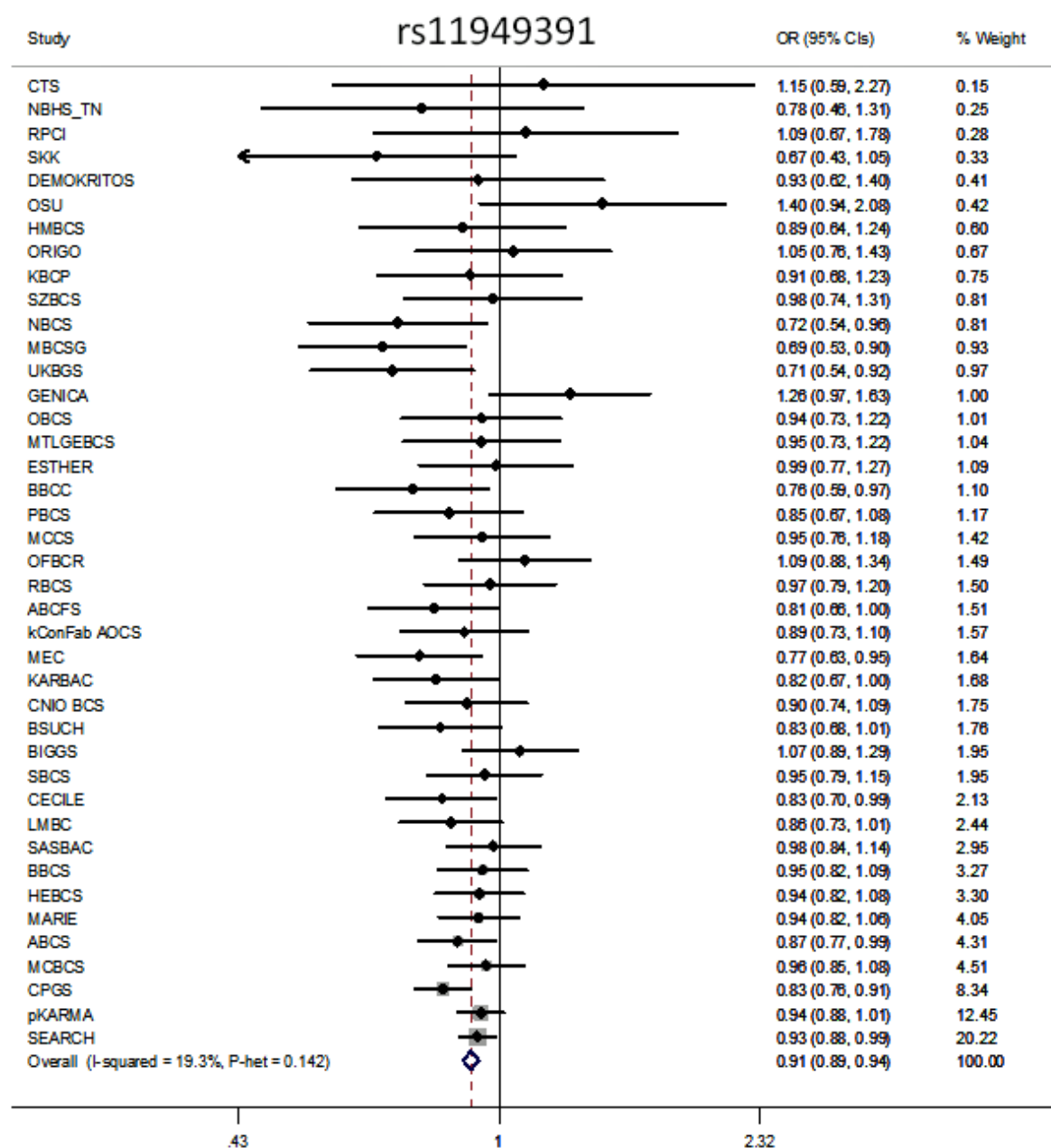


Figure S2. Forest plots for the three independently-associated SNPs marking iCHAVs 1-3.

