

Detection and replication of epistasis influencing 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 epistasis has been demonstrated in artificial gene manipulation studies in model organisms,^{4,5} and examples have been reported in other species,⁶ few convincing examples with independent replication exist for epistasis amongst natural polymorphisms in human traits.^{7,8} Its absence from empirical findings may simply be due to its 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 power and computational issues.⁹ Here we show that, using advanced computation techniques¹⁰ 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 data on 7339 gene expression levels in peripheral blood, we found 501 significant pairwise epistatic interactions between common SNPs acting on the expression levels of 238 genes ($p < 2.91 \times 10^{-16}$). Replication of these signals 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 $p < 0.05/434$. There was evidence of functional enrichment for the interacting SNPs, for instance 44 of the genetic interactions are located within 5Mb 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 natural genetic 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 re-

duced 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, when combined 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 occur amongst 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. These traits are largely heritable but on average have been shown to be less polygenic than high level phenotypes,²¹ thus many genetic effects are relatively large,²² maximising the chance at detecting epistasis, should it exist.

In our discovery dataset (Brisbane Systems Genetics Study, BSGS²²) 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 (5% significance threshold $p < 2.91 \times 10^{-16}$, 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 $\alpha \approx 0.07$, Methods, Supplementary Figure S1). Using this design we identified 501 putative genetic interactions influencing the expression levels of 238 genes (Supplementary Table S1). Of the 501 discovery interactions, 434 had available data and passed filtering (Methods) in two independent replication datasets, Fehrman¹² 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 $p < 0.05/434$, 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 ($A1$, $A2$, $D1$, $D2$, $A \times A$, $A \times D$, $D \times A$, $D \times D$) 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 interaction p -values more extreme than the 2.5% confidence interval of the quantile-quantile plot against the null hypothesis of no interactions ($p < 1.0 \times 10^{-16}$, Figure 2 and Supplementary Figure S2). Concordance of the

direction of the effect of the largest variance component was also highly significant ($p = 5.71 \times 10^{-31}$, Supplementary Table S3). The congruence of the epistatic networks in discovery and replication datasets is shown in Figure 3, demonstrating that these complex genetic patterns are common even across independent datasets. A further replication was attempted using the Centre for Health Discovery and Wellbeing (CHDWB) dataset,²³ but only 20 of the SNP pairs passed filtering because the sample size was small ($n = 139$), and likely due to insufficient power we found no evidence for replication (Supplementary Figure S6).

It should be noted that although it is a necessary step to establish the veracity of the signals from the discovery set, replication of epistasis is difficult in practice for several reasons. For example, LD between causal variants and observed markers plays an important role. Not only is the dependence on LD much greater for epistatic effects than for additive effects (Supplementary Figure S7), but when estimating epistatic variance it is more sensitive to changes in LD between observed SNPs and causal variants between independent samples when compared to additive effects (Supplementary Figure S8), and this has a direct effect on statistical power for replication. Indeed, in addition to heterogeneity between populations, sampling variance of LD r alone leads to over three fold decrease in observed variance for the most extreme case of $D \times D$ effects when compared to the average decrease for additive variance on average (Methods, Supplementary Figure S9). Therefore these results are encouraging with regards to the detection and replication of epistasis.

Though seldom the focus of association studies, SNPs with known main effects are often tested for additive \times additive genetic interactions,⁹ but our analysis shows that this is unlikely to be the most effective strategy for its detection. The majority of our discovery interactions comprised of one SNP that was significantly associated with the gene expression level in the discovery dataset, and one SNP that had no previous association²¹ (439 out of 501, Methods). Only nine interactions were between SNPs that both had known main effects while 64 were between SNPs that had no known main effects. Additionally, we observed that the largest epistatic variance component for the 501 interactions was equally divided amongst additive \times additive, additive \times dominance, dominance \times additive and dominance \times dominance at the discovery stage ($p = 0.22$ for departure from expectation). This is not surprising because the patterns of epistasis used for statistical decomposition (*i.e.* $A \times A$, $A \times D$, $D \times A$, $D \times D$) are simply convenient orthogonal parameterisations of a two locus model, and are not intended to model biological function.²⁴

Of the discovery interactions, 47 were *cis-cis* acting (both SNPs were on the same chromosome as the expression gene, median distance between interacting SNPs is 1.83Mb), 441 were *cis-trans*-acting, and 13 were *trans-trans*-acting. We observed a wide range of significant GP maps (Figure 1) but the most common pattern of epistasis that we detected involved a *trans*-SNP masking the effect of an additive *cis*-SNP. For example, MBNL1 (involved in RNA modification and regulation of splicing²⁵) has a *cis* effect at rs13069559 which in turn is controlled by 13 *trans*-SNPs and one *cis*-SNP that each exhibit a masking pattern, such

that when the *trans*-SNP is homozygous for the masking allele the decreasing allele of the *cis*-SNP no longer has an effect (Supplementary Figure S10). Each of these interactions has evidence for replication in at least one dataset and six are significantly replicated at the Bonferroni level (Supplementary Figure S3). We see similar epistatic networks involving multiple (eight or more) *trans*-acting SNPs for other gene expression levels too, for example TMEM149 (Supplementary Figure S11), NAPRT1 (Supplementary Figure S12), TRAPPC5 (Supplementary Figure S13), and CAST (Supplementary Figure S14). We observed that from pedigree analysis 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 previously²¹ (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 types²⁶ (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 states²⁷ (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 lines²⁸ (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.^{29,30}

Though we present many instances of epistasis, quantifying its relative importance to complex traits in humans remains an open question. In this study we are able to identify 238 gene expression traits with at least one significant interaction given our experiment-wide threshold, where the minimum variance explained by the epistatic effects of any interaction was 2.1% of phenotypic variance. Taking published results from a standard eQTL study²² we calculated that 1848 of the 7339 gene expression levels analysed were influenced by additive effects where the 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 Ideally

one would approach this question from a whole genome perspective³¹ but this is intractable for non-additive variance components. Nevertheless, some inference can be made from the ascertained effects in these analyses and it is evident that additive variance is overall a larger component than epistatic variance, as has been argued previously.^{2,3} Taking the additive effects detected in Powell *et al* (2012) where the minimum additive variance was 2.1%, 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 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 additive variance. There are several caveats to this comparison. Firstly, the ratio of additive to epistatic variance may differ at different minimum variance thresholds, and our estimate is determined by the threshold used. Secondly, the power of a 1 *d.f.* test exceeds that of an 8 *d.f.* test. And thirdly, the non-additive variance at causal variants is expected to be underestimated by observed SNPs in comparison to estimates for additive variance, due to differences in the rate of decay of the estimate of the genetic variance of the causal SNPs as LD decreases with the observed SNPs.

Overall, we have demonstrated that it is possible to identify and replicate epistasis in complex traits amongst common human variants, despite the relative contribution of pairwise epistasis to phenotypic variation being small. The bioinformatic analysis of the significant epistatic loci suggests that there are a large number of possible mechanisms that can lead to non-additive genetic variation. Further research into such epistatic effects may provide a useful framework to understanding molecular mechanisms and complex trait variation in greater detail. With computational techniques and data now widely available the search for epistasis in larger datasets for traits of broader interest is warranted.

Methods Summary

We searched for pairwise epistasis exhaustively in the BSGS discovery dataset,²² which comprises 846 individuals who are genotyped at 528,509 autosomal SNPs. Each individual had gene expression levels measured in peripheral blood at 47,323 probes. Only the probes that passed quality control and had significant expression in $\geq 90\%$ of individuals were used in the analysis (7,339 probes representing 6,158 RefSeq genes). Recent hardware and software¹⁰ advances that use graphics processing units (GPUs) made it possible to perform the 1.03×10^{15} statistical tests to complete this analysis. We used permutation analysis³² to calculate an experiment-wide significance threshold of $T_e = 2.91 \times 10^{-16}$ at the 5% family-wise error rate (FWER). SNP pairs were modelled for full genetic effects, including marginal additive and dominance at both SNPs plus four interaction terms. Though we could have used a less complex model to improve statistical efficiency, we deemed it important to be agnostic about the type of epistasis that might exist, and therefore chose not to over-parameterise the test.^{18,19} Because there are many large marginal effects present in these data it was necessary to perform several filtering steps to exclude SNP pairs that were

significant due to marginal effects alone. All SNP pairs with LD $r^2 > 0.1$ and $D'^2 > 0.1$ were removed to minimise the possibility of haplotype effects. All SNP pairs were required to have at least five data points in all nine genotype classes. If multiple SNP pairs were present on the same chromosomes for a particular expression trait then only the sentinel SNP pair was retained. Finally, a nested test contrasting the full genetic model against the marginal additive and dominance model was performed for each remaining SNP pair (Methods), resulting in 501 significant interactions after Bonferroni correction for multiple testing of the filtered SNPs. The significant SNP pairs were carried forward for replication in two independent datasets that used the same expression assays for analysing transcription in peripheral blood, the Fehrmann dataset¹² ($n = 1240$) and the Estonian Genome Centre University of the University of Tartu (EGCUT) dataset¹¹ ($n = 891$). Of these, 434 passed filtering in both replication datasets. A meta analysis on the interaction p -values from each replication dataset was performed to provide an overall replication statistic for each putative interaction.

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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

Figures

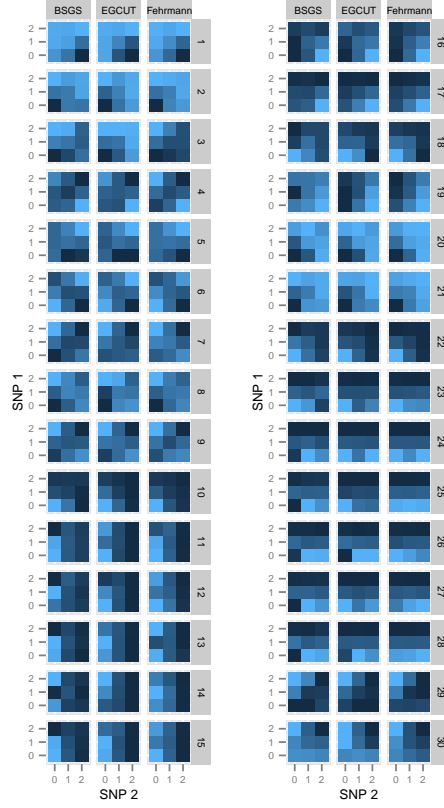


Figure 1: Replication of GP maps in two independent populations
The GP maps for each epistatic interaction that is significant at the Bonferroni level in both replication datasets are shown. Each GP map consists of nine tiles where each tile represents the expression level for that two-locus genotype 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.

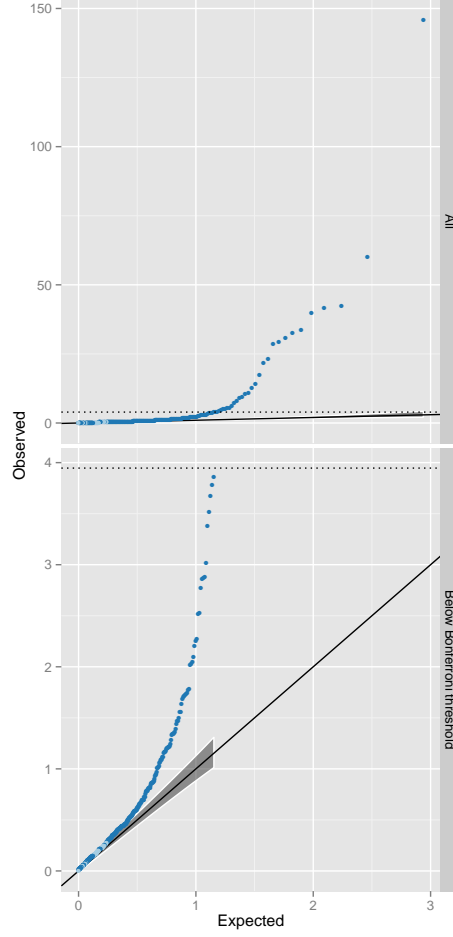


Figure 2: **Q-Q plots of interaction p -values from replication datasets** The top panel shows all 434 discovery SNPs that were tested for interactions. Observed p -values (y -axis, $-\log_{10}$ scale) are plotted against the expected p -values (x -axis, $-\log_{10}$ scale). The multiple testing correction 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.

