

1 Detection and replication of epistasis influencing
2 transcription in humans

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Abstract

Epistasis is the phenomenon whereby one polymorphism's effect on a trait depends on other polymorphisms present in the genome. The extent to which epistasis influences complex traits¹ and contributes to their variation^{2,3} is a fundamental question in evolution and human genetics. Though often demonstrated in artificial gene manipulation studies in model organisms,^{4,5} and some examples have been reported in other species,⁶ few examples exist for epistasis amongst natural polymorphisms in human traits.^{7,8} Its absence from empirical findings may simply be due to low incidence in the genetic control of complex traits,^{2,3} but an alternative view is that it has previously been too technically challenging to detect due to statistical and computational issues.⁹ Here we show that, using advanced computation¹⁰ and a gene expression study design, many instances of epistasis are found between common single nucleotide polymorphisms (SNPs). In a cohort of 846 individuals with 7339 gene expression levels measured in peripheral blood, we found 501 significant pairwise interactions between common SNPs influencing the expression of 238 genes ($p < 2.91 \times 10^{-16}$). Replication of these interactions in two independent data sets^{11,12} showed both concordance of direction of epistatic effects ($p = 5.56 \times 10^{-31}$) and enrichment of interaction p -values, with 30 being significant at a conservative threshold of $p < 0.05/501$. Forty-four of the genetic interactions are located within 2Mb of regions of known physical chromosome interactions¹³ ($p = 1.8 \times 10^{-10}$). Epistatic networks of three SNPs or more influence the expression levels of 129 genes, whereby one *cis*-acting SNP is modulated by several *trans*-acting SNPs. For example MBNL1 is influenced by an additive effect at rs13069559 which itself is masked by *trans*-SNPs on 14 different chromosomes, with nearly identical genotype-phenotype (GP) maps for each *cis-trans* interaction. This study presents the first evidence for multiple instances of segregating common polymorphisms interacting to influence human traits.

Main text

In the genetic analysis of complex traits it is usual for SNP effects to be estimated using an additive model where they are assumed to contribute independently and cumulatively to the mean of a trait. This framework has been successful in identifying thousands of associations.¹⁴ But to date, though its contribution to phenotypic variance is frequently the subject of debate,¹⁻³ there is little empirical exploration of the role that epistasis plays in the architecture of complex traits in humans.^{7,8} Beyond the prism of human association studies there is evidence for epistasis, not only at the molecular scale from artificially induced mutations⁴ but also at the evolutionary scale in fitness adaptation¹⁵ and speciation.¹⁶

Methods are now available to overcome the computational problems involved in searching for epistasis, but its detection still remains problematic due to reduced statistical power. For example, increased dependence on linkage disequilibrium (LD) between causal SNPs and observed SNPs,^{17,18} increased model

complexity in fitting interaction terms,¹⁹ and more extreme significance thresholds to account for increased multiple testing⁹ all make it more difficult to detect epistasis in comparison to additive effects. Thus, with small genetic effect sizes, as is expected in most complex traits of interest,¹⁴ the power to detect epistasis diminishes rapidly. There are two simple ways to overcome this problem. One is by using extremely large sample sizes;²⁰ another is by analysing traits that are likely to have large effect sizes among common variants. Because our focus was to ascertain the extent to which instances of epistasis arises from natural genetic variation we designed a study around the latter approach and searched for epistatic genetic effects that influence gene expression levels. Transcription levels can be measured for thousands of genes and like most complex diseases, these expression traits are typically heritable.²¹ But unlike complex diseases, genetic associations with gene expression commonly have very large effect sizes that explain large proportions of the genetic variance,²² making them good candidates to search for epistasis, should it exist.

In our discovery dataset (Brisbane Systems Genetics Study, BSGS²³) of 846 individuals genotyped at 528,509 SNPs, we used a two stage approach to identify genetic interactions. First, we exhaustively test every pair of SNPs for pairwise effects against each of 7339 expression traits in peripheral blood (1.03×10^{15} statistical tests, family-wise error rate of 5% corresponding to a significance threshold of $p < 2.91 \times 10^{-16}$, Methods). Second, we filtered the SNP pairs from stage 1 on LD and genotype class counts, and tested the remaining pairwise effects for significant interaction terms and used a Bonferroni correction for multiple testing (estimated type 1 error rate $0.05 \leq \alpha \leq 0.14$, Methods, Supplementary Figure S1). Using this design we identified 501 putative genetic interactions influencing the expression levels of 238 genes (Supplementary Table S1). We used strict quality control measures to avoid statistical associations being driven by technical artifacts (Methods). However it remains possible that unexplained technical artifacts may have led to the significant discovery interactions. Of the 501 discovery interactions, 434 had available data and passed filtering (Methods) in two independent replication datasets, Fehrmann¹² and the Estonian Genomics Centre University of Tartu (EGCUT),¹¹ in which we saw convincing evidence for replication. We used the summary statistics from the replication datasets to perform a meta analysis to obtain an independent p -value for the putative interactions, and 30 were significant after applying a Bonferroni correction for multiple testing (5% significance threshold $p < 0.05/501$, Table 1). To quantify the similarity of GP maps between the independent datasets (Figure 1) we decomposed the genetic effects of each of the SNP pairs into orthogonal additive, dominance and epistatic effects ($A1$, $A2$, $D1$, $D2$, $A1 \times A2$, $A1 \times D2$, $D1 \times A2$, $D1 \times D2$) and tested for concordance of the sign of the most significant effect (Supplementary Table S3, Methods). Sign concordance between the discovery and both replication datasets was observed in 22 out of the 30 significantly replicated interactions (expected value = 7.5 under the null hypothesis of no interactions, $p = 3.76 \times 10^{-8}$).

In addition, using the meta analysis from the replication samples only, we observed that 316 of the remaining 404 discovery SNP pairs had replication

113 interaction p -values more extreme than the 2.5% confidence interval of the
 114 quantile-quantile plot against the null hypothesis of no interactions where p -
 115 values are assumed to be uniformly distributed ($p \ll 1.0 \times 10^{-16}$, Figure 2 and
 116 Supplementary Figure S2). Concordance of the direction of the effect of the
 117 largest variance component was also highly significant ($p = 5.71 \times 10^{-31}$, Sup-
 118plementary Table S3). The congruence of the epistatic networks in discovery
 119 and replication datasets is shown in Figure 3, demonstrating that these com-
 120plex genetic patterns are common even across independent datasets. A further
 121replication was attempted using the Centre for Health Discovery and Wellbeing
 122(CHDWB) dataset,²⁴ but only 20 of the SNP pairs passed filtering because the
 123sample size was small ($n = 139$), and likely due to insufficient power we found
 124no evidence for replication (Supplementary Figure S6). It should be noted that
 125although it is a necessary step to establish the veracity of the interactions from
 126the discovery set, replication of epistasis is difficult in practice due to LD (Meth-
 127ods).

128 Though seldom the focus of association studies, SNPs with known main
 129 effects are often tested for $A \times A$ genetic interactions,⁹ but our analysis suggests
 130 this is unlikely to be the best strategy for its detection. The majority of our
 131 discovery interactions comprised of one SNP that was significantly associated
 132 with the gene expression level in the discovery dataset, and one SNP that had
 133 no previous association²² (439 out of 501, Methods). Only nine interactions
 134 were between SNPs that both had known main effects while 64 were between
 135 SNPs that had no known main effects. Additionally, we observed that the
 136 largest epistatic variance component for the 501 interactions was equally divided
 137 amongst $A \times A$, $A \times D$, $D \times A$ and $D \times D$ at the discovery stage ($p = 0.22$
 138 for departure from expectation). This is not surprising because these patterns
 139 of epistasis used for statistical decomposition are simply convenient orthogonal
 140 parameterisations of a two locus model, and are not intended to model biological
 141 function.²⁵

142 Of the discovery interactions, 26 were *cis-cis* acting (within 1Mb of the
 143 transcription start site, mean distance between SNPs was 0.53Mb), 462 were
 144 *cis-trans*-acting, and 13 were *trans-trans*-acting. We observed a wide range of
 145 significant GP maps (Figure 1) but the most common pattern of epistasis that
 146 we detected involved a *trans*-SNP masking the effect of an additive *cis*-SNP. For
 147 example, MBNL1 (involved in RNA modification and regulation of splicing²⁶)
 148 has a *cis* effect at rs13069559 which in turn is controlled by 13 *trans*-SNPs and
 149 one *cis*-SNP that each exhibit a masking pattern, such that when the *trans*-
 150 SNP is homozygous for the masking allele the decreasing allele of the *cis*-SNP
 151 no longer has an effect (Supplementary Figure S10). Each of these interac-
 152 tions has evidence for replication in at least one dataset and six are significantly
 153 replicated at the Bonferroni level (Supplementary Figure S3). We see similar
 154 epistatic networks involving multiple (eight or more) *trans*-acting SNPs for other
 155 gene expression levels too, for example TMEM149 (Supplementary Figure S11),
 156 NAPRT1 (Supplementary Figure S12), TRAPPC5 (Supplementary Figure S13),
 157 and CAST (Supplementary Figure S14). We observed that from pedigree anal-
 158 ysis these five gene expression phenotypes had non-additive variance component

estimates within the 95th percentile of the 17,994 gene expression phenotypes that were analysed previously²² (Supplementary Table S2, Methods).

In total the 501 interactions comprised 781 unique SNPs, which we analysed for functional enrichment (Methods). We tested the SNPs for cell-type specific overlap with transcriptionally active chromatin regions, tagged by histone-3-lysine-4,tri-methylation (H3K4me3) chromatin marks, in 34 cell types²⁷ (Supplementary Figure S5). There was significant enrichment for *cis*-acting SNPs in haematopoietic cell types only ($p < 1 \times 10^{-4}$ for the three tissues with the strongest enrichment after adjusting for multiple testing). However *trans*-acting SNPs did not show any tissue specific enrichment ($p > 0.1$ for all tissues). This difference between *cis* and *trans* SNPs suggests different roles in epistatic interactions where tissue specificity is provided by the *cis* SNPs. There is also enrichment for *cis*-SNPs to be localised in regions with regulatory genomic features as measured by chromatin states²⁸ (Supplementary Figure S4).

We also demonstrate physical organisation of interacting loci within the cell, suggesting a mechanism by which biological function can lead to epistatic genetic variance. It has been shown that different chromosomal regions spatially colocalise in the cell through chromatin interactions.¹³ We cross-referenced our epistatic SNPs with a map of chromosome interacting regions ($n = 96,139$) in K562 blood cell lines²⁹ (Methods) and found that 44 epistatic interactions mapped to within 5Mb ($p < 1.8 \times 10^{-10}$), (Supplementary Figure S15). Interaction of distant loci may occur through physical proximity in transcriptional factories that organise across different chromosome regions and can regulate transcription of related genes.³⁰

Quantifying the importance of epistasis in complex traits in humans remains an open question. Here we are able to identify 238 gene expression traits with at least one significant interaction given our experiment-wide threshold, where the minimum estimated variance explained by the epistatic effects of any interaction was 2.1% of phenotypic variance. Taking results from our previously published eQTL²³ we calculated that 1848 of the 7339 gene expression levels analysed were influenced by additive effects where the estimated additive variance of a locus was 2.1% or greater. Thus, we can infer that the number of instances of large additive effects is significantly greater than the number of instances of large epistatic effects.

In terms of their contribution to complex traits a more important metric might be the proportion of the variance that the epistatic loci explain.² Taking all additive effects detected in Powell *et al* (2012) that have additive variance explaining 2.1% or greater of phenotypic variance, we calculated that the proportion of total phenotypic variance of all 7339 gene expression levels explained by additive effects alone was 2.16%. By contrast, the estimated epistatic variance from the interacting SNPs detected in this study on average explain a total of 0.22% of phenotypic variance, approximately ten times lower than the estimated additive variance. There are several caveats to this comparison which we discuss in the Methods.

Overall, we have demonstrated that it is possible to identify and replicate epistasis in complex traits amongst common human variants, despite the rela-

205 tive contribution of pairwise epistasis to phenotypic variation being small. The
206 bioinformatic analysis of the significant epistatic loci suggests that there are a
207 large number of possible mechanisms that can lead to non-additive genetic varia-
208 tion. Further research into such epistatic effects may provide a useful framework
209 for understanding molecular mechanisms and complex trait variation in greater
210 detail. With computational techniques and data now widely available the search
211 for epistasis in larger datasets for traits of broader interest is warranted.

Tables

Table 1: Epistatic interactions significant at the Bonferroni level in two replication sets

	Gene (chr.)	SNP 1 (chr.)	SNP 2 (chr.)	BSGS ²	Fehrmann ³	EGCUT ³	Meta ⁴
1	ADK (10)	rs2395095 (10)	rs10824092 (10)	6.69 ¹	18.33 ¹	21.21 ¹	39.82 ¹
2	ATP13A1 (19)	rs4284750 (19)	rs873870 (19)	5.30	12.18	3.25	14.23
3	C21ORF57 (21)	rs9978658 (21)	rs11701361 (21)	9.42	6.08	16.36	21.67
4	CSTB (21)	rs9979356 (21)	rs3761385 (21)	11.99	25.20	16.72	42.27
5	CTSC (11)	rs7930237 (11)	rs556895 (11)	7.16	18.76	15.06	33.53
6	FN3KRP (17)	rs898095 (17)	rs9892064 (17)	16.16	28.24	29.39	59.95
7	GAA (17)	rs11150847 (17)	rs12602462 (17)	13.91	19.98	12.99	32.60
8	HNRPH1 (5)	rs6894268 (5)	rs4700810 (5)	15.38	8.55	3.01	10.37
9	LAX1 (1)	rs1891432 (1)	rs10900520 (1)	19.16	18.60	11.22	29.24
10	MBNL1 (3)	rs16864367 (3)	rs13079208 (3)	13.49	16.25	24.74	41.56
11	MBNL1 (3)	rs7710738 (5)	rs13069559 (3)	7.92	2.55	7.89	9.28
12	MBNL1 (3)	rs2030926 (6)	rs13069559 (3)	7.10	0.91	5.80	5.53
13	MBNL1 (3)	rs2614467 (14)	rs13069559 (3)	5.74	4.13	2.22	5.30
14	MBNL1 (3)	rs218671 (17)	rs13069559 (3)	7.63	0.62	5.82	5.23
15	MBNL1 (3)	rs11981513 (7)	rs13069559 (3)	7.71	0.43	5.36	4.58
16	MBP (18)	rs8092433 (18)	rs4890876 (18)	5.40	7.06	21.91	28.73
17	NAPRT1 (8)	rs2123758 (8)	rs3889129 (8)	8.45	15.12	16.08	30.77
18	NCL (2)	rs7563453 (2)	rs4973397 (2)	7.31	7.51	6.33	12.70
19	PRMT2 (21)	rs2839372 (21)	rs11701058 (21)	4.81	0.69	4.47	4.06
20	RPL13 (16)	rs352935 (16)	rs2965817 (16)	4.98	3.79	14.41	17.24
21	SNORD14A (11)	rs2634462 (11)	rs6486334 (11)	7.31	13.11	10.96	23.22
22	TMEM149 (19)	rs807491 (19)	rs7254601 (19)	12.16	81.55	45.78	145.78
23	TMEM149 (19)	rs8106959 (19)	rs6926382 (6)	5.80	3.06	8.80	10.72
24	TMEM149 (19)	rs8106959 (19)	rs914940 (1)	6.22	3.36	6.96	9.20
25	TMEM149 (19)	rs8106959 (19)	rs2351458 (4)	7.30	0.04	9.61	8.00
26	TMEM149 (19)	rs8106959 (19)	rs6718480 (2)	8.55	3.31	5.15	7.36
27	TMEM149 (19)	rs8106959 (19)	rs1843357 (8)	6.21	3.72	3.33	6.00
28	TMEM149 (19)	rs8106959 (19)	rs9509428 (13)	9.44	0.10	5.75	4.47
29	TRA2A (7)	rs7776572 (7)	rs11770192 (7)	8.23	3.19	1.89	4.09
30	VASP (19)	rs1264226 (19)	rs2276470 (19)	5.09	0.94	5.14	4.95