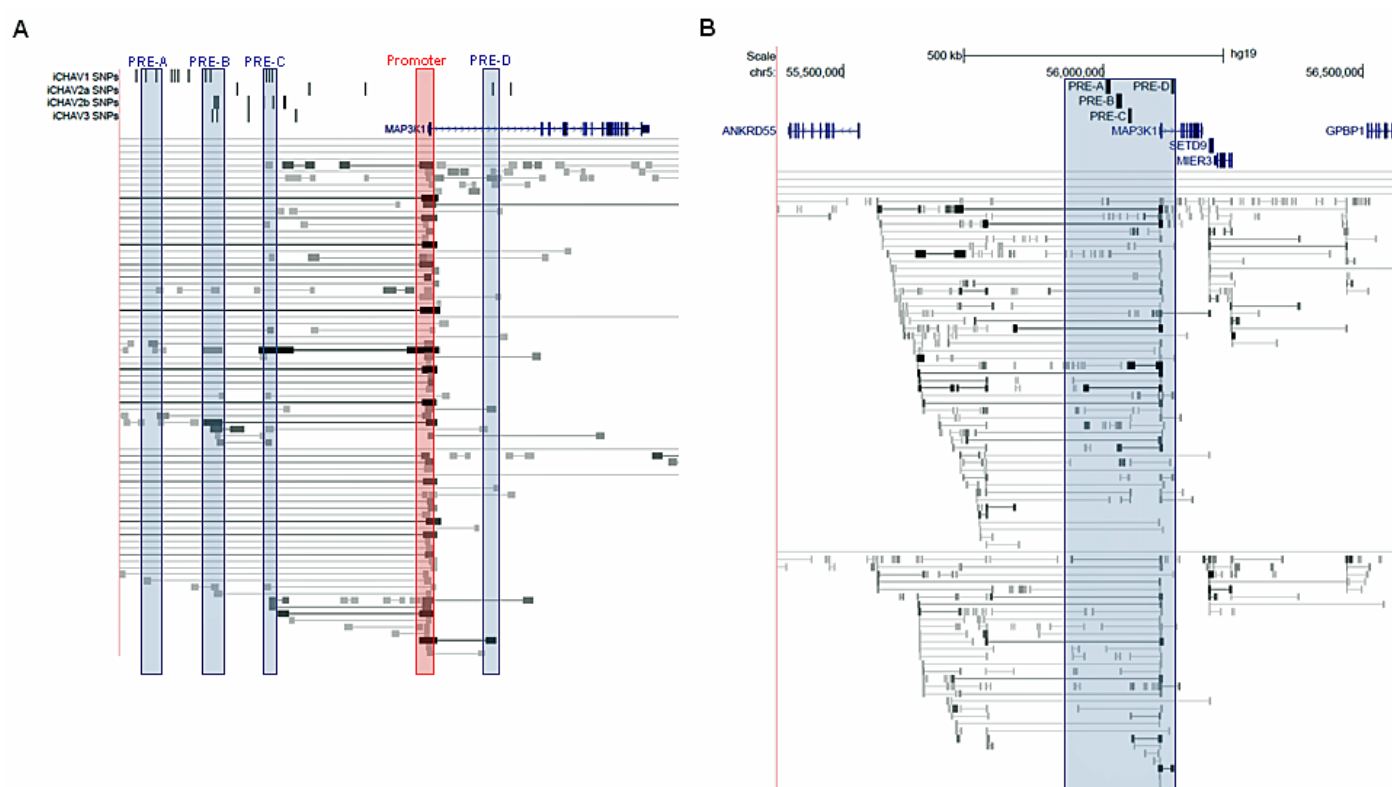


Figure S3. RNA polymerase II ChIA-PET at the 5q11.2 breast cancer risk locus in MCF7 cells shows chromatin interactions between discrete genomic regions and the *MAP3K1* promoter. ENCODE data (GEO sample accession GSM970209) was accessed using the UCSC genome browser. iCHAV variants and PREs are shown. Panel (A) shows the zoomed in region containing the PREs and *MAP3K1*. Panel (B) shows the 5q11.2 risk locus and flanking genes.