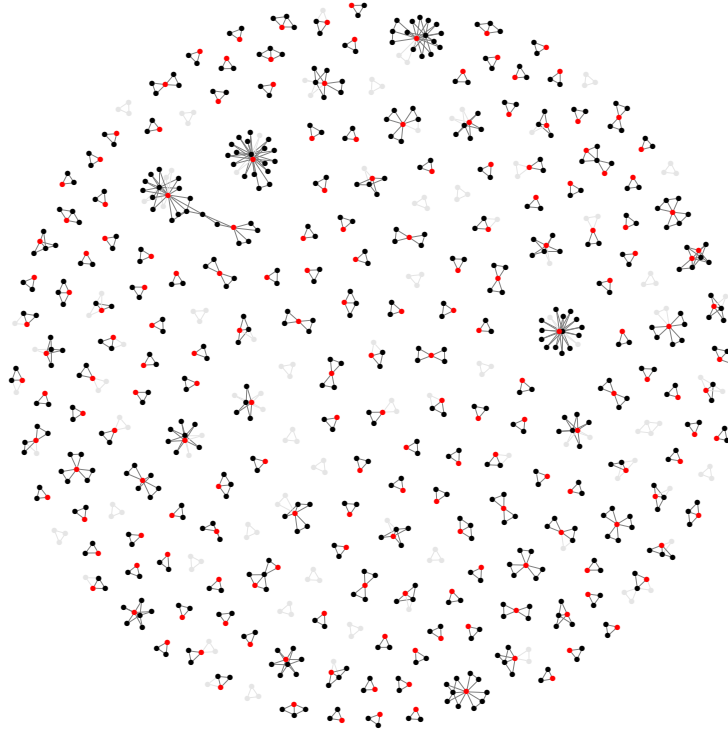


Figure 3: **Discovery and replication of epistatic networks** All 434 putative genetic interactions (edges) with data common to discovery and replication sets is shown, where black nodes represent SNPs and red nodes represent traits (gene expression probes). Three hundred and forty-five interactions 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.

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Supplementary Figures

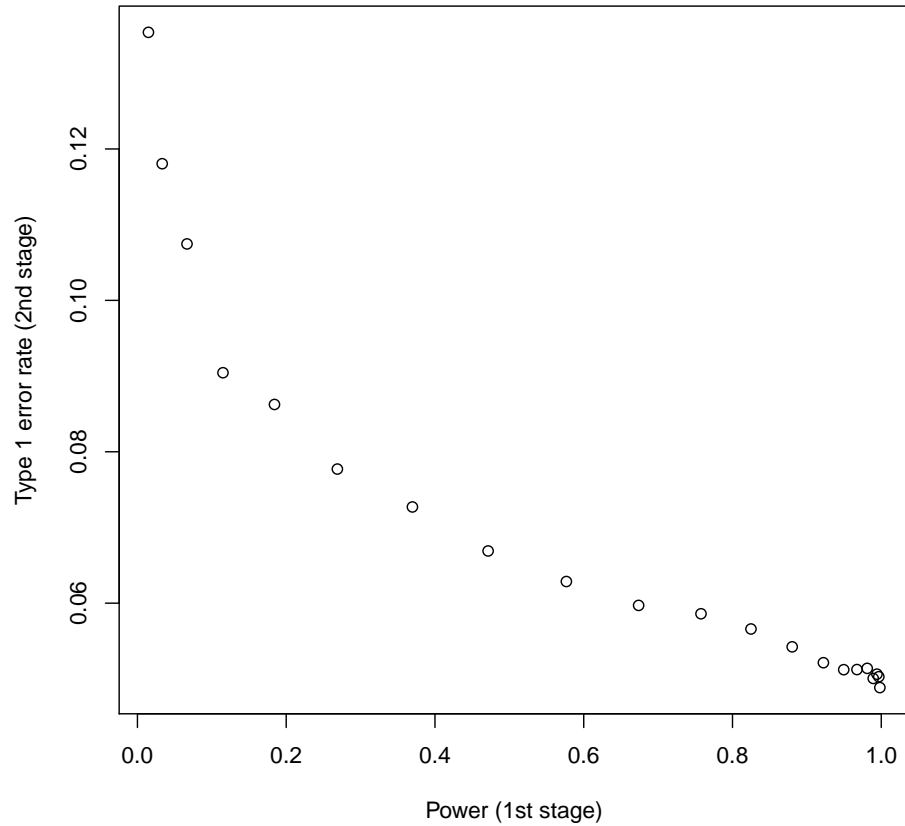


Figure S1: **Type 1 error rate of two stage design** In stage 1 SNPs are tested for full genetic effects (8 d.f.) and those that surpass a threshold for multiple testing are then tested for significant interaction terms in stage 2. These interaction p -values are then adjusted (Bonferroni) for the total 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.

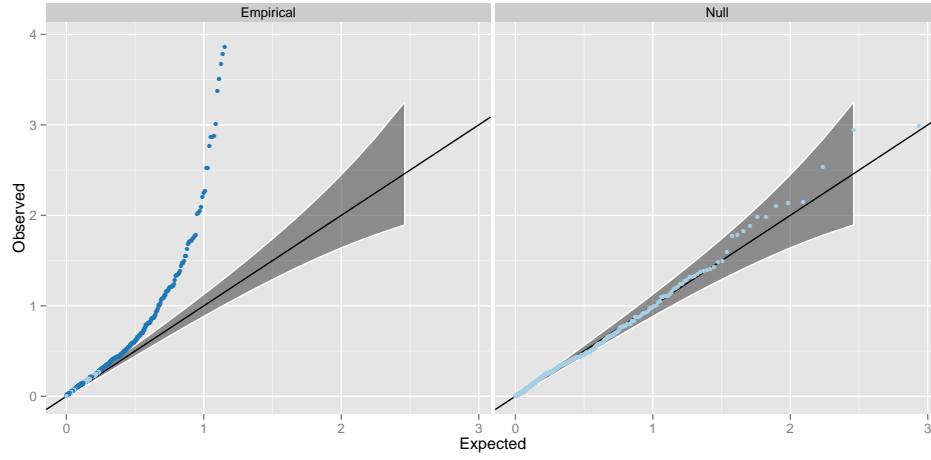


Figure S2: **Q-Q plots of interaction p -values from replication datasets, excluding the 30 points significant at the Bonferroni level** The right panel (Null) shows the interaction p -values from a meta analysis across two independent datasets on 434 randomly drawn SNP pairs. The left panel (Empirical) 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.

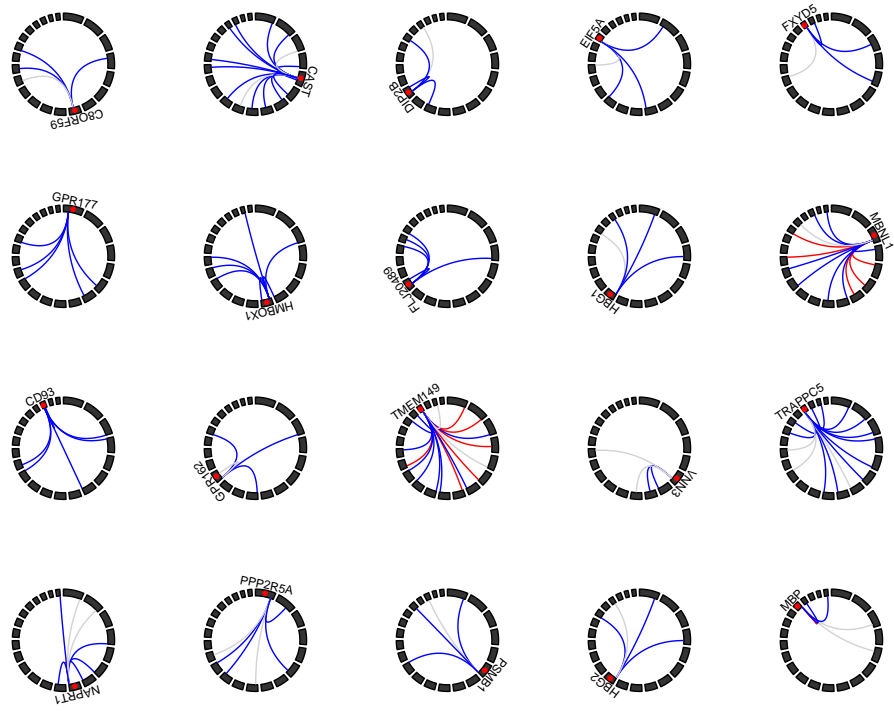


Figure S3: **Gene expression traits with four or more genetic interactions** Circle plots represent the genomic positions for SNPs (linking lines) and expression probes (red points). Chromosomes are represented by black blocks and ordered from 1 to 22 clockwise, starting from the top. Grey lines represent 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.

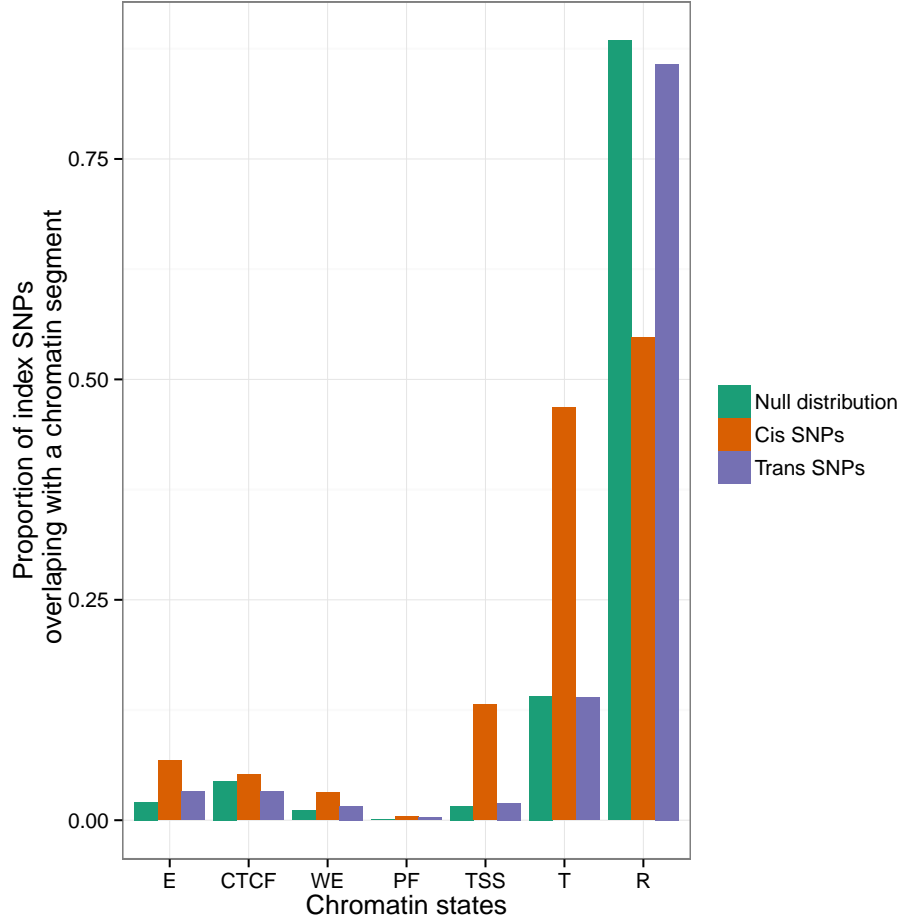


Figure S4: **Location of SNPs relative to genomic features** We used chromatin segmentation²⁷ as a method for labelling genomic features. All SNPs within 1Mb and $r^2 > 0.8$ of each *cis*- and *trans*-SNP were taken to find which genomic features (x -axis) were covered by the SNPs that compose the 501 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