¹ $-\log_{10} p$ -values for 4 *d.f.* interaction tests

² Discovery dataset

³ Independent replication dataset

⁴ Meta analysis of interaction terms between replication datasets only

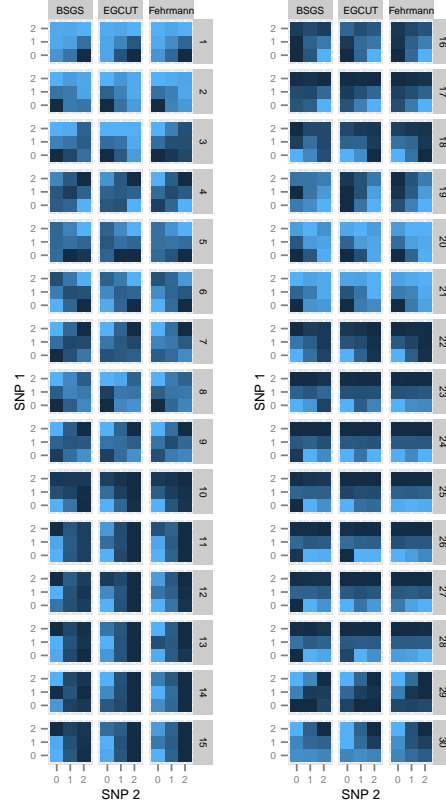


Figure 1: Replication of GP maps in two independent populations
 The GP maps for each epistatic interaction that is significant at the Bonferroni level in both replication datasets are shown. Each GP map consists of nine tiles where each tile represents the expression level for that two-locus genotype class. Phenotypes are for gene transcript levels (dark coloured tiles = high expression, light coloured tiles = low expression). Columns of GP maps are for each independent dataset. Rows of GP maps are for each of 30 significantly replicated interactions at the Bonferroni level, corresponding to the rows in Table 1. There is a clear trend of the GP maps replicating across all three datasets.

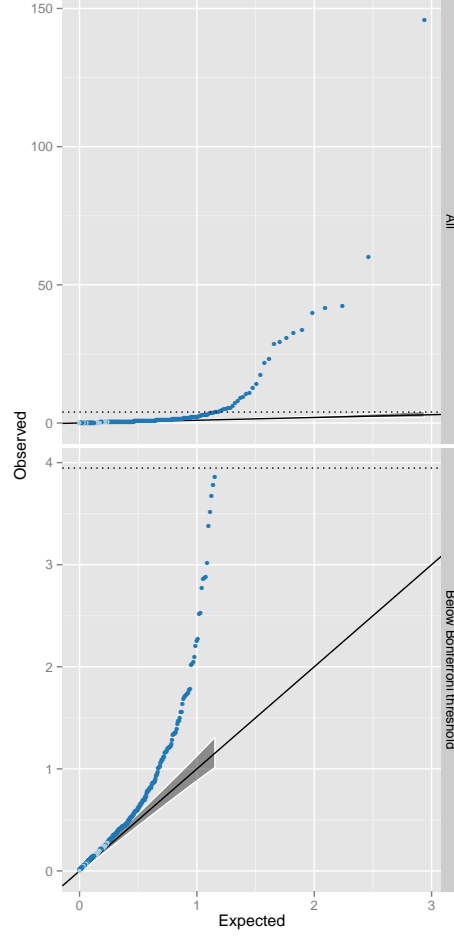


Figure 2: **Q-Q plots of interaction p -values from replication datasets**
The top panel shows all 434 discovery SNPs that were tested for interactions. Observed p -values (y -axis, $-\log_{10}$ scale) are plotted against the expected p -values (x -axis, $-\log_{10}$ scale). The multiple testing correction threshold for significance following Bonferroni correction is denoted by a dotted line. The bottom panel shows the same data as the top panel but excluding the 30 interactions that were significant at the Bonferroni level in the replication datasets. The shaded grey area represents the 5% confidence interval for the expected distribution of p -values. Dark blue points represent p -values that exceed the confidence interval, light blue are within the confidence interval.

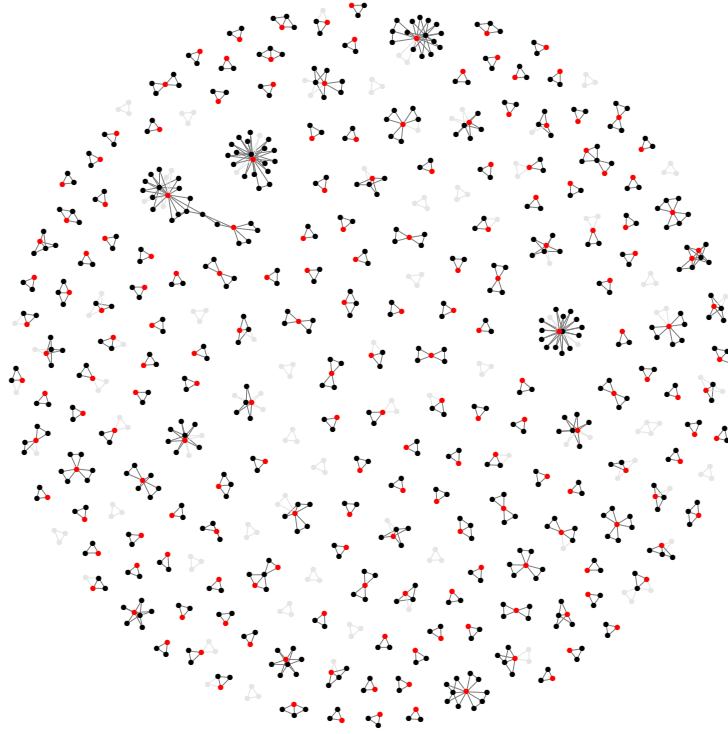


Figure 3: **Discovery and replication of epistatic networks** All 434 putative genetic interactions (edges) with data common to discovery and replication sets is shown, where black nodes represent SNPs and red nodes represent traits (gene expression probes). Three hundred and forty-five interactions had p -values exceeding the 2.5% confidence interval following meta analysis of the replication data. The remaining 89 interactions that did not replicate are depicted in grey. It is evident that a large proportion of the complex networks identified in the discovery set also exist in independent populations. An interactive version of this graph can be found here: http://kn3in.github.io/detecting_epi/

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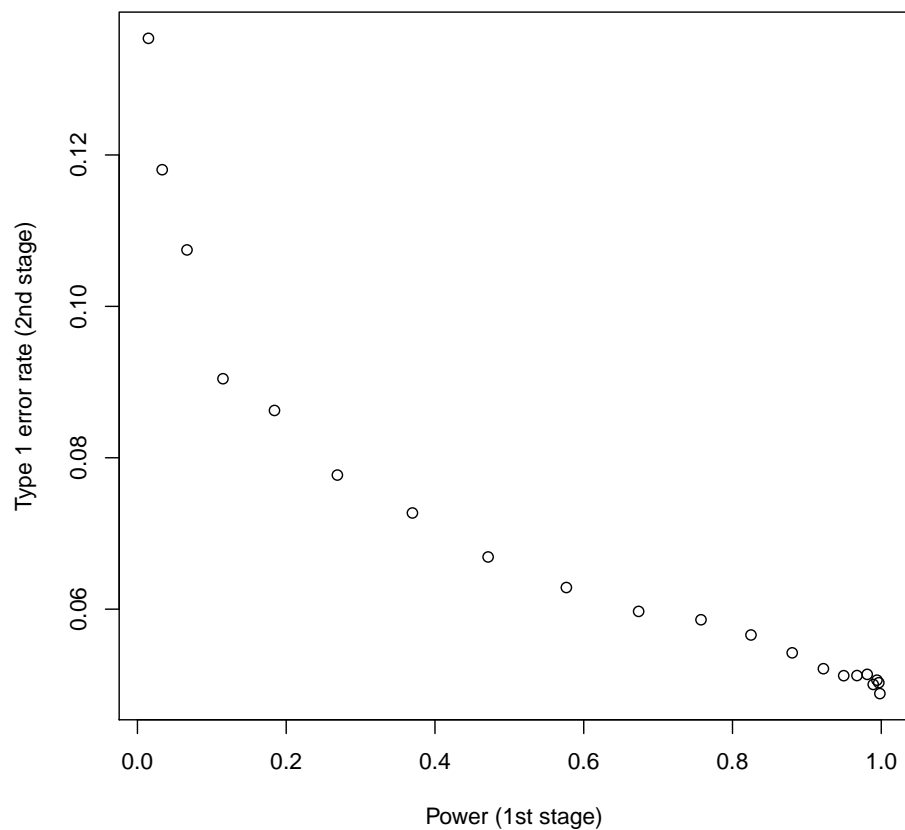


Figure S1: **Type 1 error rate of two stage design assuming a null model of one large additive effect and no epistasis** In stage 1 SNPs are tested for full genetic effects (8 d.f.) and those that surpass a threshold for multiple testing are then tested for significant interaction terms in stage 2. These interaction p -values are then adjusted (Bonferroni) for the total number of tests that passed stage 1. The type 1 error rate of this two stage design is dependent on the power, which is not known empirically.

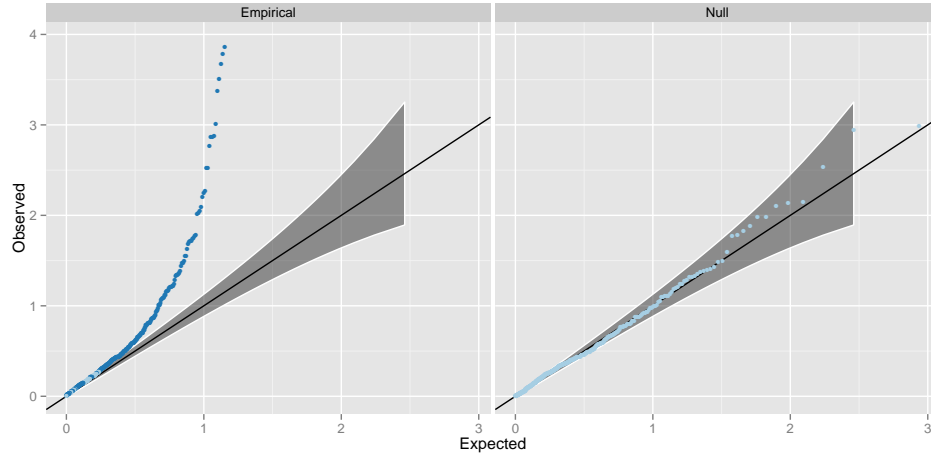


Figure S2: **Q-Q plots of interaction p -values from replication datasets, excluding the 30 points significant at the Bonferroni level** The right panel (Null) shows the interaction p -values from a meta analysis across two independent datasets on 434 randomly drawn SNP pairs. The left panel (Empirical) shows the interaction p -values from the 404 putative interactions that were not significant at the Bonferroni correction threshold. Dark blue points represent p -values that surpass the 2.5% FDR level, as in Figure 2.

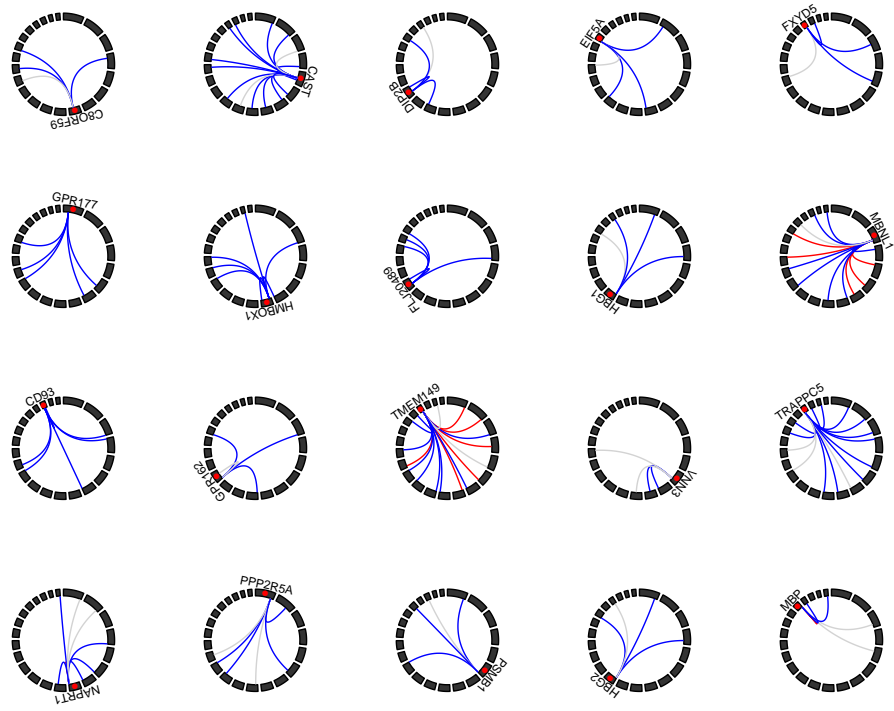


Figure S3: **Gene expression traits with four or more genetic interactions** Circle plots represent the genomic positions for SNPs (linking lines) and expression probes (red points). Chromosomes are represented by black blocks and ordered from 1 to 22 clockwise, starting from the top. Grey lines represent no evidence for replication, blue lines denote interactions that are outside the 97.5% confidence interval or the Q-Q plot (Figure 2), and red lines denote replication at the Bonferroni correction level. Most interactions are characterised as being *cis-trans* to the expression probe.

