

# Evidence that the 5p12 Variant rs10941679 Confers Susceptibility to Estrogen-Receptor-Positive Breast Cancer through *FGF10* and *MRPS30* Regulation

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Genome-wide association studies (GWASs) have revealed increased breast cancer risk associated with multiple genetic variants at 5p12. Here, we report the fine mapping of this locus using data from 104,660 subjects from 50 case-control studies in the Breast Cancer Association Consortium (BCAC). With data for 3,365 genotyped and imputed SNPs across a 1 Mb region (positions 44,394,495–45,364,167; NCBI build 37), we found evidence for at least three independent signals: the strongest signal, consisting of a single SNP rs10941679, was associated with risk of estrogen-receptor-positive (ER<sup>+</sup>) breast cancer (per-g allele OR ER<sup>+</sup> = 1.15; 95% CI 1.13–1.18; p = 8.35 × 10<sup>-30</sup>). After adjustment for rs10941679, we detected signal 2, consisting of 38 SNPs more strongly associated with ER-negative (ER<sup>-</sup>) breast cancer (lead SNP rs6864776: per-a allele OR ER<sup>-</sup> = 1.10; 95% CI 1.05–1.14; p conditional = 1.44 × 10<sup>-12</sup>), and a single signal 3 SNP (rs200229088: per-t allele OR ER<sup>+</sup> = 1.12; 95% CI 1.09–1.15; p conditional = 1.12 × 10<sup>-05</sup>). Expression quantitative trait locus analysis in normal breast tissues and breast tumors showed that the g (risk) allele of rs10941679 was associated with increased expression of *FGF10* and *MRPS30*. Functional assays demonstrated that SNP rs10941679 maps to an enhancer element that physically interacts with the *FGF10* and *MRPS30* promoter regions in breast cancer cell lines. *FGF10* is an oncogene that binds to *FGFR2* and is overexpressed in ~10% of human breast cancers, whereas *MRPS30* plays a key role in apoptosis. These data suggest that the strongest signal of association at 5p12 is mediated through coordinated activation of *FGF10* and *MRPS30*, two candidate genes for breast cancer pathogenesis.

Strong evidence for the existence of a breast cancer (MIM: 114480) susceptibility locus at 5p12 has been observed through a GWAS in Iceland (SNP rs7703618),<sup>1</sup> in the Breast Cancer Association Consortium (BCAC; SNP rs981782, 371 Kb centromeric),<sup>2</sup> and in the Cancer Genetic Markers of Susceptibility study (CGEMS; SNP rs4866929; 352 Kb

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centromeric;  $r^2 = 0.18$ ).<sup>3</sup> A subsequent study, using 22 SNPs in ~5,000 case subjects and ~33,000 control subjects of European ancestry, reported that risk at this locus could be explained by two SNPs: rs4415084 and rs10941679.<sup>4</sup> More recently, a BCAC study confirmed that rs10941679 was associated with risk of lower-grade, progesterone receptor (*PGR* [MIM: 607311])-positive breast cancer tumors.<sup>5</sup>

Here, we report the comprehensive fine-scale mapping of this locus in 104,660 subjects from 50 case-control studies participating in BCAC, including 41 studies from populations of European ancestry and nine of East Asian ancestry, and we explore the functional mechanisms underlying the associations in this region. Genotyping was conducted with the COGS array, a custom array

comprising approximately 200,000 SNPs.<sup>6</sup> After quality-control exclusions, we analyzed data from 48,155 case subjects and 43,612 control subjects of European ancestry and 6,269 case subjects and 6,624 control subjects of Asian ancestry. Estrogen receptor (*ESR1* [MIM: 133430]) status of the primary tumor was available for 27,748 European and 4,997 Asian case subjects; of these, 7,646 (22%) European and 1,623 (32%) Asian case subjects were ER<sup>-</sup>.

We examined a 1 Mb region (positions 44,394,495–45,364,167; NCBI build 37 assembly) in which the 1000 Genomes Project cataloged 1,811 variants (March 2010 Pilot version 60 CEU project data). We aimed to genotype all 628 SNPs with minor allele frequency (MAF) > 2% and correlated with rs981782 and rs10941679 at  $r^2 > 0.1$  ( $n = 424$ ), plus a set of SNPs designed to tag all remaining

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SNPs with  $r^2 > 0.9$  ( $n = 184$ ), but we managed to include 563 SNPs with a designability score (DS)  $> 0.9$  and which passed QC.<sup>6</sup> IMPUTE v.2.0 was used to impute genotypes of all known SNPs in the region using the 1000 Genome Project data (March 2012 version) as a reference panel.

Case-control analyses were conducted on 3,365 SNPs (563 genotyped and 2,776 imputed at  $r^2 > 0.3$ ). In European-ancestry women, 461 of these SNPs were associated with overall breast cancer risk, 489 with ER<sup>+</sup> and 38 with ER<sup>-</sup> breast cancer risk ( $p < 10^{-4}$ ; Table S1). SNP rs10941679 showed the strongest overall association (MAF = 0.27, per-minor (g) allele: OR = 1.12; 95% CI 1.10–1.14;  $p = 2.55 \times 10^{-26}$ ; Figure 1, Tables 1 and S1). To identify additional association signals at this region, we conducted a forward stepwise logistic regression examining SNPs with univariate  $p < 0.1$  ( $n = 1,040$ ).<sup>6</sup> The most parsimonious model included three variants: SNP1 rs10941679 (signal 1), SNP2 rs6864776 (signal 2; conditional  $p = 6.22 \times 10^{-11}$ ), and SNP3 rs200229088 (signal 3; conditional  $p = 1.12 \times 10^{-5}$ , borderline significance; Table S2). SNP1 and SNP3 are weakly correlated ( $r^2 = 0.15$ ) but SNP2 was uncorrelated with the other two ( $r^2 = 0.07$  and 0.05).

The top signal, SNP1 rs10941679, is markedly more significant than any other SNP in the locus (likelihood ratio  $> 10,000:1$ ). Hence, the most parsimonious explanation is that this SNP is causally related to risk. The next most strongly associated SNP, after adjustment for signal 1 SNP rs10941679, was rs6864776, representing signal 2 (OR per minor allele = 1.04; 95% CI 1.02–1.06;  $p = 7.84 \times 10^{-4}$ ; conditional  $p = 1.44 \times 10^{-12}$ ). Within signal 2, a further 37 SNPs correlated with rs6864776 at  $r^2 > 0.6$ , had likelihood ratios of  $<100:1$  relative to rs6864776, and hence could not be excluded from being causative

statistically (Table S2). After adjustment for both signal 1 SNP rs10941679 and signal 2 top SNP rs6864776, a single SNP remained: rs200229088 (OR overall = 1.09, 95% CI 1.07–1.12;  $p = 2.28 \times 10^{-12}$ ; conditional  $p = 1.12 \times 10^{-5}$ ). There are no other SNPs correlated with rs200229088 that could explain this association. All other SNPs were excluded from causality (likelihood ratio  $> 10,000:1$ ; Table S2). Two of the excluded variants had been previously postulated as likely causative variants<sup>4,7</sup> and so we investigated these in more depth. We found both SNPs to be partially correlated with all three signals and consequently display initially inflated effects, which are adjusted by the conditional analyses. Thus, SNP rs4415084<sup>4</sup> ( $r^2$  with signal 1 SNP rs10941679 = 0.51, with signal 2 SNP rs6864776 = 0.11, and with signal 3 SNP rs200229088 = 0.37) has odds against causality  $> 10$  million:1 versus signal 1 candidate rs10941679. Similarly, SNP rs7716600, which is an eQTL for MRPS30 expression<sup>7</sup> ( $r^2$  with SNP rs10941679 = 0.77, with SNP rs6864776 = 0.05, and with SNP rs200229088 = 0.12) has odds against causality  $>160,000:1$  versus signal 1 candidate rs10941679. These exclusions of former causal candidates highlight the need for fine-mapping studies before conducting functional analyses.

Haplotype analyses were conducted using the above three signal-representative variants, which generated eight haplotypes (Table 2). Haplotypes carrying the rare allele of signal 3 SNP rs200229088 conferred higher risks than corresponding haplotypes carrying the common allele, consistent with this allele having an independent effect. Haplotype G, carrying the minor alleles of both the signal 1 and 2 representative SNPs, is very rare and reveals that their risk alleles are negatively correlated, which is also consistent with our finding that signal 2 top SNP

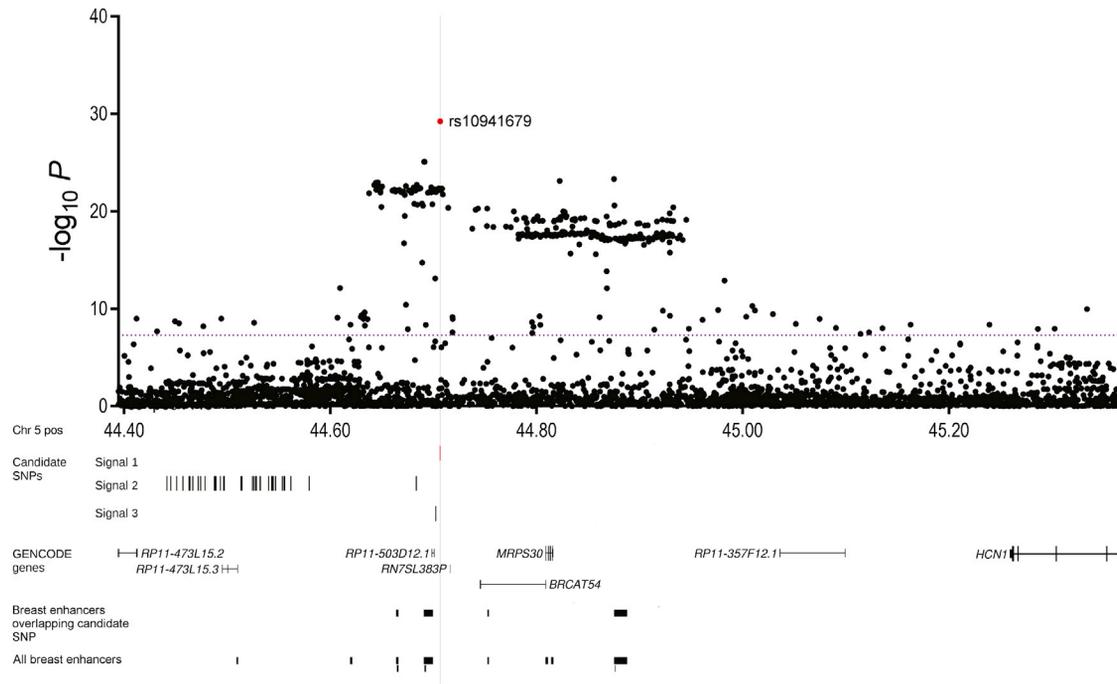
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**Figure 1. Manhattan Plot of the 5p12 Breast Cancer Susceptibility Locus**

SNPs are plotted according to their chromosomal position on the x axis and their overall p values ( $\log_{10}$  values, likelihood ratio test) from the European BCAC studies (48,155 case and 43,612 control subjects) on the y axis. The purple dotted line intersects the y axis at  $p = 10^{-8}$  and indicates genome-wide significance. Candidate SNPs in signal 1 (rs10941679), signal 2 (38 SNPs), and signal 3 (rs200229088) are shown as short vertical lines. The locations of annotated genes and putative lncRNA transcripts from GENCODE and enhancers predicted in Corradin et al.<sup>13</sup> and Hnisz et al.<sup>12</sup> from breast cancer cell lines are shown in the bottom panels.

rs6864776 increases in significance after conditioning on signal 1 SNP rs10941679 (Table 1).

We examined the associations of these three SNPs in the Asian case-control studies within BCAC. SNP1 and SNP3 both replicated in the Asian studies and the relative risk estimates with overall breast cancer were consistent with those seen in the European population: per *g*-allele OR (rs10941679) = 1.09; 95% CI 1.04–1.15;  $p = 0.0009$ , conditional  $p = 0.0859$  and per *t*-allele OR (rs200229088) = 1.09;

95% CI 1.02–1.15;  $p = 0.0065$ , conditional  $p = 0.9149$  (Table 1). SNP2 was not replicated in Asians (per *a*-allele OR = 0.94; 95% CI 0.89–1.00;  $p = 0.034$ , conditional  $p = 0.8901$ ) (Table 1).

We investigated the associations of these three signals with tumor subtypes based on ER status. SNP1 rs10941679 was largely associated with ER<sup>+</sup> breast cancer (OR ER<sup>+</sup> = 1.15; 95% CI 1.13–1.18;  $p = 8.35 \times 10^{-30}$  versus OR ER<sup>-</sup> disease = 1.04; 95% CI 1.00–1.08;  $p = 0.059$ ;

**Table 1. Associations of the Top SNPs from Each Signal with Overall Breast Cancer Risk and Breast Cancer Stratified by ER Status**

Sig	SNP	Com	Min	MAF*	OR Overall		Conditional p Value	OR ER <sup>-</sup>	p ER <sup>-</sup>	OR ER <sup>+</sup>	p ER <sup>+</sup>
					95% CI	p Overall					
<b>Europeans</b>											
1	rs10941679	A	G	0.27	1.12 (1.10–1.14)	$2.55 \times 10^{-26}$	$6.55 \times 10^{-24}$	1.04 (1–1.08)	0.059	1.15 (1.13–1.18)	$8.35 \times 10^{-30}$
2	rs6864776	G	A	0.23	1.04 (1.02–1.06)	$7.84 \times 10^{-4}$	$1.44 \times 10^{-12}$	1.10 (1.05–1.14)	$2.5 \times 10^{-5}$	1.02 (0.99–1.05)	0.08
3	rs200229088	TTG	T	0.31	1.09 (1.07–1.12)	$2.28 \times 10^{-12}$	$1.12 \times 10^{-5}$	1.03 (0.99–1.09)	0.11	1.12 (1.09–1.15)	$7.51 \times 10^{-14}$
<b>Asians</b>											
1	rs10941679	A	G	0.50	1.09 (1.04–1.15)	$9.12 \times 10^{-4}$	0.0859	1.03 (0.95–1.11)	0.53	1.11 (1.04–1.18)	$1.32 \times 10^{-3}$
2	rs6864776	G	A	0.32	0.94 (0.89–1.00)	$3.47 \times 10^{-2}$	0.8901	0.95 (0.87–1.04)	0.28	0.94 (0.89–1.00)	$6.24 \times 10^{-2}$
3	rs200229088	TTG	T	0.37	1.09 (1.02–1.15)	$6.52 \times 10^{-3}$	0.9149	1.04 (0.95–1.14)	0.43	1.08 (1.00–1.16)	$3.65 \times 10^{-2}$

Abbreviations are as follows: Com, common alleles; Min, minor alleles; MAF, minor allele frequency; OR, per-allele odds ratios (OR); 95% CI, 95% confidence intervals and 1 degree of freedom; p, significance levels for overall breast cancer are indicated in European and Asian case-control studies, and separately for ER<sup>+</sup> and ER<sup>-</sup> disease.

**Table 2. Haplotype Analysis across the BCAC Studies**

Haplotypes	rs10941679 Signal 1	rs6864776 Signal 2	rs200229088 Signal 3	Haplotype Frequency	OR	p Value
A	1	1	1	0.395440	–	–
B	1	1	2	0.120099	1.06 (1.02–1.10)	$1.49 \times 10^{-3}$
C	1	2	1	0.199599	1.10 (1.06–1.13)	$7.76 \times 10^{-11}$
D	1	2	2	0.018665	1.15 (1.04–1.27)	$5.03 \times 10^{-3}$
E	2	1	1	0.098169	1.14 (1.09–1.19)	$1.45 \times 10^{-11}$
F	2	1	2	0.154525	1.20 (1.16–1.24)	$2.72 \times 10^{-30}$
G	2	2	1	0.004248	0.91 (0.72–1.15)	$4.15 \times 10^{-1}$
H	2	2	2	0.009253	1.28 (1.10–1.48)	$1.14 \times 10^{-3}$

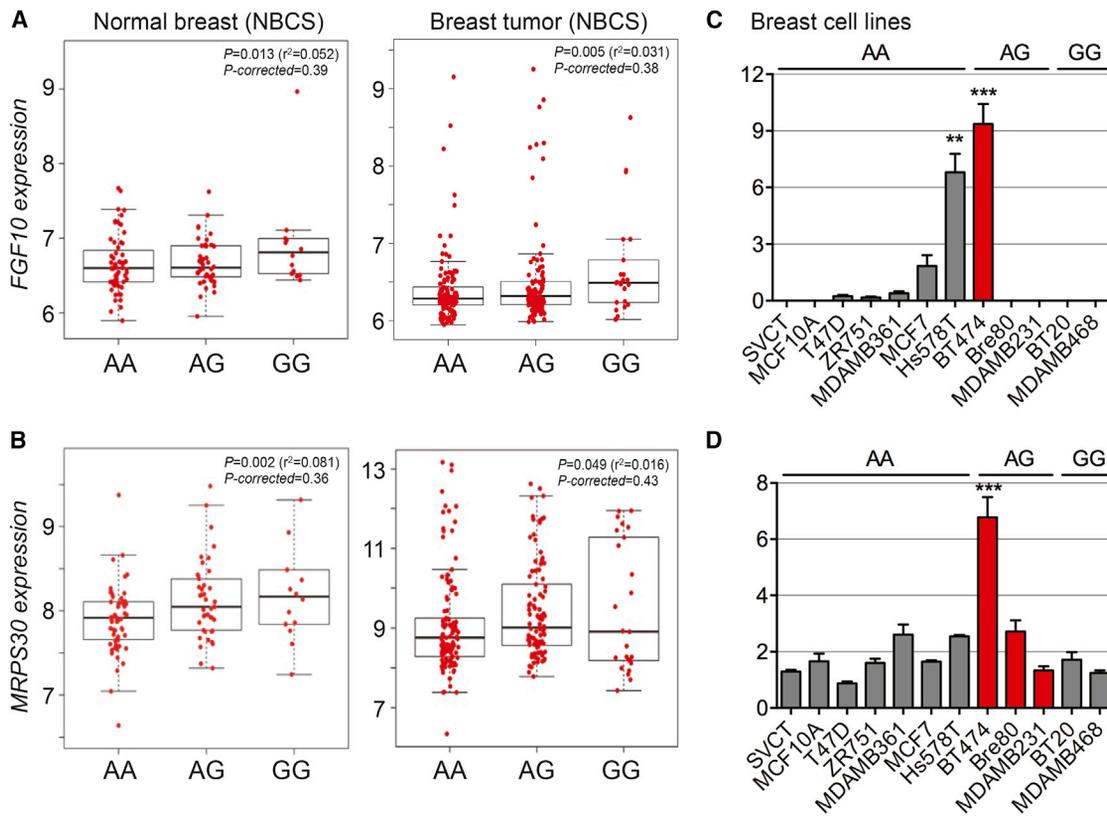
Each haplotype was compared to the ancestral haplotype carrying the common alleles of signal 1 SNP rs10941679, signal 2 SNP rs6864776, and signal 3 SNP rs200229088 (haplotype A).

p heterogeneity =  $1.5 \times 10^{-5}$ ; Table 1) as was SNP3 rs200229088 (OR ER<sup>+</sup> = 1.12; 95% CI 1.09–1.15; p =  $7.51 \times 10^{-14}$  versus OR ER<sup>−</sup> = 1.03; 95% CI 0.99–1.09; p = 0.11, p heterogeneity = 0.02). By contrast, SNP2 rs6864776 was moderately associated with ER<sup>−</sup> but not ER<sup>+</sup> tumors (OR ER<sup>−</sup> = 1.10; 95% CI 1.05–1.14; p =  $2.55 \times 10^{-5}$  versus OR ER<sup>+</sup> = 1.02; 95% CI 0.99–1.05; p = 0.08; p heterogeneity = 0.01; Table 1).

Candidate SNPs 1–3 span a 1.7 Mb region on 5p12 that includes three annotated genes—*FGF10* (MIM: 602115), *MRPS30* (MIM: 611991), and *HCN1* (MIM: 602780)—and several putative long noncoding RNAs (lncRNAs; Figure 1). To identify potential target gene(s), we examined the associations of the three lead SNPs with expression levels of genes located within 1 Mb in three different studies: (1) 116 normal breast samples and 241 breast tumors from the Norwegian Breast Cancer Study (NBCS),<sup>8</sup> (2) 93 normal and 765 breast cancer tissues from the TCGA study (germline genotype data from Affymetrix SNP 6 array were obtained from TCGA dbGAP data portal<sup>9</sup>), and (3) 183 normal breast samples from the Genotype-Tissue Expression (GTEx) project.<sup>10</sup> The SNP1 rs10941679 risk-associated *g*-allele was moderately associated with increased *FGF10* mRNA expression in NBCS normal breast (p = 0.013, p corrected = 0.39) and breast tumors (p = 0.005, p corrected = 0.38) as well as in GTEx normal breast (p corrected = 0.02; Figures 2A and S1A). The effect in TCGA was in the same direction, though not significant (normal breast p = 0.353, p corrected = 0.95 and breast tumors p = 0.057, p corrected = 0.41; Figure S1B). The *g*-allele was also associated with increased expression of *MRPS30* in the NBCS normal (p = 0.002, p corrected = 0.36) and breast tumors (p = 0.049, p corrected = 0.43), in GTEx normal breast (p corrected = 0.002), and in TCGA (normal breast p =  $6.86 \times 10^{-5}$ , p corrected =  $5.31 \times 10^{-3}$  and breast tumors p =  $7.21 \times 10^{-6}$ , p corrected =  $9.35 \times 10^{-4}$ ; Figures 2B, S1A, and S1C). No associations were observed with SNP2 rs6864776 or SNP3 variant rs200229088. We also measured endogenous levels of *FGF10*, *MRPS30*, and nearby lncRNAs *FGF10-AS1*,

*BRCAT54*, *RP11-503D12.1*, and *RP11-473L15.3* mRNA in breast cell lines homozygous (A/A or G/G) or heterozygous (A/G) for the common allele of SNP1 (Table S3, Figures 2C, 2D, S2, and S3). Total RNA from cell lines was extracted using Trizol and complementary DNA synthesized using random primers as per manufacturers' instructions. Quantitative PCR (qPCR) were performed using TaqMan assays for *FGF10* and *MRPS30* normalized against beta-glucuronidase (*GUSB* [MIM: 611499]) or with SYTO9 for lncRNAs normalized against TATA box-binding protein (*TBP* [MIM: 600075]); primers are listed in Table S4). Although the number of ER<sup>+</sup> breast cell lines carrying the risk allele was limited, *FGF10* and *MRPS30* mRNA levels were significantly higher in the BT474 heterozygous cell line (Figures 2C and 2D). *BRCAT54* was detected in the majority of cell lines but its expression appears to be genotype independent (Figure S3A). *FGF10-AS1*, *RP11-503D12.1*, and *RP11-473L15.3* transcripts were either expressed at very low levels or not detected in the cell lines analyzed (Figures S3B–S3D). Therefore, although we cannot rule out the possibility that the risk SNPs may influence local lncRNA expression, the low or absent transcript levels precluded any further evaluation.