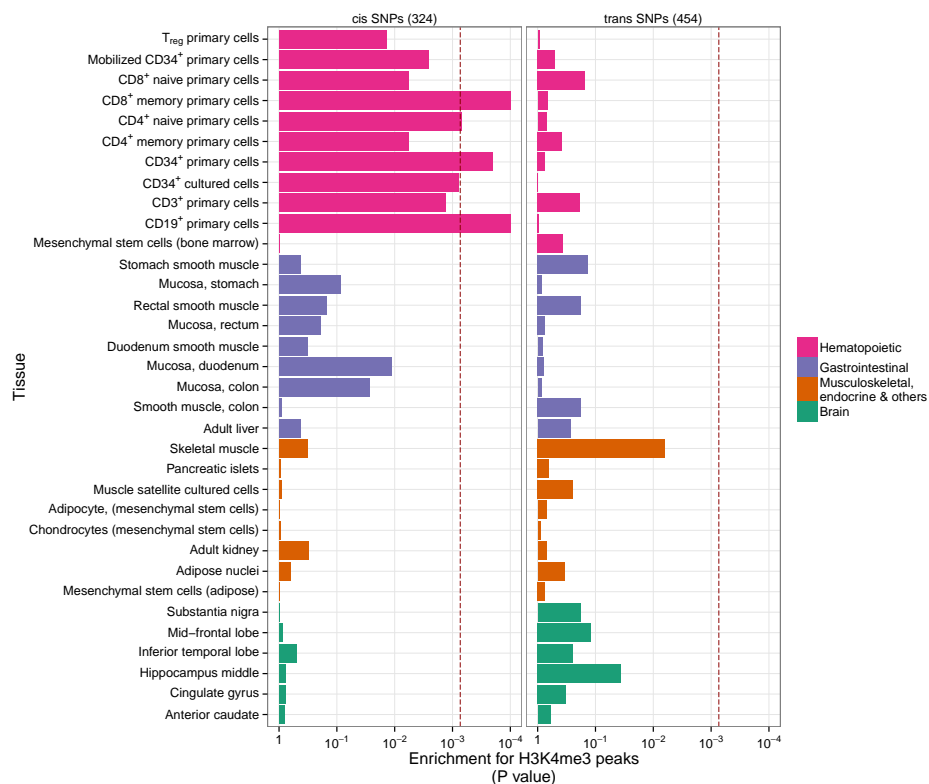


Figure S5: Tissue specific enrichment of SNPs in transcriptionally active regions The locations of transcriptional activity can be predicted by chromatin marks, assayed by H3K4me3.²⁶ Enrichment p -values are calculated using permutation analysis for 34 different cell types (y -axis) in four tissue types (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.

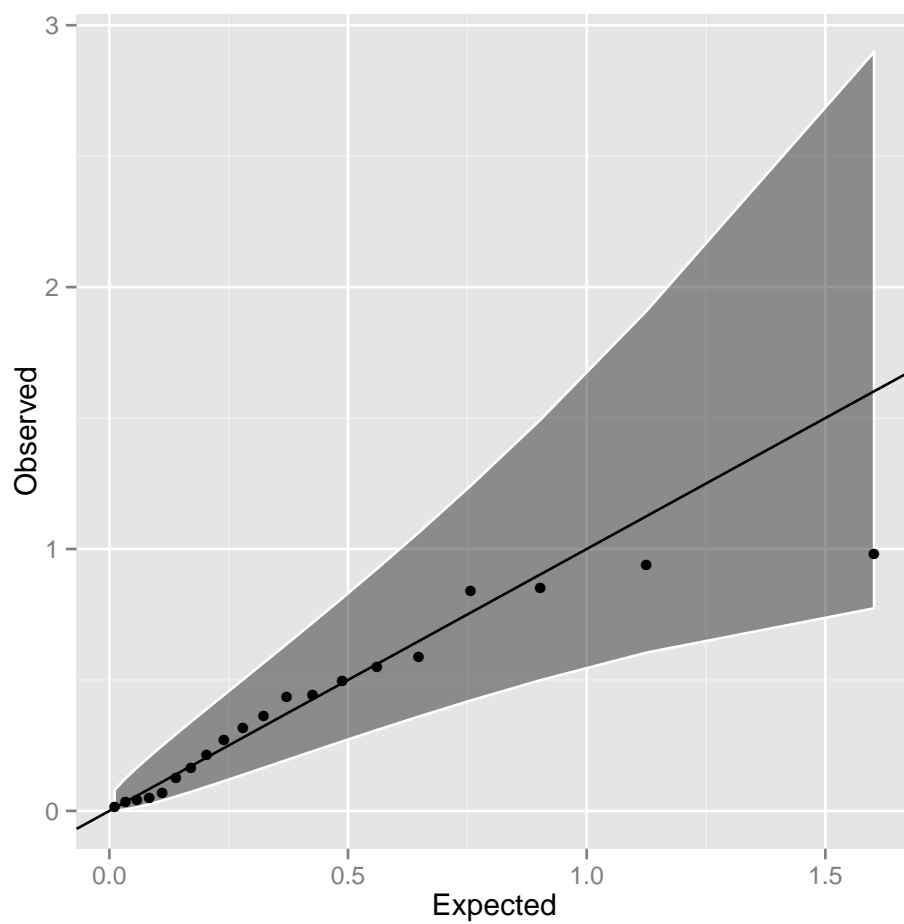


Figure S6: **Q-Q plot of interaction p -values in the CDHWB dataset**
 Twenty of the 501 discovery SNP pairs passed filtering in the CDHWB dataset (mainly due to small sample size). There is no evidence for enrichment of interaction terms, most likely due to insufficient power given the limited sample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

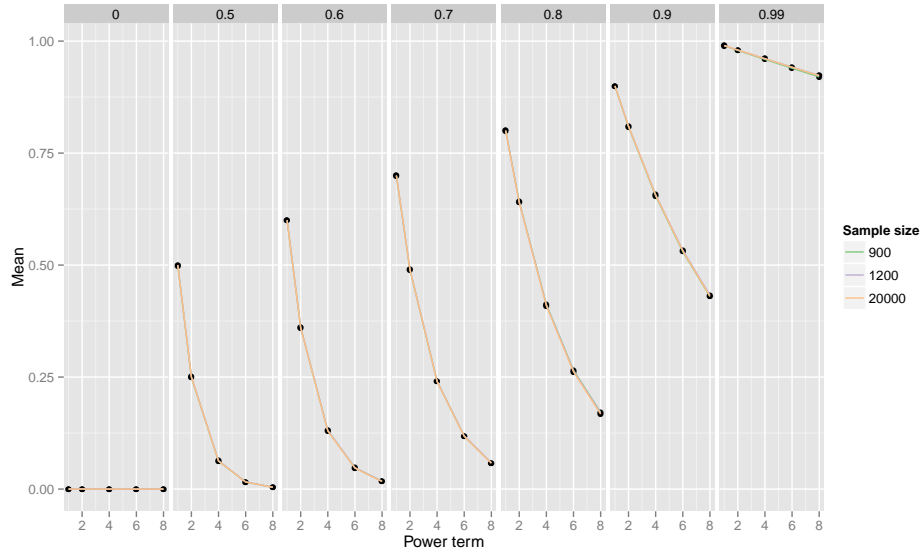


Figure S7: **Sampling mean for different power terms of population r values** Power of detection and replication of epistatic signals depends not on r^2 between causal variants and observed SNPs, but on r^4, r^6, r^8 . For a given a population value of LD r (columns of plots), plotted is the sample mean (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 .

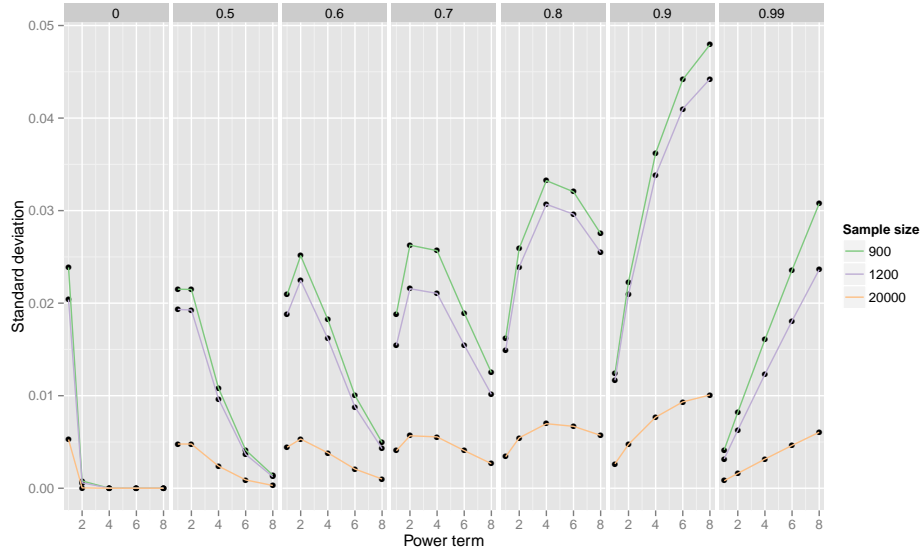


Figure S8: Sampling standard deviation for different power terms of population r values Power of detection and replication of epistatic signals depends not on r^2 between causal variants and observed SNPs, but on r^4, r^6, r^8 . For a given a population value of LD r (columns of plots), plotted is the 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.

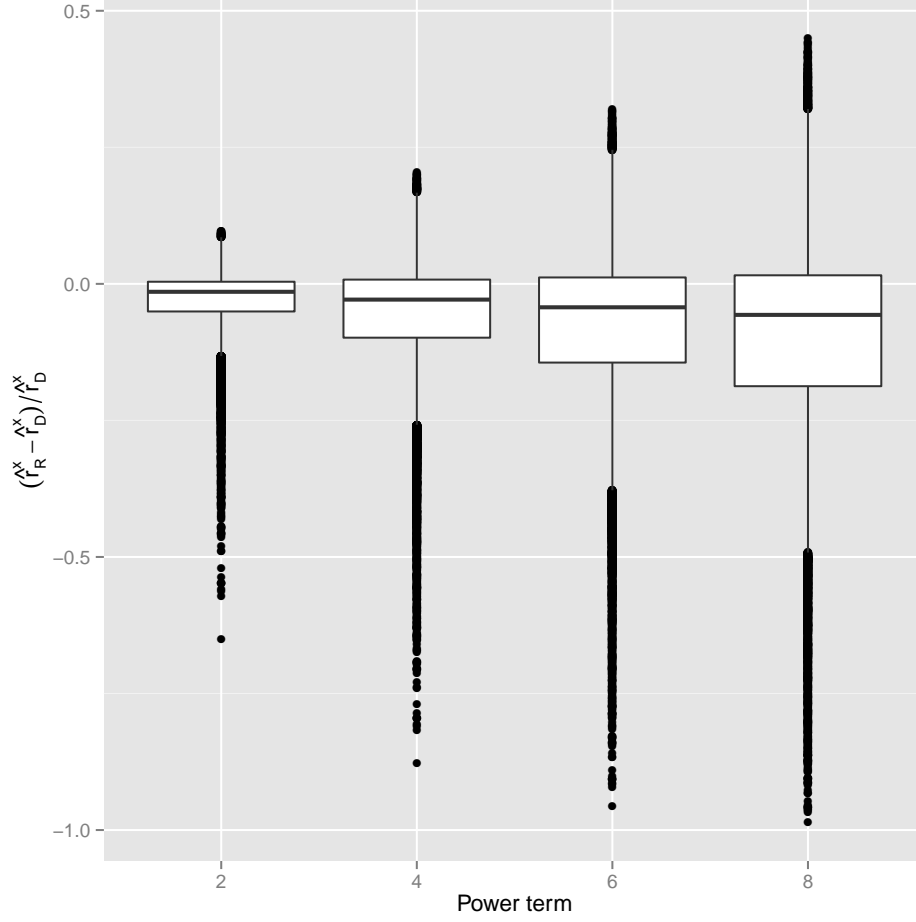


Figure S9: **Reduction in LD as estimated in replication data after ascertaining for high LD in discovery data** 100,000 “unobserved” causal variants (CVs) were tested for LD against a panel of 528,509 “observed” discovery markers (DMs). DM/CV pairs with LD $r^2 > 0.9$ were then tested in an independent 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.