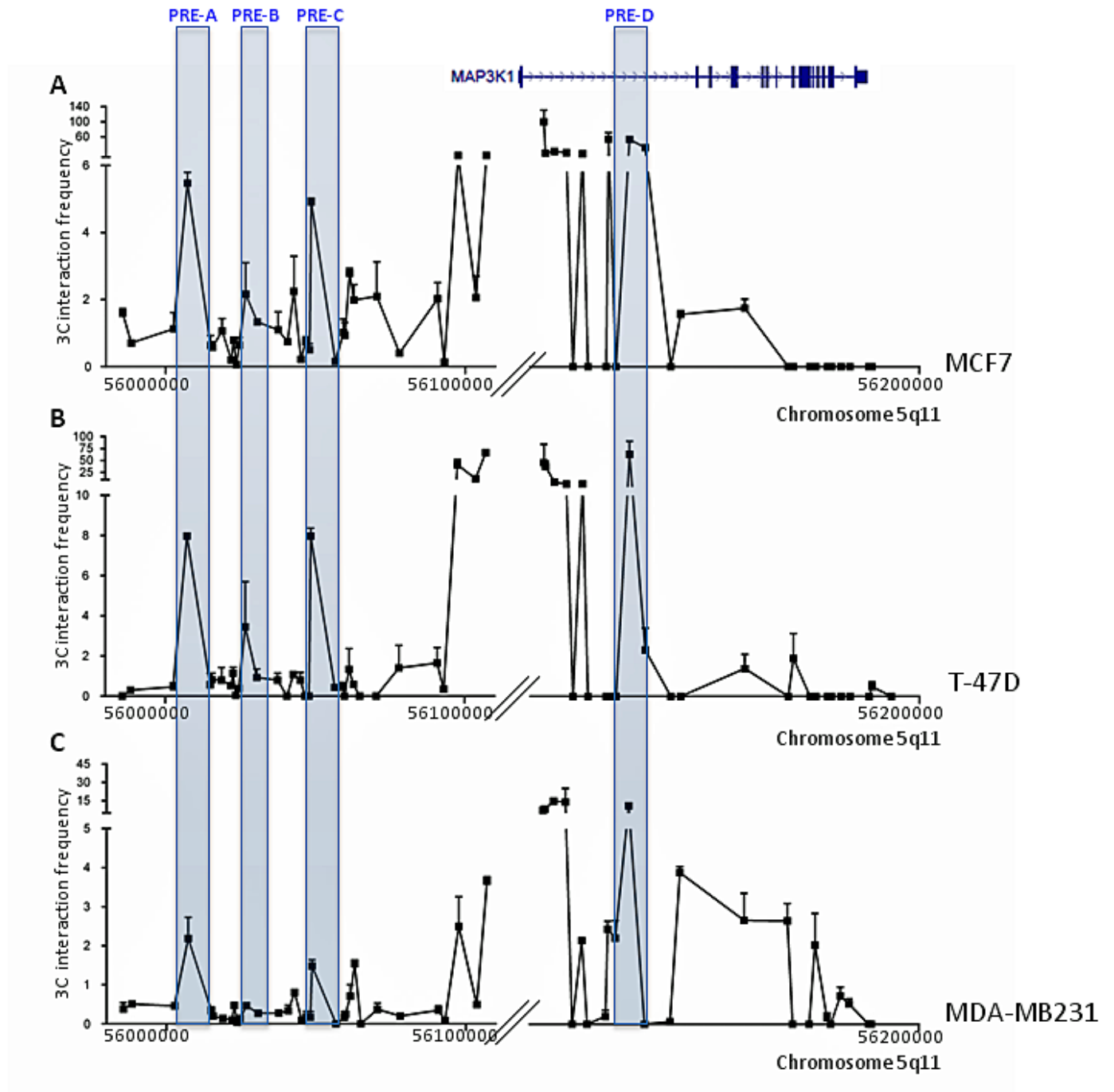


Figure S4. Chromatin interactions with the *MAP3K1* promoter in breast cancer cell lines at the 5q11.2 risk locus. 3C libraries were generated with EcoRI. Chromatin interaction frequencies were plotted at the chromosomal position of the corresponding EcoRI fragments for MCF7 (A), T-47D (B) and MDA-MB231 (C) libraries, respectively. PREs are highlighted at their chromosomal locations. A representative graph of at least two biological replicates is shown and error bars represent SD.