Candidate causal SNPs were then explored using publicly available datasets from ENCODE,<sup>11</sup> which includes information such as the location of promoter and enhancer histone marks, open chromatin, bound proteins, and altered motifs for the MCF7 breast cancer cell line, and from Hnisz et al.<sup>12</sup> and Corradin et al.<sup>13</sup> to identify the location of likely enhancers and their gene targets in a cell-specific context. Analysis of *cis* enhancer-gene interactions via PreSTIGE<sup>13</sup> showed evidence of putative regulatory elements (PREs) surrounding the top risk-associated SNPs in MCF7 breast cancer cells, but no histone-marked elements harboring a risk SNP in this cell line or in a range of cell lines and tissues analyzed in Roadmap (Figures 1 and S4). However, it is possible that certain epigenetic marks may be detected only in a specific cell subtype such as breast stem cells or in response to an external stimulus.



**Figure 2. Association of rs10941679 with *FGF10* and *MRPS30* Expression in Normal Breast Tissues, Breast Tumors, and Breast Cancer Cell Lines**

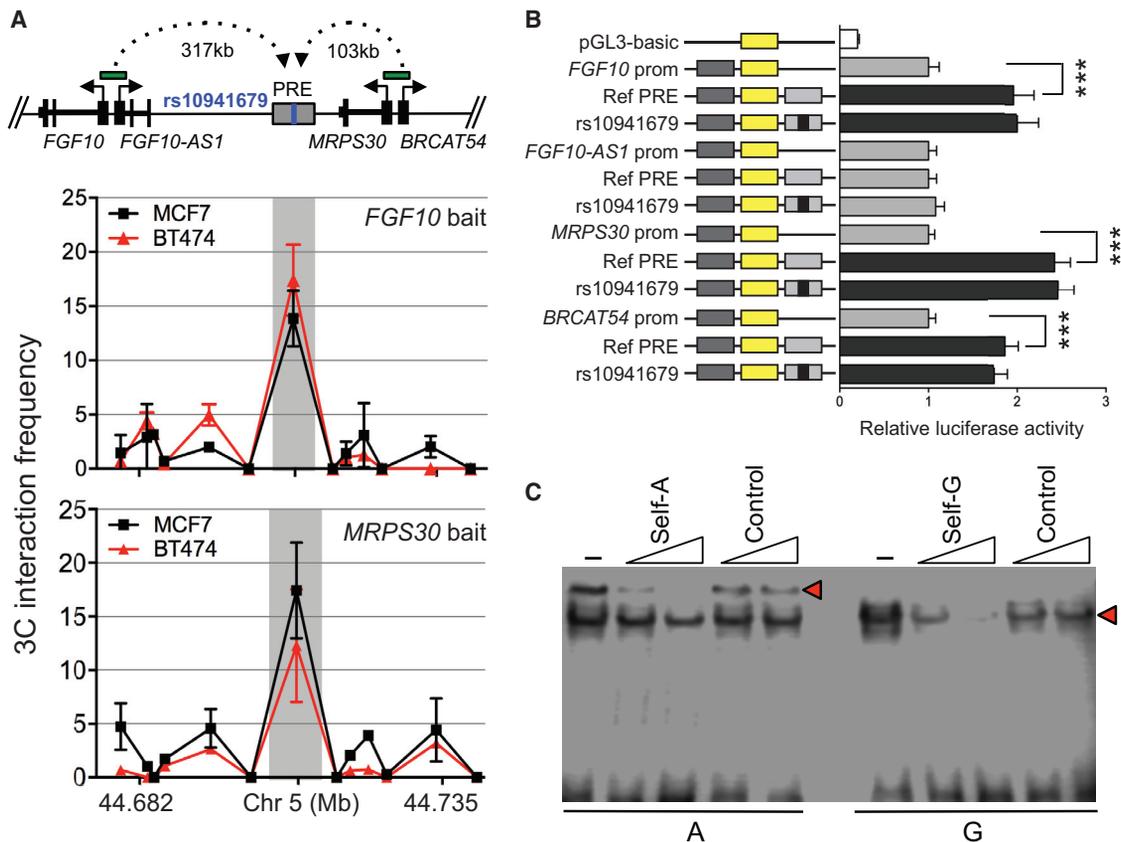
(A and B) *FGF10* (A) or *MRPS30* (B) expression in normal breast ( $n = 116$ ) or breast tumors from NBBS dataset ( $n = 241$ ). SNP genotypes are shown on the x axis and log<sub>2</sub>-normalized gene expression values on the y axis. p values are presented before and after correction for multiple testing using FDR as implemented in p.adjust function in R. Each box plot shows the median rank normalized gene expression (horizontal line), the first through third quartiles (box), and 1.5 $\times$  the interquartile range (whiskers).

(C and D) Endogenous *FGF10* (Hs00610298\_m1) (C) or *MRPS30* (Hs00169612\_m1) (D) expression measured by qPCR in untreated breast cell lines and normalized to *GUSB* (4326320E). Error bars denote SEM ( $n = 3$ ). p values were determined with a two-tailed t test. \*\* $p < 0.01$ , \*\*\* $p < 0.001$ .

To identify target gene(s), we performed chromatin conformation capture (3C) assays in ER<sup>+</sup> MCF7, BT474, and MDA-MB-361 and ER<sup>-</sup> MDA-MB-231 breast cancer cell lines and Bre80 normal breast cells (Table S5).<sup>8</sup> 3C libraries were created by cross-linking the chromatin from cell lines; DNA was then digested with EcoRI, which flanks 12 contiguous fragments that cover the PRE, and the *FGF10*, *MRPS30*, and *HCN1* promoters (Table S6); DNA was religated and decrosslinked; and qPCR with primers for the bait (gene promoters) and interactors (12 PRE fragments) was performed to detect the presence of ligation products, representing gene loops. BAC clones covering the regions of interest were used to normalize for PCR efficiency. These assays showed that the PRE containing SNP1 frequently interacted with the *FGF10* and *MRPS30* promoter regions in MCF7 and BT474 breast cancer cell lines, but only with *MRPS30* in the MDA-MB-361, MDA-MB-231, and Bre80 cell lines. This latter result was expected because *FGF10* is not expressed or expressed at very low levels in these cell lines (Figures 2C, 3A, S5, and S6). Notably, both genes share a bidirectional promoter with the lncRNAs *FGF10-AS1* and *BRCAT54*, raising the

possibility that these transcripts are also targets of the PRE (Figure 3A). No additional interactions were detected between the PRE and other annotated genes within 1 Mb of the PRE, including *HCN1* (Figure S5). To assess the potential impact of SNP1 on the identified chromatin interactions, allele-specific 3C was performed in heterozygous BT474 cell lines.<sup>8</sup> However, the sequence profiles revealed that SNP1 had no significant effect on chromatin looping (Figure S7).

The regulatory capability of the PRE, combined with the effect of SNP1, was further examined in reporter assays. Promoter-driven luciferase reporter constructs were generated by the insertion of PCR-amplified fragments containing *FGF10*, *FGF10-AS1*, *MRPS30*, or *BRCAT54* promoters into pGL3-Basic.<sup>14</sup> A 1,736-bp PRE fragment (containing either the common or minor allele of rs10941679) was then generated by PCR and cloned downstream of the modified pGL3-promoter constructs (Table S7). MCF7 and BT474 breast cancer cell lines plus Bre80 normal breast cells were transfected with the reporter plasmids and luciferase activity was measured 24 hr after transfection. To correct for any differences in transfection efficiency or



**Figure 3. Distal Regulation of *FGF10* and *MRPS30* at the 5p12 Risk Region**

(A) 3C interaction profiles between the *FGF10/FGF10AS-1* or *MRPS30/BRCAT54* bidirectional promoters and the putative regulatory element (PRE; gray bar) containing SNP rs10941679. Anchor points are set at the promoters. Graphs represent one of three independent experiments (see Figure S5B). Error bars denote SD.

(B) Luciferase reporter assays after transient transfection of ER<sup>+</sup> BT474 breast cancer cell lines. The PRE containing the major SNP allele was cloned downstream of target gene promoter-driven luciferase constructs (Ref PRE). The risk *g*-allele was engineered into the constructs and designated by the rs ID. Primers are listed in Table S7. Error bars denote 95% confidence intervals from three independent experiments. *p* values were determined by 2-way ANOVA followed by Dunnett's multiple comparisons test (\*\**p* < 0.001).

(C) EMSA for oligonucleotides containing SNP rs1094617 with the A = common allele and G = minor allele as indicated below the panel, assayed using BT474 nuclear extracts. Primers are listed in Table S8. Labels above each lane indicate inclusion of competitor oligonucleotides at 30- and 100-fold molar excess, respectively: (-) no competitor and control denotes a non-specific competitor. A red arrowhead shows a band of different mobility detected between the common and minor alleles.

cell lysate preparation, *Firefly* luciferase activity was normalized to *Renilla*. Notably, the "Ref PRE" acted as a transcriptional enhancer, leading to a 2- to 3-fold increase in *FGF10*, *MRPS30*, and *BRCAT54* promoter activity, but had no effect on the *FGF10-AS1* promoter in MCF7 and BT474 cells (Figures 3B and S8). The enhancer activity was also observed for the *MRPS30* and *BRCAT54* promoters in Bre80 cells (Figure S8). In all cell lines, inclusion of the SNP1 risk (*g*) allele had no significant effect on the PRE enhancer activity. Although this appears to rule out an effect of this SNP on transactivation, it is possible that SNP1 affects the recruitment of key proteins required for the epigenetic modification of the enhancer, which would not be observed in a reporter assay. Another possibility is that the SNP effect may be observed only under certain biological conditions such as growth factor stimulation.

To seek further evidence that SNP1 lies within an enhancer element, we performed electrophoretic mobility

shift assays (EMSAs) for both the protective (*a*) and risk (*g*) alleles.<sup>15</sup> Nuclear lysates were prepared from ER<sup>+</sup> BT474, MCF7, and MDA-MB-361 or ER<sup>-</sup> MDA-MB-231 and Hs578T cells using the NE-PER nuclear and cytoplasmic extraction reagents. Biotinylated oligonucleotide duplexes were prepared by combining sense and antisense oligonucleotides, heat annealing, and slow cooling. Duplex-bound complexes were transferred onto Zeta-Probe positively charged nylon membranes by semi-dry transfer then cross-linked onto the membranes. Membranes were processed with the LightShift Chemiluminescent EMSA kit as per the manufacturer's instructions, and signals were visualized with the C-DiGit blot scanner. For SNP1, we observed allele-specific binding by nuclear proteins only in the ER<sup>+</sup> BT474, MCF7, and MDA-MB-361 extracts (Figures 3C and S9). The protein-DNA complexes were shown to be specific, as demonstrated by increasing amounts of cold self-competitor (Figures 3C and S9 and Table S8).

Further EMSAs using competitor DNA or antibody supershifts against predicted transcription factors (TFs) suggested four proteins bound to the SNP site including FOXA1, FOXA2, CEBPB, and OCT1 (Figure S10 and Table S9). To confirm TF binding in vivo, we performed chromatin immunoprecipitation (ChIP) in heterozygous BT474 cells as previously described (Table S10).<sup>15</sup> When compared to an IgG control antibody, we observed a moderate enrichment in FOXA1 and OCT1 binding to DNA overlapping SNP rs10941679, but no difference between alleles in this cell line (Figure S11). In addition, western blot analysis indicated that FOXA1 protein expression was restricted to the ER<sup>+</sup> breast cancer cell lines analyzed, whereas OCT1 was more widely expressed (Figure S12). FOXA1 is a pioneer factor and master regulator of ER activity due to its ability to open local chromatin and recruit ER to target gene promoters.<sup>16</sup> Notably, breast cancer-associated SNPs are enriched for FOXA1 binding<sup>17</sup> and several studies have linked cooperative binding of FOXA1, ER, and OCT1 to increased gene transcription.<sup>18,19</sup> Consistent with our eQTL data, it is tempting to speculate that in specific ER<sup>+</sup> cell subtypes and/or conditions, rs10941679 alters FOXA1 affinity and OCT1 recruitment leading to target gene activation.

In conclusion, we have provided evidence for at least three independent causal SNPs with effects on the risk of breast cancer at this locus. The minor *g*-allele of signal 1 SNP rs10941679 conferred a 15% increased risk of ER<sup>+</sup> breast cancer and higher expression levels of the *MRPS30* and *FGF10* genes and was the most strongly associated SNP with *MRPS30* expression in this 1 Mb region. *MRPS30*—also called *PDCD9* (Programmed Cell Death protein 9)—encodes a mitochondrial ribosomal protein involved in apoptosis.<sup>20</sup> Although the role of mitochondria in apoptosis remains unclear, it is well established that cytochrome *c* and other pro-apoptotic proteins are released during cell death initiation.<sup>20</sup> Clearly, further investigation of the function of this protein is now merited. By contrast, *FGF10* is an extensively studied gene with compelling data suggesting its involvement in breast tumorigenesis. *FGF10* is a member of the fibroblast growth factor (FGF) family and encodes a glycoprotein that specifically binds to *FGFR2* (splice *FGFR2IIIb*) to control signaling pathways including cell differentiation, proliferation, and apoptosis.<sup>21</sup> Variants regulating *FGFR2* (MIM: 176943) have the strongest association with ER<sup>+</sup> breast cancer susceptibility identified to date.<sup>22</sup> *FGF10* is overexpressed in ~10% of human breast cancers<sup>23</sup> and increased levels of *FGF10* are highly correlated with proliferation rate of breast cancer cell lines and cancer cell invasion.<sup>24,25</sup> It signals through multiple downstream pathways including MAPK and WNT and genes such as *FGFR2*, *CCND1* (MIM: 168461), and *TGFBI* (MIM: 190180),<sup>21,24</sup> all known to play key roles in breast cancer. Therapeutic targeting of FGFs and their receptors (FGFRs) is currently a major area of drug development research, and the identification of a subgroup of individuals diag-

nosed with breast cancer with alterations in these pathways may open new avenues for personalized medicine and pathway-targeted treatments.

## Supplemental Data

Supplemental Data include Supplemental Acknowledgments, 12 figures, 10 tables, and consortia information and can be found with this article online at <http://dx.doi.org/10.1016/j.ajhg.2016.07.017>.

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## Web Resources

1000 Genomes, <http://www.1000genomes.org>  
 Cancer Cell Line Encyclopedia (CCLE), <https://portals.broadinstitute.org/ccle/home>  
 ENCODE, <https://www.encodeproject.org/>  
 GEO, <http://www.ncbi.nlm.nih.gov/geo/>  
 GTEx Portal, <http://www.gtexportal.org/home/>  
 OMIM, <http://www.omim.org/>  
 PreSTIGE, <http://genetics.case.edu/prestige/>  
 The Cancer Genome Atlas, <http://cancergenome.nih.gov/>

## References

1. Stacey, S.N., Manolescu, A., Sulem, P., Rafnar, T., Gudmundsson, J., Gudjonsson, S.A., Masson, G., Jakobsdottir, M., Thorlacius, S., Helgason, A., et al. (2007). Common variants on chromosomes 2q35 and 16q12 confer susceptibility to estrogen receptor-positive breast cancer. *Nat. Genet.* **39**, 865–869.
2. Easton, D.F., Pooley, K.A., Dunning, A.M., Pharoah, P.D., Thompson, D., Ballinger, D.G., Struwing, J.P., Morrison, J., Field, H., Luben, R., et al.; SEARCH collaborators; kConFab; AOCs Management Group (2007). Genome-wide association study identifies novel breast cancer susceptibility loci. *Nature* **447**, 1087–1093.
3. Hunter, D.J., Kraft, P., Jacobs, K.B., Cox, D.G., Yeager, M., Hankinson, S.E., Wacholder, S., Wang, Z., Welch, R., Hutchinson, A., et al. (2007). A genome-wide association study identifies alleles in *FGFR2* associated with risk of sporadic postmenopausal breast cancer. *Nat. Genet.* **39**, 870–874.
4. Stacey, S.N., Manolescu, A., Sulem, P., Thorlacius, S., Gudjonsson, S.A., Jonsson, G.F., Jakobsdottir, M., Bergthorsson, J.T., Gudmundsson, J., Aben, K.K., et al. (2008). Common variants on chromosome 5p12 confer susceptibility to estrogen receptor-positive breast cancer. *Nat. Genet.* **40**, 703–706.
5. Milne, R.L., Goode, E.L., García-Closas, M., Couch, F.J., Severi, G., Hein, R., Fredericksen, Z., Malats, N., Zamora, M.P., Arias Pérez, J.I., et al.; GENICA Network; kConFab Investigators; AOCs Group (2011). Confirmation of 5p12 as a susceptibility locus for progesterone-receptor-positive, lower grade breast cancer. *Cancer Epidemiol. Biomarkers Prev.* **20**, 2222–2231.
6. Michailidou, K., Hall, P., Gonzalez-Neira, A., Ghoussaini, M., Dennis, J., Milne, R.L., Schmidt, M.K., Chang-Claude, J., Bojesen, S.E., Bolla, M.K., et al.; Breast and Ovarian Cancer Susceptibility Collaboration; Hereditary Breast and Ovarian Cancer Research Group Netherlands (HEBON); kConFab

- Investigators; Australian Ovarian Cancer Study Group; GENICA (Gene Environment Interaction and Breast Cancer in Germany) Network (2013). Large-scale genotyping identifies 41 new loci associated with breast cancer risk. *Nat. Genet.* *45*, 353–361, e1–e2.
7. Quigley, D.A., Fiorito, E., Nord, S., Van Loo, P., Alnæs, G.G., Fleischer, T., Tost, J., Moen Volland, H.K., Tramm, T., Overgaard, J., et al. (2014). The 5p12 breast cancer susceptibility locus affects MRPS30 expression in estrogen-receptor positive tumors. *Mol. Oncol.* *8*, 273–284.
  8. Ghossaini, M., Edwards, S.L., Michailidou, K., Nord, S., Cowper-Sal Lari, R., Desai, K., Kar, S., Hillman, K.M., Kaufmann, S., Glubb, D.M., et al.; Australian Ovarian Cancer Management Group; Australian Ovarian Cancer Management Group (2014). Evidence that breast cancer risk at the 2q35 locus is mediated through IGFBP5 regulation. *Nat. Commun.* *4*, 4999.
  9. Li, Q., Seo, J.H., Stranger, B., McKenna, A., Pe'er, I., Laframboise, T., Brown, M., Tyekucheva, S., and Freedman, M.L. (2013). Integrative eQTL-based analyses reveal the biology of breast cancer risk loci. *Cell* *152*, 633–641.
  10. Consortium, G.T.; GTEx Consortium (2013). The Genotype-Tissue Expression (GTEx) project. *Nat. Genet.* *45*, 580–585.
  11. Birney, E., Stamatoyannopoulos, J.A., Dutta, A., Guigó, R., Gingeras, T.R., Margulies, E.H., Weng, Z., Snyder, M., Dermitzakis, E.T., Thurman, R.E., et al.; ENCODE Project Consortium; NISC Comparative Sequencing Program; Baylor College of Medicine Human Genome Sequencing Center; Washington University Genome Sequencing Center; Broad Institute; Children's Hospital Oakland Research Institute (2007). Identification and analysis of functional elements in 1% of the human genome by the ENCODE pilot project. *Nature* *447*, 799–816.
  12. Hnisz, D., Abraham, B.J., Lee, T.I., Lau, A., Saint-André, V., Sigova, A.A., Hoke, H.A., and Young, R.A. (2013). Super-enhancers in the control of cell identity and disease. *Cell* *155*, 934–947.
  13. Corradin, O., Saiakhova, A., Akhtar-Zaidi, B., Myeroff, L., Willis, J., Cowper-Sal Lari, R., Lupien, M., Markowitz, S., and Scacheri, P.C. (2014). Combinatorial effects of multiple enhancer variants in linkage disequilibrium dictate levels of gene expression to confer susceptibility to common traits. *Genome Res.* *24*, 1–13.
  14. Glubb, D.M., Maranian, M.J., Michailidou, K., Pooley, K.A., Meyer, K.B., Kar, S., Carlebur, S., O'Reilly, M., Betts, J.A., Hillman, K.M., et al.; GENICA Network; kConFab Investigators; Norwegian Breast Cancer Study (2015). Fine-scale mapping of the 5q11.2 breast cancer locus reveals at least three independent risk variants regulating MAP3K1. *Am. J. Hum. Genet.* *96*, 5–20.
  15. Dunning, A.M., Michailidou, K., Kuchenbaecker, K.B., Thompson, D., French, J.D., Beesley, J., Healey, C.S., Kar, S., Pooley, K.A., Lopez-Knowles, E., et al.; EMBRACE; GEMO Study Collaborators; HEBON; kConFab Investigators (2016). Breast cancer risk variants at 6q25 display different phenotype associations and regulate ESR1, RMND1 and CCDC170. *Nat. Genet.* *48*, 374–386.
  16. Hurtado, A., Holmes, K.A., Ross-Innes, C.S., Schmidt, D., and Carroll, J.S. (2011). FOXA1 is a key determinant of estrogen receptor function and endocrine response. *Nat. Genet.* *43*, 27–33.
  17. Cowper-Sal Lari, R., Zhang, X., Wright, J.B., Bailey, S.D., Cole, M.D., Eeckhoute, J., Moore, J.H., and Lupien, M. (2012). Breast cancer risk-associated SNPs modulate the affinity of chromatin for FOXA1 and alter gene expression. *Nat. Genet.* *44*, 1191–1198.
  18. Meyer, K.B., Maia, A.T., O'Reilly, M., Teschendorff, A.E., Chin, S.F., Caldas, C., and Ponder, B.A. (2008). Allele-specific up-regulation of FGFR2 increases susceptibility to breast cancer. *PLoS Biol.* *6*, e108.
  19. Belikov, S., Astrand, C., and Wrangé, O. (2009). FoxA1 binding directs chromatin structure and the functional response of a glucocorticoid receptor-regulated promoter. *Mol. Cell. Biol.* *29*, 5413–5425.
  20. Cavdar Koc, E., Ranasinghe, A., Burkhart, W., Blackburn, K., Koc, H., Moseley, A., and Spremulli, L.L. (2001). A new face on apoptosis: death-associated protein 3 and PDCD9 are mitochondrial ribosomal proteins. *FEBS Lett.* *492*, 166–170.
  21. Turner, N., and Grose, R. (2010). Fibroblast growth factor signalling: from development to cancer. *Nat. Rev. Cancer* *10*, 116–129.
  22. Meyer, K.B., O'Reilly, M., Michailidou, K., Carlebur, S., Edwards, S.L., French, J.D., Prathalingham, R., Dennis, J., Bolla, M.K., Wang, Q., et al.; GENICA Network; kConFab Investigators; Australian Ovarian Cancer Study Group (2013). Fine-scale mapping of the FGFR2 breast cancer risk locus: putative functional variants differentially bind FOXA1 and E2F1. *Am. J. Hum. Genet.* *93*, 1046–1060.
  23. Theodorou, V., Boer, M., Weigelt, B., Jonkers, J., van der Valk, M., and Hilken, J. (2004). Fgf10 is an oncogene activated by MMTV insertional mutagenesis in mouse mammary tumors and overexpressed in a subset of human breast carcinomas. *Oncogene* *23*, 6047–6055.
  24. Abolhassani, A., Riazi, G.H., Azizi, E., Amanpour, S., Muhammadnejad, S., Haddadi, M., Zekri, A., and Shirkoohi, R. (2014). FGF10: type III epithelial mesenchymal transition and invasion in breast cancer cell lines. *J. Cancer* *5*, 537–547.
  25. Chioni, A.M., and Grose, R. (2009). Negative regulation of fibroblast growth factor 10 (FGF-10) by polyoma enhancer activator 3 (PEA3). *Eur. J. Cell Biol.* *88*, 371–384.