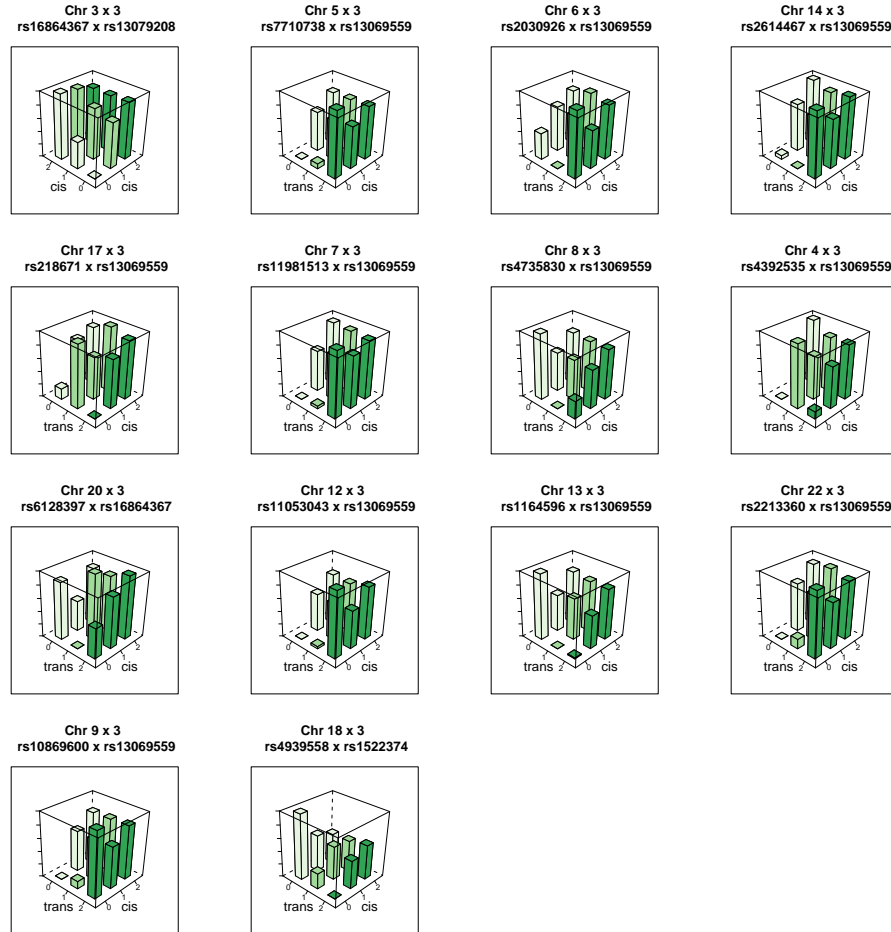


Figure S10: **Genotype-phenotype maps for 14 interactions influencing the expression of MBNL1** Each bar represents the mean phenotypic value for individuals in that genotype class. The rs13069559 SNP typically has a *cis*-additive decreasing effect on the expression of MBNL1, but in many of these interactions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

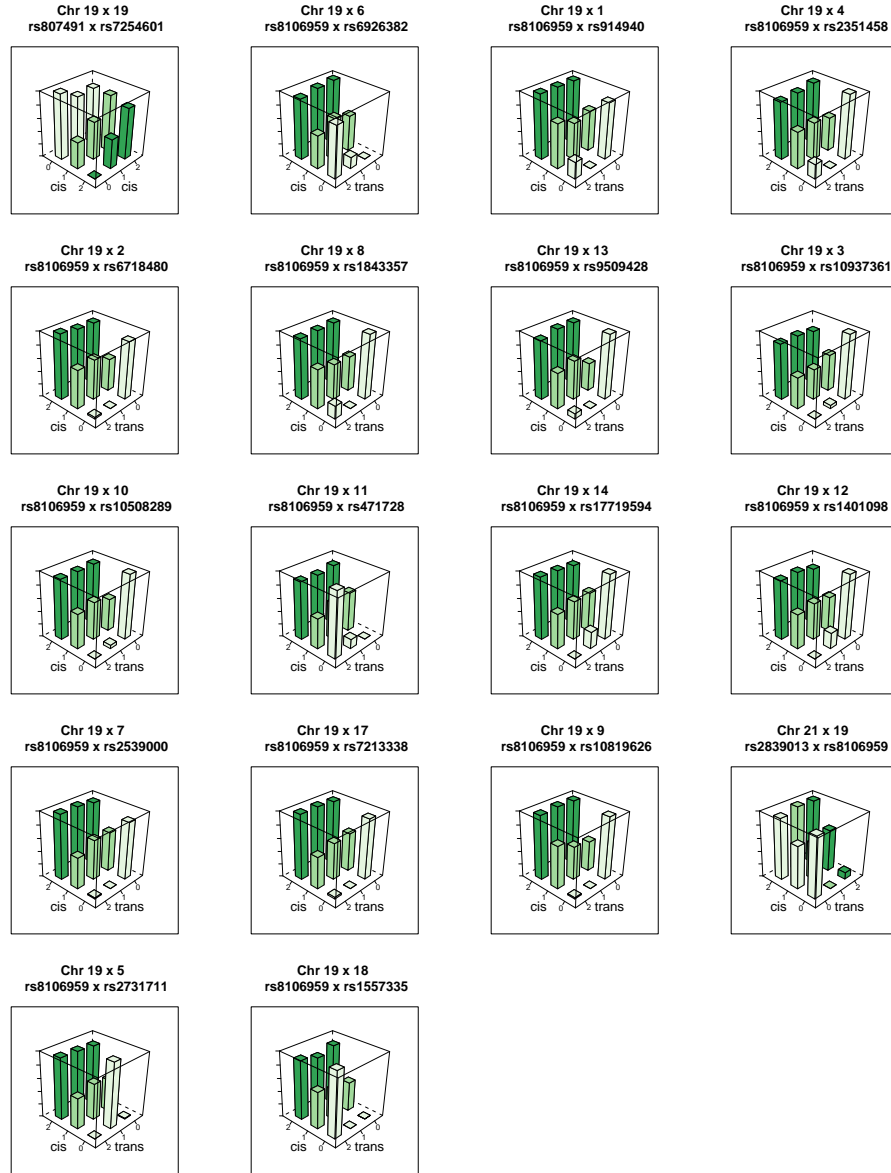


Figure S11: **Genotype-phenotype maps for 19 interactions influencing the expression of TMEM149** Each bar represents the mean phenotypic value for individuals in that genotype class. The rs13069559 SNP typically has a *cis*-additive decreasing effect on the expression of TMEM149, but in many of these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

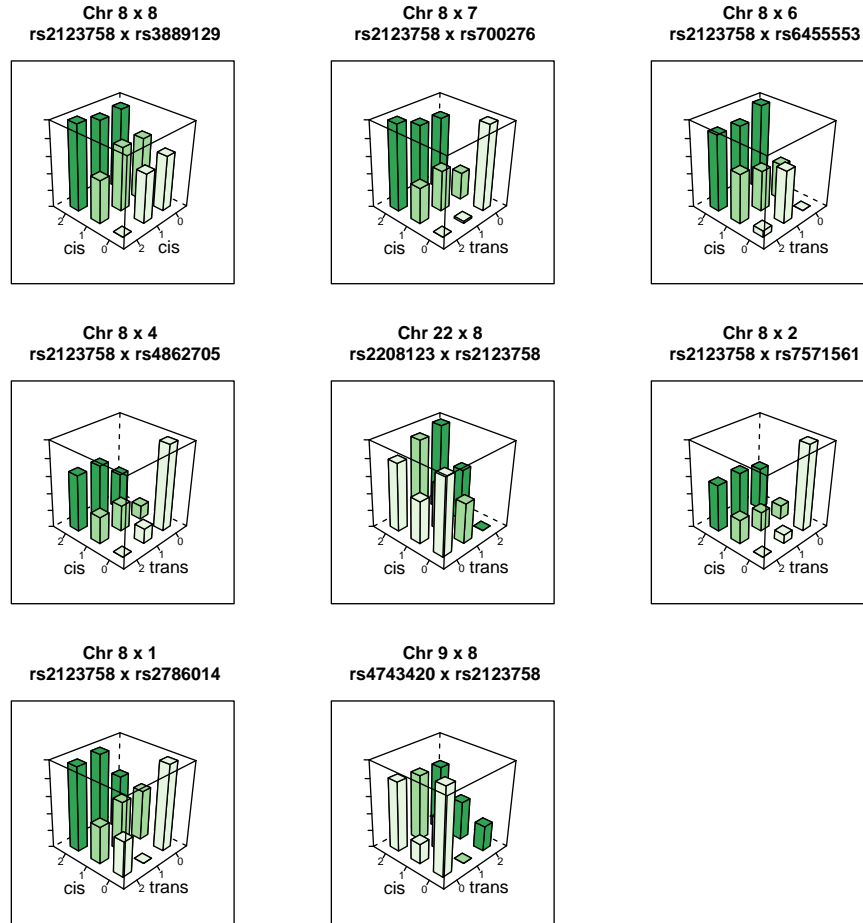


Figure S12: **Genotype-phenotype maps for 8 interactions influencing the expression of NAPRT1** Each bar represents the mean phenotypic value for individuals in that genotype class.

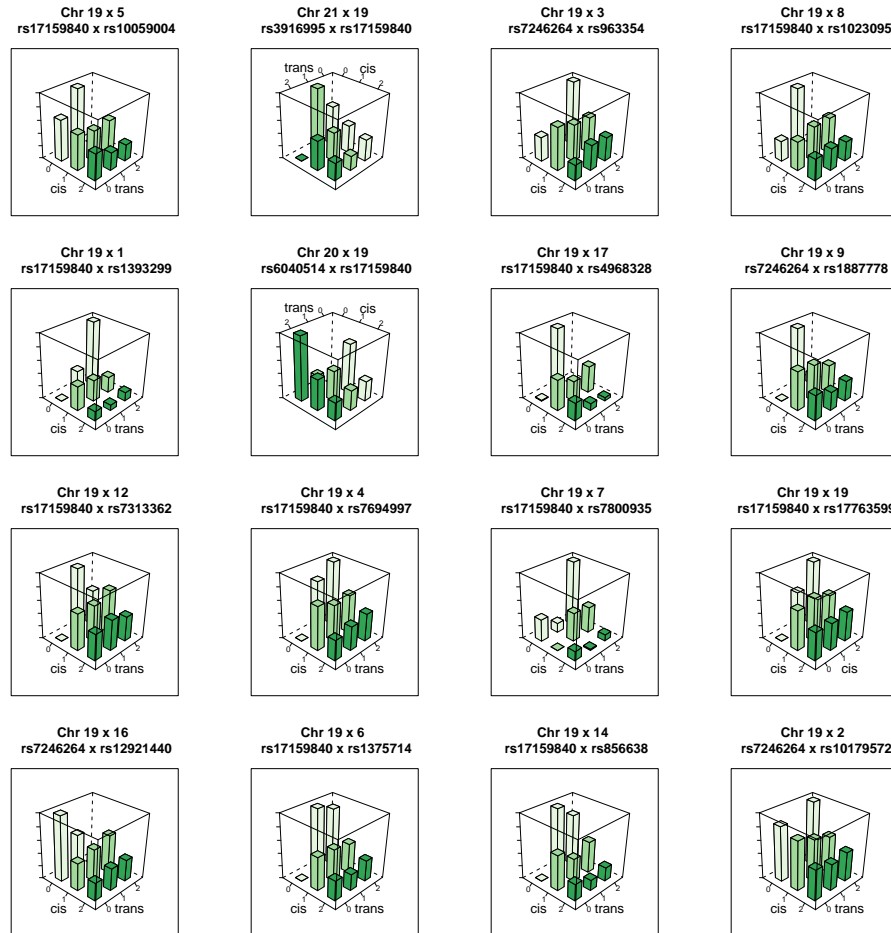


Figure S13: **Genotype-phenotype maps for 16 interactions influencing the expression of TRAPPC5** Each bar represents the mean phenotypic value for individuals in that genotype class.