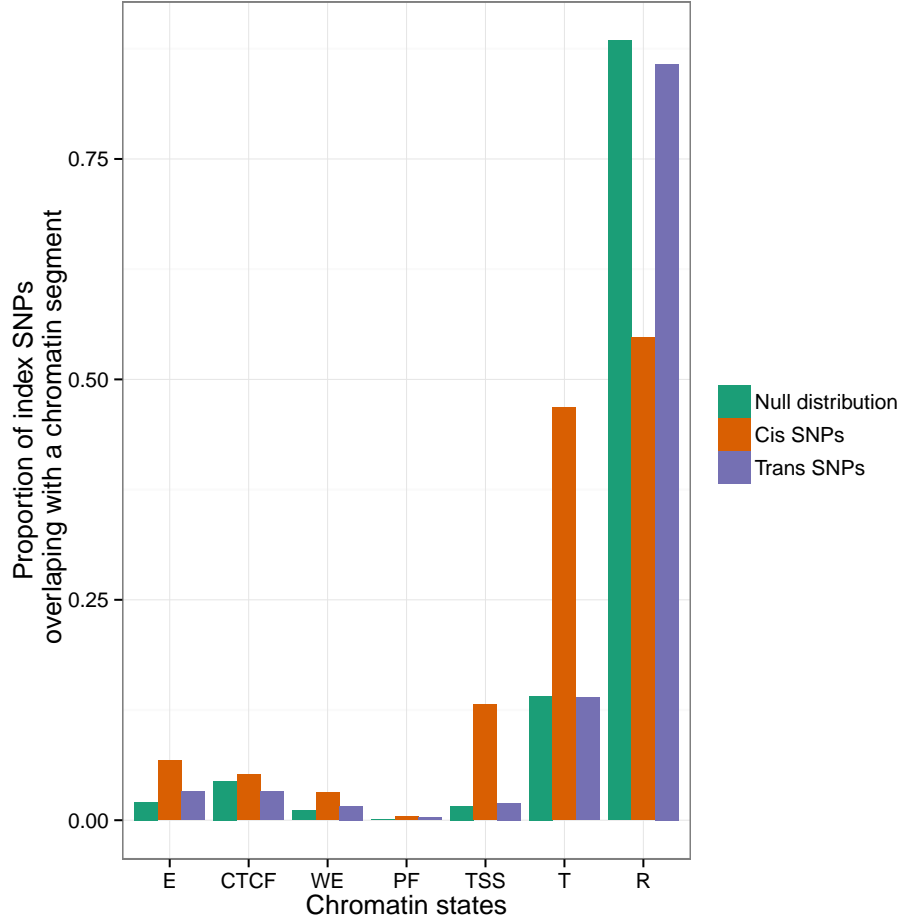


Figure S4: Location of SNPs relative to genomic features We used chromatin segmentation²⁸ as a method for labelling genomic features. All SNPs within 1Mb and $r^2 > 0.8$ of each *cis*- and *trans*-SNP were taken to find which genomic features (x -axis) were covered by the SNPs that compose the 501 significant interactions. Green bars represent the proportion (y -axis) of the 528,509 SNPs used in the analysis that fall within the range of the different genomic features. There is enrichment for *cis*-acting SNPs (red bars) in promotor regions, but *trans*-acting SNPs (blue bars) are not enriched for genomic features. The labels on the x -axis are as follows: E = Predicted enhancer, CTCF = CTCF enriched element, WE = Predicted weak enhancer or open chromatin cis regulatory element, PF = Predicted promoter flanking region, TSS = Predicted promoter region including transcriptional start site, T = Predicted transcribed region, R = Predicted Repressed or Low Activity region

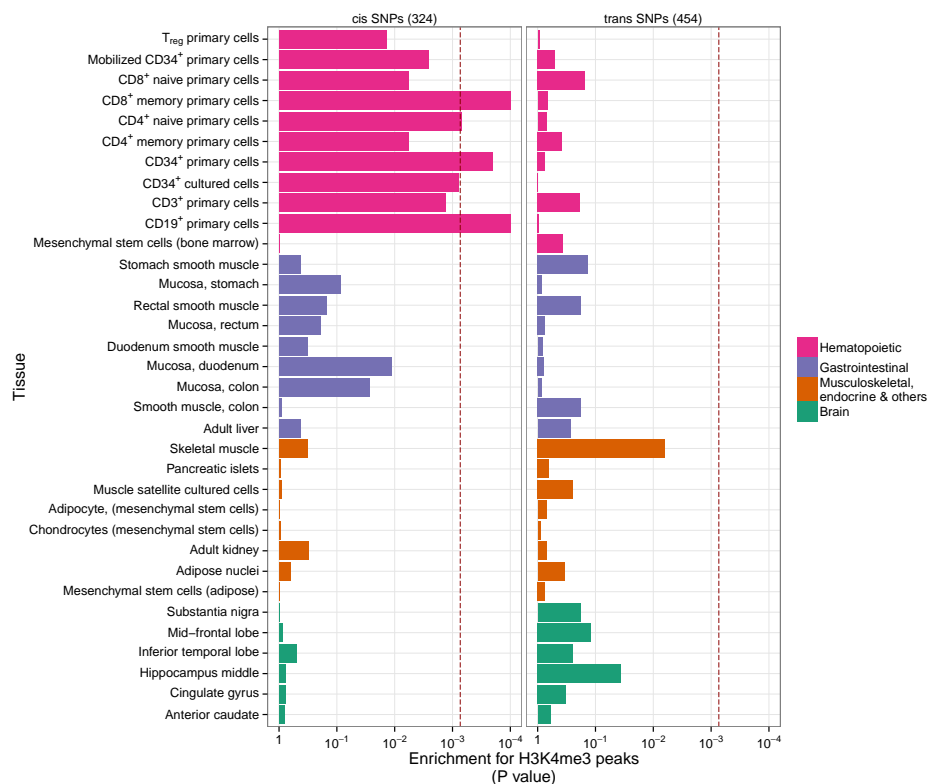


Figure S5: Tissue specific enrichment of SNPs in transcriptionally active regions The locations of transcriptional activity can be predicted by chromatin marks, assayed by H3K4me3.²⁷ Enrichment *p*-values are calculated using permutation analysis for 34 different cell types (*y*-axis) in four tissue types (Rows of boxes). The dotted red line denotes significance (Bonferroni correction for 34 cell types, *x*-axis). There is enrichment for *cis*-acting SNPs in Haematopoietic tissue types only. *Trans*-acting SNPs have no tissue specificity.

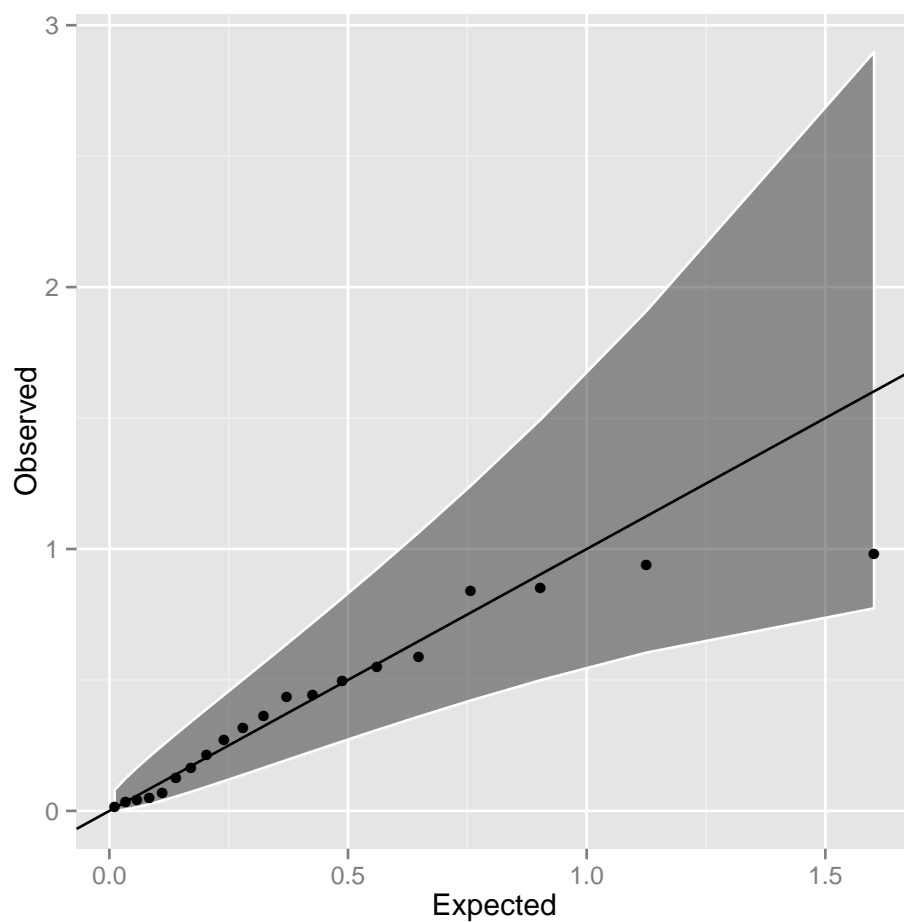


Figure S6: **Q-Q plot of interaction p -values in the CDHWB dataset**
 Twenty of the 501 discovery SNP pairs passed filtering in the CDHWB dataset (mainly due to small sample size). There is no evidence for enrichment of interaction terms, most likely due to insufficient power given the limited sample size.

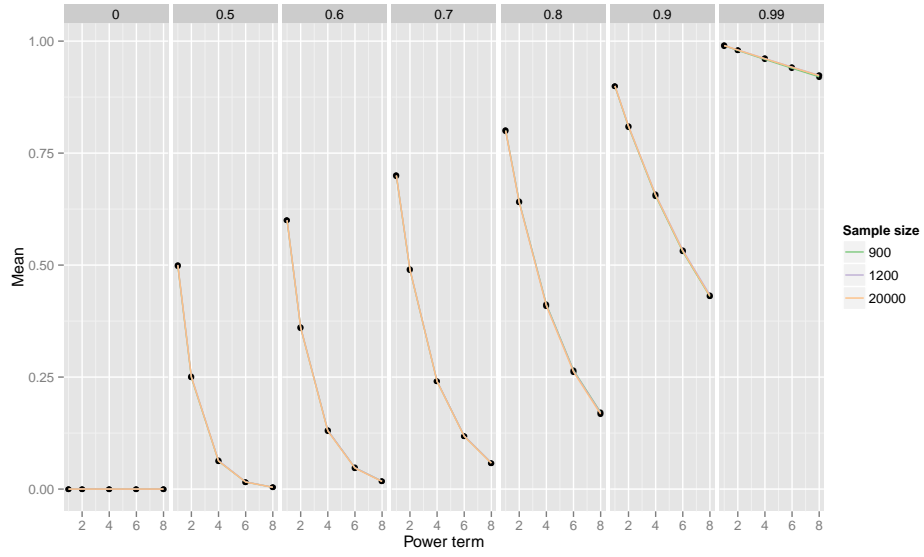


Figure S7: **Sampling mean for different power terms of population r values** Power of detection and replication of epistatic interactions depends not on r^2 between causal variants and observed SNPs, but on r^4, r^6, r^8 . For a given population value of LD r (columns of plots), plotted is the sample mean (y -axis) of \hat{r} , \hat{r}^2 (additive), \hat{r}^4 (dominance, $A \times A$), \hat{r}^6 ($A \times D$), \hat{r}^8 ($D \times D$) (x -axis) for different sample sizes (coloured lines). As true r reduces the statistical power to detect epistatic variants drops dramatically under the assumption that statistical power is proportional to higher moments of r .

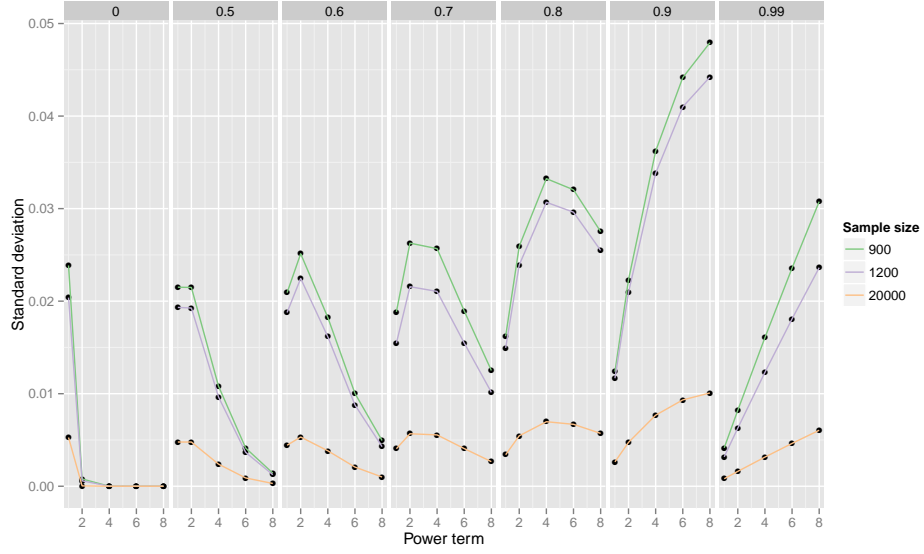


Figure S8: Sampling standard deviation for different power terms of population r values Power of detection and replication of epistatic interactions depends not on r^2 between causal variants and observed SNPs, but on r^4, r^6, r^8 . For a given a population value of LD r (columns of plots), plotted is the sampling standard deviation (y -axis) of \hat{r} , \hat{r}^2 (additive), \hat{r}^4 (dominance, $A \times A$), \hat{r}^6 ($A \times D$), \hat{r}^8 ($D \times D$) (x -axis) for different sample sizes (coloured lines). As the power term of r increases the sampling variance also increases. Supposing that there is sufficiently high r^x in the discovery sample for detection of epistasis, the replication sample is less likely to have similarly high r^x as x increases, leading to an expectation of reduced replication rates.

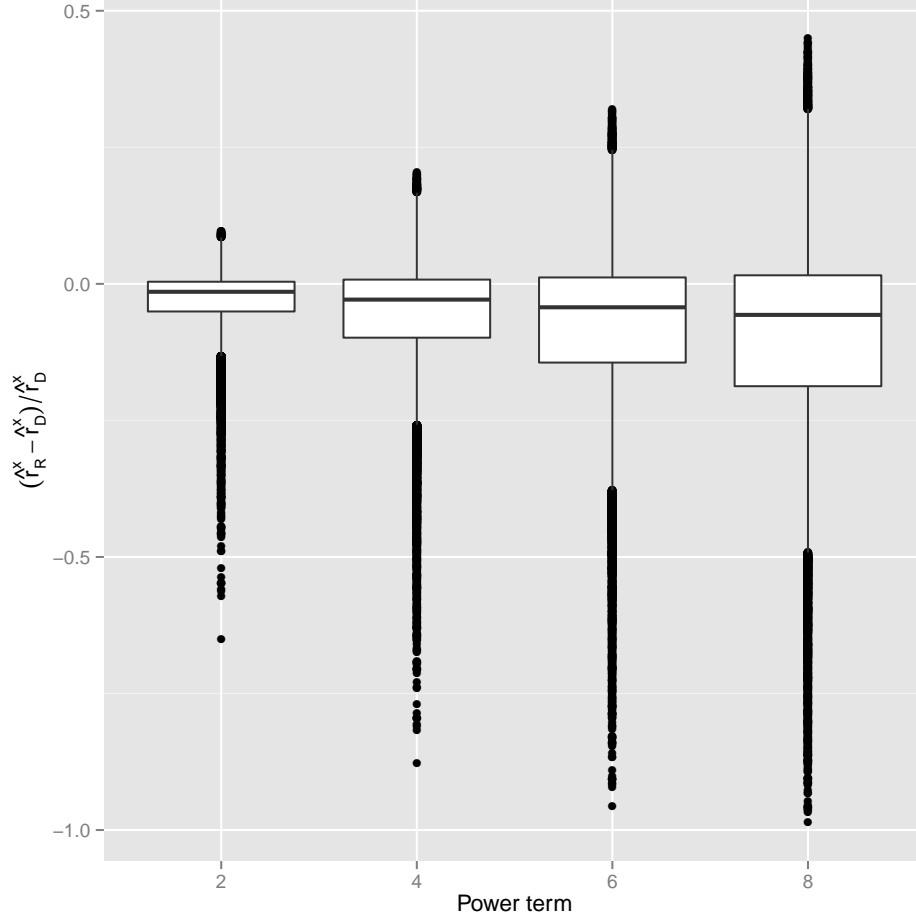


Figure S9: **Reduction in LD as estimated in replication data after ascertaining for high LD in discovery data** 100,000 “unobserved” causal variants (CVs) were tested for LD against a panel of 528,509 “observed” discovery markers (DMs). DM/CV pairs with LD $r^2 > 0.9$ were then tested in an independent sample. Simulation results of the proportional decrease between discovery and replication datasets in LD (y -axis) of $\hat{r}^2, \hat{r}^4, \hat{r}^6, \hat{r}^8$ (x -axis) are shown, where \hat{r}_D^x and \hat{r}_R^x are the sample LD measurements in the discovery and replication datasets, respectively. The average proportional decrease in the replication \hat{r}_R^x was 2.8%, 5.3%, 7.4% and 9.2% for $x = 2, 4, 6$ and 8, respectively.