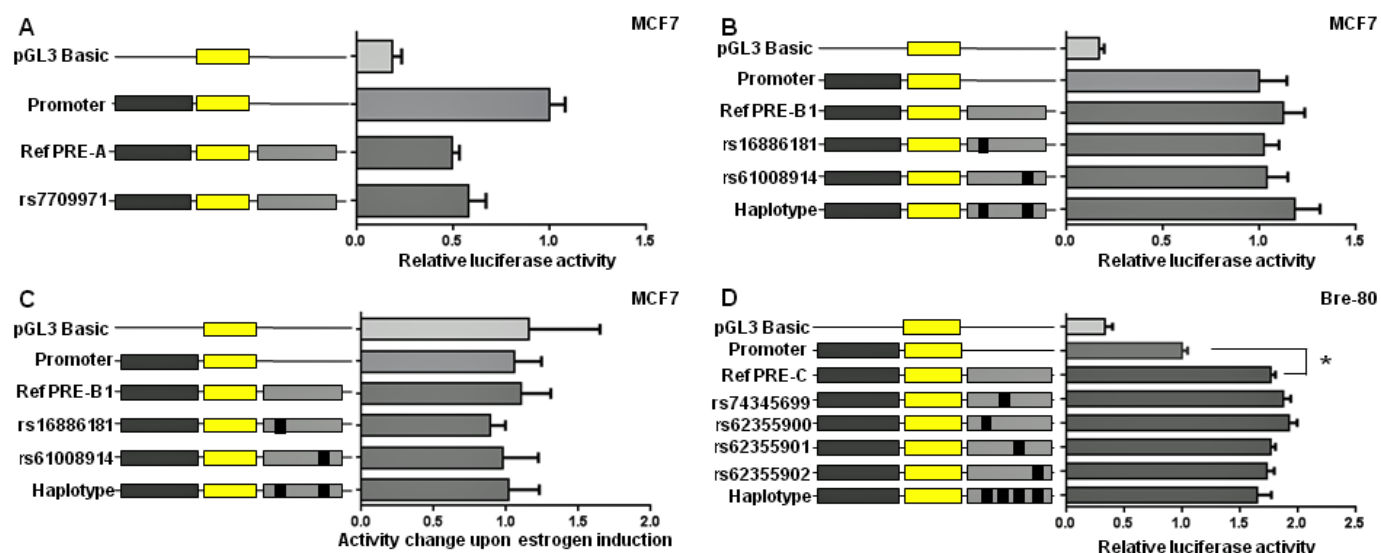


Figure S5. Luciferase reporter assays of PRE constructs containing iCHAV1 SNPs. MCF7 cells were transiently transfected with PRE-A (A) or PRE-B1 (B) and Bre-80 cells with PRE-C (D) constructs under basal conditions and assayed for luciferase activity after 24 h. Panel (C) shows results from luciferase assays after estrogen induction of MCF7 cells transfected with PRE-B1 constructs. For each reporter construct, the luciferase activity of estrogen treated cells was normalized to the activity of the corresponding vehicle treated cells. Error bars denote SEM from three experiments performed in triplicate. *P*-values were determined by repeated-measures ANOVA followed by Dunnett's multiple comparisons test (**P*<0.05).

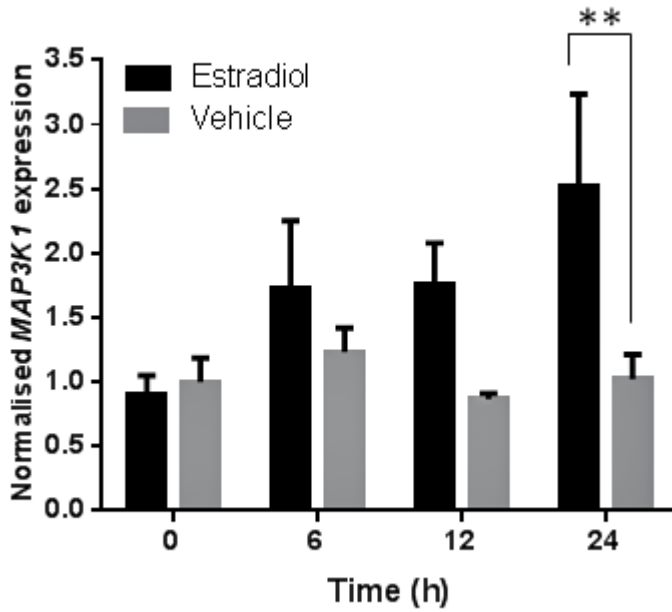


Figure S6. *MAP3K1* expression is induced in MCF7 cells after estrogen treatment. MCF7 cells were treated with either 100 nM estradiol or vehicle for 24 h. *MAP3K1* expression was measured at baseline and after 6, 12 and 24 h by qPCR and normalized using *GUS* expression as an internal control. Error bars denote SEM from three experiments performed in triplicate. Differences in expression between vehicle and estradiol treated cells were tested using a two-way repeated measures ANOVA followed by Sidak's multiple comparisons test (** $P < 0.01$).