## Supplemental Data

### Evidence that the 5p12 Variant rs10941679 Confers

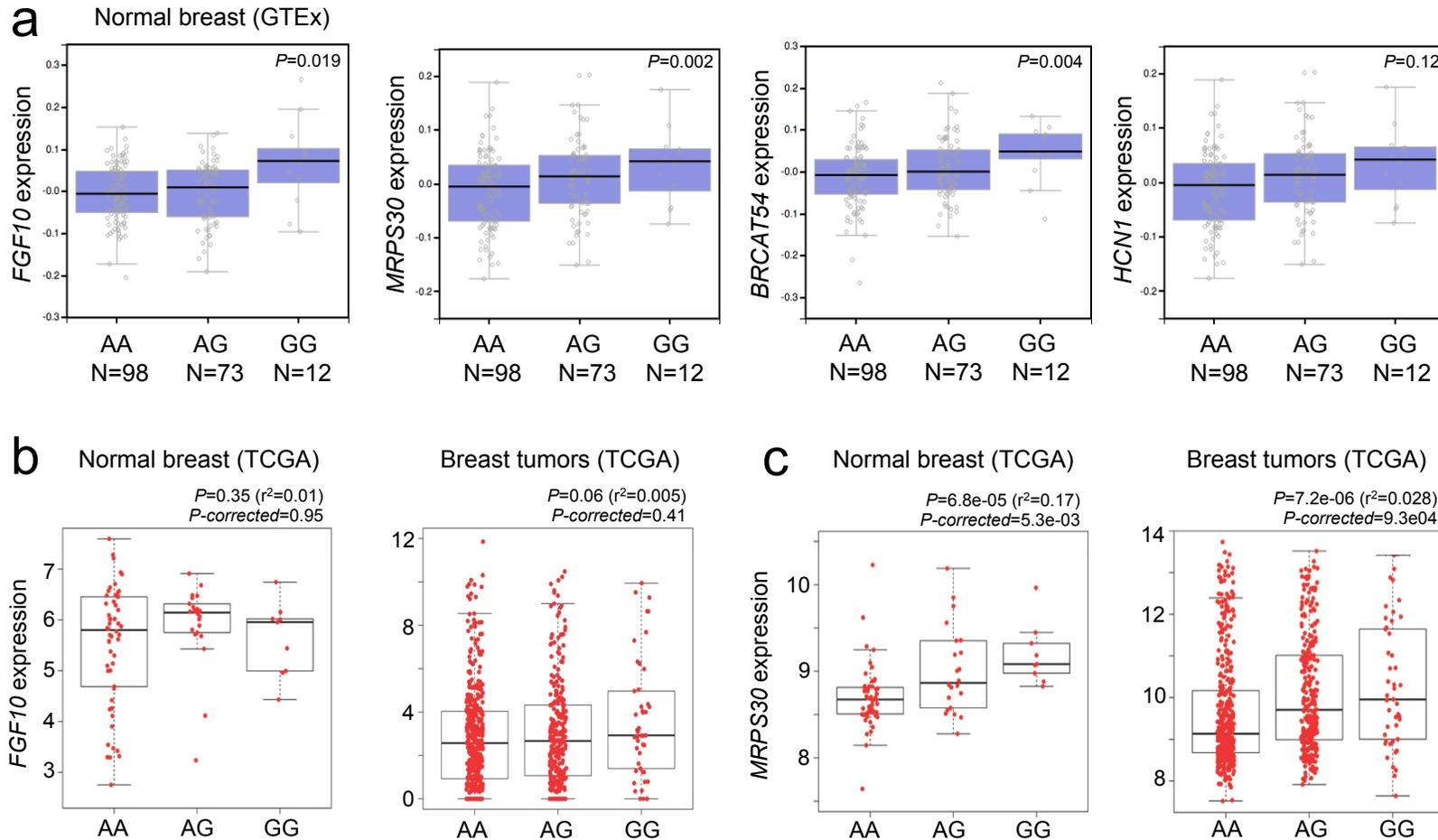
### Susceptibility to Estrogen-Receptor-Positive

### Breast Cancer through *FGF10* and *MRPS30* Regulation

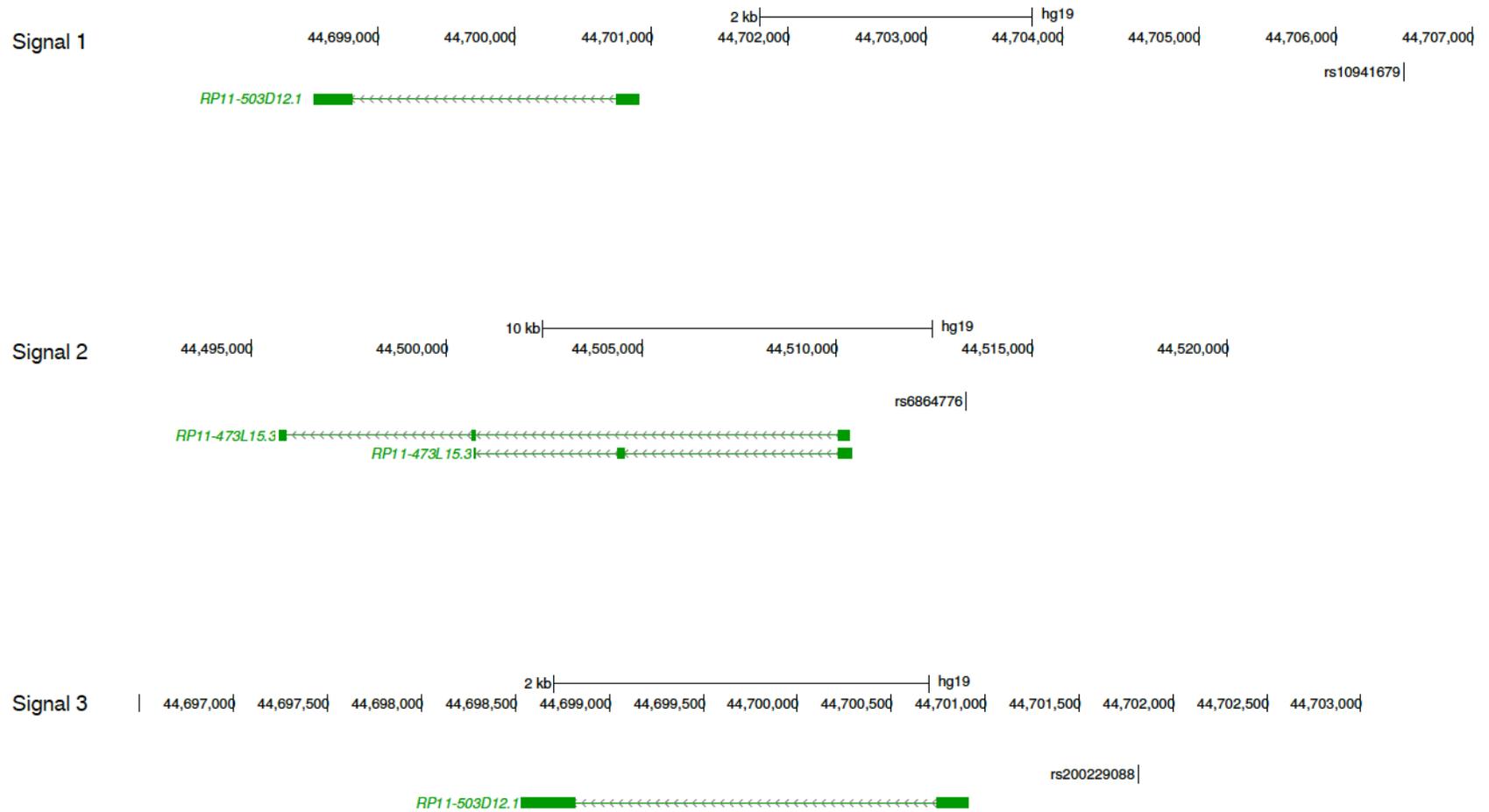
Maya Ghoussaini, Juliet D. French, Kyriaki Michailidou, Silje Nord, Jonathan Beesley, Sander Canisus, Kristine M. Hillman, Susanne Kaufmann, Haran Sivakumaran, Mahdi Moradi Marjaneh, Jason S. Lee, Joe Dennis, Manjeet K. Bolla, Qin Wang, Ed Dicks, Roger L. Milne, John L. Hopper, Melissa C. Southey, Marjanka K. Schmidt, Annegien Broeks, Kenneth Muir, Artitaya Lophatananon, Peter A. Fasching, Matthias W. Beckmann, Olivia Fletcher, Nichola Johnson, Elinor J. Sawyer, Ian Tomlinson, Barbara Burwinkel, Frederik Marme, Pascal Guénel, Thérèse Truong, Stig E. Bojesen, Henrik Flyger, Javier Benitez, Anna González-Neira, M. Rosario Alonso, Guillermo Pita, Susan L. Neuhausen, Hoda Anton-Culver, Hermann Brenner, Volker Arndt, Alfons Meindl, Rita K. Schmutzler, Hiltrud Brauch, Ute Hamann, Daniel C. Tessier, Daniel Vincent, Heli Nevanlinna, Sofia Khan, Keitaro Matsuo, Hidemi Ito, Thilo Dörk, Natalia V. Bogdanova, Annika Lindblom, Sara Margolin, Arto Mannermaa, Veli-Matti Kosma, kConFab/AOCS Investigators, Anna H. Wu, David Van Den Berg, Diether Lambrechts, Giuseppe Floris, Jenny Chang-Claude, Anja Rudolph, Paolo Radice, Monica Barile, Fergus J. Couch, Emily Hallberg, Graham G. Giles, Christopher A. Haiman, Loic Le Marchand, Mark S. Goldberg, Soo H. Teo, Cheng Har Yip, Anne-Lise Borresen-Dale, NBCS Collaborators, Wei Zheng, Qiuyin Cai, Robert Winqvist, Katri Pylkäs, Irene L. Andrulis, Peter Devilee, Rob A.E.M. Tollenaar, Montserrat García-Closas, Jonine Figueroa, Per Hall, Kamila Czene, Judith S. Brand, Hatf Darabi, Mikael Eriksson, Maartje J. Hooning, Linetta B. Koppert, Jingmei Li, Xiao-Ou Shu, Ying Zheng, Angela Cox, Simon S. Cross, Mitul Shah, Valerie Rhenius, Ji-Yeob Choi, Daehee Kang, Mikael Hartman, Kee Seng Chia, Maria Kabisch, Diana Torres, Craig Luccarini, Don M. Conroy, Anna Jakubowska, Jan Lubinski, Suleeporn Sangrajrang, Paul Brennan, Curtis Olswold, Susan Slager, Chen-Yang Shen, Ming-Feng Hou, Anthony Swerdlow, Minouk J. Schoemaker, Jacques Simard, Paul D.P. Pharoah, Vessela Kristensen, Georgia Chenevix-Trench, Douglas F. Easton, Alison M. Dunning, and Stacey L. Edwards

## **SUPPLEMENTAL ACKNOWLEDGMENTS**

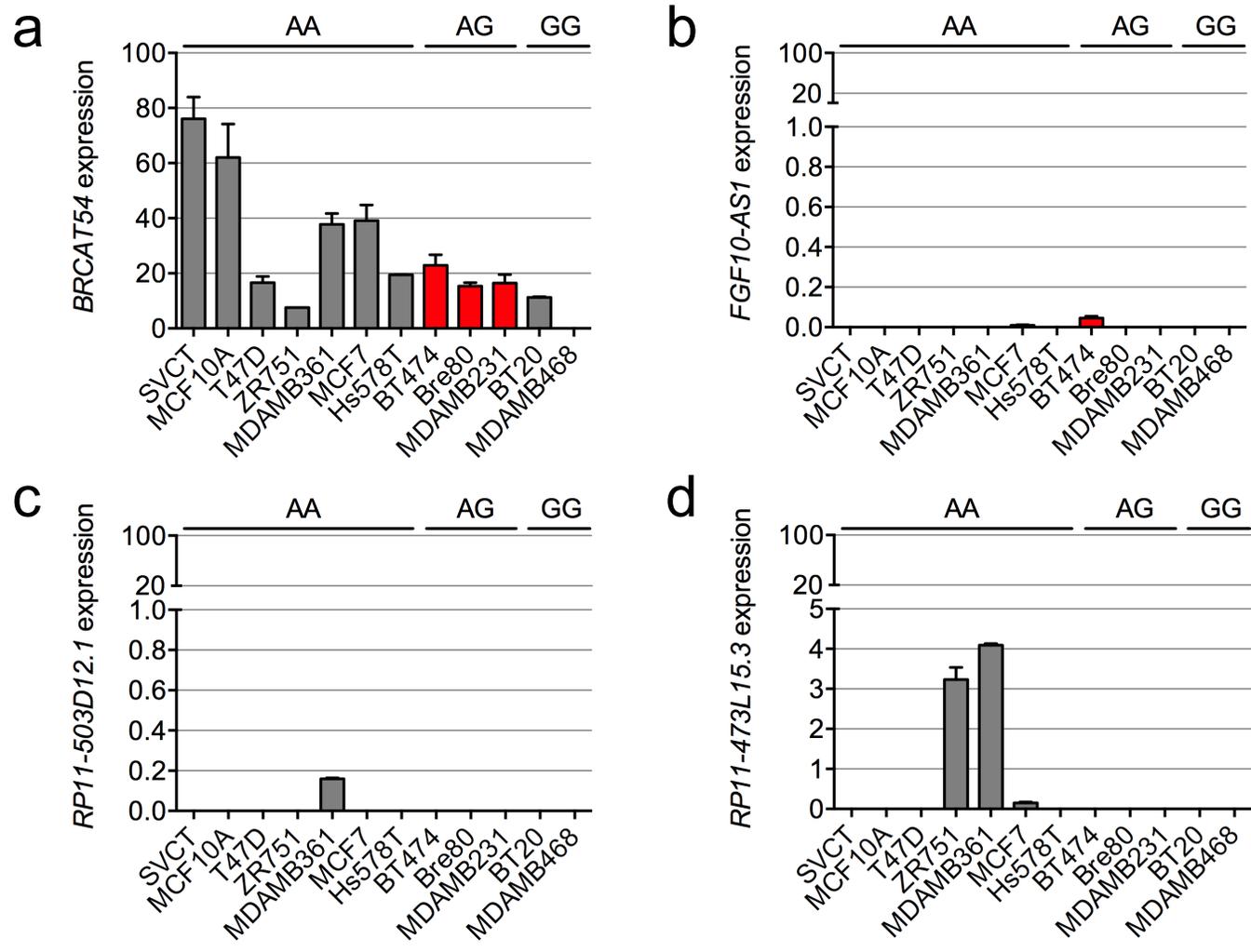
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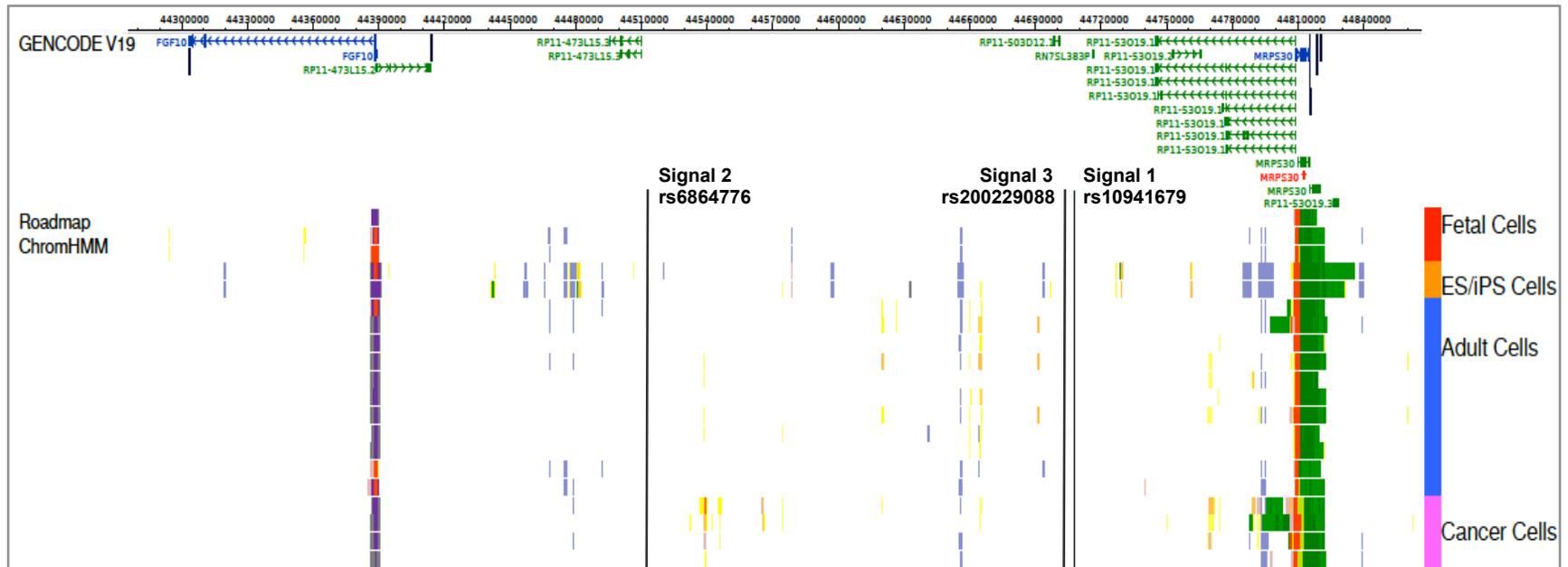
**Supplementary Figure 1. Associations of SNP rs10941679 with expression of candidate target genes. (a)** 183 normal breast samples from the GTEx database, **(b)** *FGF10* or **(c)** *MRPS30* in normal breast TCGA (n=93) and breast tumours TCGA (n=765). The x-axis of each plot corresponds to the three observed SNP genotypes and the y-axis represents log2-normalized gene expression values. *P*-values in the GTEx datasets are pre-corrected for multiple testing. *P*-values in the TCGA datasets are presented before and after correction for multiple testing using FDR as implemented in p.adjust function in R. Each box plot shows the median rank normalized gene expression (horizontal line), the first through third quartiles (box) and 1.5× the interquartile range (whiskers).



**Supplementary Figure 2. Location of putative noncoding RNAs close to the top risk SNPs.** The top risk SNPs from each signal are shown as small horizontal lines and labelled. Green boxes represent putative noncoding RNAs.

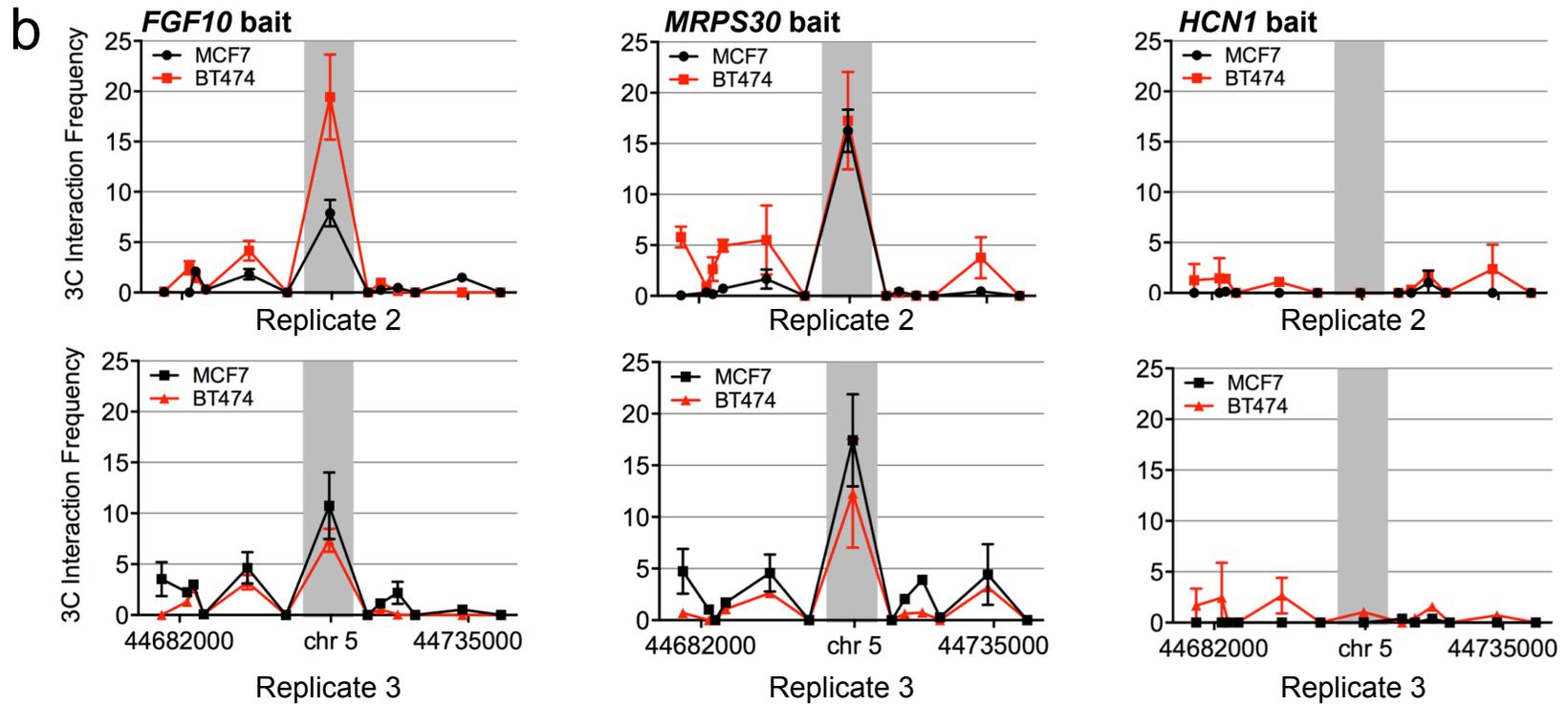
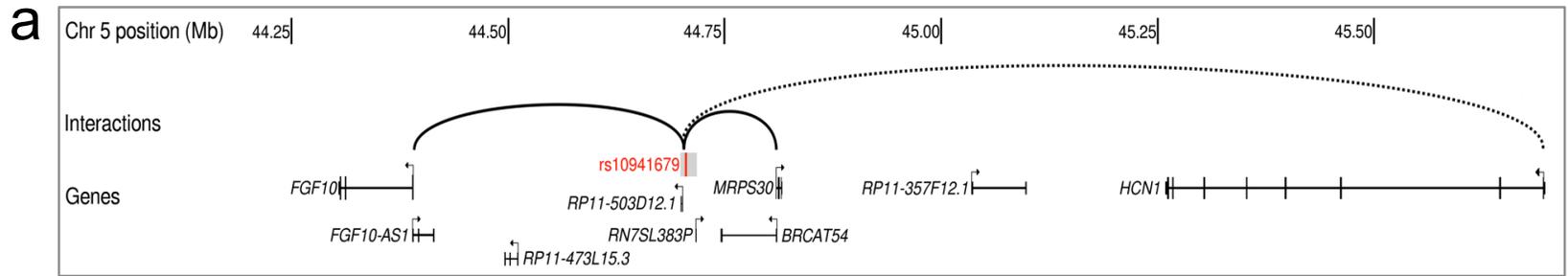