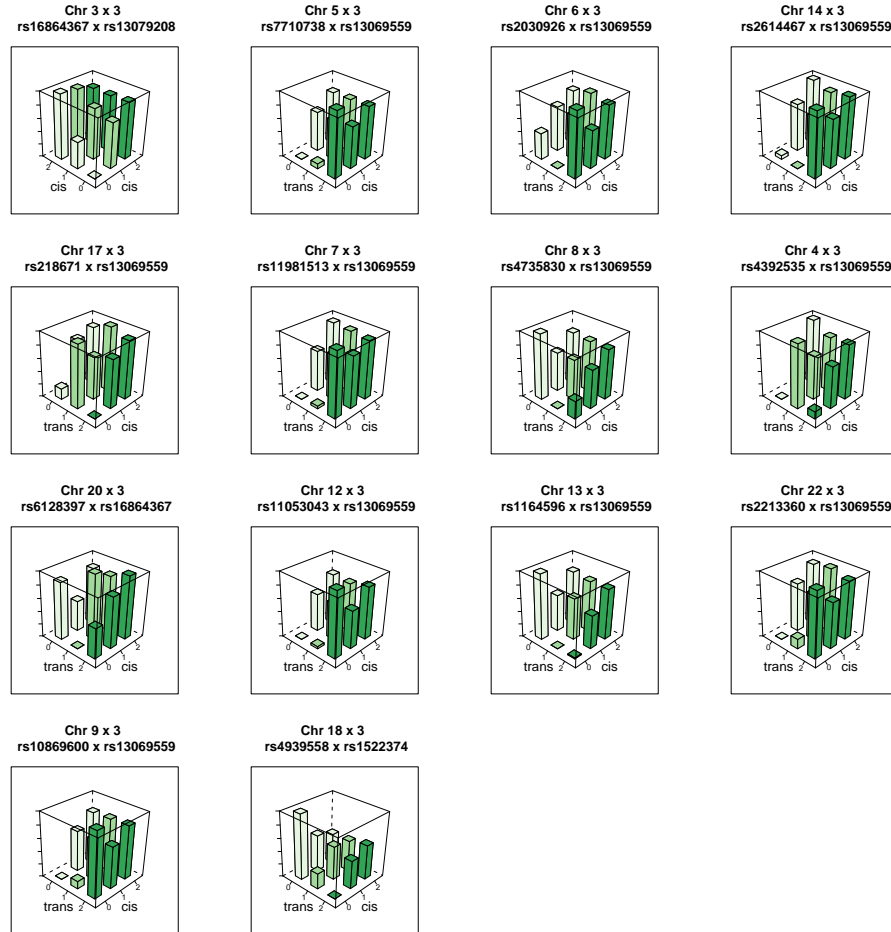


Figure S10: **Genotype-phenotype maps for 14 interactions influencing the expression of MBNL1** Each bar represents the mean phenotypic value for individuals in that genotype class. The rs13069559 SNP typically has a *cis*-additive decreasing effect on the expression of MBNL1, but in many of these interactions the *cis* effect is masked when the *trans* SNP is homozygous for the masking allele.

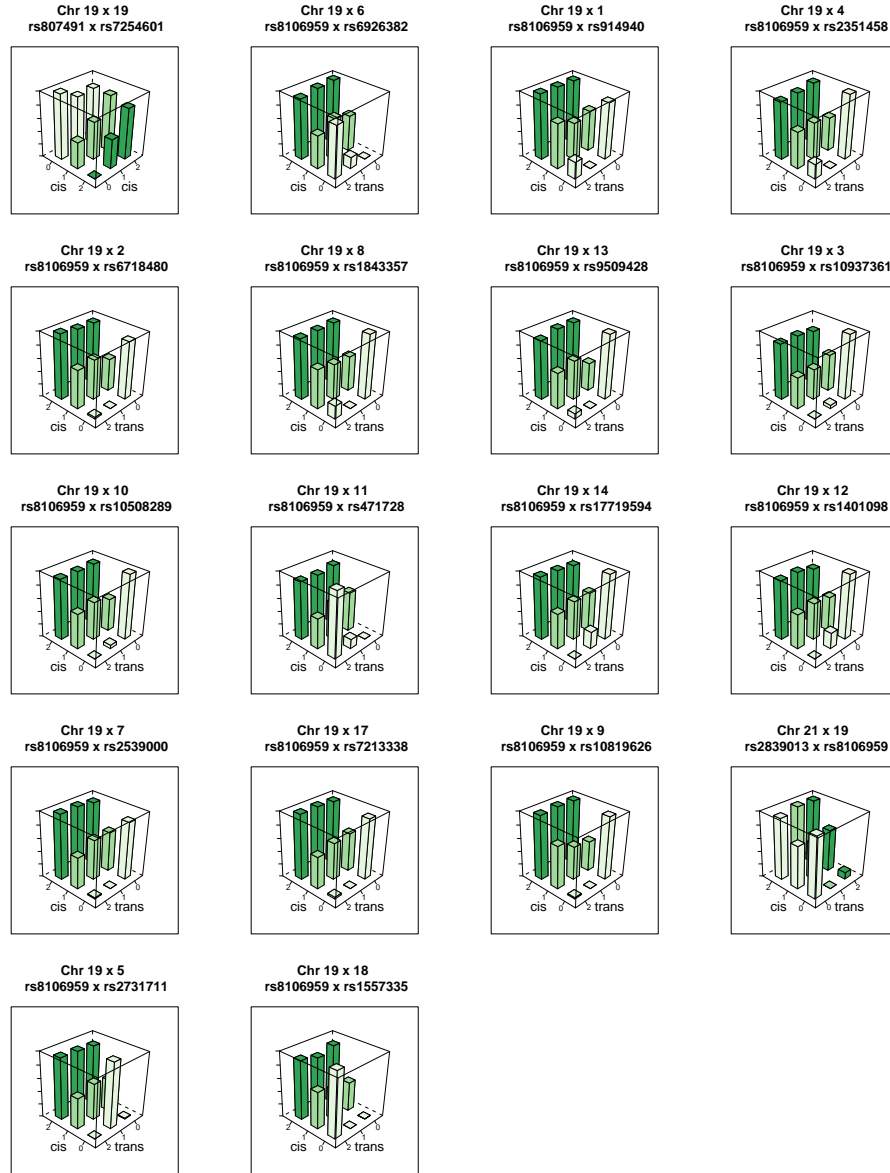


Figure S11: **Genotype-phenotype maps for 19 interactions influencing the expression of TMEM149** Each bar represents the mean phenotypic value for individuals in that genotype class. The rs13069559 SNP typically has a *cis*-additive decreasing effect on the expression of TMEM149, but in many of these interactions the *cis* effect is masked when the *trans* SNP is homozygous for the masking allele.

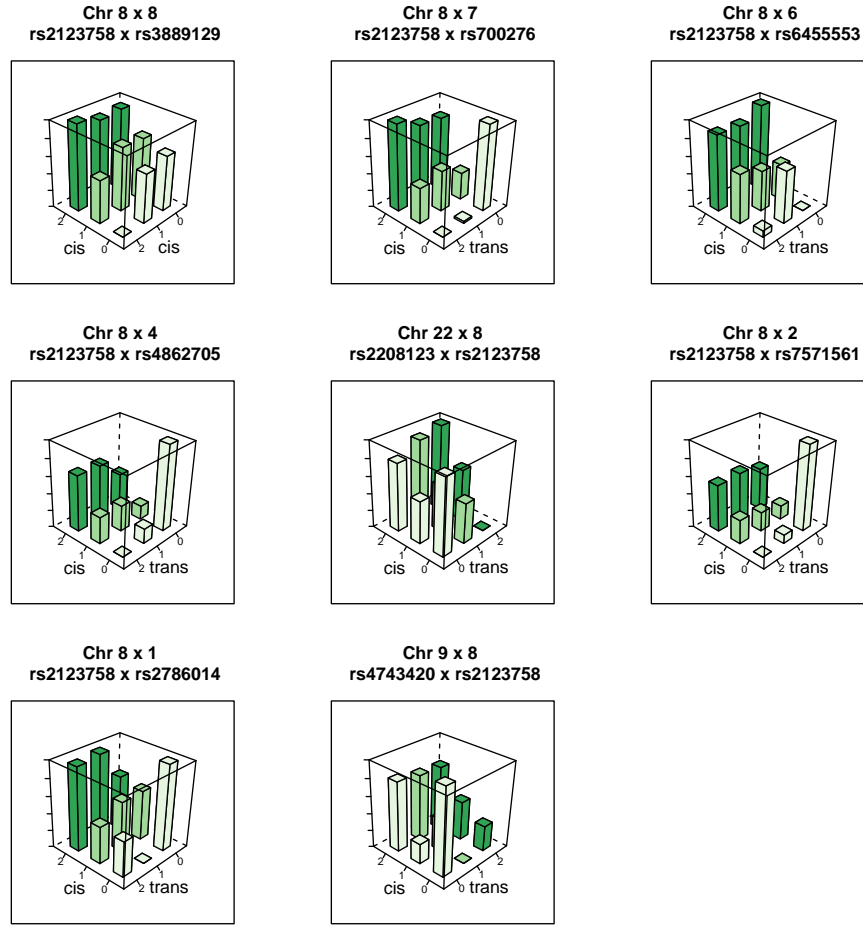


Figure S12: **Genotype-phenotype maps for 8 interactions influencing the expression of NAPRT1** Each bar represents the mean phenotypic value for individuals in that genotype class.

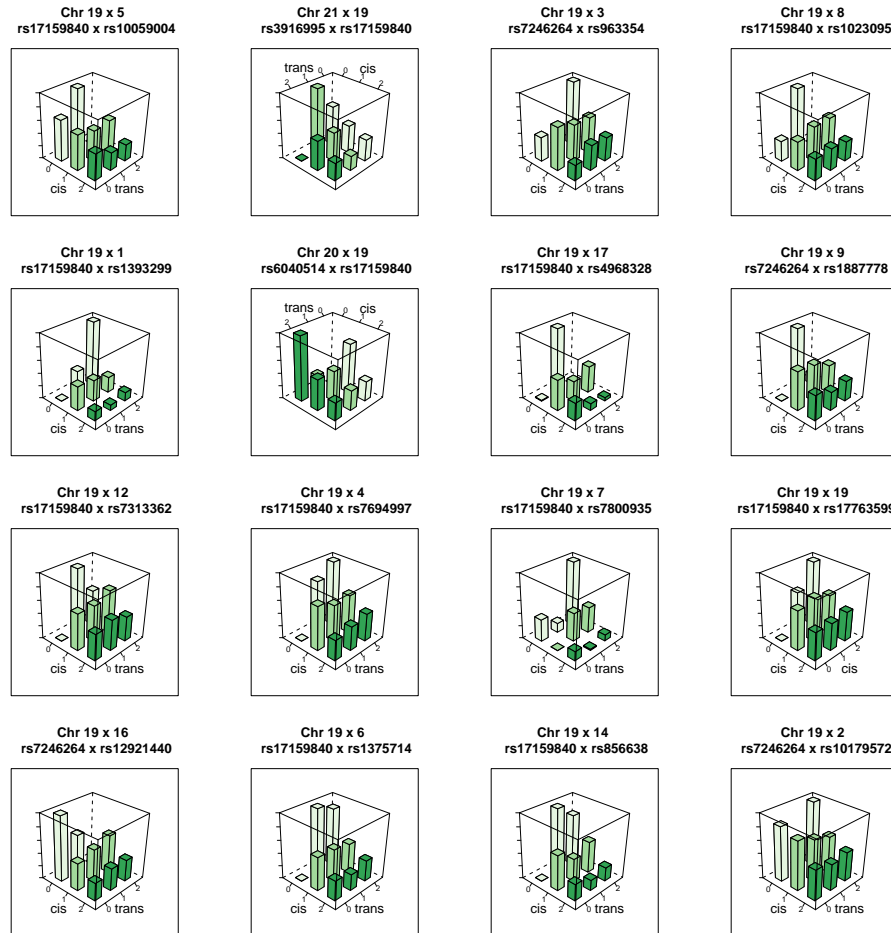


Figure S13: **Genotype-phenotype maps for 16 interactions influencing the expression of TRAPPC5** Each bar represents the mean phenotypic value for individuals in that genotype class.

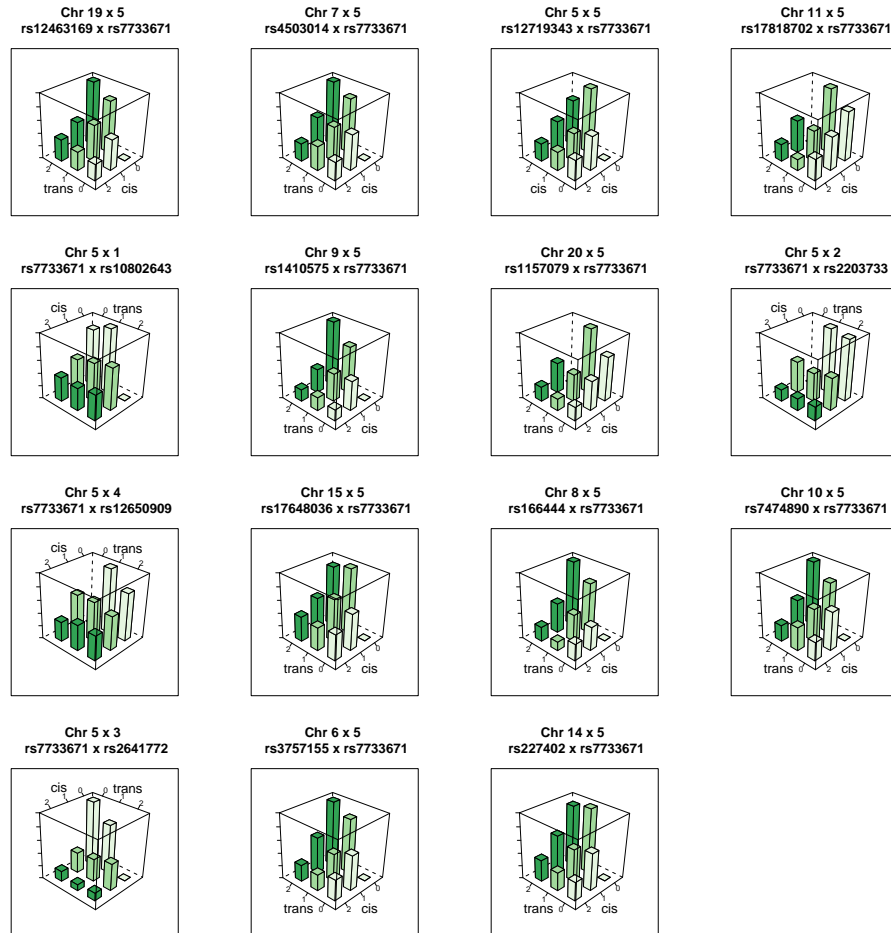


Figure S14: **Genotype-phenotype maps for 15 interactions influencing the expression of CAST** Each bar represents the mean phenotypic value for individuals in that genotype class.