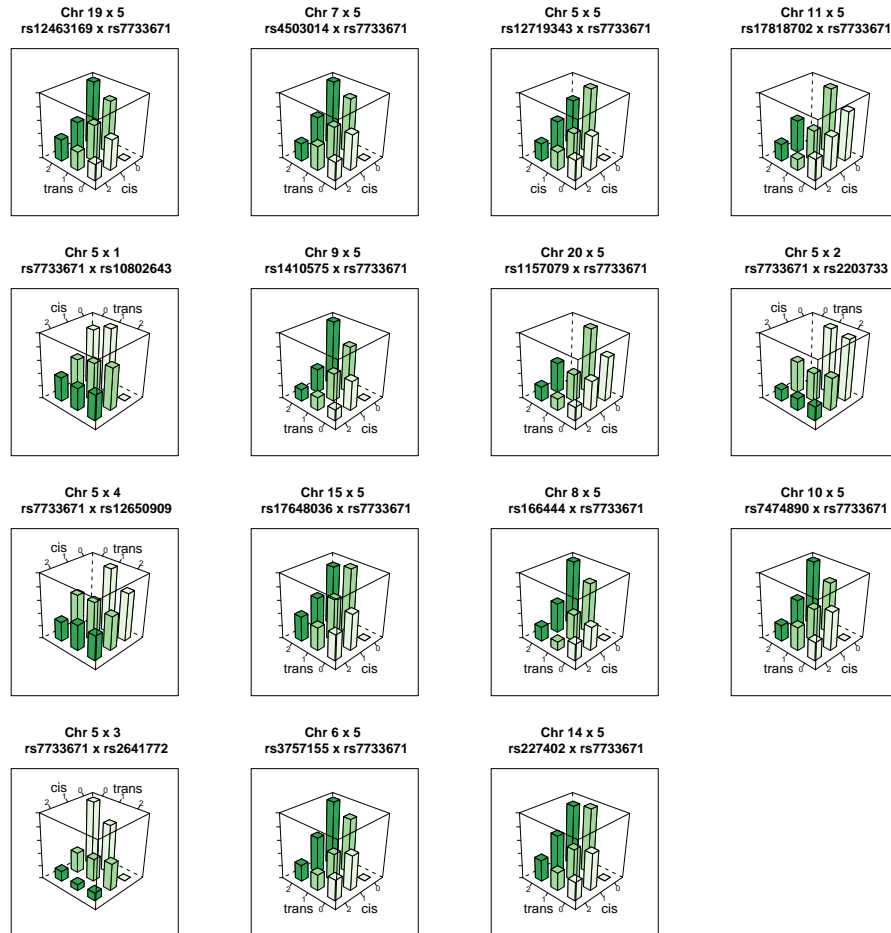


Figure S14: **Genotype-phenotype maps for 15 interactions influencing the expression of CAST** Each bar represents the mean phenotypic value for individuals in that genotype class.

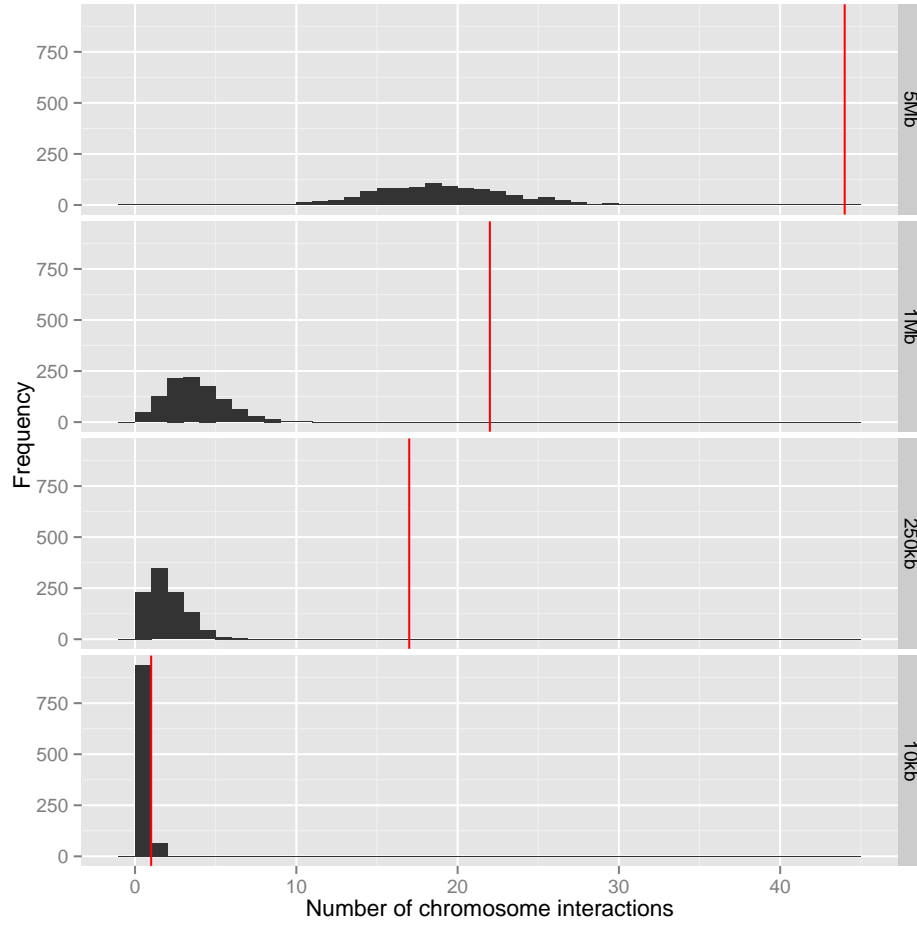


Figure S15: Number of overlaps between chromosome interactions and epistatic interactions Interacting chromosome regions may be a possible mechanism underlying epistatic interactions. The number of epistatic interactions within 20kb, 500kb, 2Mb and 10Mb of known chromosome interacting regions are shown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

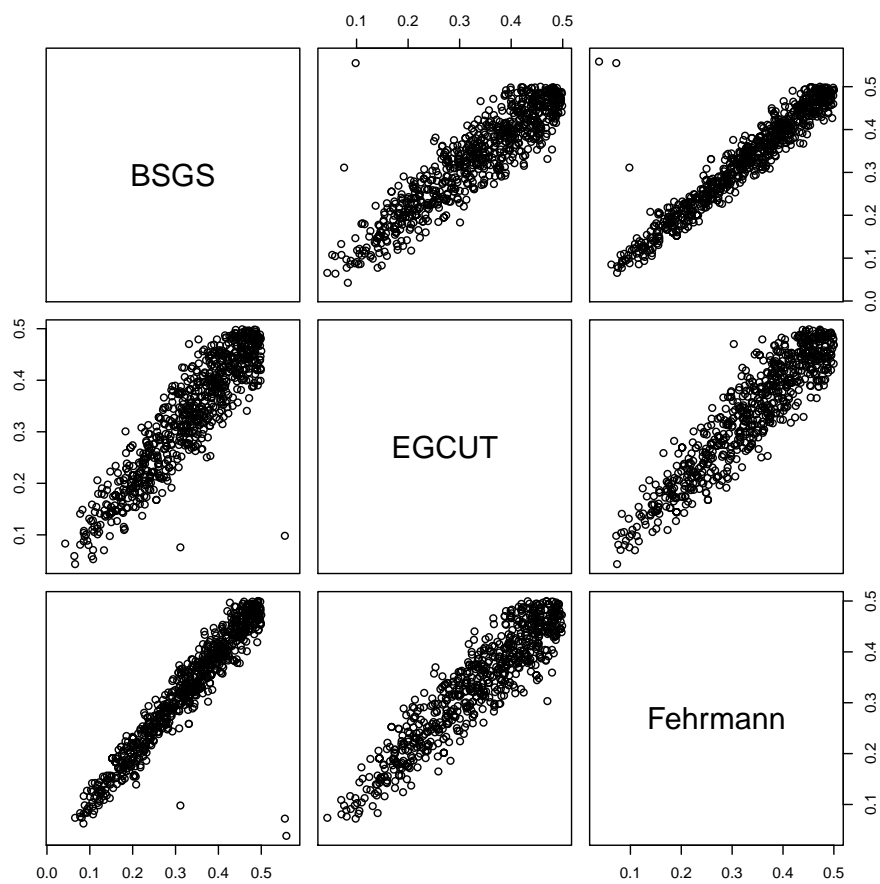


Figure S16: **Comparison of allele frequencies for 781 SNPs involved in genetic interactions across independent populations** Outliers were removed from the analysis as part of the filtering stage during replication.

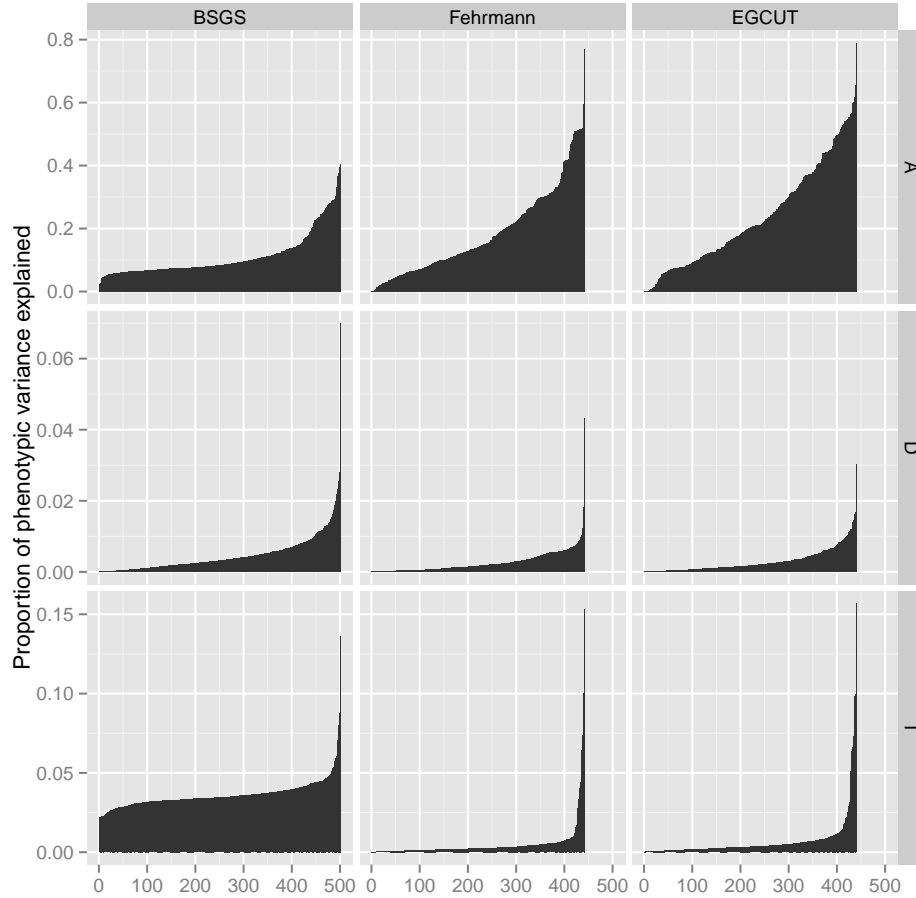


Figure S17: Comparison of variance explained by additive, dominant and epistatic effects from different cohorts How does the estimated variance decomposition change in different cohorts? The proportion of the phenotypic variance that is additive (A), dominant (D), or epistatic (I) for each putative interaction 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.