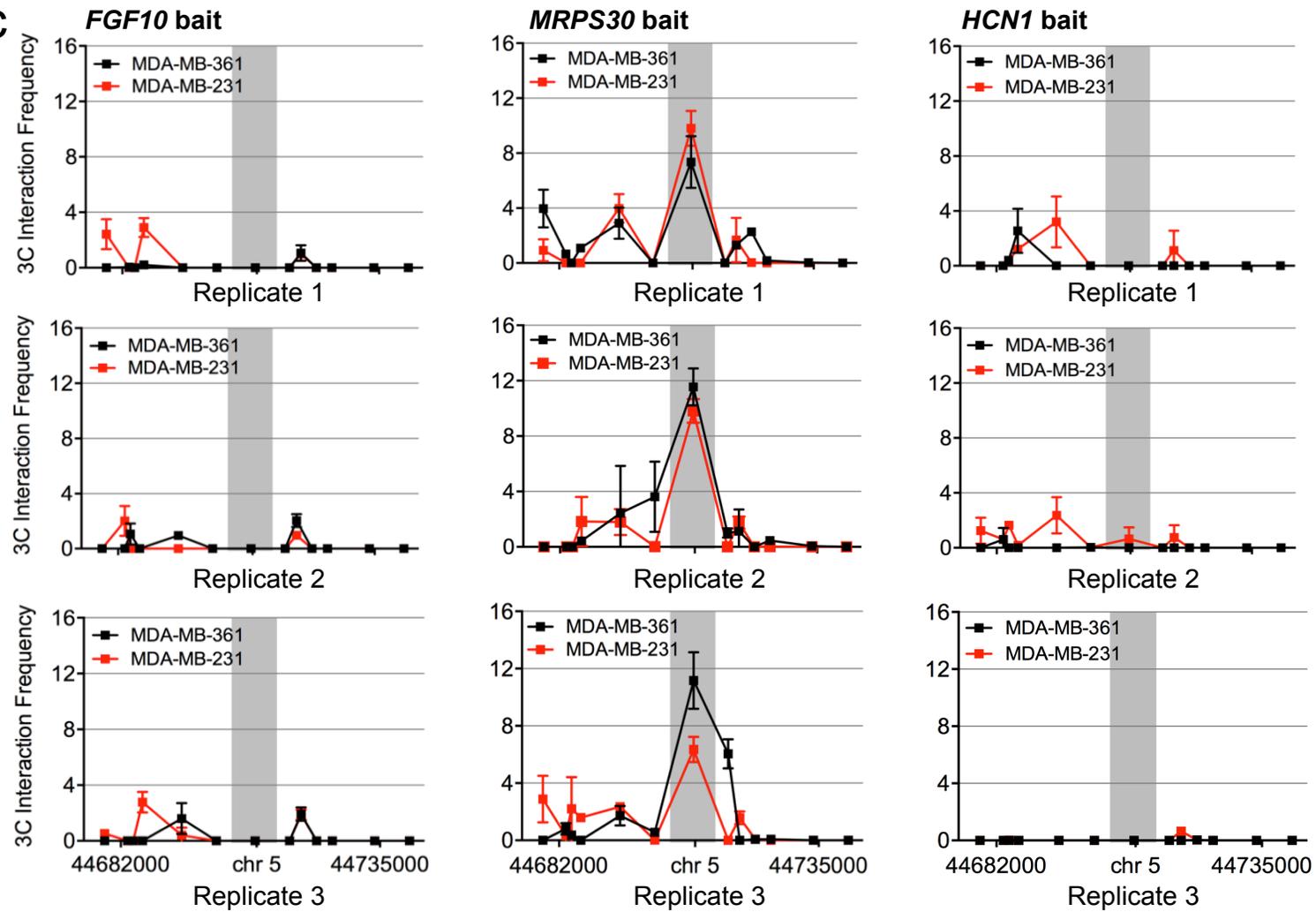
**Supplementary Figure 3.** Putative noncoding RNA expression measured by qPCR in untreated breast cell lines and normalised to *TBP*. Error bars denote SEM (n=3).

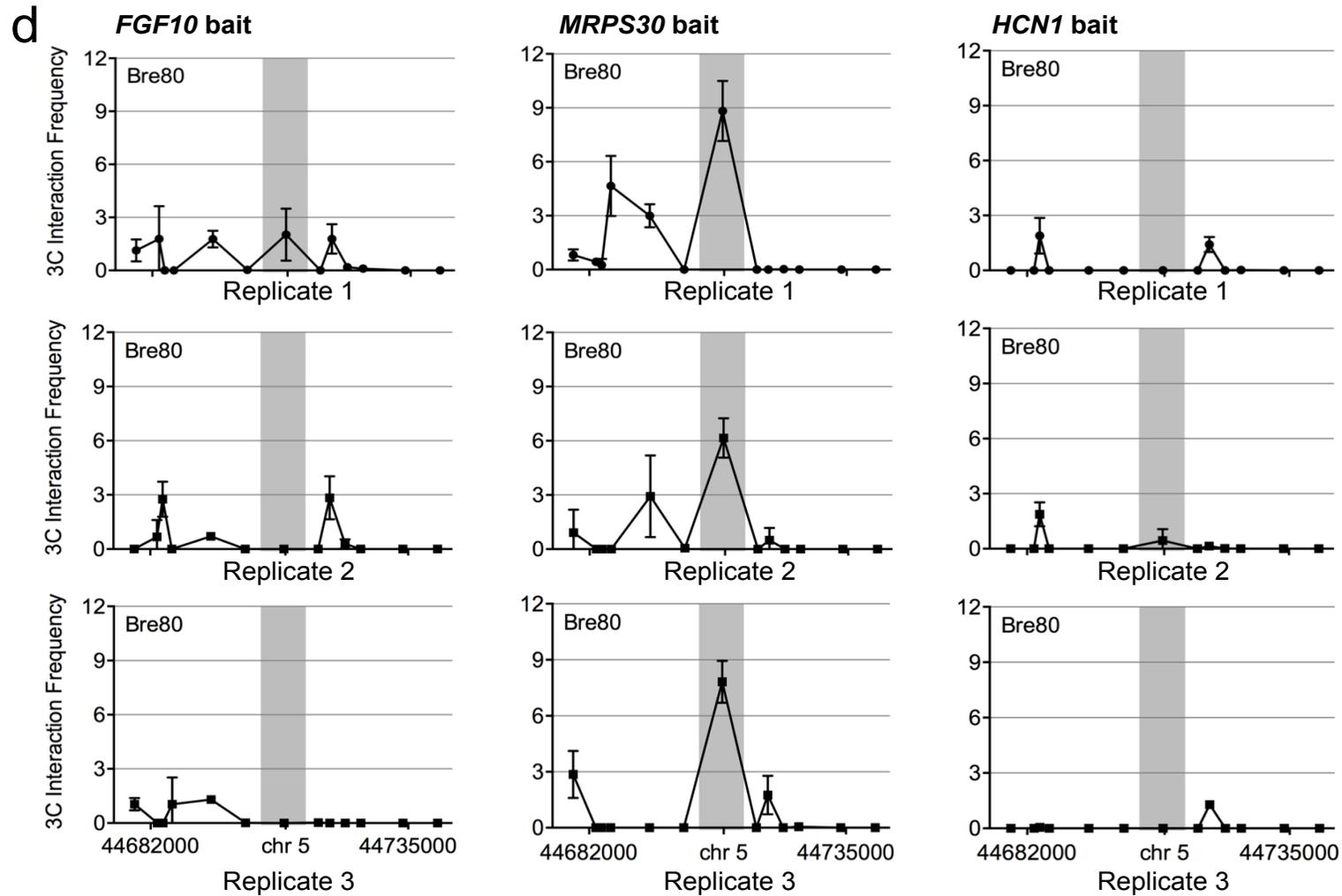


- Active TSS
- Promoter Upstream TSS
- Promoter Downstream TSS with DNase
- Promoter Downstream TSS
- Transcription 5'
- Transcription 3'
- Weak transcription
- Transcription Regulatory
- Transcription 5' Enhancer
- Transcription 3' Enhancer
- Transcription Weak Enhancer
- Active Enhancer 1
- Active Enhancer 2
- Active Enhancer Flank
- Weak Enhancer 1
- Weak Enhancer 2
- Enhancer Acetylation Only
- DNase only
- ZNF genes & repeats
- Heterochromatin
- Poised Promoter
- Bivalent Promoter
- Repressed PolyComb

**Supplementary Figure 4.** Genomic region (chr5:44276112-44866764) showing chromatin state annotations called by Roadmap Epigenomics Project Chromatin Hidden Markov Modelling (Imputed ChromHMM) relative to Gencode V19 gene models. The 25 states of chromatin segmentation depicted as colors along the genomic axis are described in the key. The position of top risk SNPs for signals 1-3 are shown by the vertical black lines.

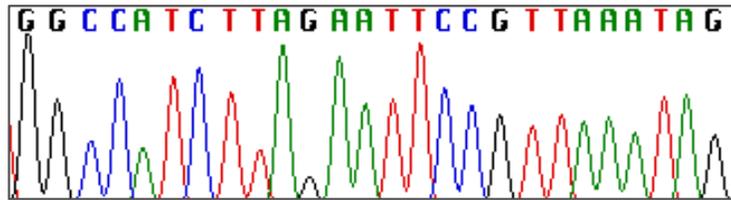


**C**

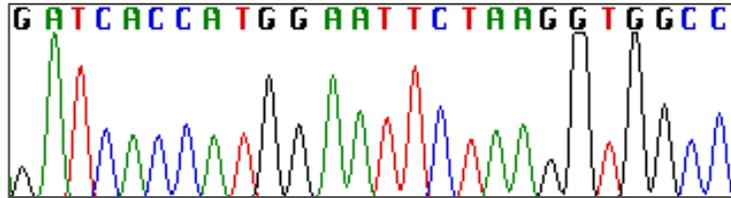


**Supplementary Figure 5. Chromatin interactions at 5p12 in breast cancer cell lines. (a)** Physical map of the region interrogated by 3C. 3C interaction profiles between the *FGF10/FGF10AS-1*, *MRPS30/BRCAT54* or *HCN1* promoters and the putative regulatory element (PRE, grey bars) containing SNP rs10941679 in **(b)** ER<sup>+</sup>/PR<sup>+</sup> MCF7 and BT474, **(c)** ER<sup>+</sup>/PR<sup>-</sup> MDA-MB-361 or ER<sup>-</sup>/PR<sup>-</sup> MDA-MB-231 breast cancer cell lines and **(d)** ER<sup>-</sup> Bre80 normal breast cells. 3C libraries were generated with *Eco*RI, with the anchor points set at the promoters. Error bars denote SD.

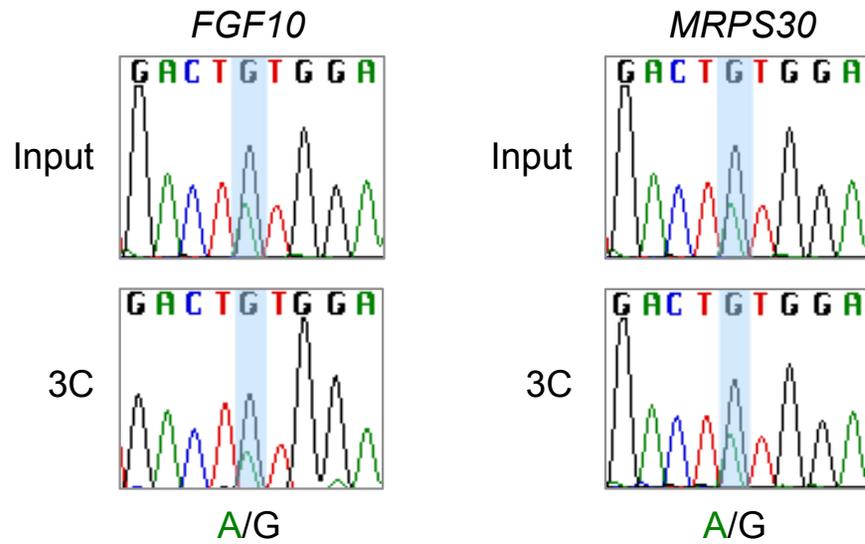
**a** 5p12 PRE ← *Eco*RI → *FGF10* promoter



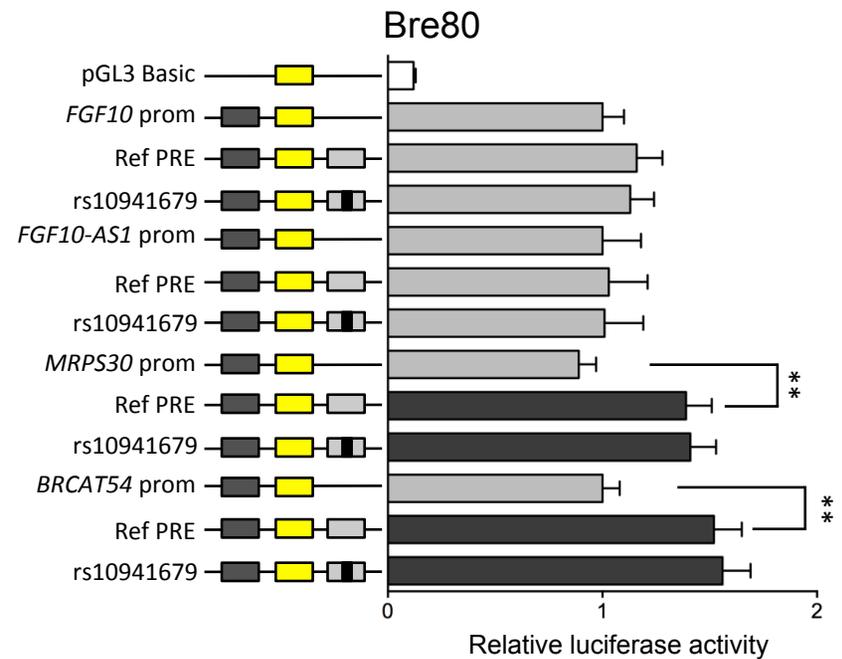
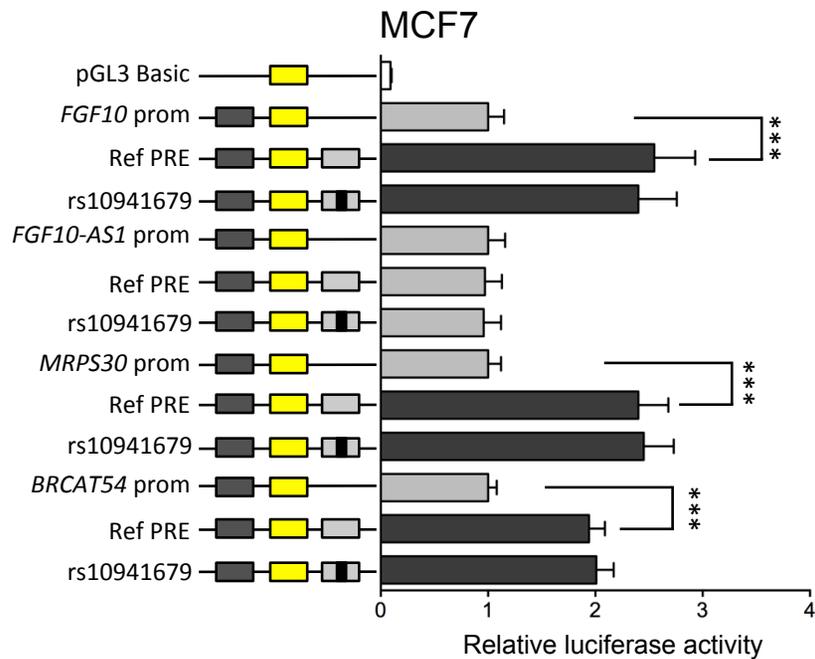
**b** *MRPS30* prom ← *Eco*RI → 5p12 PRE



**Supplementary Figure 6. DNA sequencing to confirm chromatin interactions.** Chromatogram of 3C products amplified between the (a) *FGF10* or (b) *MRPS30* promoters and the 5p12 putative regulatory element (PRE) containing SNP rs10941679.

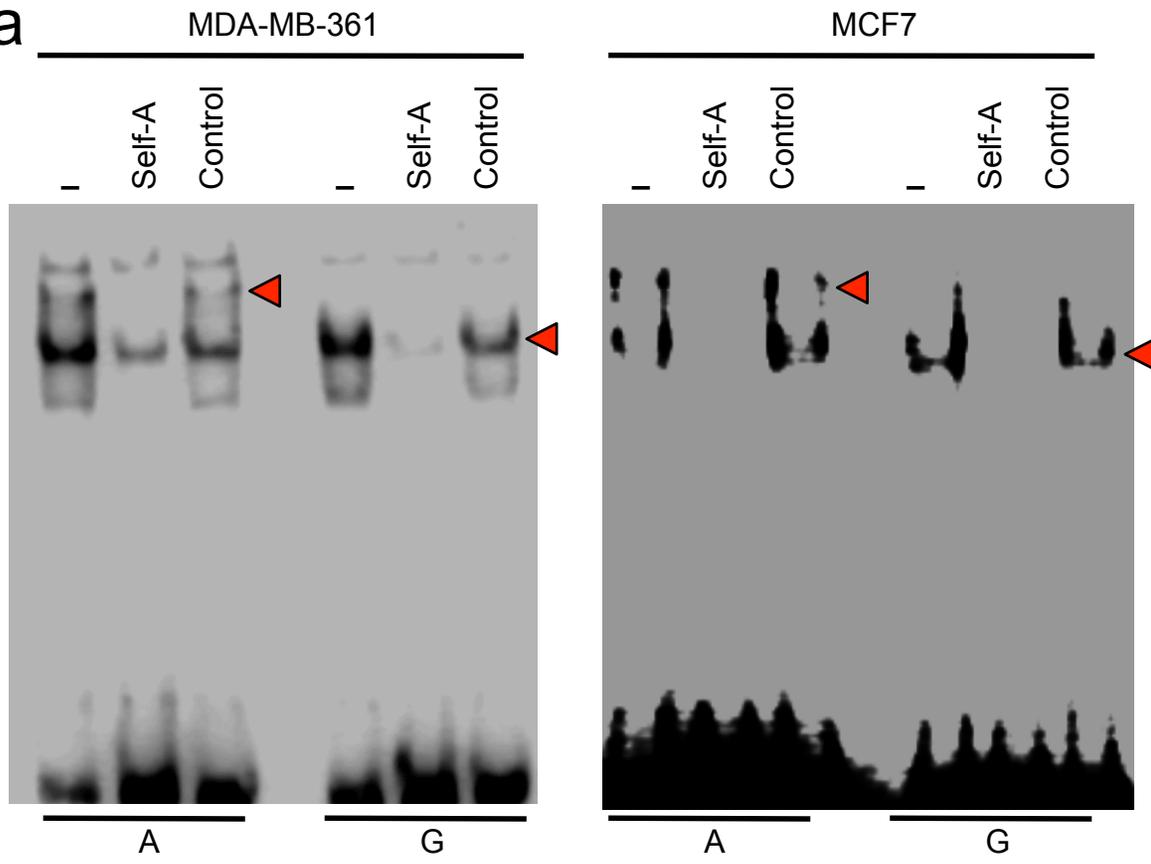


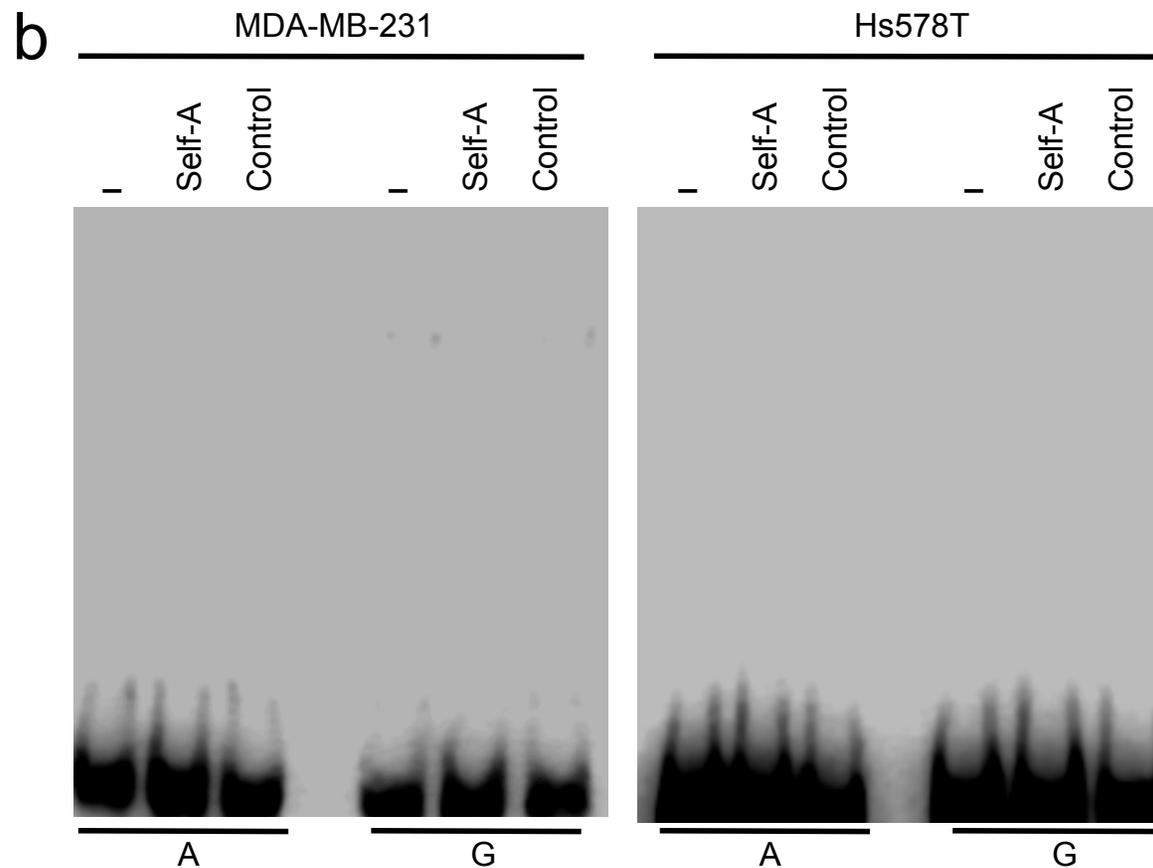
**Supplementary Figure 7. SNP rs10941679 does not influence chromatin looping between the PRE and *FGF10* or *MRPS30* promoters.** 3C followed by sequencing for the rs10941679-containing region in heterozygous BT474 ER+ breast cancer cells.