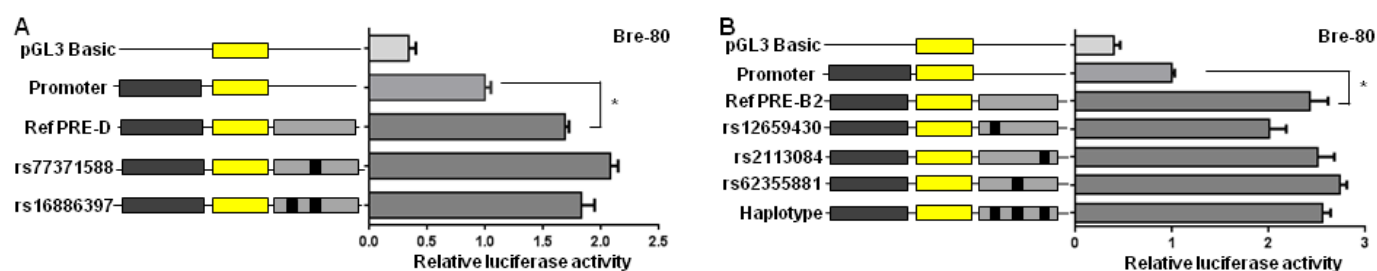


Figure S7. Luciferase reporter assays of PRE-D constructs containing iCHAV2a SNPs and PRE-B2 constructs containing iCHAV2b variants in Bre-80 cells. Bre-80 cells were transfected with PRE-D and PRE-B2 constructs. Cells were assayed for luciferase activity after 24 h. Error bars denote SEM from three experiments performed with triplicates. Statistical significance was determined by repeated-measures ANOVA followed by Dunnett's multiple comparisons test (* $P<0.05$).

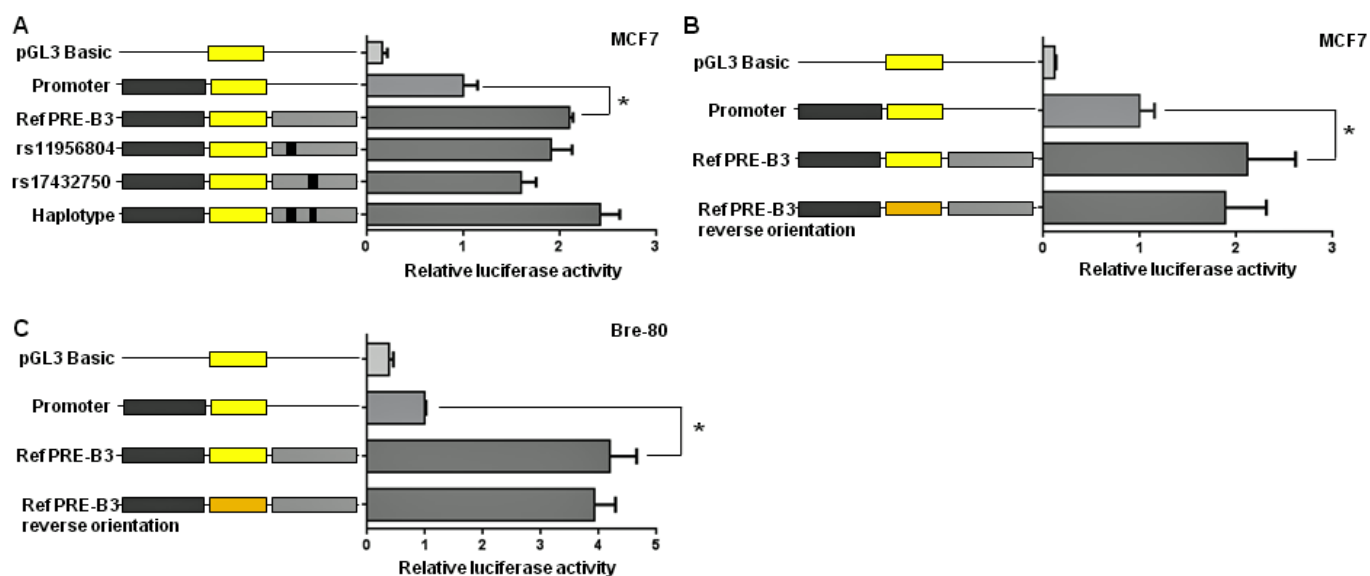


Figure S8. Luciferase reporter assays of PRE-B3 constructs containing iCHAV3 SNPs in MCF7 and Bre-80 cells. MCF7 cells were transiently transfected the PRE-B3 reference and variant luciferase constructs (A) or the PRE-B3 reference construct in the reverse orientation (B). Bre-80 cells were also transiently transfected with the reference PRE-B3 construct in the reverse orientation (C). Cells were assayed for luciferase activity after 24 h. Error bars denote SEM from three experiments performed with triplicates. *P*-values were determined by ANOVA followed by Dunnett's multiple comparisons test (**P*<0.05).