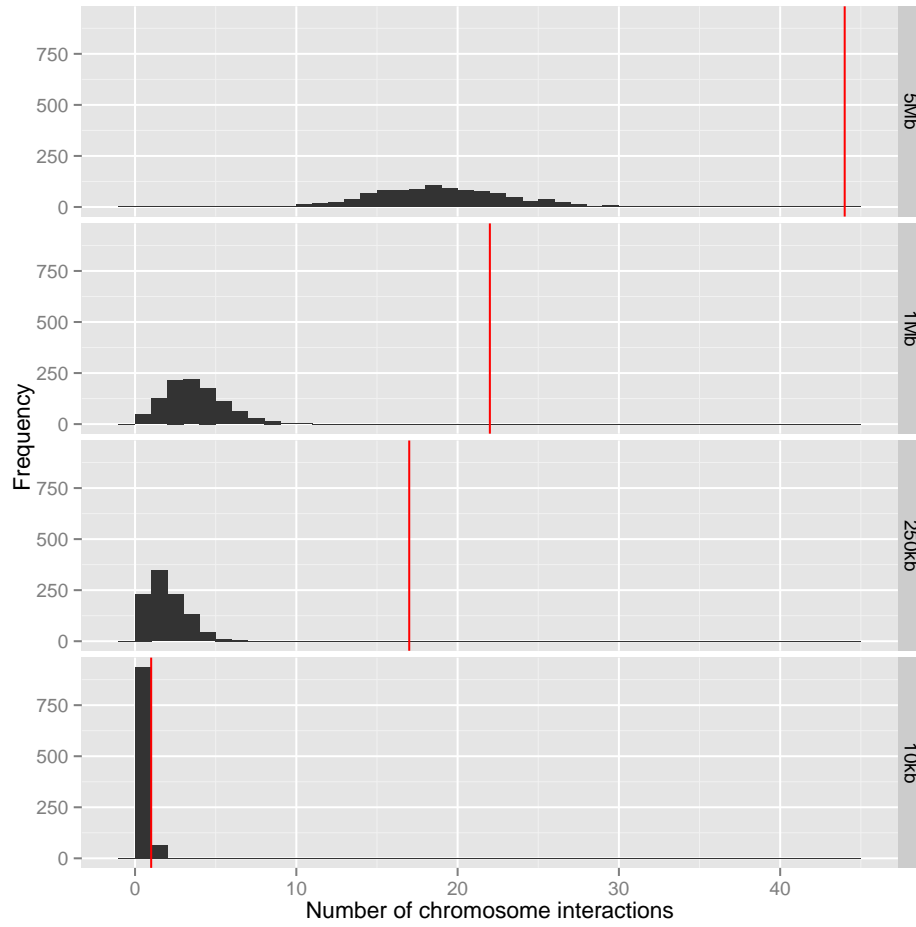


Figure S15: Number of overlaps between chromosome interactions and epistatic interactions Interacting chromosome regions may be a possible mechanism underlying epistatic interactions. The number of epistatic interactions within 20kb, 500kb, 2Mb and 10Mb of known chromosome interacting regions are shown by red vertical lines. The histograms represent the null distribution based on random sampling of 1,000 datasets for each window size.

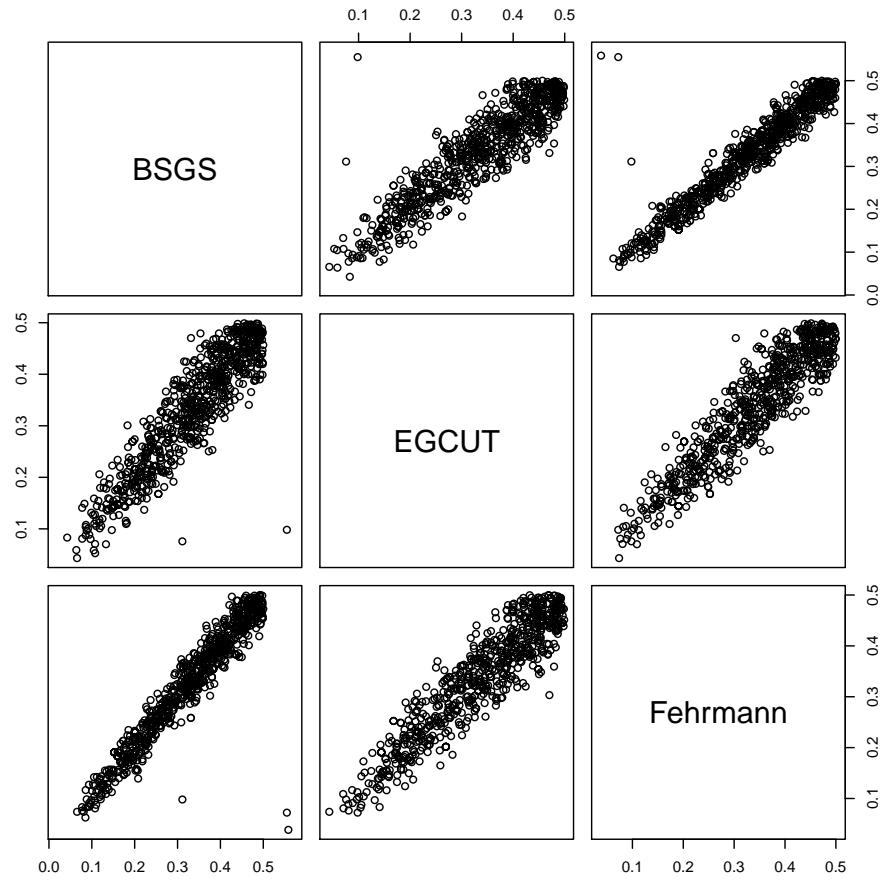


Figure S16: **Comparison of allele frequencies for 781 SNPs involved in genetic interactions across independent populations** Outliers were removed from the analysis as part of the filtering stage during replication.

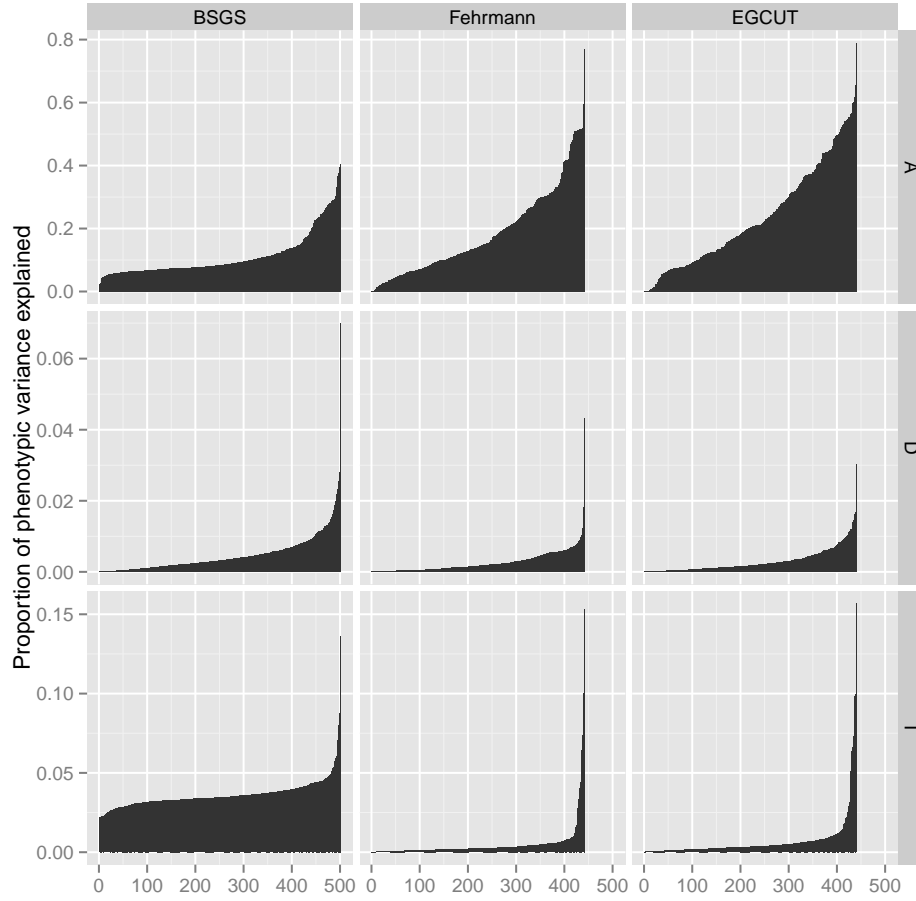


Figure S17: Comparison of variance explained by additive, dominant and epistatic effects from different cohorts How does the estimated variance decomposition change in different cohorts? The proportion of the phenotypic variance that is additive (A), dominant (D), or epistatic (I) for each putative interaction is shown on the y -axis (Note: different scales for each row). BSGS has 501 interactions whereas Fehrmann and EGCUT have 434 (x -axis). The variance estimates in each plot are ordered from lowest additive to highest. This is done independently for each cohort to depict the distribution of estimated effects.