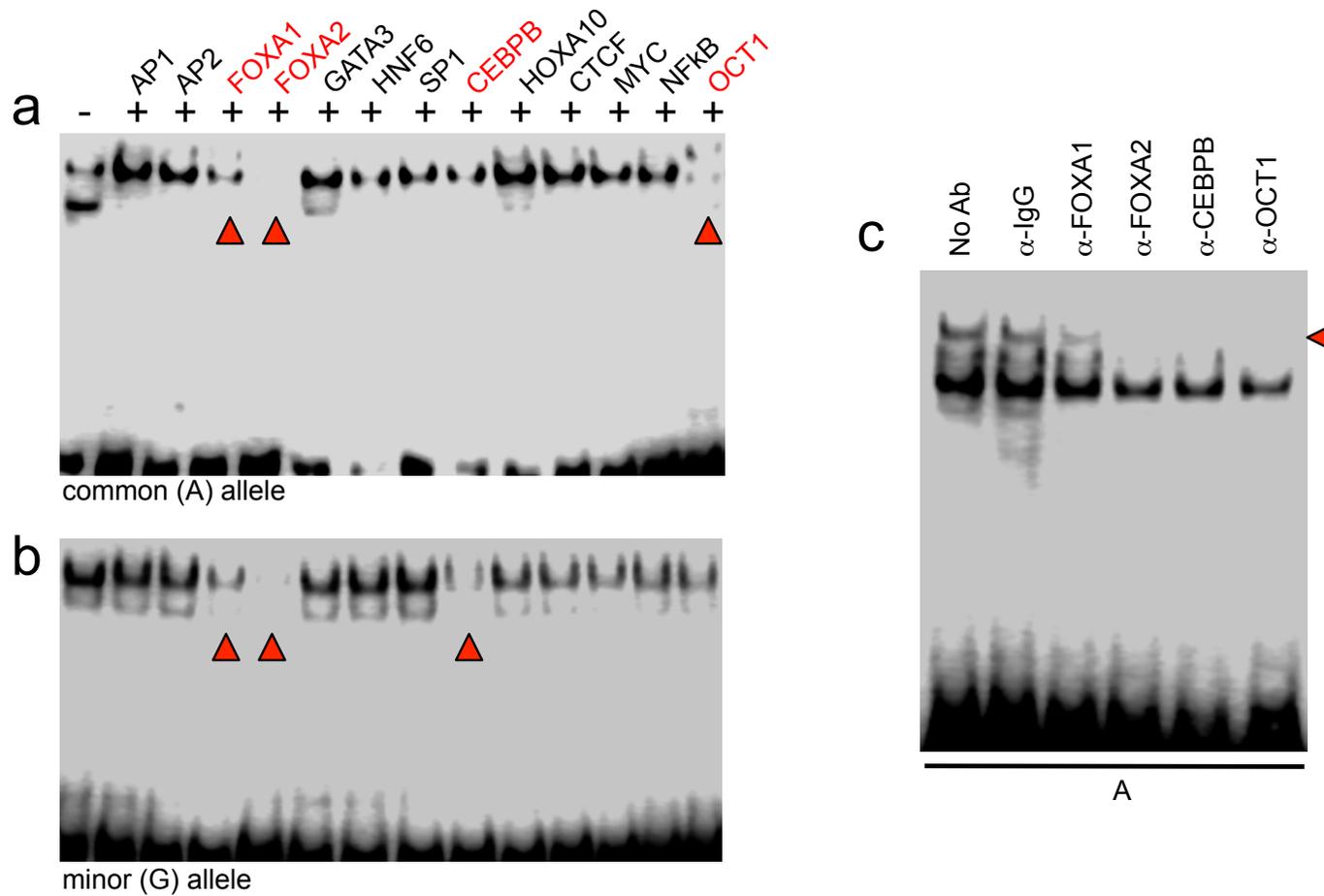
**Supplementary Figure 8. Luciferase reporter assays in ER<sup>+</sup> MCF7 and ER<sup>-</sup> Bre80 breast cell lines.** The putative regulatory element containing the major SNP allele was cloned downstream of target gene promoter-driven luciferase constructs (Ref PRE). The minor SNP allele was engineered into the constructs and is designated by the rs ID. Error bars denote 95% confidence intervals (n=3). *P*-values were determined by 2-way ANOVA followed by Dunnett's multiple comparisons test. \*\**P*<0.01, \*\*\**P*<0.001.

**a**

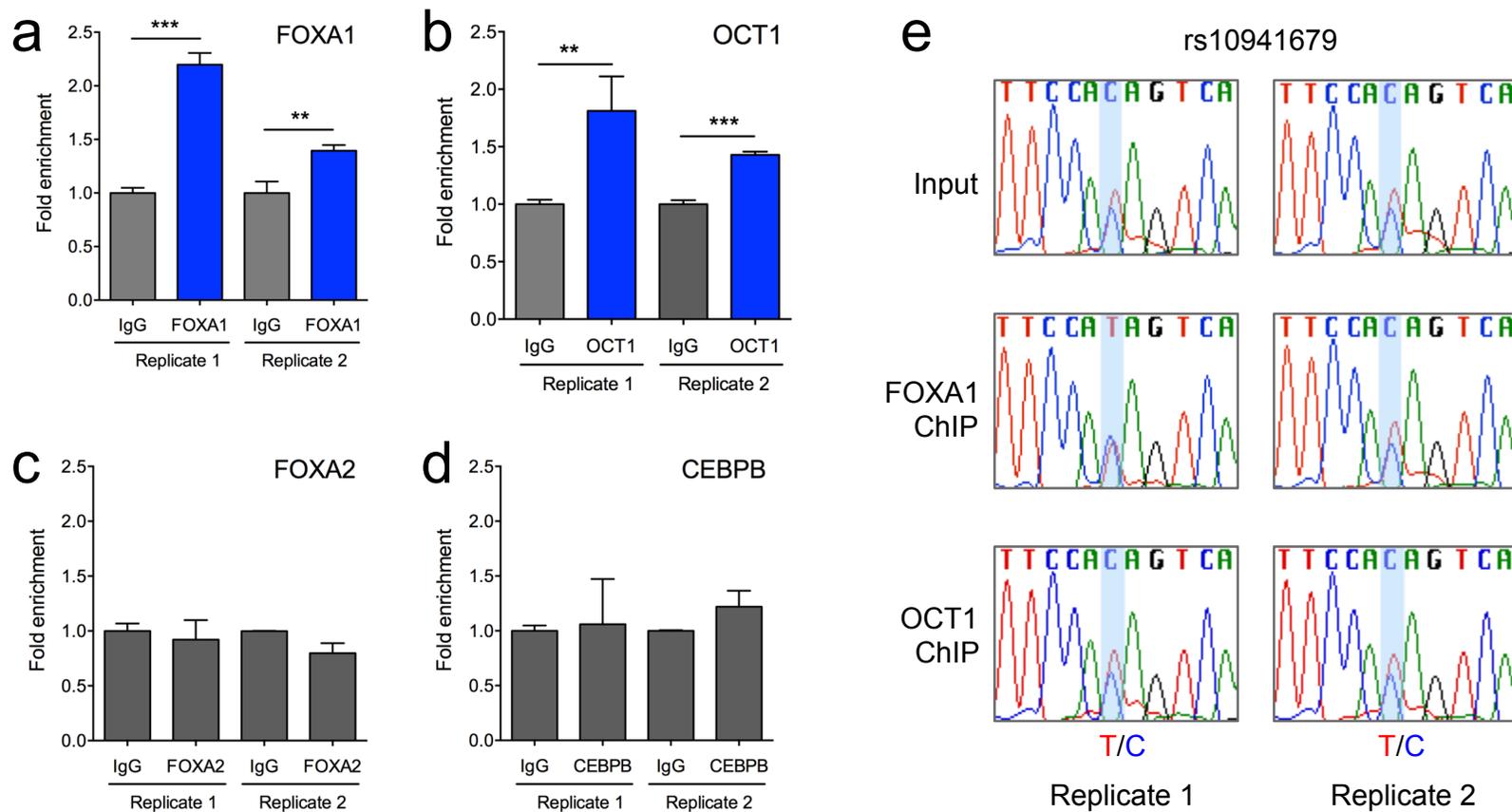




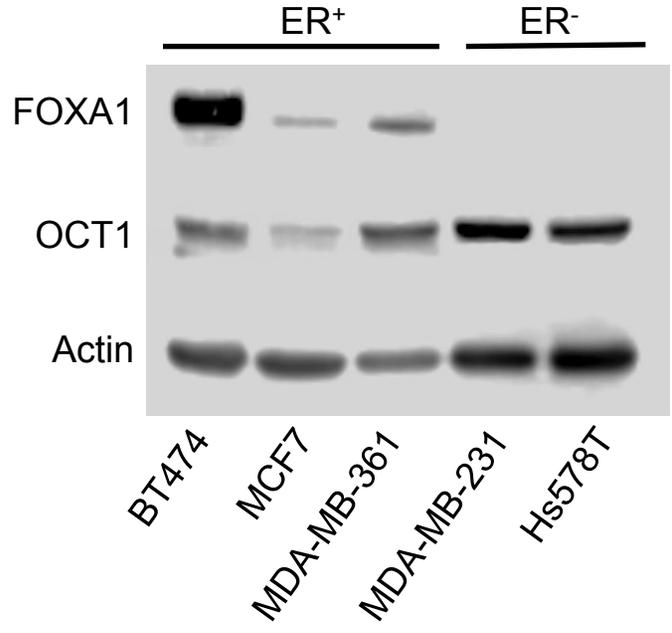
**Supplementary Figure 9. EMSAs for oligonucleotides containing SNP rs10941679.** The A=common allele and G=minor allele as indicated below the panel, assayed using (a) ER<sup>+</sup>/PR<sup>-</sup> MDA-MB-361, ER<sup>+</sup>/PR<sup>+</sup> MCF7 and (b) ER<sup>-</sup>/PR<sup>-</sup> MDA-MB-231 and Hs578T nuclear extracts. Primers are listed in **Table S8**. Labels above each lane indicate inclusion of competitor oligonucleotides at 100-fold molar excess, respectively: (-) no competitor and control denotes a non-specific competitor. Red arrowheads show bands of different mobility detected between the A and G alleles.



**Supplementary Figure 10. Competition EMSAs for SNP rs10941679 to identify candidate nuclear proteins.** Competitor oligonucleotides for predicted transcription factors (100-fold molar excess) were incubated with ER<sup>+</sup> BT474 nuclear extracts (competitor sequences are listed in **Table S9**). Red arrowheads indicate bands that were competed for complex formation on either the **(a)** common or **(b)** minor alleles. **(c)** EMSA-supershift using the common DNA duplex and 2  $\mu$ g of polyclonal antibody against FOXA1 (ab5089, abcam), FOXA2 (sc-20692, Santa Cruz Biotechnology), CEBPB (sc-150) or OCT1 (sc-232) with BT474 nuclear extracts. Rabbit IgG (sc-2027) was used as a negative control. The red arrowhead denotes supershifted complexes.



**Supplementary Figure 11. FOXA1 and OCT1 binding *in vivo*.** ChIP-qPCR results against (a) FOXA1, (b) OCT1, (c) FOXA2 or (d) CEBPB in heterozygous BT474 breast cancer cells. Error bars denote SD (n=2). A region within the second intron of *ESR1* served as a negative (Neg) control. *P*-values were determined by two-tailed t-t test. \*\**P*<0.01, \*\*\**P*<0.001. (e) Sanger sequencing of the PCR fragment generated using primers flanking SNP rs10941679 following ChIP-qPCR. Primers are listed in **Table S10**.



**Supplementary Figure 12. Western blot analysis of FOXA1 and OCT1 protein expression in breast cancer cell lines.** Actin (Sigma A2066) served as a loading control.

**Supplementary Table 1.** Strongest Associated SNPs (n=461  $P<10^{-4}$ ) with overall breast cancer risk from 41 European BCAC studies (n=104,660). \* Type refers to either genotyped or imputed SNPs.

SNP	position	Ref	Alt	EAF	OR overall	Pvalue overall	OR ER+	Pvalue ER+	OR ER-	Pvalue ER-	Type	R <sup>2</sup>
rs10941679	44706498	A	G	0.27	1.12	2.55E-26	1.15	2.44E-29	1.04	0.08	Gen	0.99
rs10462082	44800917	A	C	0.50	1.1	9.11E-22	1.11	3.40E-20	1.06	0.00	Imp	0.95
c5_pos44911206	44875449	A	G	0.48	0.91	1.67E-21	0.90	3.28E-21	0.95	0.01	Gen	1.00
rs6451770	44691395	T	G	0.39	1.1	2.50E-21	1.13	4.76E-25	1.01	0.59	Gen	1.00
rs7716600	44875005	A	C	0.77	0.9	4.13E-21	0.87	3.17E-24	0.96	0.09	Gen	1.00
rs930395	44822458	G	A	0.23	1.11	7.58E-21	1.14	4.87E-24	1.04	0.08	Gen	1.00
rs13185555	44690964	G	A	0.39	1.1	7.59E-21	1.13	5.36E-25	1.01	0.75	Imp	0.98
rs76514876	44673354	A	C	0.41	1.1	4.91E-20	1.12	5.39E-23	1.01	0.72	Imp	0.98
rs4463188	44642670	T	C	0.41	1.09	5.35E-20	1.12	8.29E-23	1.00	0.80	Gen	1.00
rs4321755	44646195	C	T	0.41	1.09	6.44E-20	1.12	6.75E-23	1.00	0.81	Gen	1.00
rs4339357	44670741	T	C	0.40	1.1	6.64E-20	1.12	9.24E-23	1.01	0.56	Imp	0.97
rs4479849	44644006	G	A	0.41	1.09	6.93E-20	1.12	4.92E-23	1.00	0.85	Gen	1.00
rs12187196	44683819	A	C	0.41	1.09	7.47E-20	1.12	8.48E-23	1.01	0.62	Gen	1.00
rs36068815	44645998	T	C	0.41	1.09	7.80E-20	1.12	5.15E-23	1.00	0.84	Gen	1.00
rs13155752	44680687	A	C	0.41	1.09	1.65E-19	1.12	7.46E-23	1.01	0.71	Imp	0.98
rs13156283	44680758	G	C	0.41	1.09	1.66E-19	1.12	7.48E-23	1.01	0.71	Imp	0.98
rs4492118	44646625	G	A	0.41	1.09	1.86E-19	1.12	1.98E-22	1.01	0.79	Gen	1.00
rs7735881	44650176	A	G	0.41	1.09	1.90E-19	1.12	1.15E-22	1.00	0.82	Gen	1.00
rs12516900	44644495	A	G	0.41	1.09	2.27E-19	1.12	2.34E-22	1.01	0.78	Gen	1.00
rs6861560	44708378	C	G	0.41	1.09	2.31E-19	1.12	7.82E-23	1.01	0.71	Imp	0.98
rs1821936	44699482	A	C	0.41	1.09	2.42E-19	1.12	1.30E-22	1.01	0.67	Imp	0.99
chr5:44648592:I	44648592	C	CTAT	0.41	1.09	2.42E-19	1.12	2.22E-22	1.01	0.76	Imp	0.98
chr5:44645330:D	44645330	AG	A	0.40	1.1	2.50E-19	1.12	3.61E-22	1.01	0.77	Imp	0.95
rs6874055	44666965	T	A	0.41	1.09	2.51E-19	1.12	1.70E-22	1.01	0.75	Imp	0.99
rs12522626	44685698	G	T	0.41	1.09	2.67E-19	1.12	1.73E-22	1.01	0.69	Gen	1.00
rs10805686	44662128	T	C	0.41	1.09	2.95E-19	1.12	2.24E-22	1.01	0.76	Gen	1.00
rs6890556	44648666	C	G	0.41	1.09	3.01E-19	1.12	3.17E-22	1.01	0.76	Gen	1.00
rs2165010	44706780	A	C	0.41	1.09	3.12E-19	1.12	1.68E-22	1.01	0.68	Imp	0.99
rs13176502	44697908	T	C	0.41	1.09	3.36E-19	1.12	2.17E-22	1.01	0.66	Imp	0.99
rs714130	44701418	T	C	0.41	1.09	3.55E-19	1.12	2.03E-22	1.01	0.68	Gen	1.00
rs1438825	44706931	G	A	0.41	1.09	3.58E-19	1.12	1.63E-22	1.01	0.72	Gen	1.00
rs10941677	44662399	G	A	0.41	1.09	3.64E-19	1.12	2.24E-22	1.01	0.77	Gen	1.00
rs1371027	44699857	T	C	0.41	1.09	3.65E-19	1.12	1.90E-22	1.01	0.69	Gen	1.00
rs2218081	44705140	C	T	0.41	1.09	3.76E-19	1.12	1.50E-22	1.01	0.74	Gen	1.00
rs4415084	44662515	C	T	0.41	1.09	3.81E-19	1.12	2.67E-22	1.01	0.78	Gen	1.00

rs1898701	44695429	A	G	0.41	1.09	3.87E-19	1.12	2.53E-22	1.01	0.67	Imp	0.99
rs2165009	44697916	C	T	0.41	1.09	4.09E-19	1.12	1.53E-22	1.01	0.72	Gen	0.99
rs4415085	44662959	C	G	0.41	1.09	4.21E-19	1.12	2.45E-22	1.00	0.82	Gen	1.00
rs6871484	44676305	T	C	0.41	1.09	4.24E-19	1.12	2.26E-22	1.01	0.75	Imp	0.99
rs920329	44702507	T	C	0.41	1.09	4.54E-19	1.12	1.71E-22	1.01	0.75	Gen	1.00
rs6863598	44668623	C	T	0.41	1.09	4.62E-19	1.12	2.67E-22	1.01	0.76	Imp	0.98
rs4419600	44678534	T	G	0.41	1.09	4.72E-19	1.12	4.27E-22	1.01	0.74	Gen	1.00
rs34762678	44705514	T	C	0.41	1.09	4.75E-19	1.12	2.22E-22	1.01	0.67	Gen	1.00
rs4571480	44687188	C	T	0.41	1.09	4.85E-19	1.12	1.64E-22	1.01	0.77	Gen	1.00
rs13156930	44698035	G	C	0.41	1.09	4.92E-19	1.12	1.76E-22	1.01	0.75	Gen	1.00
c5_pos44711954	44676197	C	T	0.41	1.09	5.20E-19	1.12	3.06E-22	1.00	0.80	Gen	1.00
chr5:44637587:I	44637587	T	TC	0.40	1.09	5.28E-19	1.12	2.86E-22	1.00	0.87	Imp	0.97
chr5:44929230:D	44929230	ATAATAT	A	0.60	0.91	5.60E-19	0.89	4.32E-21	0.98	0.44	Imp	0.92
rs10462079	44700110	G	T	0.41	1.09	5.77E-19	1.12	3.09E-22	1.01	0.69	Gen	0.99
chr5:44678911:D	44678911	TA	T	0.40	1.09	5.83E-19	1.12	3.22E-22	1.01	0.79	Imp	0.97
rs2218080	44714330	T	C	0.42	1.09	5.99E-19	1.12	2.50E-22	1.01	0.76	Imp	0.96
rs2013513	44702306	G	A	0.41	1.09	6.02E-19	1.12	3.39E-22	1.01	0.68	Imp	0.99
rs13179137	44702692	C	T	0.41	1.09	6.55E-19	1.12	2.82E-22	1.01	0.69	Gen	1.00
rs7720551	44664477	C	T	0.41	1.09	7.05E-19	1.12	4.87E-22	1.00	0.81	Imp	0.99
rs10941678	44672783	C	A	0.38	1.1	7.18E-19	1.13	7.22E-22	1.00	0.98	Imp	0.89
rs4607355	44868070	A	G	0.52	0.91	7.56E-19	0.89	3.46E-20	0.97	0.12	Imp	0.84
rs144629423	44669403	G	T	0.41	1.09	8.10E-19	1.12	4.28E-22	1.01	0.75	Imp	0.97
rs6884702	44682589	A	G	0.40	1.09	8.55E-19	1.12	3.18E-22	1.01	0.75	Imp	0.99
rs7723539	44660210	G	C	0.41	1.09	8.60E-19	1.12	5.63E-22	1.00	0.84	Imp	0.99
rs10805685	44661958	C	G	0.41	1.09	9.02E-19	1.12	5.83E-22	1.00	0.84	Imp	0.99
rs4571481	44687433	C	T	0.39	1.09	9.77E-19	1.11	4.21E-21	1.03	0.19	Gen	0.95
rs12515012	44694535	G	A	0.40	1.09	9.84E-19	1.12	4.89E-22	1.01	0.75	Imp	0.98
rs181516187	44672104	C	T	0.40	1.09	1.11E-18	1.12	6.79E-22	1.00	0.82	Imp	0.97
rs16901937	44709141	A	G	0.41	1.09	1.71E-18	1.12	7.12E-22	1.01	0.79	Gen	1.00
rs7701466	44663137	C	T	0.40	1.09	3.42E-18	1.12	1.08E-21	1.00	1.00	Imp	0.97
rs11949847	44752169	A	G	0.42	1.09	3.92E-18	1.11	1.08E-20	1.00	0.81	Gen	0.99
c5_pos44685154	44649397	A	C	0.38	1.09	4.42E-18	1.11	1.23E-20	1.02	0.32	Gen	0.98
rs56248730	44681267	A	G	0.38	1.09	4.64E-18	1.12	9.44E-21	1.02	0.32	Imp	0.98
c5_pos44720000	44684243	A	G	0.38	1.09	4.65E-18	1.12	8.11E-21	1.02	0.32	Gen	1.00
rs6867533	44827292	G	T	0.40	1.09	5.17E-18	1.11	2.05E-20	1.00	0.84	Gen	0.96
rs994793	44743247	A	G	0.42	1.09	5.41E-18	1.11	1.04E-20	1.00	0.82	Gen	1.00
rs920328	44699051	G	A	0.38	1.09	6.04E-18	1.12	7.48E-21	1.02	0.31	Gen	1.00
rs11746980	44777878	G	A	0.42	1.09	6.64E-18	1.11	2.04E-20	1.00	0.80	Gen	1.00
rs2330572	44740989	A	C	0.42	1.09	7.56E-18	1.11	1.25E-20	1.00	0.84	Gen	1.00
rs7737491	44790842	A	G	0.37	1.09	8.66E-18	1.11	4.58E-19	1.03	0.20	Imp	0.96
rs10447145	44689796	T	G	0.38	1.09	8.76E-18	1.11	1.33E-20	1.02	0.34	Imp	0.98

