

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 traits<sup>1</sup> and contributes to their variation<sup>2,3</sup> is a fundamental question in evolution and human genetics. Though often demonstrated in artificial gene manipulation studies in model organisms,<sup>4,5</sup> and some examples have been reported in other species,<sup>6</sup> few examples exist for epistasis amongst natural polymorphisms in human traits.<sup>7,8</sup> Its absence from empirical findings may simply be due to low incidence in the genetic control of complex traits,<sup>2,3</sup> but an alternative view is that it has previously been too technically challenging to detect due to statistical and computational issues.<sup>9</sup> Here we show that, using advanced computation<sup>10</sup> 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<sup>11,12</sup> 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 interactions<sup>13</sup> ( $p = 1.8 \times 10^{-10}$ ). 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.<sup>14</sup> But to date, though its contribution to phenotypic variance is frequently the subject of debate,<sup>1-3</sup> there is little empirical exploration of the role that epistasis plays in the architecture of complex traits in humans.<sup>7,8</sup> Beyond the prism of human association studies there is evidence for epistasis, not only at the molecular scale from artificially induced mutations<sup>4</sup> but also at the evolutionary scale in fitness adaptation<sup>15</sup> and speciation.<sup>16</sup>

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,<sup>17,18</sup> increased model

complexity in fitting interaction terms,<sup>19</sup> and more extreme significance thresholds to account for increased multiple testing<sup>9</sup> 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,<sup>14</sup> the power to detect epistasis diminishes rapidly. There are two simple ways to overcome this problem. One is by using extremely large sample sizes;<sup>20</sup> 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.<sup>21</sup> But unlike complex diseases, genetic associations with gene expression commonly have very large effect sizes that explain large proportions of the genetic variance,<sup>22</sup> making them good candidates to search for epistasis, should it exist.

In our discovery dataset (Brisbane Systems Genetics Study, BSGS<sup>23</sup>) 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 ( $1.03 \times 10^{15}$  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, Fehrmann<sup>12</sup> and the Estonian Genomics Centre University of Tartu (EGCUT),<sup>11</sup> 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,<sup>24</sup> 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 additive  $\times$  additive genetic interactions,<sup>9</sup> but our  
130 analysis shows that this is unlikely to be the most effective strategy for its  
131 detection. The majority of our discovery interactions comprised of one SNP  
132 that was significantly associated with the gene expression level in the discov-  
133ery dataset, and one SNP that had no previous association<sup>22</sup> (439 out of 501,  
134 Methods). Only nine interactions were between SNPs that both had known  
135 main effects while 64 were between SNPs that had no known main effects. Ad-  
136ditionally, we observed that the largest epistatic variance component for the  
137501 interactions was equally divided amongst additive  $\times$  additive, additive  $\times$   
138dominance, dominance  $\times$  additive and dominance  $\times$  dominance at the discovery  
139stage ( $p = 0.22$  for departure from expectation). This is not surprising because  
140the patterns of epistasis used for statistical decomposition (*i.e.*  $A \times A$ ,  $A \times D$ ,  
141 $D \times A$ ,  $D \times D$ ) are simply convenient orthogonal parameterisations of a two  
142locus model, and are not intended to model biological function.<sup>25</sup>

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

159 ysis these five gene expression phenotypes had non-additive variance component  
160 estimates within the 95th percentile of the 17,994 gene expression phenotypes  
161 that were analysed previously<sup>22</sup> (Supplementary Table S2, Methods).

162 In total the 501 interactions comprised 781 unique SNPs, which we analysed  
163 for functional enrichment (Methods). We tested the SNPs for cell-type specific  
164 overlap with transcriptionally active chromatin regions, tagged by histone-3-  
165 lysine-4,tri-methylation (H3K4me3) chromatin marks, in 34 cell types<sup>27</sup> (Sup-  
166plementary Figure S5). There was significant enrichment for *cis*-acting SNPs  
167 in haematopoietic cell types only ( $p < 1 \times 10^{-4}$  for the three tissues with the  
168 strongest enrichment after adjusting for multiple testing). However *trans*-acting  
169 SNPs did not show any tissue specific enrichment ( $p > 0.1$  for all tissues). This  
170 difference between *cis* and *trans* SNPs suggests different roles in epistatic in-  
171teractions where tissue specificity is provided by the *cis* SNPs. There is also  
172 enrichment for *cis*-SNPs to be localised in regions with regulatory genomic fea-  
173tures as measured by chromatin states<sup>28</sup> (Supplementary Figure S4).