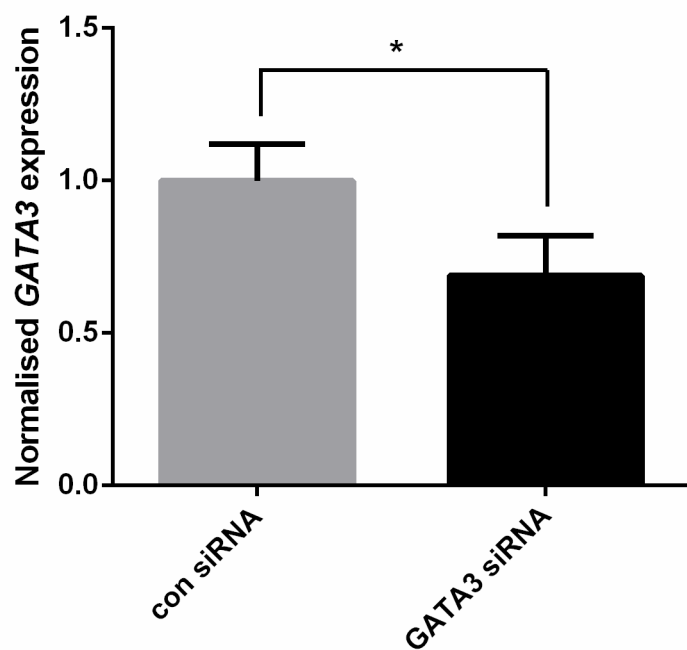


Figure S9. TaqMan qPCR assay confirming knockdown of *GATA3* in Bre80 cells. *GATA3* expression from cells transfected with non-targeting siRNA and *GATA3* is denoted by con siRNA and GATA3 siRNA, respectively. Error bars denote SEM from three experiments performed with triplicates. Statistical significance was determined by a paired t-test (* $P < 0.05$).

Supplemental tables

Table S1. 3C qPCR primers

3C primers	EcoRI site (GRCh37 chr 5 coordinates)	Sequence
Fragment 1	55,988,536	aaccttttcctgttctgcttaaggtgggtgg
Fragment 2	56,001,687	aagacagccaaagactctccaattccttctgc
Fragment 3	56,002,653	cactggctcagtcacctcattgattcagtc
Fragment 4	56,015,010	caaagaaacgagctctaagaggtgggcagc
Fragment 5	56,015,005	gaggcagatttcctggaaccttctttcc
Fragment 6	56,015,605	cctgagtctcattccctcttctgcaagagacc
Fragment 7	56,021,916	aagagtctcgtctgtcagacaaaatgtcacgg
Fragment 8	56,022,688	aagaatcactggggtgttattgctggaaagg
Fragment 9	56,023,585	ctcatggcctatgtacctcaaaggctcc
Fragment 10	56,024,547	catgtaactgctgggtttcattccagttcagg
Fragment 11	56,026,749	ggttctatgtgaagggtgccctccaaaacc
Fragment 12	56,030,623	aatccagcacctgctatgaacacctcatcc
Fragment 13	56,037,547	cctaagtcttctccagagcattttgacctgc
Fragment 14	56,040,674	atcaggaggactacaatctaggggtatgctgagg
Fragment 15	56,042,804	gcaggacttcttctcagcatcttctctttcc
Fragment 16	56,045,182	tgggactgtgggagacacacagttttgc
Fragment 17	56,046,719	agcaatcagtaatgggaaaagatgccatgc
Fragment 18	56,047,998	ctcttctgccttgttctgacctgtgtcc
Fragment 19	56,048,601	tctaggtgtgtctgcaagggcatttctgg
Fragment 20	56,056,381	gccagtgtgtttgattccaaacctctgagc
Fragment 21	56,056,646	tggaacatgctttcactaaagccttctaggaagc
Fragment 22	56,059,823	gcagtgaatgaggtgacacagatctcc
Fragment 23	56,061,410	gcgtttgaactatccagagttaacacgcatcc
Fragment 24	56,062,862	aaggtgctgtcttatgaaagaggaaaccgagg
Fragment 25	56,064,941	ggaggaaagagtaggagacggctgttcagc
Fragment 26	56,070,521	agagtaagtaatgacctgccacggcagg
Fragment 27	56,078,074	gacgaggtaggataattagtggaaccttaggc
Fragment 28	56,090,803	tcatagtgggtatggaatggtattgcattgtgc
Fragment 29	56,093,049	gtgaacacaagccgagatttcagaaggtagc
Fragment 30	56,097,598	ctatcctaccaatttgccagctaagtgattggg
Fragment 31	56,103,646	cagtttccctgatcccctattcccgc
Fragment 32	56,107,073	ttacaggaggatcttgggaaaatgtatgaaccc
Fragment 33	56,108,495	tctagccccagcactcctgcaagtattcc
Fragment 34	56,109,036	gcatgcgtgaaatcattgcaaggtttacg
Bait (promoter fragment)	56,112,866	gcttaactgaaaagggtgttctccctctgc
Fragment 37	56,113,523	gcactgtaccatggcactactgagaccaaagg
Fragment 38	56,115,580	ttgtactccagtcattcttgggttgaggg
Fragment 39	56,118,301	cggccagaccattgcttgatgtttaatagc
Fragment 40	56,119,851	aatgagagaaggcctaggaccaccttg
Fragment 41	56,121,912	cgtcatttcagcatcgtgctcagtttagtcc
Fragment 42	56,123,317	gatttacatcagagctgggacaagacagtctgg
Fragment 43	56,127,536	ttcaggttaatcagaccaactatcaggcaggc
Fragment 44	56,128,078	gaccgttccccattactgtcctaacactttgg
Fragment 45	56,129,808	aactaggatgattgaggctggatcccagacc
Fragment 46	56,132,919	cataagcagttcacataaaagacttatgggccagc