rs145342568	44672237	A	G	0.39	1.09	1.00E-17	1.12	8.13E-21	1.01	0.69	Imp	0.91
rs13160259	44828964	G	C	0.42	1.09	1.17E-17	1.11	5.45E-20	1.01	0.77	Imp	0.98
rs10072259	44834791	T	C	0.61	0.92	1.22E-17	0.90	1.39E-19	0.98	0.35	Imp	0.98
rs10512865	44823367	C	T	0.41	1.09	1.27E-17	1.11	6.61E-20	1.01	0.77	Imp	0.99
rs11958808	44945090	G	C	0.41	1.09	1.36E-17	1.11	1.25E-19	1.01	0.76	Gen	1.00
rs4866922	44932505	C	G	0.59	0.92	1.41E-17	0.90	1.11E-19	0.99	0.72	Imp	0.99
chr5:44837254:l	44837254	T	TA	0.58	0.92	1.43E-17	0.90	7.54E-20	0.99	0.77	Imp	0.99
rs1048758	44828594	G	A	0.41	1.09	1.43E-17	1.11	6.71E-20	1.00	0.80	Gen	1.00
rs1371023	44846575	C	T	0.59	0.92	1.52E-17	0.90	8.79E-20	1.00	0.79	Gen	1.00
rs10054773	44930234	C	G	0.59	0.92	1.54E-17	0.90	1.30E-19	0.99	0.71	Imp	0.98
rs10044096	44927365	A	C	0.59	0.92	1.55E-17	0.90	1.49E-19	0.99	0.71	Imp	0.99
rs7716571	44816984	G	A	0.42	1.09	1.62E-17	1.11	1.20E-19	1.01	0.78	Gen	0.96
chr5:44884873:D	44884873	TAAAC	T	0.58	0.92	1.73E-17	0.90	1.36E-19	0.99	0.73	Imp	0.98
rs2118764	44787723	C	T	0.41	1.09	1.77E-17	1.11	1.40E-19	1.00	0.80	Imp	0.98
rs10064434	44834702	G	A	0.58	0.92	1.81E-17	0.90	7.93E-20	1.00	0.81	Imp	0.98
rs930396	44856451	T	G	0.59	0.92	1.89E-17	0.90	1.58E-19	0.99	0.79	Gen	1.00
rs7380559	44837010	A	T	0.59	0.92	1.91E-17	0.90	1.02E-19	1.00	0.80	Gen	1.00
rs1821934	44789172	T	C	0.41	1.09	1.91E-17	1.11	1.63E-19	1.01	0.74	Imp	0.98
rs7711697	44780403	A	T	0.41	1.09	1.95E-17	1.11	1.24E-19	1.00	0.82	Gen	1.00
rs7711136	44823079	G	C	0.41	1.09	1.96E-17	1.11	1.10E-19	1.00	0.79	Gen	0.99
rs1438820	44797770	G	A	0.41	1.09	2.03E-17	1.11	1.20E-19	1.00	0.84	Gen	1.00
rs4866923	44932641	A	C	0.58	0.92	2.07E-17	0.90	2.19E-19	0.99	0.74	Imp	0.97
rs6896417	44802638	A	G	0.41	1.09	2.15E-17	1.11	1.56E-19	1.00	0.84	Gen	1.00
rs1061310	44820850	G	C	0.41	1.09	2.15E-17	1.11	1.16E-19	1.01	0.79	Gen	1.00
rs1438821	44858451	G	A	0.59	0.92	2.17E-17	0.90	1.82E-19	1.00	0.80	Gen	1.00
rs13177711	44796962	A	T	0.41	1.09	2.23E-17	1.11	1.69E-19	1.00	0.83	Gen	0.99
rs4629607	44857467	G	T	0.58	0.92	2.23E-17	0.90	6.97E-18	0.97	0.11	Imp	0.90
rs13159598	44805926	A	G	0.41	1.09	2.27E-17	1.11	1.61E-19	1.00	0.83	Gen	1.00
rs7730841	44823134	T	C	0.42	1.09	2.32E-17	1.11	1.24E-19	1.00	0.80	Gen	0.98
rs13362132	44858260	T	G	0.59	0.92	2.54E-17	0.90	2.03E-19	1.00	0.79	Gen	1.00
rs10043344	44926518	A	T	0.59	0.92	2.57E-17	0.90	1.50E-19	0.99	0.76	Gen	1.00
rs187822318	44671468	C	G	0.35	1.1	2.71E-17	1.12	2.72E-19	1.02	0.28	Imp	0.84
rs7720787	44817309	G	A	0.41	1.09	2.88E-17	1.11	1.52E-19	1.00	0.82	Gen	1.00
rs7705343	44879577	G	A	0.59	0.92	3.06E-17	0.90	2.70E-19	0.99	0.79	Gen	1.00
rs149708574	44825919	T	A	0.30	1.1	3.10E-17	1.12	2.35E-18	1.04	0.07	Imp	0.84
rs2877172	44790567	A	G	0.41	1.09	3.13E-17	1.11	2.26E-19	1.00	0.83	Imp	0.97
rs4518409	44870852	T	C	0.59	0.92	3.18E-17	0.90	2.75E-19	1.00	0.83	Gen	1.00
rs10077814	44916789	C	T	0.59	0.92	3.39E-17	0.90	1.88E-19	1.00	0.80	Gen	1.00
rs4329028	44872353	T	A	0.59	0.92	3.64E-17	0.90	4.23E-19	0.99	0.76	Gen	1.00
chr5:44921964:l	44921964	A	AT	0.59	0.92	4.14E-17	0.90	3.42E-19	0.99	0.78	Imp	0.96
c5_pos44933593	44897836	T	C	0.59	0.92	4.45E-17	0.90	2.78E-19	0.99	0.78	Gen	0.99

rs9292915	44871657	C	T	0.59	0.92	4.49E-17	0.90	4.55E-19	0.99	0.78	Gen	1.00
rs9292913	44870879	G	A	0.59	0.92	4.63E-17	0.90	4.50E-19	1.00	0.80	Gen	1.00
rs4412123	44876288	T	C	0.59	0.92	4.79E-17	0.90	4.15E-19	1.00	0.86	Gen	1.00
rs9790896	44900091	G	A	0.59	0.92	5.21E-17	0.90	2.76E-19	1.00	0.86	Gen	1.00
chr5:44818855:D	44818855	CCAA	C	0.38	1.09	5.37E-17	1.11	1.48E-18	1.02	0.26	Imp	0.99
rs9790879	44899885	C	T	0.59	0.92	5.65E-17	0.90	2.82E-19	1.00	0.86	Gen	1.00
rs11957920	44905086	T	C	0.59	0.92	5.72E-17	0.90	4.80E-19	1.00	0.86	Gen	1.00
rs10070339	44907882	G	A	0.59	0.92	6.40E-17	0.90	4.39E-19	1.00	0.90	Gen	1.00
rs7726586	44848822	C	T	0.62	0.92	9.03E-17	0.90	1.89E-18	0.98	0.33	Gen	1.00
rs13154781	44775027	C	T	0.39	1.09	9.35E-17	1.11	9.73E-19	1.02	0.35	Imp	0.99
rs1438827	44751956	C	A	0.39	1.09	9.41E-17	1.11	7.00E-19	1.02	0.35	Gen	1.00
rs7712949	44770345	C	T	0.39	1.09	9.90E-17	1.11	7.94E-19	1.02	0.37	Gen	1.00
chr5:44810239:D	44810239	TG	T	0.41	1.09	1.01E-16	1.11	6.06E-19	1.01	0.74	Imp	0.96
rs7708213	44848401	T	C	0.62	0.92	1.03E-16	0.90	2.52E-18	0.98	0.30	Gen	1.00
rs34237302	44852862	C	A	0.61	0.92	1.04E-16	0.90	2.61E-18	0.98	0.29	Gen	0.99
rs4440370	44853352	A	G	0.62	0.92	1.10E-16	0.90	2.63E-18	0.98	0.31	Gen	1.00
rs11747159	44737710	C	T	0.39	1.09	1.12E-16	1.11	1.12E-18	1.02	0.34	Gen	1.00
rs4492119	44855614	A	G	0.61	0.92	1.37E-16	0.90	2.83E-18	0.98	0.33	Gen	0.98
rs1371022	44847992	T	A	0.62	0.92	1.42E-16	0.90	3.56E-18	0.98	0.32	Gen	1.00
rs987394	44846378	A	G	0.62	0.92	1.49E-16	0.90	3.64E-18	0.98	0.31	Gen	1.00
rs10462081	44800665	G	A	0.38	1.09	1.58E-16	1.11	4.22E-18	1.02	0.33	Gen	1.00
rs1438819	44797846	T	A	0.38	1.09	1.60E-16	1.11	4.54E-18	1.02	0.34	Gen	1.00
rs7715731	44846844	A	C	0.62	0.92	1.69E-16	0.90	3.01E-18	0.98	0.33	Gen	1.00
rs1371025	44834233	G	A	0.62	0.92	1.70E-16	0.90	3.46E-18	0.98	0.32	Gen	0.99
rs1866406	44809945	G	C	0.39	1.09	1.74E-16	1.11	4.31E-18	1.02	0.32	Gen	0.99
rs9637797	44818181	A	G	0.39	1.09	1.74E-16	1.11	3.90E-18	1.02	0.32	Gen	1.00
rs930394	44822617	G	A	0.38	1.09	1.83E-16	1.11	3.68E-18	1.02	0.32	Gen	1.00
rs3761650	44808356	G	A	0.38	1.09	1.83E-16	1.11	4.39E-18	1.02	0.33	Gen	1.00
rs6451781	44860506	T	C	0.61	0.92	1.86E-16	0.90	6.29E-18	0.98	0.31	Gen	1.00
rs10462080	44799052	C	A	0.38	1.09	1.90E-16	1.11	5.95E-18	1.02	0.34	Imp	0.98
rs71610376	44825023	G	C	0.38	1.09	1.92E-16	1.11	3.97E-18	1.02	0.33	Imp	0.99
rs12517690	44939293	G	A	0.39	1.09	1.93E-16	1.11	5.48E-18	1.02	0.32	Gen	1.00
chr5:44816067:D	44816067	TAAAAG	T	0.38	1.09	1.96E-16	1.11	4.43E-18	1.02	0.33	Imp	0.99
rs727305	44796042	T	C	0.39	1.09	1.97E-16	1.11	4.67E-18	1.02	0.35	Gen	0.99
rs7703497	44857028	A	G	0.61	0.92	1.97E-16	0.90	5.87E-18	0.98	0.34	Gen	1.00
rs4457089	44821736	C	T	0.38	1.09	2.02E-16	1.11	4.19E-18	1.02	0.32	Gen	1.00
rs1371024	44843590	C	T	0.61	0.92	2.04E-16	0.90	4.26E-18	0.98	0.32	Gen	1.00
rs727304	44796290	G	A	0.38	1.09	2.05E-16	1.11	5.92E-18	1.02	0.34	Gen	1.00
rs10038562	44832226	C	T	0.61	0.92	2.05E-16	0.90	4.15E-18	0.98	0.32	Gen	0.99
rs13174122	44810740	T	C	0.39	1.09	2.06E-16	1.11	4.57E-18	1.02	0.32	Gen	0.99
rs7703059	44826436	G	A	0.38	1.09	2.06E-16	1.11	4.34E-18	1.02	0.34	Imp	0.99

rs2165008	44821679	G	A	0.38	1.09	2.08E-16	1.11	4.35E-18	1.02	0.32	Gen	1.00
rs7717459	44804525	C	T	0.38	1.09	2.12E-16	1.11	5.66E-18	1.02	0.34	Gen	1.00
rs4298259	44920711	G	A	0.61	0.92	2.12E-16	0.90	4.29E-18	0.98	0.33	Gen	1.00
rs11746506	44812566	C	T	0.38	1.09	2.13E-16	1.11	4.36E-18	1.02	0.32	Gen	1.00
rs12513749	44828203	G	A	0.39	1.09	2.16E-16	1.11	4.46E-18	1.02	0.34	Imp	0.99
rs13189120	44822283	A	T	0.39	1.09	2.17E-16	1.11	4.55E-18	1.02	0.32	Gen	1.00
rs6868232	44827680	G	C	0.38	1.09	2.18E-16	1.11	4.19E-18	1.02	0.34	Imp	0.99
rs16901989	44837129	C	T	0.61	0.92	2.18E-16	0.90	4.61E-18	0.98	0.32	Gen	1.00
rs13155698	44828681	A	C	0.39	1.09	2.18E-16	1.11	3.97E-18	1.02	0.33	Gen	1.00
rs6896350	44832571	C	A	0.61	0.92	2.19E-16	0.90	4.25E-18	0.98	0.33	Gen	1.00
rs1438822	44859172	C	G	0.61	0.92	2.21E-16	0.90	7.25E-18	0.98	0.32	Gen	1.00
rs729599	44842260	G	A	0.62	0.92	2.23E-16	0.90	3.96E-18	0.98	0.33	Imp	0.99
rs6451772	44785763	A	C	0.39	1.09	2.24E-16	1.11	6.13E-18	1.02	0.37	Imp	0.98
rs6872254	44803784	C	T	0.39	1.09	2.28E-16	1.11	6.90E-18	1.02	0.34	Gen	0.99
rs1837286	44858741	C	T	0.62	0.92	2.28E-16	0.90	6.49E-18	0.98	0.32	Imp	0.99
rs4373287	44862884	G	T	0.61	0.92	2.28E-16	0.90	7.68E-18	0.98	0.33	Imp	0.99
rs11741772	44814597	A	C	0.38	1.09	2.29E-16	1.11	5.23E-18	1.02	0.33	Imp	0.99
rs3761648	44808079	A	G	0.38	1.09	2.31E-16	1.11	5.47E-18	1.02	0.34	Imp	0.99
rs16901964	44783255	C	T	0.38	1.09	2.33E-16	1.11	5.86E-18	1.02	0.35	Gen	1.00
rs12651949	44798112	C	T	0.38	1.09	2.33E-16	1.11	6.20E-18	1.02	0.34	Gen	1.00
rs10462083	44806737	T	G	0.38	1.09	2.33E-16	1.11	5.65E-18	1.02	0.35	Imp	0.99
rs4596389	44836556	A	C	0.62	0.92	2.35E-16	0.90	3.59E-18	0.98	0.33	Gen	1.00
rs969679	44804052	T	C	0.38	1.09	2.35E-16	1.11	5.95E-18	1.02	0.34	Imp	0.99
rs16901990	44842740	A	C	0.62	0.92	2.35E-16	0.90	3.54E-18	0.98	0.33	Gen	1.00
rs7707315	44918805	T	C	0.61	0.92	2.38E-16	0.90	4.48E-18	0.98	0.35	Gen	1.00
rs16901965	44783358	G	A	0.38	1.09	2.39E-16	1.11	5.51E-18	1.02	0.35	Gen	1.00
rs3761649	44808164	T	C	0.38	1.09	2.40E-16	1.11	5.60E-18	1.02	0.34	Imp	0.99
rs16901963	44783102	T	A	0.39	1.09	2.40E-16	1.11	5.55E-18	1.02	0.35	Gen	1.00
rs6451775	44836788	G	A	0.62	0.92	2.41E-16	0.90	4.60E-18	0.98	0.33	Imp	0.99
rs12518851	44828231	A	G	0.38	1.09	2.44E-16	1.11	4.45E-18	1.02	0.34	Imp	0.99
rs9637796	44817919	G	A	0.38	1.09	2.45E-16	1.11	5.69E-18	1.02	0.33	Imp	0.99
rs7736952	44790379	C	T	0.39	1.09	2.49E-16	1.11	6.96E-18	1.02	0.35	Imp	0.99
rs6451783	44918293	G	A	0.61	0.92	2.51E-16	0.90	4.43E-18	0.98	0.35	Gen	1.00
rs6893319	44863729	T	G	0.62	0.92	2.53E-16	0.90	7.70E-18	0.98	0.34	Imp	0.99
rs2083243	44840147	C	T	0.62	0.92	2.55E-16	0.90	4.35E-18	0.98	0.33	Imp	0.99
rs12188871	44814004	G	A	0.39	1.09	2.55E-16	1.11	5.73E-18	1.02	0.32	Gen	1.00
rs13185174	44791791	G	A	0.38	1.09	2.56E-16	1.11	5.98E-18	1.02	0.36	Imp	0.98
rs10041518	44927406	T	C	0.61	0.92	2.57E-16	0.90	6.23E-18	0.98	0.34	Gen	1.00
rs2330620	44839822	G	A	0.62	0.92	2.58E-16	0.90	4.62E-18	0.98	0.33	Imp	0.99
rs4604199	44855716	T	C	0.62	0.92	2.59E-16	0.90	6.78E-18	0.98	0.34	Imp	0.99
rs4605791	44855712	T	G	0.62	0.92	2.60E-16	0.90	6.78E-18	0.98	0.34	Imp	0.99

rs1837285	44858490	T	G	0.62	0.92	2.60E-16	0.90	6.52E-18	0.98	0.33	Imp	0.99
rs3747479	44809162	G	C	0.39	1.09	2.60E-16	1.11	5.48E-18	1.02	0.34	Gen	1.00
rs6871052	44863317	C	T	0.62	0.92	2.63E-16	0.90	7.44E-18	0.98	0.34	Gen	1.00
rs6451784	44922331	A	G	0.58	0.92	2.63E-16	0.90	3.17E-18	0.98	0.34	Imp	0.90
rs7728431	44922679	T	C	0.61	0.92	2.65E-16	0.90	5.40E-18	0.98	0.34	Gen	1.00
rs4566804	44855739	T	C	0.62	0.92	2.66E-16	0.90	6.87E-18	0.98	0.34	Imp	0.99
rs7736092	44920995	C	T	0.61	0.92	2.66E-16	0.90	5.15E-18	0.98	0.33	Gen	1.00
rs10064437	44834721	G	C	0.61	0.92	2.69E-16	0.90	3.66E-18	0.98	0.37	Imp	0.98
rs10070928	44888557	C	G	0.60	0.92	2.72E-16	0.91	9.63E-18	0.98	0.31	Gen	0.96
rs7710952	44785791	C	T	0.38	1.09	2.74E-16	1.11	7.59E-18	1.02	0.35	Imp	0.99
rs7710978	44785864	A	G	0.38	1.09	2.75E-16	1.11	7.62E-18	1.02	0.35	Imp	0.99
rs10053247	44863959	C	T	0.62	0.92	2.78E-16	0.90	8.60E-18	0.98	0.34	Imp	0.99
rs12656984	44824702	A	G	0.38	1.09	2.79E-16	1.11	6.67E-18	1.02	0.32	Imp	0.98
rs7703618	44914579	G	A	0.61	0.92	2.83E-16	0.90	7.58E-18	0.98	0.34	Gen	1.00
rs10044321	44890760	G	A	0.61	0.92	2.85E-16	0.91	8.30E-18	0.98	0.35	Imp	0.99
rs10073945	44932134	A	G	0.61	0.92	2.86E-16	0.90	6.86E-18	0.98	0.35	Imp	0.99
chr5:44916609:D	44916609	GT	G	0.63	0.92	2.92E-16	0.90	4.50E-18	0.98	0.33	Imp	0.96
rs13179835	44908703	G	A	0.61	0.92	2.94E-16	0.90	5.90E-18	0.99	0.45	Imp	0.97
rs6871820	44919557	T	C	0.61	0.92	2.99E-16	0.90	6.36E-18	0.98	0.33	Gen	0.99
rs10038554	44927107	G	A	0.61	0.92	2.99E-16	0.90	6.52E-18	0.98	0.34	Gen	1.00
rs34325259	44936860	G	T	0.38	1.08	3.02E-16	1.10	9.54E-18	1.02	0.42	Gen	0.96
rs13183209	44803749	G	A	0.38	1.09	3.04E-16	1.11	7.72E-18	1.02	0.35	Imp	0.99
rs6451774	44793984	G	C	0.38	1.09	3.04E-16	1.10	8.79E-18	1.02	0.35	Imp	0.99
rs10063172	44930709	T	C	0.61	0.92	3.11E-16	0.90	7.55E-18	0.98	0.35	Imp	0.99
rs6451778	44857988	C	T	0.62	0.92	3.12E-16	0.90	8.11E-18	0.98	0.34	Imp	0.99
rs10039866	44925061	T	A	0.61	0.92	3.13E-16	0.90	5.21E-18	0.98	0.35	Gen	1.00
rs11948636	44881229	A	C	0.61	0.92	3.15E-16	0.91	1.00E-17	0.98	0.34	Gen	1.00
rs34115673	44892801	T	G	0.61	0.92	3.20E-16	0.91	8.96E-18	0.98	0.36	Imp	0.98
rs10035358	44892902	C	A	0.61	0.92	3.21E-16	0.91	8.71E-18	0.98	0.36	Imp	0.98
rs10072025	44843494	C	T	0.62	0.92	3.22E-16	0.90	5.21E-18	0.98	0.34	Imp	0.99
rs6875933	44786696	C	T	0.38	1.09	3.26E-16	1.11	6.99E-18	1.02	0.35	Imp	0.97
rs11750119	44786141	G	T	0.38	1.09	3.29E-16	1.11	6.37E-18	1.02	0.35	Imp	0.99
rs13154729	44892929	C	T	0.61	0.92	3.30E-16	0.91	9.07E-18	0.98	0.36	Imp	0.99
rs10057341	44931840	C	T	0.61	0.92	3.34E-16	0.90	6.56E-18	0.98	0.36	Gen	1.00
chr5:44904186:D	44904186	CACTTA	C	0.59	0.92	3.37E-16	0.90	1.97E-17	0.98	0.43	Imp	0.94
rs13153556	44928101	G	A	0.61	0.92	3.40E-16	0.90	6.16E-18	0.98	0.36	Gen	1.00
rs10059086	44872007	C	T	0.61	0.92	3.42E-16	0.91	1.13E-17	0.98	0.34	Gen	1.00
rs2330619	44796062	T	C	0.38	1.08	3.43E-16	1.10	9.62E-18	1.02	0.36	Imp	0.99
rs13168400	44883536	C	T	0.61	0.92	3.45E-16	0.91	1.22E-17	0.98	0.36	Imp	0.99
rs67274820	44782295	T	C	0.38	1.08	3.50E-16	1.11	7.18E-18	1.02	0.37	Imp	0.99
chr5:44795637:I	44795637	G	GA	0.38	1.08	3.51E-16	1.11	8.66E-18	1.02	0.36	Imp	0.99