174 We also demonstrate physical organisation of interacting loci within the cell,  
175 suggesting a mechanism by which biological function can lead to epistatic ge-  
176netic variance. It has been shown that different chromosomal regions spatially  
177 colocalise in the cell through chromatin interactions.<sup>13</sup> We cross-referenced our  
178 epistatic SNPs with a map of chromosome interacting regions ( $n = 96,139$ )  
179 in K562 blood cell lines<sup>29</sup> (Methods) and found that 44 epistatic interactions  
180 mapped to within 5Mb ( $p < 1.8 \times 10^{-10}$ ), (Supplementary Figure S15). Inter-  
181action of distant loci may occur through physical proximity in transcriptional  
182factories that organise across different chromosome regions and can regulate  
183transcription of related genes.<sup>30</sup>

184 Quantifying the importance of epistasis in complex traits in humans remains  
185 an open question. Here we are able to identify 238 gene expression traits with at  
186 least one significant interaction given our experiment-wide threshold, where the  
187 minimum estimated variance explained by the epistatic effects of any interaction  
188 was 2.1% of phenotypic variance. Taking results from our previously published  
189 eQTL<sup>23</sup> we calculated that 1848 of the 7339 gene expression levels analysed  
190 were influenced by additive effects where the estimated additive variance of a  
191 locus was 2.1% or greater. Thus, we can infer that the number of instances  
192 of large additive effects is significantly greater than the number of instances of  
193 large epistatic effects.

194 In terms of their contribution to complex traits a more important metric  
195 might be the proportion of the variance that the epistatic loci explain.<sup>2</sup> Taking  
196 all additive effects detected in Powell *et al* (2012) that have additive variance  
197 explaining 2.1% or greater of phenotypic variance, we calculated that the pro-  
198 portion of total phenotypic variance of all 7339 gene expression levels explained  
199 by additive effects alone was 2.16%. By contrast, the estimated epistatic vari-  
200 ance from the interacting SNPs detected in this study on average explain a total  
201 of 0.22% of phenotypic variance, approximately ten times lower than the esti-  
202 mated additive variance. There are several caveats to this comparison which we  
203 discuss in the Methods.

204 Overall, we have demonstrated that it is possible to identify and replicate

205 epistasis in complex traits amongst common human variants, despite the rela-  
206 tive contribution of pairwise epistasis to phenotypic variation being small. The  
207 bioinformatic analysis of the significant epistatic loci suggests that there are a  
208 large number of possible mechanisms that can lead to non-additive genetic varia-  
209 tion. Further research into such epistatic effects may provide a useful framework  
210 for understanding molecular mechanisms and complex trait variation in greater  
211 detail. With computational techniques and data now widely available the search  
212 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

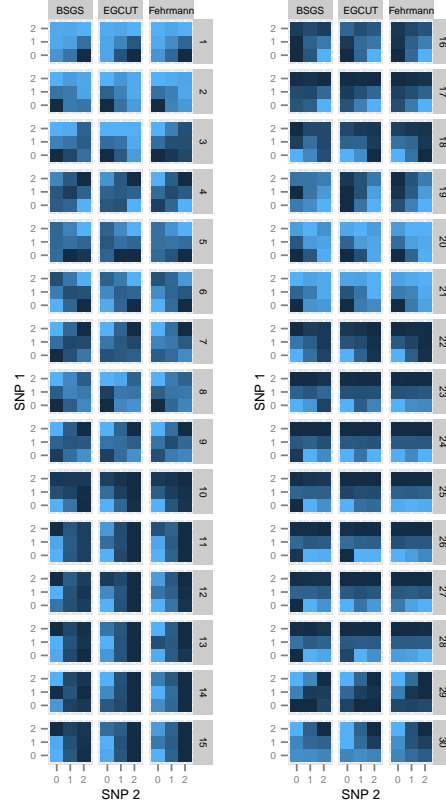
	Gene (chr.)	SNP 1 (chr.)	SNP 2 (chr.)	BSGS <sup>2</sup>	Fehrmann <sup>3</sup>	EGCUT <sup>3</sup>	Meta <sup>4</sup>
1	ADK (10)	rs2395095 (10)	rs10824092 (10)	6.69 <sup>1</sup>	18.33 <sup>1</sup>	21.21 <sup>1</sup>	39.82 <sup>1</sup>
2	ATP13A1 (19)	rs4284750 (19)	rs873870 (19)	5.30	12.18	3.25	14.23
3	C21ORF57 (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

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

<sup>2</sup> Discovery dataset

<sup>3</sup> Independent replication dataset

<sup>4</sup> 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.