Fragment 47	56,136,559	gaagcaaccaagaggaagaaaggggaactagc
Fragment 48	56,142,493	cttcacctcctgcttttctcctcacctgc
Fragment 49	56,144,772	gtaaatggagttattgtgcatgaccatgtgg
Fragment 50	56,159,585	ggacatttgagctgctcctgtttggtgg
Fragment 51	56,169,636	gaagctgacaaaaacgctcctagggtgagg
Fragment 52	56,170,833	gatgggtcatgaaagggcaggtgagg
Fragment 53	56,174,572	tgtaggctcctgggtattcatttgttctgtgg
Fragment 54	56,175,998	cctctgcaagtcataatgacttagttgcacagg
Fragment 55	56,178,759	gtctcaggatgccctccccatagtcc
Fragment 56	56,179,669	ggctcaagatgtgggaactggaactttaatgg
Fragment 57	56,181,895	cggtaggctcatttgctgagtaaatatggagc
Fragment 58	56,183,388	agatttggagctgcagccaggttgg
Fragment 59	56,188,350	ccatacatcacagctccttcatactgttgacc
Fragment 60	56,189,045	tactaagaatcagaatttgtccaacacggaagg
Fragment 61	56,193,516	gtctatgaaaaaattcatccactggatgggagg
Fragment 62	56,211,583	tttctggattctgacatgaagactgtcacc