rs10069220	44881978	G	A	0.61	0.92	3.53E-16	0.91	1.05E-17	0.98	0.33	Gen	1.00
rs4495192	44929177	T	C	0.62	0.92	3.56E-16	0.90	5.47E-18	0.98	0.35	Imp	0.99
rs13157608	44881504	C	T	0.61	0.92	3.62E-16	0.91	1.27E-17	0.98	0.36	Imp	0.99
rs10941687	44921183	G	A	0.62	0.92	3.67E-16	0.90	6.65E-18	0.98	0.35	Imp	0.99
rs10040488	44880288	G	T	0.62	0.92	3.69E-16	0.91	1.30E-17	0.98	0.36	Imp	0.99
rs9637783	44819646	T	G	0.39	1.08	3.70E-16	1.11	6.26E-18	1.02	0.37	Gen	1.00
rs4866920	44879146	G	T	0.62	0.92	3.76E-16	0.91	1.35E-17	0.98	0.36	Imp	0.99
rs10473376	44931228	C	T	0.62	0.92	3.79E-16	0.90	7.90E-18	0.98	0.36	Imp	0.99
rs4591754	44895091	G	A	0.62	0.92	3.80E-16	0.91	9.81E-18	0.98	0.36	Imp	0.99
rs7356604	44895199	T	C	0.62	0.92	3.80E-16	0.91	9.82E-18	0.98	0.36	Imp	0.99
rs12652026	44798609	C	T	0.38	1.08	3.83E-16	1.11	8.81E-18	1.02	0.36	Imp	0.99
rs4642377	44885240	A	T	0.61	0.92	3.84E-16	0.91	1.40E-17	0.98	0.35	Imp	0.99
rs6875287	44941630	C	T	0.39	1.09	3.85E-16	1.10	1.44E-17	1.02	0.32	Imp	0.98
rs10042455	44925018	A	G	0.62	0.92	3.91E-16	0.90	6.70E-18	0.98	0.35	Imp	0.99
rs13187933	44883243	T	C	0.62	0.92	3.93E-16	0.91	1.08E-17	0.98	0.36	Imp	0.99
rs6883465	44891620	T	C	0.62	0.92	4.01E-16	0.91	9.16E-18	0.98	0.37	Imp	0.99
rs4391175	44890056	A	G	0.62	0.92	4.03E-16	0.90	9.00E-18	0.98	0.37	Imp	0.99
rs10070037	44870237	A	T	0.61	0.92	4.07E-16	0.91	1.06E-17	0.98	0.37	Gen	1.00
rs7718354	44891014	T	G	0.62	0.92	4.08E-16	0.91	9.29E-18	0.98	0.37	Imp	0.99
rs7734331	44895588	G	A	0.62	0.92	4.13E-16	0.91	9.36E-18	0.98	0.37	Imp	0.99
rs11948387	44884241	T	C	0.62	0.92	4.16E-16	0.91	1.13E-17	0.98	0.36	Imp	0.99
rs6859157	44783838	G	A	0.38	1.08	4.17E-16	1.11	8.33E-18	1.02	0.37	Imp	0.99
rs11958451	44866902	A	G	0.61	0.92	4.20E-16	0.91	1.53E-17	0.98	0.35	Imp	0.99
rs4323241	44929156	T	C	0.61	0.92	4.28E-16	0.90	1.72E-17	0.98	0.33	Imp	0.97
rs10060878	44899424	T	C	0.61	0.92	4.37E-16	0.91	8.78E-18	0.98	0.36	Gen	1.00
rs6894324	44867336	G	C	0.62	0.92	4.37E-16	0.91	1.38E-17	0.98	0.35	Imp	0.99
rs10040082	44865854	T	C	0.62	0.92	4.43E-16	0.91	1.50E-17	0.98	0.34	Gen	1.00
rs9292914	44871381	A	C	0.61	0.92	4.45E-16	0.91	1.23E-17	0.98	0.37	Gen	1.00
rs10065638	44866162	T	C	0.61	0.92	4.48E-16	0.91	1.39E-17	0.98	0.34	Gen	0.98
rs10473377	44931246	T	C	0.62	0.92	4.49E-16	0.90	8.71E-18	0.98	0.35	Imp	0.99
rs34501299	44913079	T	G	0.61	0.92	4.50E-16	0.91	1.16E-17	0.98	0.38	Gen	1.00
rs4395640	44869100	T	C	0.62	0.92	4.53E-16	0.91	1.47E-17	0.98	0.35	Gen	1.00
rs7708686	44922838	C	G	0.62	0.92	4.69E-16	0.90	7.26E-18	0.98	0.35	Imp	0.99
rs7708506	44922704	C	G	0.62	0.92	4.73E-16	0.90	7.31E-18	0.98	0.35	Imp	0.99
rs12109710	44929928	A	G	0.62	0.92	4.74E-16	0.90	9.42E-18	0.98	0.35	Imp	0.98
rs6868779	44886000	A	G	0.61	0.92	4.80E-16	0.91	1.42E-17	0.98	0.36	Gen	1.00
rs13356086	44887679	T	C	0.62	0.92	4.84E-16	0.91	1.08E-17	0.98	0.37	Imp	0.98
rs11951760	44872172	G	A	0.62	0.92	4.97E-16	0.91	1.18E-17	0.98	0.39	Imp	0.99
rs4866784	44901131	T	C	0.61	0.92	5.05E-16	0.91	1.01E-17	0.98	0.38	Gen	1.00
rs10057521	44865986	C	T	0.60	0.92	5.07E-16	0.91	1.51E-17	0.98	0.36	Gen	0.95
rs4129642	44898129	G	T	0.61	0.92	5.12E-16	0.91	9.43E-18	0.98	0.36	Gen	1.00

rs6881563	44912853	C	T	0.61	0.92	5.28E-16	0.91	1.38E-17	0.98	0.38	Gen	1.00
rs4866783	44876507	C	T	0.62	0.92	5.62E-16	0.91	1.38E-17	0.98	0.39	Imp	0.98
rs7708705	44922925	A	G	0.62	0.92	5.67E-16	0.90	7.88E-18	0.98	0.35	Imp	0.98
rs10039173	44903376	C	T	0.61	0.92	5.74E-16	0.91	1.03E-17	0.98	0.41	Gen	1.00
rs6870136	44910662	G	A	0.61	0.92	5.76E-16	0.91	1.41E-17	0.98	0.40	Gen	1.00
rs10060645	44908199	C	T	0.61	0.92	6.23E-16	0.91	1.57E-17	0.98	0.40	Imp	0.99
rs10045264	44886432	G	A	0.61	0.92	6.24E-16	0.91	1.93E-17	0.98	0.36	Gen	1.00
rs6880275	44908935	T	C	0.62	0.92	7.75E-16	0.91	1.48E-17	0.98	0.41	Imp	0.99
rs9791056	44903891	T	C	0.61	0.92	8.09E-16	0.91	1.25E-17	0.98	0.41	Gen	1.00
rs9791059	44900447	A	C	0.62	0.92	8.13E-16	0.90	8.88E-18	0.98	0.42	Imp	0.98
chr5:44904190:D	44904190	TAACTC	T	0.61	0.92	8.31E-16	0.91	1.55E-17	0.98	0.41	Imp	0.99
chr5:44842792:D	44842792	CT	C	0.56	0.92	8.78E-16	0.90	3.54E-19	1.00	0.97	Imp	0.94
chr5:44842794:D	44842794	CCAGTT	C	0.56	0.92	8.80E-16	0.90	3.56E-19	1.00	0.97	Imp	0.94
chr5:44842796:D	44842796	AGTTC	A	0.56	0.92	8.80E-16	0.90	3.55E-19	1.00	0.97	Imp	0.94
rs4457088	44900954	T	A	0.62	0.92	8.85E-16	0.91	1.29E-17	0.98	0.41	Imp	0.99
chr5:44782479:D	44782479	AT	A	0.38	1.08	8.87E-16	1.10	1.30E-17	1.02	0.37	Imp	0.98
rs4866921	44917137	T	C	0.62	0.92	9.83E-16	0.91	1.34E-17	0.98	0.41	Imp	0.99
rs9791164	44904030	A	G	0.62	0.92	9.92E-16	0.91	1.70E-17	0.98	0.42	Imp	0.99
rs6874167	44916674	G	A	0.62	0.92	1.01E-15	0.91	1.49E-17	0.98	0.41	Imp	0.99
rs4360054	44929004	A	G	0.61	0.92	1.10E-15	0.91	2.52E-17	0.98	0.41	Imp	0.98
rs6895062	44916492	T	C	0.62	0.92	1.12E-15	0.91	1.84E-17	0.98	0.40	Imp	0.98
chr5:44852861:D	44852861	AC	A	0.61	0.92	1.59E-15	0.91	6.38E-17	0.98	0.31	Imp	0.96
rs7449277	44841457	T	A	0.60	0.92	2.06E-15	0.91	7.34E-17	0.98	0.35	Imp	0.95
chr5:44885826:I	44885826	A	AAATT	0.61	0.92	2.29E-15	0.90	1.66E-17	0.99	0.53	Imp	0.96
rs7702464	44826259	A	C	0.27	1.11	2.32E-15	1.13	6.87E-17	1.03	0.20	Imp	0.74
chr5:44928978:D	44928978	ACCCTC	A	0.55	0.92	2.80E-15	0.90	3.15E-17	0.99	0.58	Imp	0.88
chr5:44921956:I	44921956	T	TA	0.61	0.92	3.19E-15	0.90	3.29E-17	0.99	0.54	Imp	0.95
rs74724331	44758161	G	A	0.40	1.08	3.28E-15	1.10	7.61E-17	1.01	0.47	Imp	0.95
rs13178923	44929285	T	A	0.57	0.92	1.03E-14	0.91	4.38E-15	0.98	0.34	Imp	0.87
rs12658334	44689131	G	A	0.39	1.09	2.72E-14	1.12	3.65E-16	1.02	0.50	Imp	0.73
chr5:44832897:I	44832897	A	ATGTT	0.59	0.93	5.26E-14	0.91	6.86E-16	0.99	0.79	Imp	0.94
rs183946926	44688992	G	A	0.37	1.09	1.05E-13	1.12	1.55E-16	1.03	0.25	Imp	0.69
chr5:44868315:D	44868315	AAT	A	0.48	0.93	1.26E-12	0.91	1.39E-13	0.97	0.13	Imp	0.79
rs200229088	44701817	TTG	T	0.31	1.09	2.28E-12	1.12	1.26E-13	1.04	0.15	Imp	0.65
rs4562047	44868093	C	T	0.48	0.93	1.52E-11	0.91	7.18E-14	0.99	0.68	Imp	0.77
rs2067980	44982317	A	G	0.16	1.09	2.51E-11	1.12	4.35E-13	1.04	0.17	Gen	1.00
rs73093976	44609392	C	T	0.17	1.09	6.67E-11	1.12	8.34E-13	1.02	0.36	Imp	0.95
rs147039293	44673324	C	T	0.14	0.92	3.98E-10	0.90	2.66E-10	0.93	0.01	Imp	0.96
rs75209549	44629545	A	G	0.14	1.09	6.64E-10	1.11	1.25E-10	1.02	0.53	Imp	0.89
rs112234443	44922557	C	T	0.05	1.16	1.10E-09	1.19	5.72E-10	1.07	0.14	Imp	0.90
rs76001691	44633373	T	C	0.05	1.16	1.14E-09	1.18	4.23E-09	1.09	0.07	Imp	0.82

rs4549535	44630291	A	G	0.05	1.15	1.58E-09	1.17	9.53E-10	1.08	0.09	Imp	0.93
rs6880469	44975989	A	G	0.35	1.06	1.83E-09	1.08	2.19E-10	1.01	0.57	Imp	0.98
rs6451787	45009269	T	G	0.17	1.09	2.53E-09	1.11	2.10E-10	1.03	0.23	Imp	0.84
rs28705196	44631795	T	C	0.05	1.14	4.04E-09	1.17	1.01E-09	1.06	0.18	Imp	0.97
chr5:44633359:l	44633359	T	TTC	0.05	1.14	4.39E-09	1.17	1.36E-09	1.06	0.17	Imp	0.95
chr5:44633372:l	44633372	T	TC	0.05	1.14	4.39E-09	1.17	1.36E-09	1.06	0.17	Imp	0.95
rs5004228	44632008	A	G	0.05	1.14	4.63E-09	1.16	2.03E-09	1.07	0.13	Gen	1.00
rs116443643	44606969	C	A	0.15	0.92	5.30E-09	0.91	1.78E-09	0.95	0.04	Imp	0.95
rs13357090	44632713	C	T	0.05	1.14	7.05E-09	1.16	3.72E-09	1.07	0.12	Imp	0.96
chr5:44681918:l	44681918	C	CA	0.04	0.84	7.16E-09	0.82	3.71E-08	0.88	0.03	Imp	0.58
c5_pos44839303	44803546	A	G	0.14	0.92	7.49E-09	0.91	2.36E-08	0.93	0.01	Gen	1.00
rs112754768	44718764	A	G	0.05	1.15	8.65E-09	1.18	6.28E-09	1.06	0.19	Imp	0.88
rs34692501	45029217	A	G	0.13	1.09	9.49E-09	1.11	7.72E-10	1.03	0.30	Gen	1.00
rs10941712	45917605	C	T	0.23	0.93	9.61E-09	0.91	1.42E-10	0.95	0.02	Imp	0.78
rs11750654	44494303	T	C	0.15	0.92	1.13E-08	0.91	2.84E-09	0.95	0.04	Imp	0.95
rs11750655	44494312	T	C	0.15	0.92	1.13E-08	0.91	2.84E-09	0.95	0.04	Imp	0.95
rs3935086	44960923	T	A	0.33	1.06	1.14E-08	1.07	2.76E-09	1.01	0.64	Gen	1.00
rs79670114	44861045	A	G	0.04	1.16	1.24E-08	1.19	2.52E-09	1.06	0.23	Imp	0.88
rs11742346	45003293	C	T	0.13	1.09	1.29E-08	1.11	1.45E-09	1.03	0.24	Imp	0.94
rs12520604	44476872	A	C	0.04	1.16	1.34E-08	1.20	4.72E-09	1.03	0.56	Imp	0.91
c5_pos44832736	44796979	C	T	0.14	0.92	1.36E-08	0.91	4.34E-08	0.93	0.01	Gen	0.99
c5_pos44671855	44636098	G	C	0.20	0.93	1.41E-08	0.92	6.26E-09	0.96	0.10	Gen	1.00
rs187108781	44619502	A	G	0.15	0.92	1.59E-08	0.91	6.61E-09	0.95	0.04	Imp	0.95
chr5:44795636:l	44795636	A	AG	0.28	1.07	1.68E-08	1.09	9.96E-10	1.00	0.87	Imp	0.85
c5_pos45369617	45333860	T	C	0.26	0.94	1.71E-08	0.92	1.56E-10	0.96	0.05	Gen	1.00
rs12516986	44526154	A	G	0.04	1.15	1.80E-08	1.18	6.09E-09	1.03	0.50	Gen	1.00
rs7702731	44914285	A	G	0.68	0.94	2.80E-08	0.94	2.63E-08	0.99	0.71	Gen	0.98
rs75036127	44802928	A	T	0.04	1.15	3.47E-08	1.19	5.58E-09	1.05	0.33	Imp	0.88
rs72748037	44449376	G	A	0.15	0.93	3.51E-08	0.91	1.02E-08	0.95	0.04	Imp	0.97
rs13183434	45074633	G	A	0.13	1.08	3.67E-08	1.11	1.91E-09	1.04	0.21	Gen	1.00
chr5:45943429:D	45943429	CTCT	C	0.16	0.92	3.72E-08	0.89	1.09E-09	0.92	0.01	Imp	0.70
rs11745472	44929256	A	T	0.12	1.11	3.83E-08	1.13	5.18E-08	1.03	0.39	Imp	0.61
rs72748026	44431924	C	G	0.14	0.92	3.86E-08	0.91	3.78E-09	0.95	0.05	Imp	0.95
rs17343002	44853593	G	C	0.31	0.94	4.36E-08	0.94	9.13E-08	0.96	0.06	Gen	1.00
c5_pos44489027	44453270	T	G	0.15	0.93	4.89E-08	0.91	1.65E-08	0.95	0.04	Gen	1.00
rs11738503	45011742	A	G	0.16	1.08	5.09E-08	1.10	4.71E-09	1.04	0.19	Imp	0.87
rs11741260	44412065	G	A	0.14	0.93	5.14E-08	0.91	5.32E-09	0.95	0.05	Gen	1.00
rs11948186	45051434	A	G	0.34	1.06	5.17E-08	1.07	7.10E-09	1.01	0.69	Imp	0.97
rs7701656	44947702	T	C	0.13	1.08	5.71E-08	1.11	2.64E-09	1.00	0.90	Gen	1.00
chr5:44692568:l	44692568	G	GT	0.21	0.93	6.86E-08	0.92	8.13E-08	0.96	0.09	Imp	0.90
rs78797445	44795797	G	A	0.19	0.94	8.81E-08	0.93	1.30E-07	0.96	0.08	Imp	0.97

chr5:45162687:D	45162687	AGATCT	A	0.14	0.93	1.03E-07	0.91	2.86E-08	0.93	0.01	Imp	0.95
c5_pos44754116	44718359	G	A	0.19	0.94	1.03E-07	0.93	1.25E-07	0.95	0.05	Gen	1.00
rs111385188	44675187	G	A	0.04	1.16	1.22E-07	1.19	2.96E-08	1.07	0.22	Imp	0.85
rs62372990	45988038	G	A	0.06	1.14	1.26E-07	1.15	3.53E-07	1.07	0.12	Imp	0.76
c5_pos44754266	44718509	G	A	0.07	1.11	1.27E-07	1.15	5.01E-10	1.04	0.30	Gen	1.00
rs7293402	45990381	G	A	0.06	1.13	1.40E-07	1.15	4.28E-07	1.07	0.13	Imp	0.76
rs147872430	45239336	C	T	0.14	0.93	1.52E-07	0.91	3.35E-08	0.93	0.01	Imp	0.96
chr5:44945030:I	44945030	T	TA	0.19	0.93	1.76E-07	0.93	3.34E-07	0.96	0.08	Imp	0.92
rs10051592	45090306	T	G	0.33	1.06	1.83E-07	1.07	1.67E-08	1.01	0.79	Gen	1.00
c5_pos45171325	45135568	G	C	0.14	0.93	1.97E-07	0.91	4.97E-08	0.93	0.01	Gen	1.00
chr5:44518580:I	44518580	C	CT	0.21	0.94	2.25E-07	0.93	2.21E-06	0.93	0.00	Imp	0.87
rs11743309	45122388	A	G	0.14	0.93	2.27E-07	0.92	8.71E-08	0.93	0.00	Gen	1.00
c5_pos45322045	45286288	T	A	0.14	0.93	2.61E-07	0.92	6.68E-08	0.94	0.01	Gen	0.99
c5_pos45028737	44992980	G	T	0.14	0.93	3.09E-07	0.92	1.57E-06	0.94	0.02	Gen	1.00
c5_pos45149948	45114191	C	T	0.14	0.93	3.74E-07	0.92	1.15E-07	0.93	0.01	Gen	1.00
rs9763350	45882593	T	C	0.26	1.07	4.25E-07	1.08	6.84E-07	1.08	0.00	Imp	0.71
rs11740651	45302476	C	A	0.14	0.93	7.20E-07	0.91	7.54E-08	0.94	0.03	Imp	0.91
chr5:44976998:I	44976998	G	GTGAT	0.11	1.08	9.83E-07	1.11	5.80E-08	1.02	0.44	Imp	0.93
rs13172124	45160377	C	T	0.46	0.95	1.13E-06	0.94	4.25E-07	0.98	0.22	Imp	0.96
rs13186320	44302177	A	T	0.47	1.05	1.19E-06	1.07	1.46E-07	1.04	0.05	Imp	0.86
rs72750030	45903837	G	A	0.11	0.92	1.26E-06	0.89	1.33E-07	0.92	0.02	Imp	0.70
rs72751936	45984433	G	A	0.11	0.91	1.33E-06	0.88	2.66E-07	0.90	0.01	Imp	0.59
c5_pos45135048	45099291	G	A	0.09	0.92	1.53E-06	0.91	7.00E-06	0.92	0.01	Gen	1.00
rs147517548	44711323	C	T	0.01	1.3	1.55E-06	1.37	5.35E-07	1.18	0.11	Imp	0.68
rs13156720	44823321	A	G	0.02	1.19	1.79E-06	1.25	1.18E-07	1.08	0.28	Imp	0.77
c5_pos44444935	44409178	C	A	0.06	0.91	1.90E-06	0.89	8.08E-07	0.95	0.18	Gen	1.00
rs4866929	45266589	A	G	0.47	0.95	1.98E-06	0.94	5.79E-07	0.98	0.21	Gen	1.00
chr5:45210850:D	45210850	CTATAA	C	0.45	0.95	2.02E-06	0.94	7.09E-07	0.97	0.17	Imp	0.92
rs11959880	45958226	G	A	0.09	0.9	2.06E-06	0.89	2.01E-06	0.90	0.01	Imp	0.58
rs17268417	44700201	T	C	0.05	1.12	2.08E-06	1.15	4.60E-07	1.08	0.07	Gen	0.95
chr5:45210851:D	45210851	TATAAA	T	0.45	0.95	2.31E-06	0.94	7.39E-07	0.97	0.16	Imp	0.92
chr5:44349569:D	44349569	TATCAGA	T	0.43	1.06	2.33E-06	1.06	3.88E-06	1.03	0.22	Imp	0.70
rs11738948	44999799	G	C	0.20	0.94	2.45E-06	0.94	1.32E-05	0.95	0.05	Imp	0.96
rs72750027	45899603	G	A	0.01	1.43	2.55E-06	1.46	1.45E-05	1.42	0.01	Imp	0.66
rs72765759	45846620	C	T	0.11	0.92	2.91E-06	0.90	2.28E-07	0.94	0.06	Imp	0.73
rs981782	45285718	A	C	0.47	0.96	3.13E-06	0.95	8.27E-07	0.98	0.30	Imp	0.97
rs137877813	45084544	G	C	0.14	0.94	3.13E-06	0.93	4.96E-06	0.94	0.02	Imp	0.96
rs138117035	45938300	T	C	0.10	0.91	3.23E-06	0.89	7.08E-07	0.93	0.05	Imp	0.61
rs145696903	45933347	G	A	0.10	0.91	3.38E-06	0.89	9.01E-07	0.92	0.04	Imp	0.61
chr5:44302175:D	44302175	CA	C	0.51	1.05	3.46E-06	1.06	3.54E-07	1.03	0.17	Imp	0.94
rs75442098	44498735	C	A	0.03	0.86	3.56E-06	0.84	1.40E-05	0.91	0.15	Imp	0.65