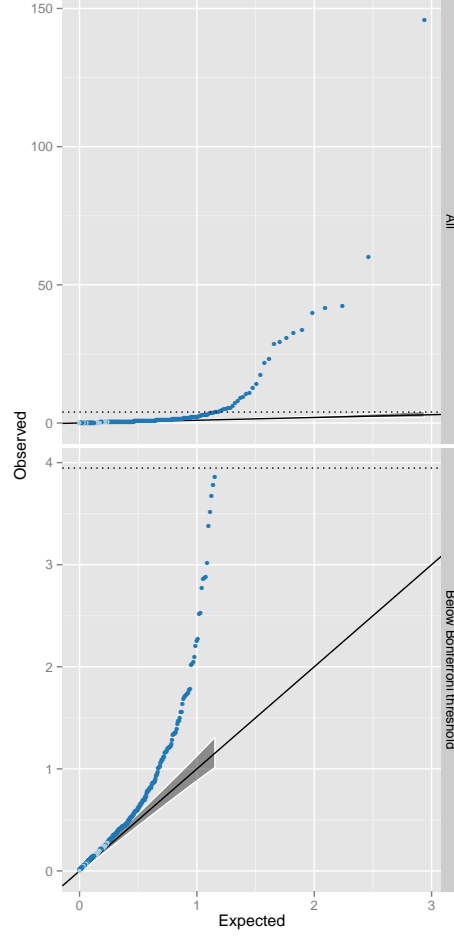


Figure 2: **Q-Q plots of interaction  $p$ -values from replication datasets** The top panel shows all 434 discovery SNPs that were tested for interactions. Observed  $p$ -values ( $y$ -axis,  $-\log_{10}$  scale) are plotted against the expected  $p$ -values ( $x$ -axis,  $-\log_{10}$  scale). The multiple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

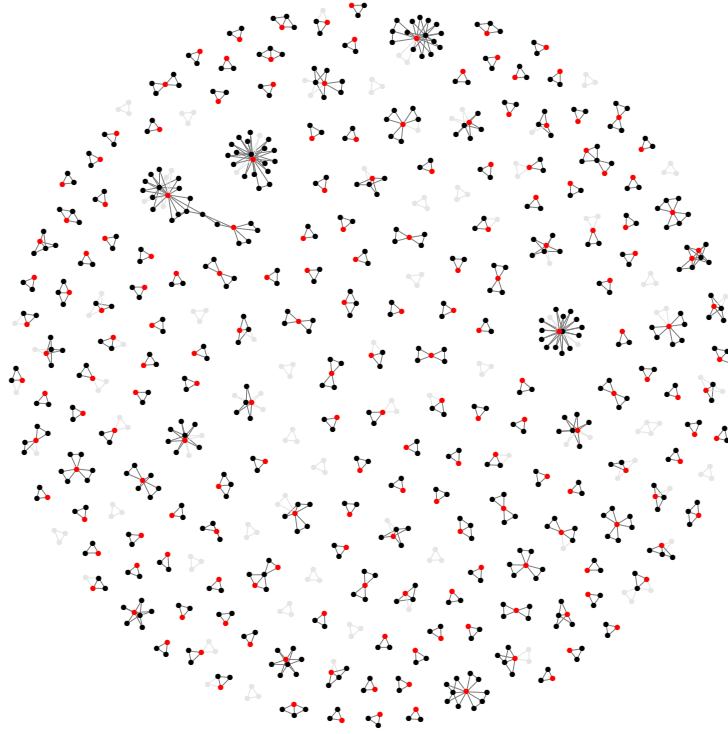


Figure 3: **Discovery and replication of epistatic networks** All 434 putative genetic interactions (edges) with data common to discovery and replication sets is shown, where black nodes represent SNPs and red nodes represent traits (gene expression probes). Three hundred and forty-five interactions 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/](http://kn3in.github.io/detecting_epi/)