291 **Supplementary Tables**

Table S1 – continued from previous page

Gene ID ^a	Expression trait	Chr.	rs ID	Chr.	SNP 1	Pos/Mb ^c	Association ^d	rs ID	Chr.	SNP 2	Pos/Mb ^c	Association ^d	BSGS ^e	Interaction statistic ^f	-log ₁₀ p-values	EGCUT ^g	Meta ^g	Distance / Mb ^h
C8ORF59	ILMN.1653205	8	rs8051751	16	7188323	7188323	C9ORF72	rs2890452	8	86102223	86102223	C8ORF59	5.79	1.39	0.18	0.18	0.87	
C9ORF72	ILMN.1741881	9	rs10122802	9	27356780	27356780	C9ORF72	rs2526698	1	242029101	242029101	CABC1	6.36	0.96	0.01	0.01	0.37	
CABC1	ILMN.1731064	10	rs12765847	10	4333308	4333308	INPP5E	rs7338725	1	227174210	227174210	CABC1	6.36	0.94	0.00	0.00	0.34	
CARD9	ILMN.1712532	9	rs4266703	9	138289825	138289825	INPP5E	rs684040	5	82128660	82128660	INPP5E	5.81	0.99	0.86	0.42	0.42	
CARD9	ILMN.1712532	9	rs4266703	11	6026061	6026061	INPP5E	rs4077315	9	1392660486	1392660486	INPP5E	7.61	0.23	0.96	0.62	0.75	
CARD9	ILMN.1712532	9	rs4266703	20	17737878	17737878	INPP5E	rs7733671	5	96000269	96000269	CABC1	7.77	0.77	0.96	0.62	0.75	
CABC1	ILMN.1717234	5	rs1329169	5	17321369	17321369	CABC1	rs7733671	5	96000269	96000269	CABC1	7.03	0.02	2.85	1.62	1.62	
CABC1	ILMN.1717234	5	rs12599264	16	81840122	81840122	CABC1	rs7733671	5	96000269	96000269	CABC1	7.03	0.02	2.85	1.62	1.62	
CABC1	ILMN.1717234	5	rs12719343	5	125369113	125369113	CABC1	rs7733671	5	96000269	96000269	CABC1	7.68	0.36	1.37	1.20	1.20	29.369
CABC1	ILMN.1717234	5	rs1410575	5	78255630	78255630	CABC1	rs7733671	5	96000269	96000269	CABC1	6.55	0.13	1.34	0.78	0.78	
CABC1	ILMN.1717234	5	rs166444	5	78392770	78392770	CABC1	rs7733671	5	96000269	96000269	CABC1	7.01	0.27	0.52	0.37	0.37	
CABC1	ILMN.1717234	5	rs17648036	15	27321111	27321111	CABC1	rs7733671	5	96000269	96000269	CABC1	7.81	0.97	0.03	0.41	0.41	
CABC1	ILMN.1717234	5	rs17818702	11	86107920	86107920	CABC1	rs7733671	5	96000269	96000269	CABC1	6.62	1.15	0.59	1.09	1.09	
CABC1	ILMN.1717234	5	rs227402	14	70496867	70496867	CABC1	rs7733671	5	96000269	96000269	CABC1	6.12	0.11	0.01	0.01	0.01	
CABC1	ILMN.1717234	5	rs282124	21	15166804	15166804	CABC1	rs7733671	5	96000269	96000269	CABC1	6.87	0.07	0.33	0.12	0.12	
CABC1	ILMN.1717234	5	rs3757155	6	136458593	136458593	CABC1	rs7733671	5	96000269	96000269	CABC1	7.24	0.72	0.33	0.12	0.12	
CABC1	ILMN.1717234	5	rs503014	7	31149140	31149140	CABC1	rs7733671	5	96000269	96000269	CABC1	5.88	0.92	1.56	1.72	1.72	
CABC1	ILMN.1717234	5	rs747890	10	59590078	59590078	CABC1	rs7733671	5	96000269	96000269	CABC1	6.74	0.49	0.12	0.23	0.23	
CABC1	ILMN.1717234	5	rs7733671	5	96000269	96000269	CABC1	rs10802643	1	238120177	238120177	CABC1	7.42	0.75	0.78	0.93	0.93	
CABC1	ILMN.1717234	5	rs7733671	5	96000269	96000269	CABC1	rs12650909	4	170192890	170192890	CABC1	7.42	0.23	0.78	0.50	0.50	
CABC1	ILMN.1717234	5	rs7733671	5	96000269	96000269	CABC1	rs2203733	2	224093101	224093101	CABC1	6.97	0.22	0.87	0.54	0.54	
CABC1	ILMN.1717234	5	rs7733671	5	96000269	96000269	CABC1	rs2641772	3	195531841	195531841	CABC1	6.03	0.19	0.26	0.15	0.15	
CABC1	ILMN.1651705	11	rs872311	18	66175386	66175386	CDC88B	rs11032695	11	34447586	34447586	CDC88B	6.41	0.26	0.30	0.32	0.32	
CDC88B	ILMN.1772208	11	rs2353203	11	17099980	17099980	CDC88B	rs541207	11	64125142	64125142	CDC88B	5.68	0.33	0.37	0.31	0.31	
CDC88B	ILMN.1772208	11	rs694739	11	64097233	64097233	CDC88B	rs12771349	10	96989193	96989193	CDC88B	5.62	0.23	0.18	0.14	0.14	
CDC88B	ILMN.1784863	7	rs3211834	7	80280117	80280117	CDC88B	rs1254900	2	85816334	85816334	VAMP8	6.93	0.15	0.01	0.02	0.02	
CDC88B	ILMN.1800540	1	rs750801	11	76033374	76033374	CDC88B	rs6700168	1	207502534	207502534	CD55	5.09	0.08	0.03	0.02	0.02	
CDC88B	ILMN.1800540	1	rs1884655	20	23074375	23074375	CDC88B	rs10255470	7	157182040	157182040	CD55	6.06	1.74	0.24	1.20	1.20	
CDC88B	ILMN.1704730	20	rs1884655	20	23074375	23074375	CDC88B	rs4696726	4	7992632	7992632	CD55	5.71	0.13	0.80	0.42	0.42	
CDC88B	ILMN.1704730	20	rs1884655	20	23074375	23074375	CDC88B	rs7622580	3	196721395	196721395	CD55	5.56	0.04	0.27	0.08	0.08	
CDC88B	ILMN.1704730	20	rs1884655	20	23074375	23074375	CDC88B	rs838875	12	125145394	125145394	CD55	6.31	0.24	1.67	1.16	1.16	
CDC88B	ILMN.1704730	20	rs1884655	20	23074375	23074375	CDC88B	rs9576388	13	38434472	38434472	CD55	7.88	0.71	0.22	0.45	0.45	
CDC88B	ILMN.1704730	20	rs2868504	20	37717578	37717578	CDC88B	rs1884655	20	23074375	23074375	CD55	5.71	0.64	0.75	0.81	0.81	14.697
CDC88B	ILMN.1704730	20	rs4813479	20	23076914	23076914	CDC88B	rs10925747	1	238899903	238899903	CD55	7.43	0.72	0.20	0.44	0.44	
CDC88B	ILMN.1704730	20	rs4813479	20	23076914	23076914	CDC88B	rs2873420	18	136500554	136500554	CD55	7.02	0.92	0.02	0.36	0.36	
CDC88B	ILMN.1704730	20	rs4813479	20	23076914	23076914	CDC88B	rs4328531	18	74439542	74439542	CD55	6.13	0.47	1.28	0.67	0.67	
CDC88B	ILMN.1704730	20	rs4813479	20	23076914	23076914	CDC88B	rs7726482	17	77264482	77264482	CD55	6.08	0.21	0.14	0.11	0.11	15.781
CDC88B	ILMN.1704730	20	rs4813479	14	104162263	104162263	CDC88B	rs7324744	13	115008038	115008038	CD55	5.46	0.95	0.07	0.45	0.45	
CDC88B	ILMN.2339796	13	rs861544	14	46614102	46614102	CDC88B	rs11655031	17	30833162	30833162	CD55	5.47	0.90	0.12	0.48	0.48	
CDC88B	ILMN.1730928	17	rs9905940	17	51956250	51956250	CDC88B	rs4803481	19	42066556	42066556	CD55	6.15	2.16	0.16	1.44	1.44	
CDC88B	ILMN.1745949	19	rs200609	18	42066556	42066556	CDC88B	rs2421050	5	158943044	158943044	CD55	6.67	0.16	0.14	0.12	0.12	
CDC88B	ILMN.1745949	19	rs4803481	18	42066556	42066556	CDC88B	rs13132719	4	180265266	180265266	CD55	5.75	0.15	0.24	0.12	0.12	
CDC88B	ILMN.1703754	18	rs6505780	18	13069782	13069782	CDC88B	rs13079012	3	134247706	134247706	CD55	6.36	0.23	0.10	0.09	0.09	
CDC88B	ILMN.2389845	3	rs3825569	14	101350298	101350298	CDC88B	rs772788	2	235248562	235248562	CD55	5.65	0.72	0.20	0.44	0.44	
CDC88B	ILMN.2359945	16	rs8192935	16	55861794	55861794	CDC88B	rs2695290	12	102087844	102087844	CD55	5.74	0.92	0.02	0.36	0.36	
CDC88B	ILMN.2359945	16	rs8192935	13	38838122	38838122	CDC88B	rs867578	11	81937002	81937002	CD55	4.75	0.07	1.28	0.67	0.67	
CDC88B	ILMN.2202940	12	rs591967	13	102277782	102277782	CDC88B	rs7313235	12	10132283	10132283	CD55	5.55	0.74	0.36	0.73	0.73	
CDC88B	ILMN.2202940	12	rs591967	16	84471642	84471642	CDC88B	rs3903088	10	134236688	134236688	CD55	7.54	0.95	0.07	0.45	0.45	
CDC88B	ILMN.1663142	12	rs429790	16	101350646	101350646	CDC88B	rs6863172	5	175595960	175595960	CD55	5.55	0.27	0.07	0.02	0.02	
CDC88B	ILMN.2403228	12	rs305054	11	963929337	963929337	CDC88B	rs169130	16	63121080	63121080	CD55	7.56	0.07	0.07	0.02	0.02	
CDC88B	ILMN.1674609	5	rs17129799	19	1047161	1047161	CDC88B	rs7366017	13	67713633	67713633	CD55	6.33	1.92	0.28	1.39	1.39	
CDC88B	ILMN.1770280	19	rs3752237	19	1047161	1047161	CDC88B	rs14536016	4	61738094	61738094	CD55	6.34	0.10	0.01	0.01	0.01	
CDC88B	ILMN.1653435	8	rs4353645	8	145369555	145369555	CDC88B	rs14536016	4	61738094	61738094	CD55	6.34	0.10	0.01	0.01	0.01	

0.23 Continued on next page

Gene ID ^a	Expression trait	Chr.	rs ID	Chr.	SNP 1	Pos/Mb ^c	Association ^d	rs ID	Chr.	SNP 2	Pos/Mb ^c	Association ^d	BSGS ^e	Interaction statistic ^f – Fehrmann ^g	– log ₁₀ p-values	Metag ^h	Distance / Mb
CPV1	ILMN.1682928	7	rs2835998	21	39202070	4	188859080	rs245884	7	29188475	CPV1	5.55	0.19	0.03	0.04		
CPV1	ILMN.1813256	2	rs2131290	4	188859080	4	188859080	rs1531133	7	46843631	CPV1	5.47	0.28	0.10	0.12		
CRLS1	ILMN.1737685	20	rs16139887	20	5986234	20	5986234	rs14773927	25	62406408	CRLS1	6.18	0.10	0.36	0.15		
CS1B	ILMN.1761797	21	rs9979356	21	45230974	21	45230974	rs3761385	25	45198355	CS1B	11.99	25.20	16.72	42.27	0.033	
CTNNA1	ILMN.1804854	5	rs2494943	18	69500505	18	69500505	rs176382	5	138226767	CTNNA1	5.74	0.02	0.41	0.11		
CTSC	ILMN.1866347	11	rs9457684	11	88139983	11	88139983	rs7079264	10	108679892	CTSC	5.67	0.92	0.74	1.03		
CTSC	ILMN.2242463	11	rs7572236	22	26250645	22	26250645	rs7128352	11	88073757	CTSC	5.84	0.49	0.80	0.73		
CTSC	ILMN.2242463	11	rs7930237	11	88117962	11	88117962	rs556895	7	16	18.76	15.06	33.53	0.040			
CTSC	ILMN.1651886	10	rs7108734	11	11456027	11	11456027	rs12784396	10	102027407	CTSC	5.42	0.21	0.01	0.03		
CYBRD1	ILMN.1712305	4	rs2592948	4	129994690	4	129994690	rs88427	2	172368120	CYBRD1	5.89	0.23	0.53	0.34		
CYBRD1	ILMN.1712305	4	rs7852475	9	140698856	9	140698856	rs88427	2	172368120	CYBRD1	5.68	0.20	0.02	0.04		
CYBRD1	ILMN.2087692	2	rs11257679	10	12318284	10	12318284	rs88427	2	172368120	CYBRD1	5.81	0.39	1.87	1.47		
CYBRD1	ILMN.2087692	2	rs36137908	20	23344590	20	23344590	rs88427	2	172368120	CYBRD1	5.53	0.05	0.83	0.36		
CYBRD1	ILMN.2087692	2	rs88427	20	172368120	20	172368120	rs88427	2	160112881	CYBRD1	5.85	0.87	0.10	0.44		
CYBRD1	ILMN.1704985	3	rs6021982	20	36571928	20	36571928	rs7591849	2	219650616	CYBRD1	5.42	0.29	0.86	0.60		
DAB2	ILMN.1218428	5	rs7778910	7	10451383	7	10451383	rs835223	5	39381357	DAB2	5.44	0.48	0.41	0.44		
DAB2	ILMN.1811648	17	rs9600173	17	43411688	17	43411688	rs1343244	6	82076988	DAB2	9.12	0.00	0.58	0.12		
DDT	ILMN.1690982	22	rs5760102	22	24248761	22	24248761	rs2378341	3	187475208	DDT	5.62	0.64	0.25	0.42		
DDT	ILMN.1779001	9	rs4937097	13	125962645	13	125962645	rs7042042	7	32451144	DDT	5.31	0.61	0.29	0.41		
DDT	ILMN.1779001	9	rs4937097	13	125962645	13	125962645	rs219515	7	88204888	DDT	5.47	0.08	0.41	0.16		
DDT	ILMN.1783996	1	rs10120023	9	137810259	9	137810259	rs10120023	9	137810259	DDT	6.39	0.77	0.02	0.29		
DDT	ILMN.1783996	1	rs123633827	11	109703727	11	109703727	rs7566044	2	169960422	DDT	6.00	0.06	1.17	0.58		
DHR9	ILMN.1733998	12	rs1519956	12	89468283	12	89468283	rs7566044	2	169960422	DHR9	6.48	0.37	0			

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Table S1 – continued from previous page

Gene ID ^a	Probe ID ^b	Expression trait		SNP 1			SNP 2			Interaction statistic ^f / -log ₁₀ p-values			Distance / Mb ^h		
		rs ID	Chr.	Pos/Mb ^c	Association ^d	rs ID	Chr.	Pos/Mb ^c	Association ^d	BSGS ^e	Fehrmann ^f	EGCUT ^g			
FE22	ILMN-1739586	2	rs2356400	19	44321776			rs13406184	2	36791226	FE22	5.78	0.14	0.33	0.16
FE22	ILMN-1739586	2	rs969010	4	159963132			rs11691600	2	36810133	FE22	6.59	0.14	0.28	0.14
FGD2	ILMN-2115005	6	rs4803848	19	46203030			rs31486	6	37001267	FGD2	5.69	0.12	0.25	0.11
FLJ20489	ILMN-2115005	12	rs9024634	10	133943931			rs831489	6	36999652	FLJ20489	5.49	1.20	0.11	0.66
FLJ20489	ILMN-1778144	12	rs17615703	12	117036766	FLJ20489		rs3782908	12	48169526	FLJ20489	5.81	0.06	0.70	0.29
FLJ20489	ILMN-1778144	12	rs472408	12	4569326			rs17615703	12	16769951	FLJ20489	5.73	0.03	0.11	0.02
FLJ20489	ILMN-1778144	12	rs472408	12	97033126			rs3782908	12	48169526	FLJ20489	6.49	0.31	0.47	0.36
FLJ20489	ILMN-1778144	12	rs204135	16	50626195			rs3782908	12	48169526	FLJ20489	6.04	0.38	0.17	0.21
FLJ20489	ILMN-1763663	16	rs9325634	21	43818790			rs3782908	12	48169526	FLJ20489	6.04	0.38	0.17	0.21
FLJ43093	ILMN-1763663	16	rs17112712	14	107276627			rs6906101	6	36667610	FLJ43093	5.48	0.14	0.95	0.53
FLJ43093	ILMN-212450	6	rs6906101	14	36667610	FLJ43093		rs13214069	6	32705248	FLJ43093	5.44	0.39	0.06	0.13
FN3KRP	ILMN-212450	6	rs6906101	14	36667610			rs9892064	17	80829703		5.48	0.00	0.64	0.18
FUCA1	ILMN-1652333	17	rs898095	17	80890638			rs9892064	17	80829703	FUCA1	16.16	28.24	29.39	59.95
FXD5	ILMN-1752728	17	rs4971478	17	80890638			rs788175	13	98328559		6.41	0.01	0.30	0.06
FXD5	ILMN-2309848	19	rs1633921	19	35695200			rs2285515	13	98328559	FXD5	3.70	0.09	0.41	0.17
FXD5	ILMN-2309848	19	rs17398183	20	55609148			rs11739594	5	141709563		6.58	0.03	0.48	0.15
FXD5	ILMN-2309848	19	rs2285515	19	35660450	FXD5		rs13067700	3	95331048		5.70	0.07	0.17	0.05
FXD5	ILMN-2309848	19	rs2285515	19	35660450	FXD5		rs17036504	2	47567329		6.00	0.09	0.09	0.51
G3BP2	ILMN-2309848	4	rs2285515	19	35660450	FXD5		rs1553985	4	7654604		6.10	0.28	0.08	0.04
G3BP2	ILMN-231758	4	rs10230232	17	29390239			rs12602462	17	78146016		13.91	19.98	12.99	32.60
GAA	ILMN-2410783	17	rs11150847	17	78153130			rs10902506	12	132678089		5.65	0.11	0.39	0.17
GAA	ILMN-2410783	17	rs8068856	17	78100731	GAA		rs7605821	2	2335655228		5.85	0.01	0.78	0.28
GAPT	ILMN-1675191	5	rs10070522	5	57786110	GAPT		rs10070522	5	57786110	GAPT	5.72	0.26	0.11	0.11
GAPT	ILMN-1675191	5	rs20838717	10	128038717			rs2950520	7	99827148	GATS	5.47	0.83	0.63	0.87
GATS	ILMN-1699631	7	rs1147447	14	66460742			rs2950520	7	99827148	GATS	5.47	0.83	0.63	0.87
GATS	ILMN-1699631	7	rs2425256	20	35056572			rs2197465	14	48572632		6.22	0.38	0.35	0.33
GDPD3	ILMN-1774901	16	rs3809624	16	30102802	GDPD3		rs1015111	4	128972357		5.86	0.55	0.09	0.24
GDPD3	ILMN-1774901	16	rs7204270	16	30102802	GDPD3		rs7577283	4	128972357		5.78	0.02	0.45	0.13
GDPD3	ILMN-1774901	16	rs7204270	16	30102802	GDPD3		rs7960552	12	111164237	GNLY	5.72	0.36	0.46	0.39
GNLY	ILMN-17906962	2	rs145072	13	110899955			rs2707210	12	6902002	GPR162	5.49	0.25	0.03	0.06
GNLY	ILMN-3239426	12	rs71986426	16	26084476			rs4740848	9	6554558	GPR162	5.07	0.25	0.06	0.07
GNLY	ILMN-3239426	12	rs1860563	16	6478898			rs9827054	3	188880113	GPR177	5.45	0.72	0.67	0.81
GPR162	ILMN-1730816	12	rs2272500	12	79685913			rs12065581	1	68732819	GPR177	5.76	0.17	0.40	0.22
GPR162	ILMN-1730816	12	rs2707210	12	6902002	GPR162		rs12065581	1	68732819	GPR177	5.43	0.79	1.43	1.50
GPR162	ILMN-1730816	12	rs2707210	12	6902002	GPR162		rs12065581	1	68732819	GPR177	5.43	0.79	1.43	1.50
GPR177	ILMN-1660549	1	rs11057383	12	124369421			rs12065581	1	68732819	GPR177	6.04	0.95	0.21	0.60
GPR177	ILMN-1660549	1	rs12527241	6	120468039			rs12065581	1	68732819	GPR177	5.86	0.24	0.34	0.23
GPR177	ILMN-1660549	1	rs12532999	9	127939793			rs12065581	1	68732819	GPR177	5.86	0.24	0.34	0.23
GPR177	ILMN-1660549	1	rs225613	16	11169683			rs12065581	1	68732819	GPR177	5.86	0.24	0.34	0.23
GPR177	ILMN-1660549	1	rs9575097	13	82986268			rs12065581	1	68732819	GPR177	5.86	0.24	0.34	0.23
GPR177	ILMN-2283325	1	rs6566669	18	70506011			rs12065581	1	68732819	GPR177	5.86	0.24	0.34	0.23
GPR177	ILMN-2283325	1	rs9290426	6	371399932			rs4965745	15	101508261	GPR177	5.86	0.24	0.34	0.23
GSDMB	ILMN-2347193	17	rs11557467	17	38028634	GSDMB		rs11101992	1	110266754	GSTM1	6.11	0.27	1.14	0.79
GSTM1	ILMN-2391861	13	rs12248673	10	53192833			rs11101992	1	110266754	GSTM1	5.91	0.27	1.14	0.79
GSTM1	ILMN-2391861	13	rs1547574	13	85345257			rs3754446	6	110253241	GSTM1	6.77	0.36	0.66	0.65
GSTM2	ILMN-2201580	1	rs6432807	13	96159560			rs4533333	2	77919015		6.36	0.52	0.27	0.31
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs6497007	15	58677017		6.52	0.27	0.31	0.23
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs9883949	21	19532546		6.52	0.27	0.31	0.23
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979			rs2855039	11	5271671	HBG2	5.70			