rs16902086	45285752	A	G	0.34	1.05	3.60E-06	1.06	6.45E-07	1.02	0.34	Gen	1.00
rs72762052	45427937	C	T	0.13	0.93	3.83E-06	0.91	6.81E-07	0.93	0.03	Imp	0.81
rs72750038	45912279	T	C	0.10	0.91	3.87E-06	0.89	1.15E-06	0.92	0.02	Imp	0.65
rs74833952	44961224	G	A	0.02	0.76	4.76E-06	0.75	2.64E-05	0.80	0.06	Imp	0.42
rs11747840	44984579	C	T	0.21	0.95	4.97E-06	0.94	1.74E-05	0.96	0.08	Imp	0.94
rs72751917	45949866	G	A	0.01	1.4	5.21E-06	1.40	6.02E-05	1.38	0.02	Imp	0.51
rs11750845	44373060	C	T	0.51	1.05	5.39E-06	1.06	4.54E-07	1.02	0.27	Gen	1.00
rs72759922	45005236	C	T	0.21	0.95	6.67E-06	0.94	1.09E-05	0.97	0.16	Imp	0.94
rs72763959	45707287	C	A	0.01	1.33	6.84E-06	1.36	3.85E-05	1.28	0.03	Imp	0.69
rs35190075	44776604	C	G	0.02	1.18	7.49E-06	1.25	5.20E-07	1.11	0.15	Imp	0.76
rs185330077	45366544	G	A	0.01	1.48	7.60E-06	1.69	9.92E-08	0.94	0.70	Imp	0.39
rs182455548	45746470	G	A	0.01	1.33	7.67E-06	1.35	3.70E-05	1.28	0.04	Imp	0.69
rs72763975	45767075	G	A	0.01	1.33	7.78E-06	1.35	3.66E-05	1.28	0.04	Imp	0.69
c5_pos44897529	44861772	T	C	0.04	1.11	8.41E-06	1.14	9.56E-07	1.05	0.26	Gen	1.00
rs72763992	45806419	A	T	0.01	1.33	8.60E-06	1.36	3.32E-05	1.28	0.04	Imp	0.67
rs148456234	45922475	C	T	0.11	0.92	8.93E-06	0.90	1.29E-06	0.93	0.05	Imp	0.65
rs72750006	45862423	A	G	0.01	1.3	8.95E-06	1.35	1.57E-05	1.24	0.05	Imp	0.63
rs72762048	45408899	T	C	0.12	0.93	9.74E-06	0.91	2.49E-06	0.93	0.03	Imp	0.80
rs5003385	44907327	A	C	0.04	1.11	1.01E-05	1.14	9.29E-07	1.05	0.27	Gen	1.00
rs191491730	45160378	G	A	0.00	1.6	1.03E-05	1.83	4.17E-07	1.21	0.37	Imp	0.59
rs148415520	45317307	G	A	0.45	0.96	1.13E-05	0.95	6.59E-06	0.97	0.12	Imp	0.91
rs76810418	45882608	G	A	0.25	1.06	1.21E-05	1.07	1.71E-05	1.06	0.02	Imp	0.74
chr5:44606887:D	44606887	AT	A	0.15	0.94	1.28E-05	0.93	2.00E-05	0.97	0.26	Imp	0.95
rs147763247	45313905	G	C	0.45	0.96	1.32E-05	0.95	6.73E-06	0.97	0.12	Imp	0.91
rs10473395	45882607	T	C	0.25	1.06	1.52E-05	1.07	2.30E-05	1.06	0.02	Imp	0.74
chr5:45799799:D	45799799	AT	A	0.04	1.12	1.59E-05	1.13	5.54E-05	1.06	0.26	Imp	0.81
rs192741232	44838737	A	G	0.01	1.3	1.59E-05	1.38	4.74E-06	1.14	0.27	Imp	0.75
rs150276216	44889423	T	C	0.04	1.11	1.71E-05	1.14	2.26E-06	1.06	0.26	Imp	0.97
rs191712805	45795107	A	G	0.01	1.33	1.72E-05	1.36	5.89E-05	1.28	0.05	Imp	0.61
rs4613718	44649944	C	T	0.60	1.04	1.92E-05	1.06	5.73E-07	0.98	0.22	Gen	1.00
rs114416420	45951524	G	T	0.10	0.91	2.00E-05	0.90	3.06E-05	0.90	0.01	Imp	0.56
c5_pos44663080	44627323	C	T	0.21	0.95	2.02E-05	0.95	4.76E-05	0.95	0.04	Gen	1.00
rs180724159	45562846	T	C	0.00	1.53	2.04E-05	1.71	1.34E-06	1.20	0.36	Imp	0.72
rs181673846	45837084	C	A	0.01	1.34	2.08E-05	1.38	4.81E-05	1.29	0.05	Imp	0.69
chr5:45853664:D	45853664	CAT	C	0.02	1.22	2.15E-05	1.22	1.25E-04	1.10	0.30	Imp	0.58
rs139424826	45522979	C	A	0.12	0.93	2.44E-05	0.90	1.23E-06	0.94	0.10	Imp	0.68
rs6859397	45136050	T	C	0.33	1.04	2.79E-05	1.06	4.14E-06	1.00	0.83	Imp	0.95
rs4492120	45187804	G	A	0.33	1.04	2.82E-05	1.06	2.90E-06	1.00	0.81	Gen	1.00
chr5:44349567:D	44349567	TATAT	T	0.38	1.05	2.86E-05	1.06	2.14E-05	1.02	0.43	Imp	0.69
rs10035564	45252500	A	G	0.34	1.04	2.98E-05	1.06	2.91E-06	1.01	0.72	Gen	0.96
rs75513092	44544909	G	T	0.21	0.95	3.22E-05	0.95	6.48E-05	0.96	0.06	Imp	0.98

rs6867827	45317216	G	T	0.17	1.06	3.36E-05	1.06	5.26E-05	1.04	0.11	Imp	0.97
rs139928702	45938421	A	C	0.05	1.12	3.47E-05	1.13	1.02E-04	1.08	0.18	Imp	0.62
rs11748830	44265513	C	T	0.16	0.94	3.70E-05	0.94	3.87E-05	0.94	0.02	Imp	0.92
rs17268006	44538881	C	T	0.21	0.95	4.19E-05	0.95	9.20E-05	0.96	0.05	Gen	0.99
chr5:45686224:D	45686224	TA	T	0.07	1.12	4.22E-05	1.09	5.64E-03	1.15	0.01	Imp	0.50
chr5:44816722:D	44816722	ACTT	A	0.04	1.11	4.29E-05	1.14	7.24E-06	1.07	0.18	Imp	0.95
rs72759920	44998794	T	G	0.04	1.11	5.13E-05	1.13	1.88E-05	1.06	0.21	Imp	0.92
rs72759916	44990010	T	C	0.04	1.11	6.09E-05	1.13	1.69E-05	1.06	0.24	Imp	0.92
rs62370579	45792329	A	T	0.03	1.15	6.40E-05	1.15	7.86E-04	1.10	0.17	Imp	0.77
chr5:45843775:D	45843775	TTA	T	0.04	1.12	6.78E-05	1.13	2.15E-04	1.06	0.29	Imp	0.84
rs181701912	44888831	C	T	0.11	0.92	6.92E-05	0.90	2.49E-05	0.96	0.36	Imp	0.51
rs10078961	44888856	T	C	0.29	0.95	7.00E-05	0.93	8.16E-06	0.99	0.73	Imp	0.63
chr5:45379590:D	45379590	CA	C	0.19	1.05	7.02E-05	1.06	7.32E-05	1.04	0.15	Imp	0.97
rs75946047	44454153	C	T	0.03	1.14	7.34E-05	1.20	6.92E-06	1.08	0.26	Imp	0.81
rs62372989	45987296	A	G	0.03	1.14	7.45E-05	1.15	3.22E-04	1.07	0.31	Imp	0.70
rs12187122	44577879	A	G	0.03	1.13	7.77E-05	1.16	3.14E-05	1.01	0.93	Imp	0.91
rs6451798	45387854	C	T	0.19	1.05	7.93E-05	1.06	8.43E-05	1.03	0.16	Gen	1.00
rs62370581	45793718	G	A	0.03	1.12	8.32E-05	1.12	3.67E-04	1.06	0.28	Imp	0.91
rs1816683	44701827	G	A	0.57	1.04	8.34E-05	1.06	2.73E-06	0.98	0.30	Imp	0.85
rs75964308	45946479	G	C	0.04	1.12	8.48E-05	1.13	2.36E-04	1.06	0.33	Imp	0.77
rs1501362	45378207	C	T	0.19	1.05	8.52E-05	1.06	7.26E-05	1.03	0.18	Imp	0.98
rs141075872	45790426	A	T	0.03	1.13	8.55E-05	1.13	7.34E-04	1.07	0.25	Imp	0.82
rs7709262	45342044	A	G	0.19	1.05	8.69E-05	1.06	8.62E-05	1.03	0.18	Imp	0.98
rs144163740	45376455	G	T	0.36	0.95	8.87E-05	0.94	3.12E-05	0.98	0.52	Imp	0.61
rs62370604	45843007	T	C	0.03	1.12	9.55E-05	1.13	3.17E-04	1.06	0.32	Imp	0.86
rs74859464	45951751	C	A	0.03	1.12	9.58E-05	1.13	2.33E-04	1.06	0.33	Imp	0.83
rs6874127	45305615	A	G	0.19	1.05	9.63E-05	1.06	9.12E-05	1.03	0.17	Gen	1.00
rs75212852	45805592	C	T	0.03	1.12	9.76E-05	1.12	4.25E-04	1.06	0.29	Imp	0.91
chr5:45804917:I	45804917	T	TA	0.03	1.12	9.79E-05	1.12	4.27E-04	1.06	0.29	Imp	0.91
c5_pos45103929	45068172	G	A	0.04	1.1	9.85E-05	1.12	1.27E-04	1.05	0.31	Gen	1.00

**Supplementary Table 2.** The three independent signals showing the lead SNP in each signal together with the correlated SNPs that could not be excluded from causality.

SNP	Position	Ref/Alt	MAF	Imputation r2	OR (95% CI)	Single SNP p_value	P-value conditional on signal 1 top hit	P-value conditional on signal 1 + 2	cor_rs10941679	cor_rs6864776	cor_rs200229088
<b>Signal 1</b>											
rs10941679	44706498	A/G	0.27	0.99	1.12(1.1-1.15)	2.60E-26	NA	NA	1.00	0.07	0.14
<b>Signal 2</b>											
rs6864776	44513304	G/A	0.23	0.97	1.04(1.02-1.06)	0.00078	6.22E-11	NA	0.07	1.00	0.05
chr5.44527739.l	44527739	ATACT/A	0.24	0.96	1.04(1.02-1.07)	0.00044	6.40E-11	NA	0.06	0.98	0.05
rs4634356	44553611	C/T	0.23	0.99	1.04(1.02-1.06)	0.00076	7.89E-11	NA	0.07	0.99	0.06
rs1905192	44531538	A/T	0.23	0.99	1.04(1.02-1.06)	0.00077	8.76E-11	NA	0.07	0.99	0.06
rs4866905	44555867	C/T	0.23	1.00	1.04(1.02-1.06)	0.00086	1.02E-10	NA	0.07	0.98	0.06
rs1482663	44543102	A/G	0.23	1.00	1.04(1.02-1.06)	0.00085	1.04E-10	NA	0.07	0.99	0.06
chr5.44496660.l	44496660	AG/A	0.23	0.98	1.04(1.02-1.06)	0.00087	1.04E-10	NA	0.06	0.99	0.05
rs7710996	44514350	G/A	0.23	0.99	1.04(1.02-1.06)	0.00085	1.07E-10	NA	0.06	0.99	0.05
rs6451763	44527841	C/A	0.23	0.99	1.04(1.02-1.06)	0.00087	1.09E-10	NA	0.07	0.99	0.05
c5_pos44562807	44527050	A/C	0.23	1.00	1.04(1.02-1.06)	0.00087	1.15E-10	NA	0.07	0.99	0.06
rs1351633	44543851	C/T	0.23	0.99	1.04(1.02-1.06)	0.00094	1.18E-10	NA	0.07	0.99	0.06
rs1384453	44496114	T/C	0.23	0.99	1.04(1.02-1.06)	0.00091	1.20E-10	NA	0.06	0.99	0.05
rs1482665	44546628	G/A	0.23	1.00	1.04(1.02-1.06)	0.00093	1.20E-10	NA	0.07	0.99	0.06
rs983940	44544136	G/A	0.23	0.99	1.04(1.02-1.06)	0.00095	1.21E-10	NA	0.07	0.99	0.06
rs6897963	44554781	C/A	0.23	1.00	1.04(1.02-1.06)	0.00098	1.33E-10	NA	0.07	0.98	0.06
rs1384454	44492998	C/T	0.23	0.99	1.04(1.02-1.06)	0.00098	1.43E-10	NA	0.06	0.99	0.05
rs10079222	44561473	G/C	0.23	1.00	1.04(1.02-1.06)	0.001	1.45E-10	NA	0.07	0.98	0.06
rs7736427	44524276	A/G	0.25	0.93	1.04(1.02-1.07)	0.00028	1.48E-10	NA	0.05	0.94	0.05
rs10512860	44539953	G/C	0.23	1.00	1.04(1.02-1.06)	0.001	1.50E-10	NA	0.07	0.99	0.05
rs4866776	44525589	T/A	0.23	1.00	1.04(1.02-1.06)	0.001	1.52E-10	NA	0.07	0.99	0.05
rs1482690	44488840	T/A	0.24	0.98	1.04(1.02-1.06)	0.0011	1.91E-10	NA	0.06	0.99	0.05
rs12516346	44488156	A/C	0.23	0.99	1.04(1.01-1.06)	0.0016	2.39E-10	NA	0.07	0.97	0.06
rs1482684	44478742	G/A	0.23	1.00	1.04(1.01-1.06)	0.0013	2.41E-10	NA	0.06	0.99	0.05
chr5.44496659.l	44496659	TA/T	0.25	0.93	1.04(1.01-1.06)	0.0017	2.41E-10	NA	0.07	0.95	0.05
rs1482691	44487477	C/A	0.23	0.99	1.04(1.01-1.06)	0.0013	2.52E-10	NA	0.06	0.99	0.05
rs7724859	44471395	T/C	0.23	1.00	1.04(1.01-1.06)	0.0014	2.84E-10	NA	0.06	0.99	0.05
rs2128430	44462926	T/C	0.23	0.99	1.04(1.01-1.06)	0.0014	3.03E-10	NA	0.06	0.99	0.05
rs7707044	44471392	A/T	0.23	1.00	1.04(1.01-1.06)	0.0015	3.15E-10	NA	0.06	0.99	0.05
rs1905191	44456782	C/T	0.23	0.98	1.04(1.01-1.06)	0.0015	3.21E-10	NA	0.06	0.99	0.05

rs1120718	44466578	T/C	0.23	1.00	1.04(1.01-1.06)	0.0017	3.86E-10	NA	0.06	0.99	0.05
rs4866899	44444857	C/T	0.23	0.98	1.04(1.01-1.06)	0.0017	4.49E-10	NA	0.06	0.98	0.05
rs7712213	44451269	A/C	0.23	0.98	1.04(1.01-1.06)	0.002	5.30E-10	NA	0.07	0.98	0.05
rs6451762	44463186	C/T	0.24	0.96	1.04(1.01-1.06)	0.0023	7.36E-10	NA	0.06	0.98	0.05
rs7703171	44441175	C/T	0.23	0.95	1.04(1.01-1.06)	0.0018	8.24E-10	NA	0.06	0.95	0.05
rs6879342	44473605	A/G	0.24	0.98	1.03(1.01-1.06)	0.0067	3.57E-09	NA	0.07	0.96	0.06
<b>Signal 3</b>											
rs200229088	44701817	TTG/T	0.31	0.65	1.09(1.07-1.12)	2.30E-12	5.61E-04	1.12E-05	0.14	0.05	1.00

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**Supplementary Table 3.** ER/PR status, genotype and copy number at 5p12 of breast cell lines used in functional assays.

	<b>ER<sup>a</sup> status</b>	<b>PR<sup>b</sup> status</b>	<b>Signal 1 rs10941679</b>	<b>Signal 2 rs6864776</b>	<b>Signal 3 rs200229088</b>	<b>Copy Number<sup>c</sup></b>	<b>Haplotype</b>
SVCT	-	-	AA	GG	TTG/T	Not det	A/B
MCF10A	-	-	AA	GG	TTG/T	Not det	A/B
T47D	+	+	AA	AA	TTG/T	Gain	C/D
ZR751	+	+	AA	AA	TTG/T	Normal	C/D
MDA-MB-361	+	-	AA	AA	TTG/T	Gain	C/D
MCF7	+	+	AA	AG	TTG/T	Gain	Not det
Hs578T	-	-	AA	AG	TTG/T	Gain	Not det
BT474	+	+	AG	AG	TTG/T	Gain	Not det
Bre80	-	-	AG	AG	TTG/T	Not det	Not det
MDA-MB-231	-	-	AG	AG	TTG/T	Gain	Not det
BT-20	-	-	GG	AA	TTG/T	Gain	G/H
MDA-MB-468	-	-	GG	AA	TTG/T	Gain	G/H

<sup>a</sup>Estrogen receptor-alpha, <sup>b</sup>Progesterone receptor, <sup>c</sup>Copy number status from array CGH data (segmentation of normalized log<sub>2</sub> ratios) generated by the Cancer Cell Line Encyclopedia (CCLE), Not det = Not determined.

**Supplementary Table 4.** Oligonucleotides used for qRT-PCR of putative noncoding RNAs.

<b>qRT Primer</b>	<b>Sequence (5' to 3')</b>
FGF10-AS1FOR	TGCACCAACATCCATAACTCCTCG
FGF10-AS1REV	CTTTGTGAAGGAAAGCTGGACAGG
BRCAT54FOR	TGTGTGCTTAGACCTGACGCTGG
BRCAT54REV	CTCCTCACATGGCAGGAAGAAGG
RP11-503D12.1FOR	CAAGGTCAGTTGGCATTGTTCTGG
RP11-503D12.1REV	CAACCACCATGTCAAGGAACTCTAGC
RP11-473L15.3FOR	CACATGCATTGGCTATGACCTTGC
RP11-473L15.3REV	GCTGGCCATAAAGCTTCAATGTCC
TATA-binding proteinFOR	CATGGATCAGAACAACAGCCTGC
TATA-binding proteinREV	TTGTGAGAGTCTGTGAGTGGAAGAGC

**Supplementary Table 5.** Cell lines used in the study.

<b>Cell line<sup>1,2</sup></b>	<b>Source</b>
SVCT	ECACC 94122105
MCF10A	ATCC CRL-10317
T47D	ATCC HTB-133
ZR751	ATCC CRL-1500
MDA-MB-361	Gift from S. Lankani, UQCCR, AUS
MCF7	ATCC HTB-22
Hs578T	ATCC HTB-126
BT474	ATCC HTB-20
Bre80	Gift from R. Reddel, CMRI, AUS
MDA-MB-231	ATCC HTB-26
BT-20	ATCC HTB-19
MDA-MB-468	ATCC HTB-132

<sup>1</sup> Cell lines used in experiments were within passages 15-30.

<sup>2</sup> Cell lines were routinely tested for mycoplasma and short tandem repeat (STR) profiled.

**Supplementary Table 6.** Oligonucleotides used in 3C assays.

<b>3C Primer (<i>Eco</i>RI)</b>	<b>Sequence (5' to 3')</b>
FGF10 promoter bait	GCTGCTCTTCTGTACAGCGTGATGACAAGAGG
MRPS30 promoter bait	CCTTCACAGAAGTAAGCAACATGAGGGAAGC
HCN1 promoter bait	TCCCCTGGCACCTGCAAGTATGTGC
PRE Fragment 1	CTAAGACTGTCTCACCATGGAGCAACTCATGC
PRE Fragment 2	TGGCTTCCTGCCACTTGTTTCCTAATGC
PRE Fragment 3	CTCCTTGCTGTGAGAAGGTTGCACATCC
PRE Fragment 4	AGCACAAGAGCAGCCTCTACTGGGATATGC
PRE Fragment 5	TCCAAGTGAGTCAGTCTCCTGCCTCAGG
PRE Fragment 6	TCACTTGCTTTCTAGATTTGTGATTCTGCTTTCC
PRE Fragment 7	GTCCAGAGTCAAGTGGAGGAGATAACTCAAGGG
PRE Fragment 8	GAGGAATCAGGCAATCTTCATAAATATGGCTTACC
PRE Fragment 9	GGAATGCAAAGTGCAGCTATTGCTCTGC
PRE Fragment 10	GAAAATTTGGCATCATCCTTAGTGCACAGTAGG
PRE Fragment 11	CTGTGTAAGTTACAGAGCCATGATCATTGGTGG
PRE Fragment 12	GCCACATCCCTAGATGCCATGGTCC

**Supplementary Table 7.** Oligonucleotides used in cloning luciferase constructs.

Primer	Sequence (5' to 3')
FGF10 promoter FOR	GATCGCCATAAAGTGCGTTTGC
FGF10 promoter REV	CTGGAAGGGTAAGACCCGATGC
FGF10-AS1 promoter FOR	CCCAGGCATAATTTACGCTGAGG
FGF10-AS1 promoter REV	GCATTCACTTCTGGCCAGATCC
MRPS30 promoter FOR	ACTTTCCATGGCTCTGACTCAGTCC
MRPS30 promoter REV	CGGACCCAGAGGTCAACTTAAGC
BRCAT54 promoter FOR	CCTATCGTGAACGGTTATTGTGAGC
BRCAT54 promoter REV	AGTTCCTGGCAAAGGAGGTTGC
5p12 enhancer FOR	GACTCTGTATCTTCACGCATTTCCAGG
5p12 enhancer REV	TTAGAGACCAGAAATGTGGGCAAAGG
rs10941679 mut FOR	GCTTTTTATTGACTGTTGGAAAGAACACAGC
rs10941679 mut REV	GCTGTGTTCTTTCCACAGTCAATAAAAAGC

**Supplementary Table 8.** Oligonucleotides used in EMSAs.

SNP	allele <sup>a</sup>	Sequence (5' to 3') <sup>b</sup>
rs10941679	com	<sup>BIO</sup> GCTTTTTATTGACTATGGAAAGAACACAGC
rs10941679	min	<sup>BIO</sup> GCTTTTTATTGACTGTTGGAAAGAACACAGC
Neg control		AGAGATTGCCTGACGTCAGAGAGCTAG

<sup>a</sup> com: common allele, min: minor allele

<sup>b</sup> BIO: 5' biotinylation (present on both the sense and antisense strands of the duplex)

**Supplementary Table 9.** EMSA competitor duplexes and their target DNA binding proteins.

<b>Competition Target</b>	<b>Sequence (5' to 3')</b>
AP1	CGCTTGATGACTCAGCCGGAA
AP2	GATCGAACTGACCGCCCGCGGCCCGT
CEBP	TGCAGATTGCGCAATCTGCA
CTCF	AAGAAACCGCTAGGGGGCCTACT
FOXA1	CTGGTCTTAAAGGTGTTTACCTTGTCTGAT
FOXA2	GTTGACTAAGTCAATAATCAGAATCAG
GATA3	CACTTGATAACAGAAAGTGATAACTCT
HNF6	GATTCATATTGATTTCAAAA
HOXA10	GCATTCAGAAGGTTATAGCTTT
MYC	TCAGACCACGTGGTCGGG
NFKB	AGTTGAGGGGACTTTCCCAGGC
OCT1	TGTCGAATGCAAATCACTAGAA
SP1	ATTCGATCGGGGCGGGGCGAGC

**Supplementary Table 10.** Oligonucleotides used for ChIP-qPCR analyses.

<b>Name</b>	<b>Sequence (5' to 3')</b>
ChIP rs10941679 FOR	TTGAATCAGATGTGTAGTGCTTCC
ChIP rs10941679 REV	AACCTGAATGCAGCAAAATAGC
ChIP negcontrol FOR	CTAAGGACGAGATGCACATGG
ChIP negcontrol REV	TCAAGTTTCCAACACTCCAACAGG

## **CONSORTIA**

For members of kConFab, see <http://www.kconfab.org/Organisation/Members.aspx>; for AOCS, see [http://www.aocstudy.org/org\\_coll.asp](http://www.aocstudy.org/org_coll.asp).

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