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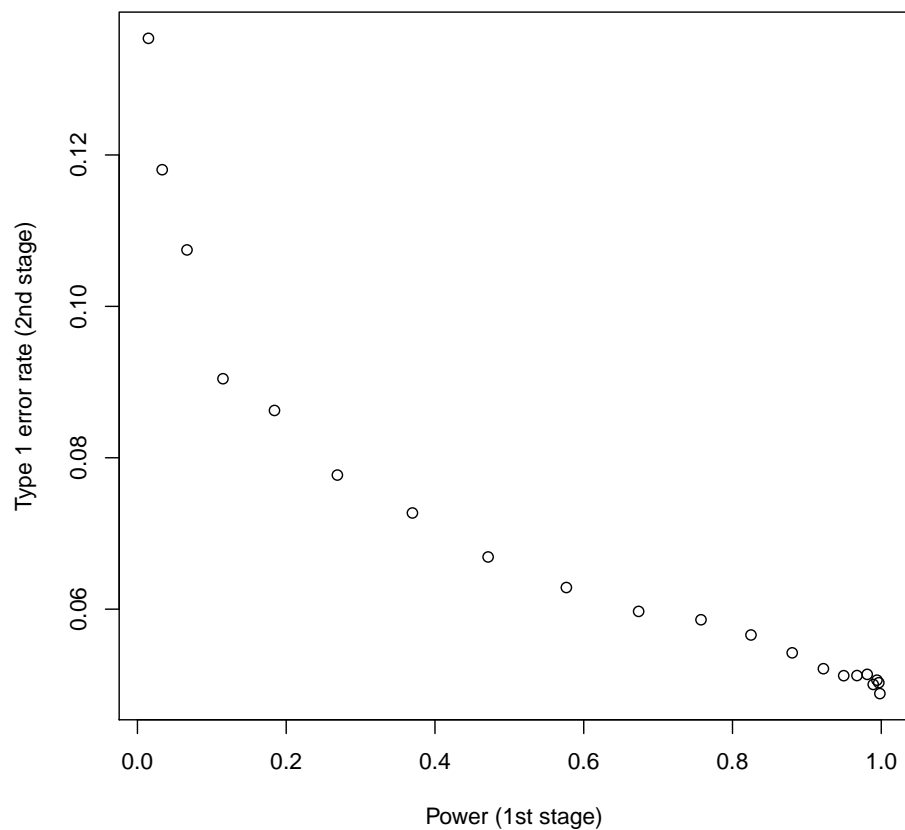


Figure S1: **Type 1 error rate of two stage design assuming a null model of one large additive effect and no epistasis** In stage 1 SNPs are tested for full genetic effects (8 d.f.) and those that surpass a threshold for multiple testing are then tested for significant interaction terms in stage 2. These 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.

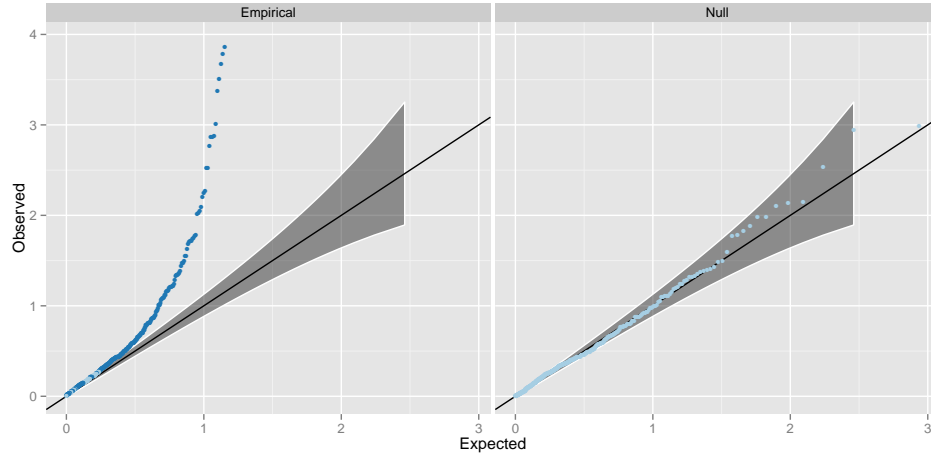


Figure S2: **Q-Q plots of interaction  $p$ -values from replication datasets, excluding the 30 points significant at the Bonferroni level** The right panel (Null) shows the interaction  $p$ -values from a meta analysis across two independent datasets on 434 randomly drawn SNP pairs. The left panel (Empirical)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