Supplementary Tables

Table S1 – continued from previous page

Expression trait			SNP 1			SNP 2			Interaction statistic / -log ₁₀ p-values			Distance / Mb		
Gene ID ^a	Probe ID ^b	Chr.	rs ID	Chr.	Pos / Mb ^c	Association ^d	rs ID	Chr.	Pos / Mb ^c	Association ^d	BSGS ^e		Fehrmann ^f	EGCUT ^g
CBORF59	ILMN_1633205	8	rs8051751	16	7188323		rs2896452	8	86102223	CBORF59	5.79	1.39	0.18	0.87
CBORF72	ILMN_1741881	9	rs10122902	9	27556780	C9ORF72	rs2526698	9	242029101		6.36	0.96	0.01	0.37
CAC1	ILMN_1731064	1	rs12765847	10	4353908		rs3738725	1	221714210	CAC1	6.36	0.94	0.00	0.34
CARD9	ILMN_1712532	9	rs4260763	9	139289825	INPP5E	rs684040	1	82128660		5.81			
CAST	ILMN_1717234	5	rs4573661	11	6026661		rs4077515	9	139266496	INPP5E	6.61	0.09	0.86	0.42
CAST	ILMN_1717234	5	rs11540769	20	6778978		rs7733671	5	96000269	CAST	7.07	0.23	0.96	0.62
CAST	ILMN_1717234	5	rs12463169	19	17321669		rs7733671	5	96000269	CAST	5.73	0.02	2.85	1.75
CAST	ILMN_1717234	5	rs12599264	16	81840122		rs7733671	5	96000269	CAST	7.00			
CAST	ILMN_1717234	5	rs12719343	5	125369113		rs7733671	5	96000269	CAST	7.68	0.36	1.57	29.369
CAST	ILMN_1717234	5	rs1410575	9	78255630		rs7733671	5	96000269	CAST	6.55	0.13	1.34	0.78
CAST	ILMN_1717234	5	rs166444	8	78392770		rs7733671	5	96000269	CAST	7.01	0.27	0.52	0.37
CAST	ILMN_1717234	5	rs17648036	15	27311111		rs7733671	5	96000269	CAST	7.81	0.97	0.03	0.41
CAST	ILMN_1717234	5	rs17818702	11	86107920		rs7733671	5	96000269	CAST	6.62	1.15	0.59	1.09
CAST	ILMN_1717234	5	rs227402	14	70496867		rs7733671	5	96000269	CAST	6.12	0.11	0.01	0.01
CAST	ILMN_1717234	5	rs2822124	21	15166804		rs7733671	5	96000269	CAST	6.87			
CAST	ILMN_1717234	5	rs3757155	6	136458593		rs7733671	5	96000269	CAST	7.24	0.07	0.33	0.12
CAST	ILMN_1717234	5	rs4503014	7	31149140		rs7733671	5	96000269	CAST	5.88	0.92	1.56	1.72
CAST	ILMN_1717234	5	rs7474890	10	59590078		rs7733671	5	96000269	CAST	6.74	0.49	0.12	0.23
CAST	ILMN_1717234	5	rs7733671	5	96000269	CAST	rs10802643	1	238120177		7.42	0.75	0.78	0.93
CAST	ILMN_1717234	5	rs7733671	5	96000269	CAST	rs12650909	2	170129890		7.42	0.23	0.87	0.54
CAST	ILMN_1717234	5	rs7733671	5	96000269	CAST	rs2203733	2	224093101		6.07	0.22	0.78	0.50
CAST	ILMN_1717234	5	rs7733671	5	96000269	CAST	rs2641772	3	195531841		6.93	0.19	0.26	0.15
CAT	ILMN_1651705	11	rs872311	18	66175386		rs11032695	11	34447586	CAT	6.41	0.26	0.30	0.22
CCDC88B	ILMN_1722208	11	rs2352303	19	17099980		rs541207	11	64125142	CCDC88B	5.68	0.33	0.37	0.31
CCDC88B	ILMN_1772208	11	rs694739	16	64097233	CCDC88B	rs12771349	10	96989193		5.62	0.23	0.18	0.14
CD36	ILMN_1784863	7	rs3211834	11	76033374		rs1254900	2	85816334	CD36	6.93	0.15	0.01	0.02
CD55	ILMN_1800540	1	rs750801	11	80283117		rs10255470	7	207502534	CD55	5.09	0.03	0.03	0.02
CD93	ILMN_1704730	20	rs1884655	20	23074375	CD93	rs6700168	1	207502534	VAMP8	6.06	1.74	0.24	1.20
CD93	ILMN_1704730	20	rs1884655	20	23074375	CD93	rs4696726	4	7992632		5.71	0.13	0.80	0.42
CD93	ILMN_1704730	20	rs1884655	20	23074375	CD93	rs7622580	3	196721395		5.56	0.04	0.27	0.08
CD93	ILMN_1704730	20	rs1884655	20	23074375	CD93	rs838875	12	125145394		6.31	0.24	1.67	1.16
CD93	ILMN_1704730	20	rs1884655	20	23074375	CD93	rs9576388	13	38434472		7.88	0.71	0.22	0.45
CD93	ILMN_1704730	20	rs2868504	20	37771578		rs1884655	20	23074375	CD93	5.71	0.64	0.75	0.81
CD93	ILMN_1704730	20	rs4813479	20	23076914	CD93	rs10925747	1	238899903		7.43			
CD93	ILMN_1704730	20	rs4813479	20	23076914	CD93	rs2873420	8	136500554		7.02			
CD93	ILMN_1704730	20	rs4813479	20	23076914	CD93	rs4295531	18	74439542		6.13			
CD93	ILMN_1704730	20	rs4813479	20	23076914	CD93	rs7489981	17	77264482		6.08			
CD93	ILMN_1704730	20	rs861544	14	104162263		rs7324744	13	115008038	CD93	5.46	0.21	0.14	0.11
CDK5R1	ILMN_23309796	13	rs9059590	17	46614102	HOXB2	rs11655031	17	30833162	CDK5R1	5.47	0.95	0.07	0.45
CEACAM21	ILMN_1745949	19	rs200690	20	51956250		rs4803481	19	42066556	CEACAM21	6.15	0.90	0.12	0.48
CEACAM21	ILMN_1745949	19	rs4803481	19	42066556		rs2421050	5	158943044	CEACAM21	6.67	2.16	0.16	1.44
CEACAM21	ILMN_1703754	18	rs6505780	18	33699782	CEACAM21	rs13132719	4	180265266		5.75	0.15	0.24	0.12
CEP192	ILMN_1787808	3	rs3825569	14	101350298	CEP192	rs13079012	3	134247706	ANAPC13	6.36	0.23	0.10	0.09
CEP63	ILMN_2359945	16	rs8192935	16	55861794	CES1	rs772788	2	235248562		5.65			
CHPT1	ILMN_2202940	12	rs591967	13	38388122		rs2695290	12	102087844	CHPT1	5.74	0.72	0.20	0.44
CHPT1	ILMN_2202940	12	rs6539014	12	102277782		rs867578	11	81937002	CHPT1	4.75	0.92	0.02	0.36
CLEC12A	ILMN_1663142	12	rs429790	16	84471642		rs7313235	12	10132283	CLEC12A	5.55	0.07	1.28	0.67
CLEC12A	ILMN_2403228	12	rs7305054	11	10156646		rs3903088	10	134236688	CLEC12A	7.54	0.95	0.36	0.73
CLEC12A	ILMN_1674609	5	rs17129799	11	96929337		rs6863172	5	175595960	CLTB	5.55		0.27	
CLTB	ILMN_1770290	19	rs3752237	19	1047161	ABCA7	rs169130	16	63121080	CLTB	7.56	0.07	0.07	0.02
CNN2	ILMN_1770290	19	rs3752237	19	1047161	ABCA7	rs7336017	13	67713633	CNN2	6.33	1.92	0.28	1.39
CPFL	ILMN_1682928	8	rs4336345	8	145569535	ABCA7	rs1455268	4	61738094	CPFL	6.34	0.10	0.01	0.01
CPVL	ILMN_1654545	7	rs12596791	16	26115562		rs2455884	7	29188475	CPVL	5.74	0.06	0.57	0.23

Continued on next page

Gene ID ^a			Expression trait			SNP 1			SNP 2			Interaction statistic ^c - Log ₁₀ p-values			Distance / Mb		
rs ID	Chr.	Probe ID ^b	rs ID	Chr.	Pos / Mb ^c	Association ^d	rs ID	Chr.	Pos / Mb ^c	Association ^d	BSGS ^e	Fehrmann ^f	EGCUT ^g	Metag ^h			
PEZ2	2	ILMN_1739586	rs2356400	19	43221776		rs13406184	2	36791226	PEZ2	5.78	0.14	0.33	0.16			
PEZ2	2	ILMN_1739586	rs969010	4	15963132		rs11691600	2	36791226	PEZ2	6.59	0.14	0.28	0.14			
FCGD2	6	ILMN_2115005	rs4803848	19	46205050		rs831486	6	37901267	FCGD2	5.69	0.12	0.25	0.11			
FCGD2	6	ILMN_2115005	rs902634	10	133943951		rs831489	6	36999682	FCGD2	5.49	1.20	0.21	0.66			
FLJ20489	12	ILMN_1778144	rs17615703	12	117036766		rs3872908	12	48169526	FLJ20489	5.81	0.06	0.70	0.29		68.867	
FLJ20489	12	ILMN_1778144	rs3872908	12	48169526	FLJ20489	rs897511	4	167695661		5.53	0.03	0.11	0.02			
FLJ20489	12	ILMN_1778144	rs4792199	17	7992118		rs3872908	12	48169526	FLJ20489	5.74	0.19	0.02	0.04			
FLJ20489	12	ILMN_1778144	rs4984440	15	97033129		rs3872908	12	48169526	FLJ20489	6.49	0.31	0.47	0.36			
FLJ20489	12	ILMN_1778144	rs7204135	16	50626195		rs3872908	12	48169526	FLJ20489	6.90	0.38	0.17	0.21			
FLJ20718	18	ILMN_1763663	rs9325634	21	43818790		rs278197	16	50106594	FLJ20718	6.04	0.14	0.95	0.53			
FLJ43093	6	ILMN_2123450	rs1112712	14	107276627		rs6906101	6	36667610	FLJ43093	5.48	0.39	0.06	0.13		3.962	
FLJ43093	6	ILMN_2123450	rs6906101	6	36667610		rs13214069	6	32705248		5.44	0.00	0.64	0.18		0.063	
FN3KRP1	17	ILMN_1652333	rs898095	17	80380638		rs9892064	17	80827903		16.16	28.24	29.39	59.95			
FXFD5	19	ILMN_2309848	rs4971478	2	1346063		rs12744386	1	24168019	FUCA1	6.41	0.01	0.30	0.06			
FXFD5	19	ILMN_2309848	rs1633921	20	35659200		rs788178	13	98328559		3.70	0.09	0.41	0.17			
FXFD5	19	ILMN_2309848	rs1739183	19	5609148	FXFD5	rs2285515	19	35660450	FXFD5	6.58	0.03	0.48	0.15			
FXFD5	19	ILMN_2309848	rs2285515	19	35660450	FXFD5	rs11739594	5	141709563		5.70	0.07	0.17	0.05			
FXFD5	19	ILMN_2309848	rs2285515	19	35660450	FXFD5	rs13067700	3	95331048		6.00	0.09	0.09	0.51			
FXFD5	19	ILMN_2309848	rs2285515	19	35660450	FXFD5	rs17036504	2	47567329		6.10	0.28	0.08	0.37			
G3BP2	4	ILMN_2381768	rs10230232	7	29390239		rs1553985	4	76554604		5.19	0.08	0.37	0.14			
GAA	17	ILMN_2410783	rs11150847	17	78151330		rs12602462	17	13216016		13.91	19.98	12.99	32.60		0.007	
GAA	17	ILMN_2410783	rs8068856	17	78100731		rs10902506	12	782738089		5.65	0.11	0.39	0.17			
GAP	5	ILMN_1675191	rs10070522	5	57786117	GAPT	rs7605821	2	235659228		5.85	0.01	0.78	0.28			
GAPT	5	ILMN_1675191	rs7082031	10	128038717	GAPT	rs10070522	5	57786110	GAPT	5.72	0.26	0.11	0.11			
GATS	7	ILMN_1699631	rs1147447	14													

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Table S1 – continued from previous page

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Table S1 – continued from previous page