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Table S1 – continued from previous page

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Table S1 – continued from previous page

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Table S1 – continued from previous page

Gene ID ^a	Probe ID ^b		SNP 1		SNP 2		Interaction statistic ^c / -log ₁₀ p-values				Distance / Mb ^d				
	Chr.	rs ID	Chr.	Pos / Mb ^c	Association ^d	Chr.	Pos / Mb ^c	Association ^d	BSGS ^e	Fehrmann ^f		EGCUT ^g	Meta ^g		
REBE	1	rs4982958	14	24987865		1	rs301819	1	8501786	REBE	5.66	0.61	1.23	1.17	
REBE	1	rs7697290	4	135248366		1	rs301819	1	8501786	REBE	5.74	0.14	0.10	0.06	
REBE	1	rs11085829	19	13174312		1	rs301819	1	8501786	REBE	5.74	0.21	0.33	0.21	
REBE	1	rs3852011	3	112844086		1	rs301819	1	8501786	REBE	5.71	0.08	0.60	0.26	
RNASE6	14	rs11628398	14	8106521	RNASE6	13	rs7324365	13	100601327		5.48	0.42	0.21	0.26	
RNASE6	14	rs6003134	19	8106521		14	rs11628398	14	21182800	RNASE6	5.11	0.09	0.22	0.08	
RNF167	17	rs238230	17	4875566		13	rs4848457	13	54668512		4.37				
RNF167	17	rs400688	17	4839930	RNF167	3	rs11706900	3	36348968		5.59	0.71	0.46	0.64	
RNFEP	1	rs1107121	17	67153386		1	rs2819365	1	201983242		6.27	0.11	0.30	0.13	
RNFEP	1	rs8071611	17	67153386		1	rs2819365	1	201983242		4.32	1.48	0.32	1.28	
RPL13	16	rs352935	16	89648580		16	rs2965817	16	89513234		4.98	3.79	14.41	17.24	
RPL23AP7	2	rs1401202	16	80320056		2	rs4849261	2	114450028	RPL23AP7	5.55	0.13	0.73	0.38	
RPL36AL	14	rs3007033	14	50103816	RPL36AL	9	rs17495030	9	138035083		5.46	0.09	0.06	0.02	
RPL36AL	14	rs4009028	14	50020817		6	rs1502991	6	66137260		5.86	0.32	0.20	0.19	
RPL8	8	rs2958482	8	145984615	RPL8	1	rs1619856	1	234585790		4.59	0.10	0.37	0.15	
RPL8	8	rs4143674	20	4741304		8	rs2958482	8	145984615	RPL8	4.33	0.13	0.45	0.22	
SEC13	3	rs4889214	16	80913946		3	rs696221	3	10342876	SEC13	6.48				
SEC13	3	rs17085428	5	95388015		3	rs7695	1	156147326	SEC13	5.70	0.22	1.73	1.17	
SES3	11	rs12147460	14	104412137		11	rs684856	11	94906111	SES3	5.50	0.02	0.51	0.15	
SES3	11	rs355391	15	46591793	SES3	11	rs684856	11	94906111	SES3	5.67	0.31	0.06	0.10	
SES3	11	rs684856	15	46591793		8	rs7004947	8	134606425	SES3	5.60	0.21	0.51	0.31	
SH3BGL2	6	rs10838191	11	43893658		3	rs1354034	3	56849749	PPBP	5.52	0.70	0.12	0.35	
SH3BGL2	6	rs2545385	5	68283979		3	rs1354034	3	56849749	PPBP	5.97	0.50	0.51	0.30	
SH3BGL2	6	rs6845304	4	88280502		3	rs1354034	3	56849749	PPBP	5.23	0.32	0.71	0.33	
SH3BGL2	6	rs1034120	21	18196922		9	rs117455517	9	131785369	SH3BGL2	7.40	0.22	0.18	0.13	
SIRPG	20	rs1535883	20	1612819	SIRPG	4	rs6842739	4	60489510		5.74	0.29	0.18	0.17	
SLC22A18	11	rs11673260	19	52181798		11	rs367035	11	2923826	SLC22A18	5.47	0.09	0.24	0.09	
SLC22A18	11	rs367035	11	2923826	SLC22A18	7	rs3110874	7	153224179		5.70	0.15	0.10	0.06	
SLC22A18	11	rs367035	11	2923826	SLC22A18	2	rs6777203	2	241678528		6.15	0.39	0.13	0.19	
SLC41A3	3	rs012136	11	29236743		3	rs6777203	3	125801067	SLC41A3	5.88	1.10	0.82	1.24	
SLC45A4	8	rs698508	8	142337734	SLC45A4	5	rs7701916	5	174598073		5.95	0.86	0.07	0.40	
SLC46A3	13	rs948505	17	5502091		13	rs7701916	13	29259349	SLC46A3	5.52	0.09	0.58	0.26	
SMG7	1	rs8035259	15	97403923		1	rs10911353	1	183489203	SMG7	6.52	0.17	0.09	0.06	
SMG7	1	rs8035259	15	97403923		2	rs116777815	2	65800982		5.68	0.39	0.62	0.32	
SNHG8	20	rs116777815	20	4161500	SMOX	4	rs705832	4	119225940	SNHG8	4.11				
SNHG8	20	rs116777815	20	4161500		11	rs214097	11	17291499	SNHG8	6.00	0.29	1.03	0.72	
SNORD14A	4	rs1509420	15	46250108		11	rs214097	11	17291499	SNORD14A	7.31	13.11	10.96	23.22	
SNORD14A	4	rs2634462	11	17339197		11	rs6486334	11	17015557		6.08				
SNORD89	2	rs2045863	11	115929241		2	rs750783	2	101889306	SNORD89	5.96				
SNORD89	2	rs11605822	11	122986326		2	rs750783	2	101889306	SNORD89	6.33				
SNORD89	2	rs2135064	5	26778066		16	rs7185362	16	81888905	SNORD89	6.45				
SNUPN	15	rs1346466	21	46376528	SNUPN	3	rs1472075	3	193706323		5.59	0.13	1.41	0.83	
SNUPN	15	rs1346466	21	46376528	SNUPN	15	rs4774580	15	45652086	SPATA5L1	5.44	0.34	0.00	0.06	
SPATA5L1	15	rs131620	19	41117869		15	rs4774580	15	45652086	SPATA5L1	5.44				
STARD10	11	rs2221406	13	90174526		11	rs1006620	11	72509713		5.44				
STARD10	11	rs4073164	14	104947517		7	rs17685	7	75616105	STYXL1	5.65	0.67	0.12	0.33	
STYXL1	7	rs939294	4	180439236	SULF2	14	rs939294	4	180439236	STYXL1	5.88	0.57	0.17	0.31	
SULT1A4	16	rs1463965	20	74332954		16	rs3785354	16	28550667	TUFM	5.51	0.46	0.24	0.30	
SULT1A4	16	rs2436657	21	40119768		16	rs3785354	16	28550667	TUFM	7.05	0.01	0.05	0.00	
SURF6	9	rs6099626	20	56013994		9	rs3118663	9	136281753	SURF6	5.83				
SYTL2	13	rs1375719	13	103410782		13	rs485485	13	85495269	SYTL2	6.14	0.26	0.16	0.14	
THBS3	1	rs1939875	13	95422867		1	rs4072037	1	155162067	THBS3	5.47	0.28	0.31	0.24	
THBS3	1	rs1939875	13	95422867		1	rs4072037	1	155162067	THBS3	5.55	0.03	0.15	0.03	
THBS3	1	rs8014956	14	20687978		1	rs20449805	1	155194980	THBS3	5.65	0.31	0.76	0.55	
TIPRL	21	rs2823245	21	16745523		1	rs1320993	1	168154599	TIPRL	5.22	0.07	0.40	0.15	

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	SNP 2	Interaction statistic /	—	og ₁₀ p-values

Gene ID ^a			Expression trait		SNP 1			SNP 2			Interaction statistic ^b - log ₁₀ p-values			Distance / Mb ^c	
Gene ID ^a	Chr.	rs ID	Chr.	Pos/Mb ^c	Association ^d	rs ID	Chr.	Pos/Mb ^c	Association ^d	BSGS ^e	Fehrmann ^f	EGCUT ^g	Metag ^h		
TNME14	1	rs19404003	11	132389627		rs1725246	7	44581986	TNME14	5.70	0.06	1.34	0.70		
TNME14	1	rs2839013	21	47248091		rs8106959	19	36219525	TNME14	8.11	0.16	0.48	0.26		
TNME14	1	rs5762235	22	27925288		rs8106959	19	36219525	TNME14	6.79					
TNME14	1	rs6900518	20	45207005		rs8106959	19	36219525	TNME14	11.09					
TNME14	1	rs807491	19	36268923	SNX26	rs7254601	19	36147315	TNME14	12.16	81.55	45.78	145.78	0.122	
TNME14	1	rs8106959	19	36219525	TNME14	rs10508289	10	4799159		8.12	1.55	3.09	3.67		
TNME14	1	rs8106959	19	36219525	TNME14	rs10819626	9	133025756		8.02	0.40	0.99	0.80		
TNME14	1	rs8106959	19	36219525	TNME14	rs10819626	9	133025756		8.39	3.61	1.18	3.78		
TNME14	1	rs8106959	19	36219525	TNME14	rs14101098	12	12884559		7.37	2.41	1.00	2.52		
TNME14	1	rs8106959	19	36219525	TNME14	rs1557335	18	64268976		6.95	0.08	0.07	0.77	2.87	
TNME14	1	rs8106959	19	36219525	TNME14	rs17719594	14	90932598		6.93	3.06	0.77	2.87		
TNME14	1	rs8106959	19	36219525	TNME14	rs1843357	8	13822381		6.21	3.72	3.33	6.00		
TNME14	1	rs8106959	19	36219525	TNME14	rs2351458	4	113317583		7.30	0.04	9.61	8.00		
TNME14	1	rs8106959	19	36219525	TNME14	rs2539000	7	147619772		6.70	1.57	1.52	2.27		
TNME14	1	rs8106959	19	36219525	TNME14	rs2731711	5	171792273		5.92	0.19	0.33	0.19		
TNME14	1	rs8106959	19	36219525	TNME14	rs471728	11	121959460		8.89	0.90	3.62	3.51		
TNME14	1	rs8106959	19	36219525	TNME14	rs6718480	2	233879066		8.55	3.31	5.15	7.36		
TNME14	1	rs8106959	19	36219525	TNME14	rs6926382	6	161683974		5.80	3.06	8.80	10.72		
TNME14	1	rs8106959	19	36219525	TNME14	rs7213338	17	80357420		5.49	0.07	3.14	2.10		
TNME14	1	rs8106959	19	36219525	TNME14	rs914940	1	242889492		6.22	3.36	6.96	9.20		
TNME14	1	rs8106959	19	36219525	TNME14	rs9509428	13	21473952		9.44	0.10	5.75	4.47		
TNME14	1	rs8106959	19	36219525	TNME14	rs1449226	11	656845	TNME163A	5.79	0.64	0.12	0.32		
TNME14	1	rs8106959	19	36219525	TNME14	rs4963126	11	656845	TNME180	5.61	0.11	0.15	0.37		
TNME14	1	rs8106959	19	36219525	TNME14	rs10488630	7	128593948	IRF 5	5.52	1.03	0.17	0.62		
TNME14	1	rs8106959	19	36219525	TNME14	rs10488630	7	128593948	IRF 5	8.23	3.19	1.89	4.09	0.031	
TNME14	1	rs8106959	19	36219525	TNME14	rs11770192	7	23498358		5.61	0.28	0.40	0.29		
TNME14	1	rs8106959	19	36219525	TNME14	rs3916581	11	11888787	TRAPPC4	5.52	0.93	0.01	0.36	12.131	
TNME14	1	rs8106959	19	36219525	TNME14	rs3916581	11	11888787	TRAPPC4	5.97	0.21	1.60	1.07		
TNME14	1	rs8106959	19	36219525	TNME14	rs10059004	5	166970604		6.92	0.37	0.87	0.68		
TNME14	1	rs8106959	19	36219525	TNME14	rs1023095	8	132022957		7.79	0.12	0.18	0.08		
TNME14	1	rs8106959	19	36219525	TNME14	rs1375714	6	156404902		6.43	0.63	0.47	0.59		
TNME14	1	rs8106959	19	36219525	TNME14	rs1393299	1	242329791		6.38	0.21	0.24	0.16		
TNME14	1	rs8106959	19	36219525	TNME14	rs17663399	19	2369415		6.38	0.21	0.24	0.16		
TNME14	1	rs8106959	19	36219525	TNME14	rs4963828	17	57493457		7.51	0.50	0.38	0.44		
TNME14	1	rs8106959	19	36219525	TNME14	rs7513392	12	12743332		7.98	0.53	0.34	0.38		
TNME14	1	rs8106959	19	36219525	TNME14	rs778194	12	9941781		5.86	0.23	0.30	0.25		
TNME14	1	rs8106959	19	36219525	TNME14	rs7800935	7	145600926		6.27	0.15	0.33	0.16		
TNME14	1	rs8106959	19	36219525	TNME14	rs856638	14	85493550		6.73	0.24	0.07	0.08		
TNME14	1	rs8106959	19	36219525	TNME14	rs17159840	19	7758194	TRAPPC5	7.58	0.85	0.78	1.01		
TNME14	1	rs8106959	19	36219525	TNME14	rs17159840	19	7758194	TRAPPC5	7.73	0.51	0.55	0.56		
TNME14	1	rs8106959	19	36219525	TNME14	rs17159840	19	7758194	TRAPPC5	8.10	0.02	0.02	0.02		
TNME14	1	rs8106959	19	36219525	TNME14	rs10179572	2	228504503		6.71	0.14	0.26	0.13		
TNME14	1	rs8106959	19	36219525	TNME14	rs12921440	16	30408765		7.34	0.08	0.86	0.40		
TNME14	1	rs8106959	19	36219525	TNME14	rs1887778	9	134635088	RAPGEF1	7.05	0.08	0.60	0.69		
TNME14	1	rs8106959	19	36219525	TNME14	rs963354	3	157393770		7.41	0.36	0.90	0.69		
TNME14	1	rs8106959	19	36219525	TNME14	rs2395771	6	41264577	TREM1	5.42	0.11	0.25	0.11		
TNME14	1	rs8106959	19	36219525	TNME14	rs2395771	6	41264577	TREM1	5.92	1.20	1.23	1.69		
TNME14	1	rs8106959	19	36219525	TNME14	rs2032447	6	26043469	TRIM38	6.46	0.04	0.07	0.39		
TNME14	1	rs8106959	19	36219525	TNME14	rs10748526	10	82273079	TSPAN14	6.00	0.07	0.18	0.06		
TNME14	1	rs8106959	19	36219525	TNME14	rs12800998	11	2317951	TSPAN32	5.01	0.04	0.04	0.06		
TNME14	1	rs8106959	19	36219525	TNME14	rs620607	6	137947208		5.51	0.51	0.51	0.51	45.345	
TNME14	1	rs8106959	19	36219525	TNME14	rs620607	6	137947208		6.34	0.07	0.18	0.06		
TNME14	1	rs8106959	19	36219525	TNME14	rs1198819	22	238746880		6.13					
TNME14	1	rs8106959	19	36219525	TNME14	rs4783126	16	85147633							