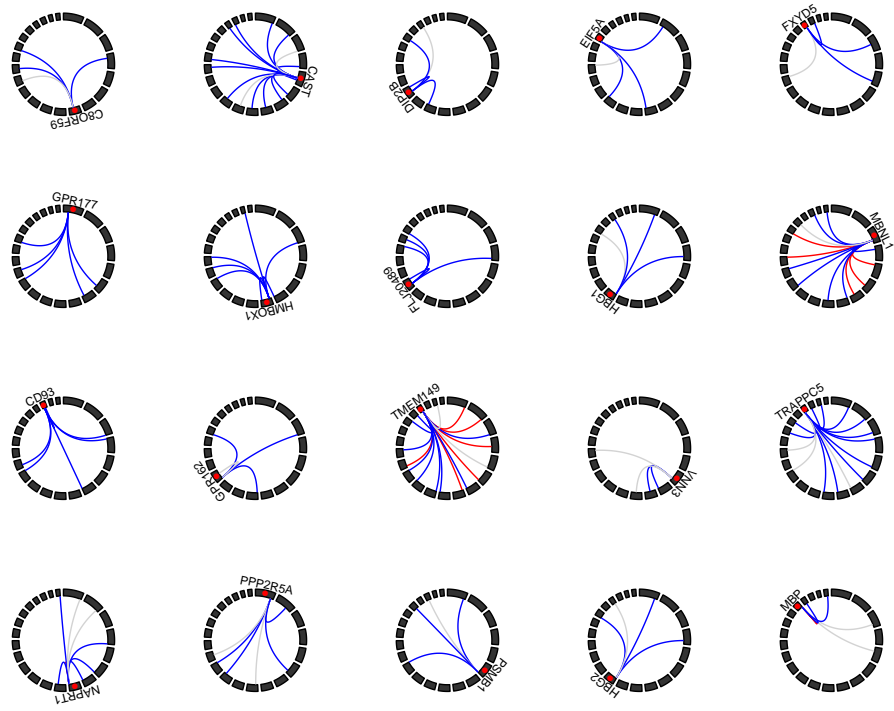
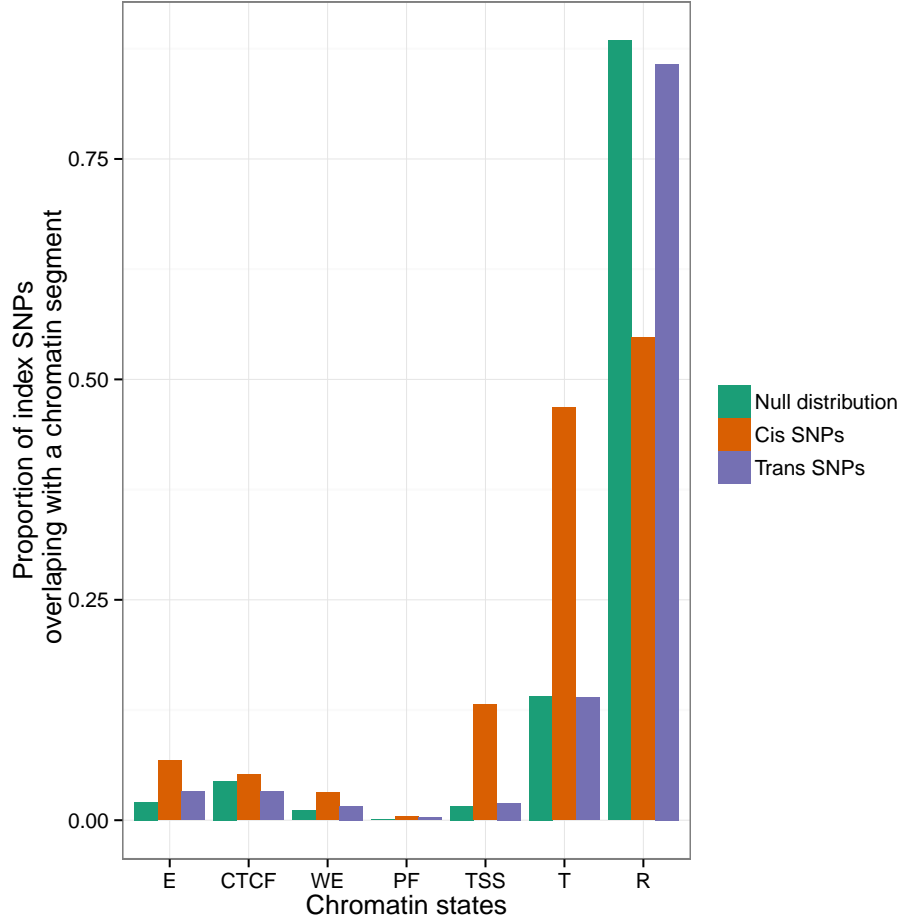