Table S2. Reporter gene construct cloning and overlapping mutagenesis primers

Primer	Sequence
MAP3K1 promoter forward	<u>acgcgtccac</u> ctctctgcagtaacatag
MAP3K1 promoter reverse	<u>aagcttgcttc</u> ctctcggcaatctcg
rs74762363 forward	ccagtttcaaccactg <a>ac aaacagtcaaaaatcag
rs74762363 reverse	ctgatttttgactgtttgtcagtgggtgaaactgg
PRE-A forward	<u>accggtgctca</u> actagccaaagcactcc
PRE-A reverse	<u>cctgcaggggg</u> caagtctaaagcagtgtgg
rs7709971 forward	cagttcagccactgtgg <a>a agcagtgagatctggag
rs7709971 reverse	ctccagatctcactgcttccacagtggctgaactg
PRE-B3 forward	<u>accggtgga</u> agatgggacaagccttgatgc
PRE-B3 reverse	<u>cctgcagggga</u> agaacaacctgtctcaatgatgg
rs17432750 forward	caacctggattcttctact <a>a tcacacaagtcagg
rs17432750 reverse	cctgacttgtgtgattagtgaagaatccagggtg
rs11956804 forward	gctgagcagtttatcttgtcatatt <a>a gtaggatgaatg
rs11956804 reverse	cattcatctctactaaatatgacaaagataaactgctcagc
PRE2c forward	<u>accggtaaa</u> agatagctttcaaagg
PRE2c reverse	<u>gcgtcgacc</u> atagttacttcaaagg
rs12659430 forward	<u>catgcatttg</u> atgtgtcctataaaaag
rs12659430 reverse	<u>ctttatagg</u> acacatccaaatgcatg
rs62355881 forward	<u>aaagtca</u> cggtgcttctggtagc
rs62355881 reverse	<u>gctaccaga</u> agcatccgtgacttt
rs2113084 forward	<u>taaattt</u> gtggcatgcaaataattaaaac
rs2113084 forward	<u>gttttaata</u> tttgcatagccacaaattta
PRE-C forward	<u>accggtcca</u> agtttcatgcatggctctgtgg
PRE-C reverse	<u>cctgcaggg</u> tggcctcttccagtacagtgg
rs74345699 forward	ggctcacgcctgtaatctcagcactttgg
rs74345699 reverse	ccaaagtgtgagattacaggcgtgagcc
rs62355900 forward	tgggggagcatccgaggtggatgaag
rs62355900 reverse	cttcatccacctcggatgctcccca
rs62355901 forward	ggagatcaagaccatcccggtagcacg
rs62355901 reverse	cgtgctagccggatggtcttgatctcc
rs62355902 forward	tgggcaacagagcgagactccatctcaaaaaaca
rs62355902 reverse	tgtttttgagatggagtctcgctctgttgccca
PRE-D forward	<u>accggtggc</u> cattatagcagtgtctttgc
PRE-D reverse	<u>cctgcaggg</u> gctgatgcctagtagtcaattaagc
rs77371588 forward	ctgggcagtgaagccgtcttccagtg
rs77371588 reverse	cactggaagacgggtcactgcccag

Underlined sequences correspond to restriction enzyme recognition sites and bases highlighted in red denote SNP loci.

Table S3. Primer pairs used in ChIP-RT-PCR and sequence confirmation

Primer	Sequence	Ref or comment
rs17432750-for1	GGCCATCTGTTTTACCAACC	RT-PCR
rs17432750-rev1	ATTTGCACATGCCTTTCTCC	RT-PCR
rs17432750-for2	CAATGCAAATCTTCCTTGCTT	RT-PCR & sequencing
rs17432750-rev2	TGGGAAGGAGTCGTTGAGTT	RT-PCR & sequencing
ER- α -ENH-for	TGTAGGCTAGTTTTGTTTAACGATTTTT	Cancer Research 2007; 67:6477-6483
ER- α -ENH-rev	GGTGATGGGAGAATTGCTTAGAA	as above
<i>CCND1</i> -for	TGCCACACACCAGTGACTTT	Genes Dev. 2006; 20:2513- 2526.
<i>CCND1</i> -rev	ACAGCCAGAAGCTCCAAAAA	as above

Table S6. Associations of the iCHAV representative SNPs with breast cancer risk in Asian and African American studies

SNP	Position	Asian Studies						African American Studies					
		MAF	Imp r^2	P-trend	OR	LCI	UCI	MAF	Imp r^2	P-trend	OR	LCI	UCI
iCHAV1 rs62355902	56053723	0.38	0.94	3.30E-02	1.06	1.00	1.12	0.09	0.98	2.30E-01	1.14	0.92	1.36
iCHAV2a rs113317823	56087883	0.13	0.88	1.40E-05	1.19	1.11	1.27	0.06	0.78	7.80E-01	1.04	0.77	1.31
iCHAV2b rs62355899	56050465	0.26	1.00	7.46E-01	0.99	0.93	1.05	0.04	1.00	9.60E-01	0.99	0.72	1.36
iCHAV2c rs7721581	56087883	0.56	1.00	3.80E-01	0.98	0.93	1.03	0.06	0.78	7.78E-01	1.04	0.78	1.38
iCHAV3 rs11949391	56045081	0.05	1.00	2.60E-02	0.87	0.75	0.99	0.10	1.00	1.20E-01	0.84	0.62	1.06

Table S8

[Click here to download Supplemental Movies and Spreadsheets: Table S8.xlsx](#)