Gene ID ^a	Chr.	Expression trait		SNP 1		SNP 2		Interaction statistic ^f / -log ₁₀ p-values		MetaSig ^g	Distance / Mb ^h		
		rs ID	Chr.	Pos/Mb ^c	Association ^d	rs ID	Chr.	Pos/Mb ^c	Association ^d			BSGS ^e	Fehrmann ^f
MBNL1	3	rs4392435	4	41513423		rs13069559	3	152187431	MBNL1	8.39	0.02	4.33	3.02
MBNL1	3	rs4735850	8	895841		rs15069559	3	152187431	MBNL1	6.74	0.32	4.21	3.38
MBNL1	3	rs4935958	18	46278091		rs1522374	3	15223530		7.72	0.03	0.27	0.07
MBNL1	3	rs6128397	20	57263132		rs10804367	3	152234166		7.22	1.34	1.15	1.73
MBNL1	3	rs710738	20	57263132		rs13069559	3	152187431	MBNL1	7.92	2.55	7.89	9.28
MBNL1	3	rs710738	20	57263132		rs10804367	3	152234166	MBP	7.40	0.03	0.23	0.07
MBP	18	rs30568	22	43210981		rs2051344	18	74715653	MBP	5.56	0.02	0.26	0.27
MBP	18	rs30568	22	43210981		rs1125539	3	155204939		5.79	0.02	0.76	0.27
MBP	18	rs2051344	18	74715653	MBP	rs1125539	3	155204939		6.03	0.15	0.50	0.26
MBP	18	rs2051344	18	74715653	MBP	rs2051344	18	74715653	MBP	5.82	0.03	0.47	0.14
MBP	18	rs4805021	19	33436367		rs4808076	19	74747424		5.40	7.06	3.13	1.71
MBP	18	rs8002433	18	74747424		rs966396	1	12050634	MBP	4.63	1.13	1.33	1.33
MBP	13	rs7989895	13	109401737		rs4846085	4	171860973	MBP	5.71	0.61	0.25	0.41
MBP	13	rs7989895	13	109401737		rs11725347	1	171860973	MBP	5.07	0.13	0.30	0.14
MBP	13	rs7989895	13	109401737		rs12718598	7	50428445	MBP	5.57	0.07	1.02	0.05
MBP	13	rs7989895	13	109401737		rs12718598	7	50428445	MBP	5.57	0.05	0.08	0.02
MBP	13	rs7989895	13	109401737		rs2660665	8	137526799	MBP	4.17	0.05	0.27	0.40
MBP	13	rs7989895	13	109401737		rs4147592	7	1565600146	MBP	5.45	0.57	0.02	0.02
MBP	13	rs7989895	13	109401737		rs11771552	7	1547087616	MBP	5.90	0.01	0.23	0.15
MBP	13	rs7989895	13	109401737		rs1805	11	118076069	MBP	5.64	0.97	1.08	1.35
MBP	13	rs7989895	13	109401737		rs750495	5	1782046	MBP	6.89	0.34	0.18	0.19
MBP	13	rs7989895	13	109401737		rs2863095	10	102746503	MBP	5.71	0.26	0.18	0.19
MBP	13	rs7989895	13	109401737		rs3811188	14	42194916	MBP	6.56	0.14	0.44	0.22
MBP	13	rs7989895	13	109401737		rs722269	6	42158596	MBP	7.48	0.46	0.70	0.64
MBP	13	rs7989895	13	109401737		rs2395803	6	42158596	MBP	6.85	0.31	0.63	0.46
MBP	13	rs7989895	13	109401737		rs13217993	6	4216401	MBP	6.21	0.41	0.25	0.28
MBP	13	rs7989895	13	109401737		rs12431444	14	42068689	MBP	5.18	1.87	1.87	2.86
MTMR15	15	rs7178375	15	21521718	MTMR10	rs11160227	14	95514596		6.31	0.46	0.52	0.50
MTMR15	15	rs7178375	15	21521718	MTMR10	rs4973801	21	29363604		5.83	0.11	0.50	0.23
MTMR15	15	rs459498	21	42795027	MTMR10	rs8130120	21	29363604	MYBPC3	6.78	0.29	0.92	0.65
MTMR15	21	rs459498	21	42795027	MTMR10	rs1317149	11	4748685	MYBPC3	5.56	0.13	0.46	0.23
MTMR15	21	rs459498	21	42795027	MTMR10	rs124681	11	47529947	MYBPC3	5.71	0.04	0.08	0.02
MTMR15	13	rs10134030	13	109550561	MYOM1	rs2737422	16	87580855	MYBPC3	6.02	0.74	0.15	0.40
MTMR15	13	rs7322768	13	109550561	MYOM1	rs2737422	16	87580855	MYBPC3	6.02	0.74	0.15	0.40
MTMR15	16	rs4798075	16	87580855	MYOM1	rs11649236	16	86324237	MYBPC3	5.54	0.20	0.59	1.77
MTMR15	16	rs12444224	16	87580855	MYOM1	rs6826085	4	76870229	MYBPC3	5.65	0.20	0.03	0.30
MTMR15	4	rs2070757	4	74638723	MYOM1	rs6826085	4	76870229	MYBPC3	5.46	0.27	0.43	0.30
MTMR15	22	rs2071856	22	37770630	MYOM1	rs6826085	4	76870229	MYBPC3	6.08	0.07	0.48	0.18
MTMR15	8	rs21233678	8	144663661	NAPRT1	rs2786014	1	234897243	MYBPC3	8.45	15.12	16.08	30.77
MTMR15	8	rs21233678	8	144663661	NAPRT1	rs3889129	9	93911111	MYBPC3	5.62	1.27	0.19	0.81
MTMR15	8	rs21233678	8	144663661	NAPRT1	rs4862705	4	187445552	MYBPC3	6.12	0.87	0.76	1.01
MTMR15	8	rs21233678	8	144663661	NAPRT1	rs6455553	6	167811764	MYBPC3	6.06	1.10	2.58	2.77
MTMR15	8	rs21233678	8	144663661	NAPRT1	rs700276	7	146189057	MYBPC3	6.83	0.13	0.47	0.23
MTMR15	8	rs21233678	8	144663661	NAPRT1	rs7571561	2	213386267	MYBPC3	6.60	0.29	0.88	0.63
MTMR15	22	rs2208123	22	48214812	NAPRT1	rs12123758	8	144663661	NAPRT1	5.50	0.12	0.17	0.08
MTMR15	8	rs2208123	8	103488089	NAPRT1	rs12123758	8	144663661	NAPRT1	5.58	0.67	1.10	0.40
MTMR15	8	rs4743420	8	50882019	NAPRT1	rs930280	9	98391111	NAPRT1	5.58	0.82	1.10	0.40
MTMR15	19	rs1405655	19	50882019	NAPRT1	rs930280	9	98391111	NAPRT1	5.58	0.82	1.10	0.40
MTMR15	19	rs1405655	19	50882019	NAPRT1	rs10852406	10	93976932	NAPRT1	7.38	2.11	0.44	0.71
MTMR15	19	rs1405655	19	50882019	NAPRT1	rs7077137	2	234721287	NAPRT1	7.38	2.11	0.44	0.71
MTMR15	19	rs1405655	19	50882019	NAPRT1	rs4797384	17	23228670	NAPRT1	7.38	2.11	0.44	0.71
MTMR15	19	rs1405655	19	50882019	NAPRT1	rs4797384	17	23228670	NAPRT1	7.38	2.11	0.44	0.71
MTMR15	22	rs1096857	22	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878	3	183141003	NMT2	6.84	0.43	0.34	0.32
MTMR15	16	rs1096857	16	15239498	NMT2	rs12400878							

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Table S1 – continued from previous page

Gene ID ^a		Expression trait		SNP 1			SNP 2			Interaction statistic / -log ₁₀ p-values			Distance / Mb ^b
rs ID	Chr.	rs ID	Chr.	Pos/Mb ^c	Association ^d	rs ID	Chr.	Pos/Mb ^c	Association ^d	BSGS ^e	Fehrmann ^f	EGCUT ^g	Meta ^g
NRBF2	10	rs6025645	20	56157341		rs7923609	10	65133822	NRBF2	5.45			
NRBF2	10	rs6517815	21	19819016		rs7923609	10	65133822	NRBF2	6.11			
NRBF2	10	rs4852124	2	240680022		rs6588415	1	52334047	NRBF2	6.13			
NRDI	8	rs5017351	11	25453452		rs1005901	8	21964378	NUDT18	5.44	0.47	0.05	0.17
NUDT18	8	rs11613438	12	113480510		rs1047944	6	163997407	NUDT18	8.59	1.27	1.35	2.03
OAS1	12	rs1334483	12	113480510		rs2721133	12	113409260	OAS1	4.13	4.12	0.81	3.86
OAS1	12	rs1334483	12	113480510		rs2721133	12	113409260	OAS1	4.38	0.87	0.46	0.76
OPIN	10	rs7192613	16	74286646		rs17312962	10	13169066	OPIN	5.64	0.42	0.06	0.14
OSBPL5	21	rs2829679	21	26625433		rs998639	11	3149249	OSBPL5	5.00	0.36	0.00	0.07
OSTF1	9	rs17780195	17	70624189		rs2273770	9	77755469	OSTF1	5.42	0.16	0.87	0.49
OSTF1	9	rs2273770	9	77755469	OSTF1	rs7775469	9	77755469	OSTF1	5.42	1.20	0.08	0.62
OVGP1	1	rs10802822	1	240132968		rs7718088	5	179590952	OVGP1	5.43	0.13	1.48	0.88
OVGP1	1	rs347351	1	102148107		rs1264898	1	111992823	OVGP1	6.04	0.25	1.21	0.82
PAM	5	rs28092	5	140149795	PAM	rs7846000	1	40139553	HPCAL4	5.59	0.66	0.44	0.59
PCYOX1L	5	rs2438490	5	14826162	PCYOX1L	rs2731939	3	21395989	HPCAL4	6.20	0.19	0.26	0.16
PEX5	12	rs10444467	12	128052636		rs4329748	12	7364442	PEX5	5.85	0.09	0.71	0.32
PEX5	12	rs7495797	12	27246462		rs4329748	12	7364442	PEX5	5.74	0.34	0.09	0.13
PFAAF5	13	rs131969	22	49151303		rs7328733	13	33126737	PFAAF5	5.64	0.87	0.36	0.67
PGLYRP1	19	rs12982353	19	46529456		rs1263806	14	21982957	PFAAF5	6.51	0.03	0.65	0.24
PHCA	11	rs493642	11	123097386	PGLYRP1	rs10736812	11	76708086	PHCA	5.51	0.36	0.90	0.70
PISD	22	rs4141404	22	31675185	PIK3IP1	rs2065841	1	61728597	PISD	5.60	0.20	0.01	0.03
PISD	22	rs470072	22	32263151	PISD	rs10498313	14	30398876	PISD	5.23	0.02	0.87	0.33
PISD	22	rs615752	22	332334931	PISD	rs1465754	22	32097775	PISD	7.11	0.00	1.19	0.48
PISD	22	rs715572	22	332334931	PISD	rs6518754	22	32097775	PISD	4.12	0.05	0.42	0.15
PNKD	16	rs6869411	16	155781604		rs928046	9	140487108	PNKD	6.35	0.16	0.04	0.04
PNPLA7	9	rs11639998	16	45271109		rs4735800	11	35619930	PNPLA7	5.15	0.31	0.78	0.56
PPFIBP2	11	rs911019	20	49668255		rs1156875	14	35619930	PPFIBP2	4.44	0.29	0.33	0.26
PPFIBP2	14	rs12914603	15	58350896		rs12120009	1	212447167	PPFIBP2	5.81	0.12	0.42	0.19
PPFIBP2	14	rs10930170	2	163999467		rs12120009	1	212447167	PPFIBP2	5.63	0.72	0.48	0.66
PPF2R5A	1	rs12433255	12	123595064		rs12120009	1	212447167	PPF2R5A	5.72	0.08	0.95	0.46
PPF2R5A	1	rs1889083	13	66222691		rs12120009	1	212447167	PPF2R5A	5.61	0.36	0.13	0.17
PPF2R5A	1	rs682334	11	107417238		rs12120009	1	212447167	PPF2R5A	5.65	1.69	0.28	1.21
PPF2R5A	1	rs757871	6	135030045		rs12120009	1	212447167	PPF2R5A	5.95	0.37	0.06	0.12
PPF2R5A	1	rs7871178	9	271048475		rs12120009	1	212447167	PPF2R5A	5.72	0.16	0.30	0.16
PRKDX5	11	rs8019823	16	95040482		rs11600990	11	64082807	PRKDX5	6.43	0.81	0.14	0.44
PRKDX5	11	rs2188355	16	23867776		rs11600990	11	64082807	PRKDX5	7.34	0.53	0.11	0.25
PRMT2	21	rs1029231	21	47931653	C21ORF57	rs958127	18	31497346	PRMT2	5.60	0.19	0.03	0.04
PRMT2	21	rs2839372	21	48063862	C21ORF57	rs1171058	21	47776382	PRMT2	4.81	0.69	4.47	4.06
PSMB1	6	rs3862607	17	121774705		rs13207114	6	170877444	PSMB1	5.79	0.00	0.26	0.04
PSMB1	6	rs4890648	18	43983954		rs6928843	6	170890384	PSMB1	5.14	0.44	0.21	0.27
PSMB1	6	rs6060930	20	30347832		rs9295415	6	170823379	PSMB1	5.44	1.95	0.64	1.78
PSMB1	6	rs6928843	6	170800384	PSMB1	rs2769689	1	225797957	PSMB1	5.42	1.18	0.32	0.86
PSMB1	6	rs7299749	12	131727816		rs13207114	6	170877444	PSMB1	5.40	0.03	0.48	0.15
PWPI	12	rs2353567	14	95478823		rs11036212	11	5221825	PTDSS1	5.00	0.03	0.32	0.15
PWPI	12	rs4969205	17	76598123		rs11036212	11	5221825	PTDSS1	5.90	0.80	0.08	0.38
PWPI	12	rs631562	11	126852438		rs11036212	11	5221825	PTDSS1	5.70	0.02	0.40	0.11
QDPR	4	rs4946705	6	106348246		rs10020773	4	17526682	QDPR	5.75	1.03	1.25	1.55
RAB31P	12	rs241730	22	33375704		rs7305307	12	70235726	RAB31P	6.55	0.25	0.08	0.09
RAB31P	12	rs241730	22	33375704		rs7305307	12	70235726	RAB31P	6.42	0.28	0.84	0.59
RAB31P	19	rs1075728	19	42462788	RAB31P	rs7951028	11	120161117	RAB31P	5.23	0.03	0.31	0.08
RBL2	16	rs6931702	16	53526551	AKTIP	rs1863464	15	26988488	RBL2	5.40	0.38	0.37	0.47
RCN1	11	rs10879131	12	41147155		rs4922579	11	32136436	RCN1	5.43	0.58	0.37	0.47
RCN1	11	rs4922579	11	32136436	RCN1	rs1166957	8	141177468	RCN1	4.32	0.41	0.09	0.17
RCN1	11	rs4922579	11	32136436	RCN1	rs1341899	1	102740645	RCN1	5.40	0.04	0.26	0.07