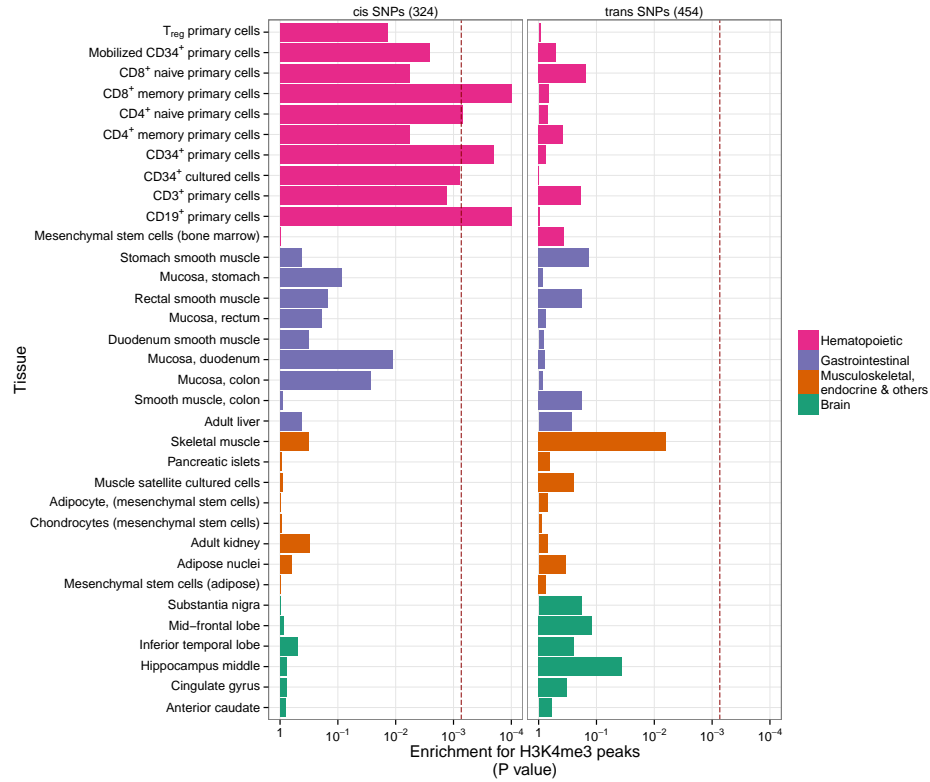