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Table S1 – continued from previous page

Expression trait			SNP 1			SNP 2			Interaction statistic ^f / -log ₁₀ p-values			Distance / Mb ^h		
Gene ID ^a	Probe ID ^b	Chr.	rs ID	Chr.	Pos/Mb ^c	Association ^d	rs ID	Chr.	SNP-2	Association ^d	BSGS ^e	Fehrmann ^f	EGCUT ^g	Meta ^g
UBASH3A	LMN-2338348	21	rs1893592	21	43855067	UBASH3A	rs7201194	16	83600397		5.91	0.59	0.42	0.52
UBASH3A	LMN-2338348	21	rs1893592	21	43855067	UBASH3A	rs7201194	16	83600397		6.01	0.48	1.29	1.10
USP36	LMN-1697227	17	rs2279308	17	76794981	USP36	rs7225546	17	75151717		5.71	0.03	0.14	0.03
VASP	LMN-1743646	19	rs1264226	19	40063167		rs2276470	19	45974668		5.09	0.94	5.14	4.95
VNN2	LMN-1678939	6	rs10435352	7	103252718		rs1883613	6	133077063	VNN2	5.64	0.84	0.15	0.46
VNN2	LMN-1678939	6	rs10435352	7	103252718		rs1883613	6	133077063	VNN2	5.44	0.39	0.69	0.57
VNN2	LMN-1678939	6	rs134447	22	49927332		rs1883617	6	133072650	VNN2	5.72			
VNN2	LMN-1678939	6	rs134447	22	49927332		rs1883617	6	133072650	VNN2	5.77	0.33	0.19	0.19
VNN3	LMN-1678939	6	rs216495	11	16834510		rs1883617	6	133072650	VNN2	6.44	0.16	0.74	0.41
VNN3	LMN-1678939	6	rs10278073	7	151662184		rs2267952	6	133067782	VNN3	5.74	0.23	0.48	0.31
VNN3	LMN-1804935	6	rs1443946	8	73006453		rs2267952	6	133067782	VNN3	6.44	0.31	0.17	0.17
VNN3	LMN-1804935	6	rs348462	9	75547169		rs2267952	6	133067782	VNN3	5.82	0.03	0.19	0.04
VNN3	LMN-1804935	6	rs7157055	14	83262064		rs2267952	6	133067782	VNN3	6.12	0.73	1.15	1.21
VNN3	LMN-2387680	6	rs2823165	21	5694253		rs2267952	6	133067782	VNN3	4.83	0.46	0.05	0.16
VNN3	LMN-2387680	6	rs9596457	13	51692548		rs2267952	6	133067782	VNN3	5.60	0.53	0.54	0.57
VSTM1	LMN-1763455	19	rs9596457	19	54553697	VSTM1	rs4532100	18	71024750		5.71	0.48	0.17	0.26
VSTM1	LMN-1763455	19	rs10500316	19	54553697	VSTM1	rs7895870	10	123098249		5.71	0.48	0.17	0.26
VSTM1	LMN-1763455	19	rs10500316	19	54553697	VSTM1	rs7895870	10	123098249		5.88	0.81	1.38	1.47
VSTM1	LMN-1763455	19	rs9628570	22	30261219		rs10500316	19	54553697	VSTM1	5.88	0.19	0.13	0.09
WDR48	LMN-1762103	3	rs1388935	3	188927822		rs6778963	3	39091812	WDR48	5.88	0.19	0.13	0.09
WDR48	LMN-1762103	3	rs1887778	9	134635088		rs883349	3	39067925	WDR48	6.34	0.57	1.35	1.22
WDR48	LMN-1762103	3	rs9554833	13	102624790		rs7619193	3	39044116	WDR48	5.85	0.18	0.61	0.35
WDR6	LMN-1669484	3	rs12362253	11	123371708		rs7619193	3	39044116	WDR6	4.86	1.64	1.43	2.25
XAF1	LMN-2330573	17	rs1535031	21	36773170	XAF1	rs11715581	3	49194331		5.86	2.38	0.17	1.63
ZEP00	LMN-1684628	16	rs906446	21	37040648		rs12591171	15	68173945		5.48	0.09	0.36	0.15
ZNF500	LMN-1700238	16	rs4282793	22	48283177		rs1829968	16	93173945	ZFP90	5.79	0.67	0.27	0.46
ZYX	LMN-1701875	7	rs6056281	20	89353122		rs2290560	16	4799041	ZNF500	5.29	0.67	0.27	0.46
ZYX	LMN-1701875	7	rs6056281	20	89353122		rs2242601	7	143093824	ZYX	6.04	0.26	0.01	0.05

^a Phenotypes are expression levels of RefSeq Genes^b Illumina probe ID used to measure gene expression^c Physical SNP position in base pairs (HG19)^d RefSeq Gene ID of gene expression level that is influenced by the SNP (BSGS discovery dataset, significance threshold = 1.29 × 10⁻¹¹)^e Interaction - log₁₀ p-value from discovery dataset^f Interaction - log₁₀ p-value from replication dataset^g Interaction - log₁₀ p-value from meta analysis of replication datasets only^h Distance in Mb between interacting SNPs for *cis-cis* acting SNP pairsⁱ p-values are absent if the interaction did not pass the QC filtering in the replication dataset^j Meta analysis p-values are absent if the interaction did not pass the QC filtering in either replication dataset

Table S2: **Estimation of additive and non-additive variance components from pedigree information** Taken from previous analysis in Powell et al 2013²²

Gene	Probe	Additive		Non-additive	
		Variance	s.e.	Variance	s.e.
NAPRT1	ILMN_1710752	0.37	0.03	0.14	0.05
TMEM149	ILMN_1786426	0.41	0.04	0.09	0.04
MBNL1	ILMN_2313158	0.18	0.03	0.11	0.04
TRAPPC5	ILMN_2372639	0.32	0.04	0.13	0.05
CAST	ILMN_1717234	0.31	0.03	0.10	0.04

Table S3: **Concordance of sign of epistatic variance components between discovery and replication datasets**

Test	Interactions ^a	Dataset	n^b	Expected ^c	Observed ^d	p -value
1 ^e	All	EGCUT	434	217.00	306	6.69×10^{-18}
		Fehrmann	434	217.00	278	5.04×10^{-9}
		Both	434	108.50	221	5.56×10^{-31}
	Significant	EGCUT	30	15.00	25	3.25×10^{-4}
		Fehrmann	30	15.00	24	1.43×10^{-3}
		Both	30	7.50	22	3.76×10^{-8}
2 ^f	All	EGCUT	434	54.25	92	4.22×10^{-7}
		Fehrmann	434	54.25	79	6.18×10^{-4}
		Both	434	6.78	30	2.55×10^{-11}
	Significant	EGCUT	30	3.75	19	9.46×10^{-11}
		Fehrmann	30	3.75	19	9.46×10^{-11}
		Both	30	0.47	18	2.23×10^{-25}
3 ^g	All	EGCUT	1133	566.50	775	7.10×10^{-36}
		Fehrmann	1133	566.50	726	1.90×10^{-21}
		Both	1133	283.25	562	1.39×10^{-70}
	Significant	EGCUT	73	36.50	55	1.69×10^{-5}
		Fehrmann	73	36.50	55	1.69×10^{-5}
		Both	73	18.25	46	7.86×10^{-12}

^a “All” denotes 434 discovery interactions and “Significant” denotes 30 interactions with significant replication p -values

^b Number of tests for concordance

^c Expected number of concordant cases under the null hypothesis of no interactions

^d Observed number of concordant cases

^e The sign of the most significant epistatic variance component in discovery is the same as the corresponding variance component in the replication data.

^f The largest epistatic variance component in the discovery is the same as in the replication with the same sign in both.

^g The sign of all epistatic variance components in the discovery with $p < 0.05$ are the same as the corresponding variance components in the replication data.

Table S4: **Concordance of sign of epistatic variance components between discovery and replication datasets using test 4**

Interactions ^a	Dataset	n^b	0 ^c	1 ^c	2 ^c	3 ^c	4 ^c	p
Expected ^d	-	-	0.06	0.25	0.38	0.25	0.06	-
All	EGCUT	434	0.06	0.22	0.41	0.23	0.08	0.194
All	Fehrman	434	0.07	0.22	0.39	0.24	0.08	0.385
All	Combined	868	0.07	0.22	0.40	0.23	0.08	0.0448
Significant	EGCUT	30	0.07	0.03	0.30	0.33	0.27	4.72×10^{-4}
Significant	Fehrman	30	0.03	0.07	0.33	0.27	0.30	6.69×10^{-4}
Significant	Combined	60	0.05	0.05	0.32	0.30	0.28	5.49×10^{-8}

^a “All” denotes 434 discovery interactions and “Significant” denotes 30 interactions with significant replication p -values.

^b Number of tests for concordance.

^c Proportion of tests that have 0, 1, 2, 3 or 4 concordant signs between discovery and replication.

^d Expected proportion of concordant signs under the null hypothesis of no epistasis.

Table S5: Details on linkage disequilibrium and relative positions of all discovery interactions with SNPs on the same chromosome

Chr	Gene	SNP 1	SNP 2	Position 1	Position 2	Distance / Mb	R^2	D'
19	TMEM149	rs807491	rs7254601	36268923	36147315	0.122	0.000	0.001
17	FN3KRP	rs898095	rs9892064	80890638	80827903	0.063	0.063	0.088
21	CSTB	rs9979356	rs3761385	45230974	45198355	0.033	0.041	0.066
3	MBNL1	rs16864367	rs13079208	152234166	152116652	0.118	0.041	0.117
10	ADK	rs2395095	rs10824092	76446305	75929517	0.517	0.013	0.020
11	CTSC	rs7930237	rs556895	88117962	88077479	0.040	0.012	0.045
17	GAA	rs11150847	rs12602462	78153130	78146016	0.007	0.000	0.001
8	NAPRT1	rs2123758	rs3889129	144663661	144613680	0.050	0.053	0.060
1	LAX1	rs1891432	rs10900520	203877662	203780591	0.097	0.065	0.106
18	MBP	rs8092433	rs4890876	74747424	74732087	0.015	0.035	0.053
11	SNORD14A	rs2634462	rs6486334	17339127	17015557	0.324	0.008	0.012
21	C21ORF57	rs9978658	rs11701361	48027084	47764477	0.263	0.032	0.065
16	RPL13	rs352935	rs2965817	89648580	89513234	0.135	0.054	0.060
19	ATP13A1	rs4284750	rs873870	19810050	19738554	0.071	0.008	0.015
2	NCL	rs7563453	rs4973397	232301670	232291471	0.010	0.027	0.029
5	HNRPH1	rs6894268	rs4700810	179032488	178991794	0.041	0.000	0.001
19	VASP	rs1264226	rs2276470	46063167	45974668	0.088	0.018	0.022
7	TRA2A	rs7776572	rs11770192	23528927	23498358	0.031	0.064	0.064
21	PRMT2	rs2839372	rs11701058	48063862	47776382	0.287	0.100	0.122
12	OAS1	rs13311	rs2072133	113448652	113409260	0.039	0.002	0.016
16	N4BP1	rs12444224	rs11649236	87580855	48632478	38.948	0.007	0.021
5	CAST	rs12719343	rs7733671	125369113	96000269	29.369	0.001	0.001
7	DNAJB6	rs2286842	rs3779589	157216093	157163614	0.052	0.005	0.006
1	OVGP1	rs10802822	rs1264898	240132968	111992823	128.140	0.008	0.030
20	CD93	rs2868504	rs1884655	37771578	23074375	14.697	0.000	0.002
11	PHCA	rs493642	rs10736812	123097386	76708086	46.389	0.002	0.008
21	MX1	rs459498	rs8130120	42795027	29363604	13.431	0.000	0.000
16	AKTIP	rs2896940	rs13332406	57721127	53489705	4.231	0.000	0.001
17	CDK5R1	rs9905940	rs11655031	46614102	30833162	15.781	0.000	0.000
2	CYBRD1	rs888427	rs7591849	172368120	160112881	12.255	0.000	0.000
8	HMBOX1	rs587639	rs7837237	132725731	28876221	103.850	0.001	0.001
11	TRAPPC4	rs1793823	rs3916581	131018917	118887887	12.131	0.001	0.002
12	PEX5	rs10444467	rs4329748	128052636	7364442	120.688	0.000	0.000
12	FLJ20489	rs17615703	rs3782908	117036766	48169526	68.867	0.001	0.002
16	PRKCB1	rs2188355	rs10492793	23867776	12639800	11.228	0.000	0.000
14	MRPL52	rs1950857	rs3811188	26710271	23299135	3.411	0.002	0.004
17	C17ORF60	rs9907897	rs7405659	63502633	59874129	3.629	0.004	0.011
6	FLJ43093	rs6906101	rs13214069	36667610	32705248	3.962	0.000	0.000
19	TRAPPC5	rs17159840	rs17763599	7758194	2369415	5.389	0.000	0.000
22	PISD	rs715572	rs6518754	33234931	32097775	1.137	0.001	0.003
12	DIP2B	rs871257	rs12427378	117994348	51074199	66.920	0.001	0.001
12	GPR162	rs2272500	rs2707210	79685913	6902002	72.784	0.003	0.005
17	USP36	rs2279308	rs7225546	76794981	75151717	1.643	0.000	0.000