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Table S1 – continued from previous page

Gene ID ^a			Expression trait			SNP 1			SNP 2			Interaction statistic ^f			BSCS ^e			-log ₁₀ p-values			Distance / Mb ^b		
Gene	ID ^a	Chr.	rs ID	Chr.	Pos/Mb ^c	Association ^d	rs ID	Chr.	Pos/Mb ^c	Association ^d	rs ID	Chr.	Pos/Mb ^c	F _{max}	F _{max} ²	ECGUT ^g	Meta ^g	ECGUT ^g	Meta ^g	ECGUT ^g	Meta ^g		
RENE	ILMN_1802830	1	rs4982958	14	24987865		rs301819	1	8501786	RENE	rs301819	1	8501786	RENE	5.66	0.61	1.23	1.17					
RENE	ILMN_1802838	1	rs7697290	4	135248366		rs301819	1	8501786	RENE	rs301819	1	8501786	RENE	5.74	0.14	0.10	0.06					
RENE	ILMN_1802840	1	rs11085629	19	13174312		rs301819	1	8501786	RENE	rs301819	1	8501786	RENE	5.12	0.21	0.33	0.21					
RENE	ILMN_1802842	1	rs301819	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	RENE	5.71	0.08	0.60	0.26					
RENE	ILMN_1802844	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802846	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802848	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802850	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802852	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802854	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802856	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802858	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802860	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802862	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802864	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802866	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802868	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802870	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802872	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802874	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802876	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802878	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802880	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802882	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802884	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802886	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802888	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802890	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802892	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802894	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802896	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802898	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802900	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802902	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802904	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802906	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802908	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802910	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802912	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802914	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802916	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802918	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802920	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802922	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802924	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802926	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802928	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802930	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802932	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802934	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802936	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802938	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802940	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802942	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802944	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802946	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1802948	1	rs10285098	14	21182850	RNASE6	rs10285098	13	100601327	RNASE6	rs10285098	13	100601327	RNASE6	5.48	0.42	0.21	0.26					
RENE	ILMN_1																						

Continued on next page

Table S1 – continued from previous page

Gene ID ^a		Expression trait		SNP 1			SNP 2			Interaction statistic / -log ₁₀ p-values					
TMEMD4	Probe ID ^b	Chr.	rs ID	Chr.	Pos/Mb ^c	Association ^d	rs ID	Chr.	Pos/Mb ^c	Association ^d	BSGS ^e	Fehrmann ^f	EGCUT ^g	Meta ^g	Distance / Mb ^h
TMEM149	ILMN-17804148	7	rs19340400	11	132389627		rs17725246	7	44581986	TMEMD4	3.70	0.06	1.34	0.70	
	ILMN-1780426	19	rs28390113	21	47248981		rs8106939	19	36219525	TMEM149	8.11	0.16	0.48	0.26	
	ILMN-1780426	19	rs5762235	22	27925288		rs8106939	19	36219525	TMEM149	6.79				
	ILMN-1780426	19	rs6090518	20	43207005		rs8106939	19	36219525	TMEM149	11.09	0.76			
	ILMN-1780426	19	rs807491	19	36268923	SNX26	rs7254600	19	36147315	TMEM149	12.16	81.55	45.78	145.78	0.122
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs10508289	10	36147315	TMEM149	8.12	1.55	3.09	3.07	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs10508289	10	36147315	TMEM149	8.02	0.40	0.99	0.80	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs10937361	3	188350436	TMEM149	8.39	3.61	1.18	3.78	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs1401098	12	128884559	TMEM149	7.37	2.41	1.00	2.52	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs1557335	18	64268976	TMEM149	6.95	0.08	0.07	0.03	
TMEM149	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs17719594	14	90932398	TMEM149	6.93	3.06	0.77	2.87	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs1843357	8	13822381	TMEM149	6.21	3.72	3.33	6.00	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs2351458	4	113317583	TMEM149	7.30	0.04	9.61	8.00	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs2539000	7	147619772	TMEM149	6.70	1.57	1.52	2.27	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs2731711	5	171792273	TMEM149	6.70	0.07	3.14	2.10	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs4711728	11	129595460	TMEM149	5.92	0.19	0.33	0.19	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs6718480	2	233879066	TMEM149	8.89	0.90	3.62	3.51	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs6926382	2	161683974	TMEM149	8.55	3.31	5.15	7.36	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs7213338	17	80357420	TMEM149	5.80	3.06	8.80	10.72	
	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs914940	1	242889492	TMEM149	6.22	3.36	6.96	9.20	
TMEM63A	ILMN-1780426	19	rs8106939	19	36219525	TMEM149	rs9509428	13	21473952	TMEM63A	9.44	0.10	5.75	4.47	
	ILMN-1780426	19	rs1254086	13	72890603		rs4149226	1	226027323	TMEM63A	5.60				
	ILMN-1780426	11	rs1548475	9	58058246		rs4963126	11	65845	TMEM80	5.79	0.64	0.12	0.32	
	ILMN-1780426	7	rs1537146	9	4859303		rs10488630	7	128593948	IRF5	5.61	0.11	0.15	0.07	
	ILMN-1780426	7	rs199793	20	22287303		rs10488630	7	128593948	IRF5	5.52	1.03	0.17	0.62	
	ILMN-1780426	7	rs7776572	20	23528927		rs11770192	7	23498358	IRF5	8.23	3.19	1.89	4.09	
	ILMN-1780426	13	rs13531675	11	113531675		rs3916581	11	11888787	TRAPPC4	5.61	0.28	0.40	0.29	
	ILMN-1780426	19	rs17159840	19	7758194	TRAPPC5	rs10050904	5	166970604	TRAPPC4	5.52	0.93	0.01	0.36	
	ILMN-1780426	19	rs17159840	19	7758194	TRAPPC5	rs1023095	8	132022957	TRAPPC4	6.92	0.37	1.60	1.07	12.131
	ILMN-1780426	19	rs17159840	19	7758194	TRAPPC5	rs1375714	6	156404902	TRAPPC4	7.79	0.12	0.18	0.07	
TMEM149	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs1393299	1	24329791	TRAPPC5	6.43	0.63	0.47	0.59	
	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs17763599	19	2369415	TRAPPC5	6.38	0.21	0.24	0.16	
	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs4968328	17	57495457	TRAPPC5	6.51	0.50	0.38	0.44	
	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs7313362	12	129644342	TRAPPC5	7.08	0.04	0.65	0.25	
	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs7694997	4	9947811	TRAPPC5	5.86	0.20	0.36	0.22	
	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs7800935	7	146690926	TRAPPC5	6.27	0.15	0.33	0.16	
	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs856638	14	85439550	TRAPPC5	6.73	0.24	0.07	0.08	
	ILMN-2372639	19	rs380708	22	22740855		rs17159840	19	7758194	TRAPPC5	7.58				
	ILMN-2372639	19	rs3916995	19	45128455		rs17159840	19	7758194	TRAPPC5	7.73	0.85	0.78	1.01	
	ILMN-2372639	19	rs6040514	20	11272861		rs10179572	2	228504503	TRAPPC5	8.10	0.51	0.55	0.56	
TMEM149	ILMN-2372639	19	rs7246264	19	7762978		rs10179572	2	228504503	TRAPPC5	6.71	0.14	0.02	0.02	
	ILMN-2372639	19	rs7246264	19	7762978		rs12921440	16	30408705	TRAPPC5	7.34	0.14	0.26	0.13	
	ILMN-2372639	19	rs7246264	19	7762978		rs1887778	3	134635088	TRAPPC5	7.05	0.08	0.86	0.40	
	ILMN-2372639	19	rs7246264	19	7762978		rs963354	3	157393770	TRAPPC5	7.41	0.36	0.90	0.69	
	ILMN-2372639	19	rs10862975	12	85749398		rs2395771	6	41264577	TRAPPC5	5.42	0.11	0.25	0.11	
	ILMN-2372639	6	rs12412964	7	158850412		rs2032447	6	26044369	TRAPPC5	5.92	1.20	1.23	1.69	
	ILMN-2372639	17	rs2527180	7	158850412		rs10748526	10	82273079	TRAPPC5	6.46	0.04	0.91	0.39	
	ILMN-2372639	19	rs908726	17	27194634	MYBPC3	rs12800998	11	2317951	TRAPPC5	6.00	0.07	0.18	0.06	
	ILMN-1718621	11	rs10838738	11	47663049	TSPAN32	rs12800998	11	2317951	TSPAN32	5.01				
	ILMN-1697971	6	rs2527180	7	158850412	TSPAN32	rs12800998	11	2317951	TSPAN32	5.01				
TMEM149	ILMN-3232126	22	rs1403240	22	50971266	EGGF1	rs6206007	6	137947208	TSPAN32	5.34				
	ILMN-3232126	22	rs1403240	22	50971266	EGGF1	rs1198819	2	238746880	TSPAN32	6.34				
	ILMN-3232126	22	rs1403240	22	50971266	EGGF1	rs4783126	16	85147633	TSPAN32	6.13				

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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	0.17	1.35	1.22
WDR48	LMN-1762103	3	rs1887778	9	134635088		rs883349	3	39067925	WDR48	6.34	0.59	1.63	1.43
WDR48	LMN-1762103	3	rs9554833	13	102624790		rs7619193	3	39044131	WDR48	5.85	0.18	0.61	0.35
WDR6	LMN-1669484	3	rs12362253	11	123371708		rs7619193	3	39044131	WDR6	4.86	1.64	1.43	2.25
XAF1	LMN-2330573	17	rs1535031	21	9673170	XAF1	rs11715581	3	49194331		5.48	2.38	0.17	1.63
ZEP00	LMN-2370573	17	rs12362253	17	6673170		rs12591171	15	68179799		5.48	0.09	0.36	0.15
ZFP90	LMN-1684628	16	rs909446	21	37040648		rs18128968	16	93573945	ZFP90	5.79	0.09	0.36	0.15
ZNF500	LMN-1700238	16	rs4283723	22	48283177		rs2290560	16	4799041	ZNF500	5.29	0.67	0.27	0.46
ZYX	LMN-1701875	7	rs6056281	20	8935312		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 2013²¹

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 *cis-cis* interactions

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