Figure S3: **Gene expression traits with four or more genetic interactions** Circle plots represent the genomic positions for SNPs (linking lines) and expression probes (red points). Chromosomes are represented by black blocks and ordered from 1 to 22 clockwise, starting from the top. Grey lines represent 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 segmentation<sup>28</sup> 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.<sup>27</sup> 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.

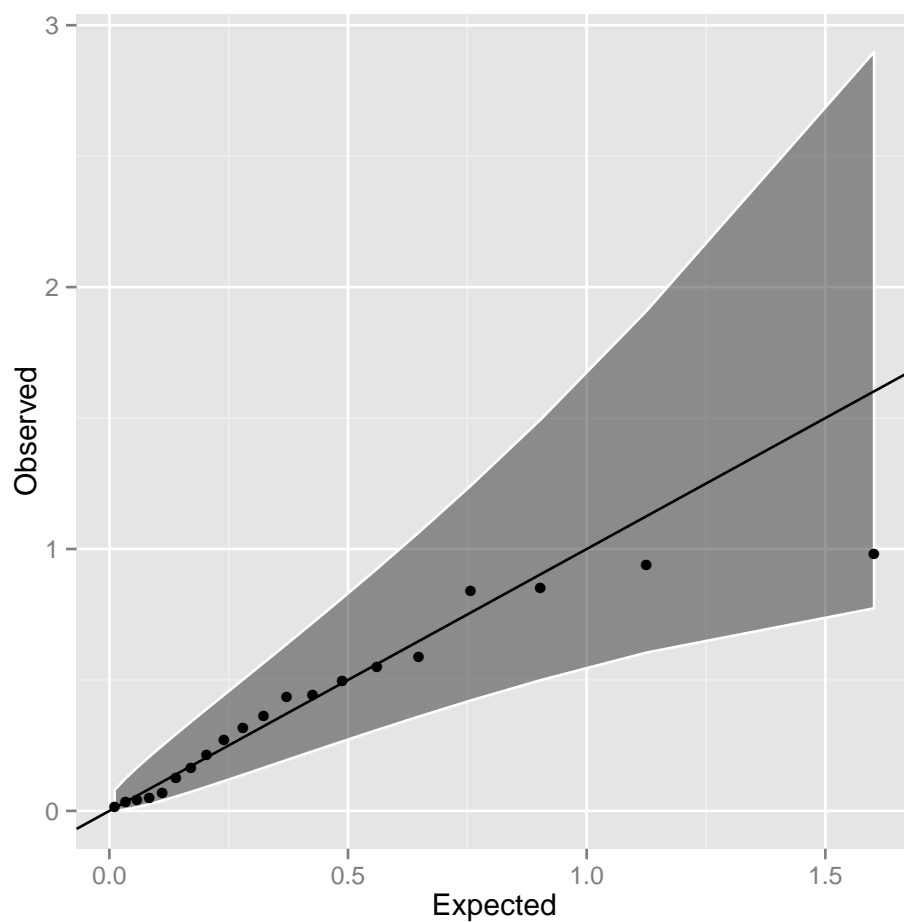


Figure S6: **Q-Q plot of interaction  $p$ -values in the CDHWB dataset**  
 Twenty of the 501 discovery SNP pairs passed filtering in the CDHWB dataset (mainly due to small sample size). There is no evidence for enrichment of interaction terms, most likely due to insufficient power given the limited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

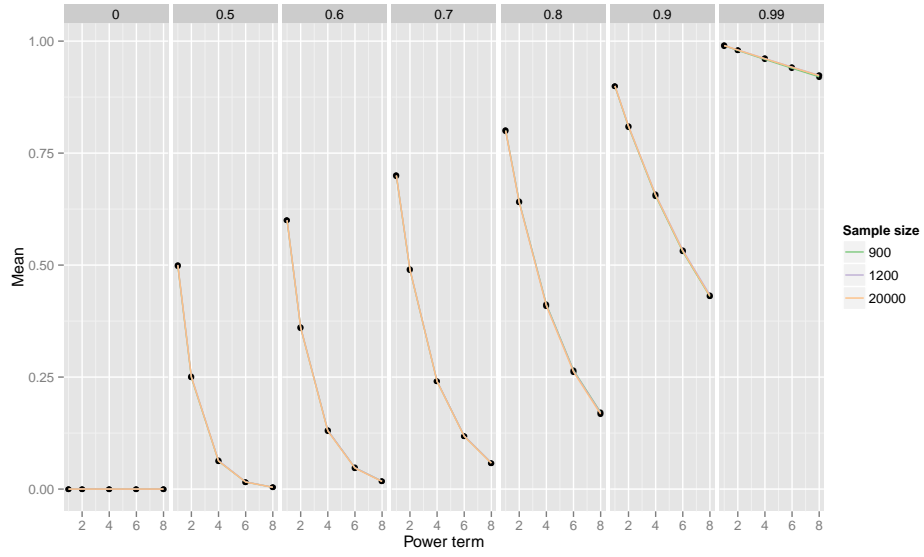
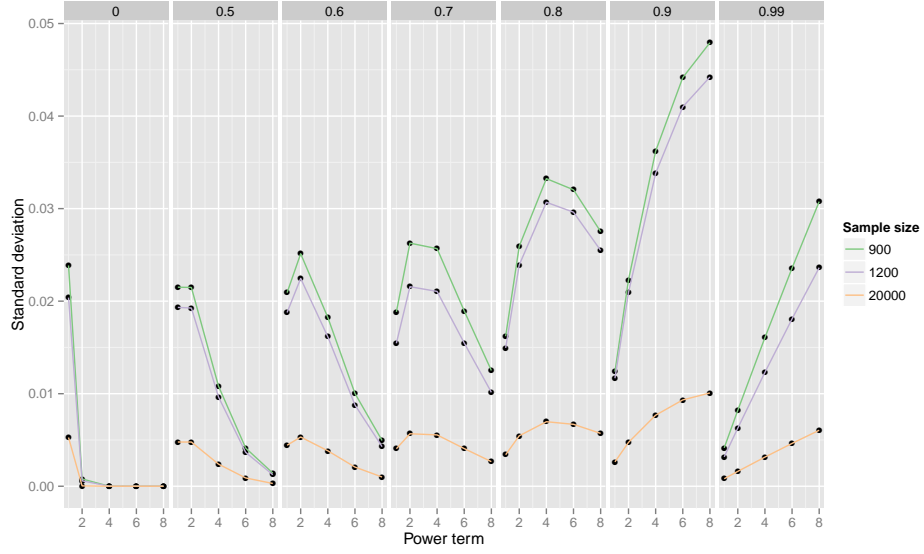


Figure S7: **Sampling mean for different power terms of population  $r$  values** Power of detection and replication of epistatic interactions depends not on  $r^2$  between causal variants and observed SNPs, but on  $r^4, r^6, r^8$ . For a given population value of LD  $r$  (columns of plots), plotted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

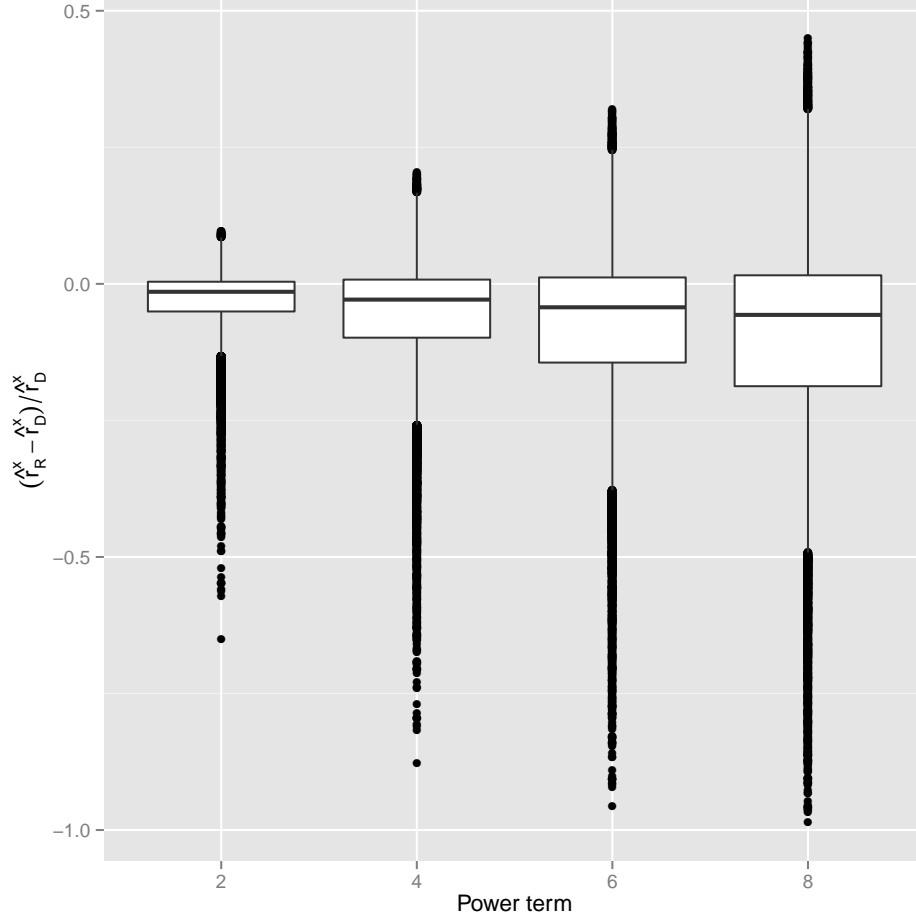


Figure S9: **Reduction in LD as estimated in replication data after ascertaining for high LD in discovery data** 100,000 “unobserved” causal variants (CVs) were tested for LD against a panel of 528,509 “observed” discovery markers (DMs). DM/CV pairs with LD  $r^2 > 0.9$  were then tested in an independent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

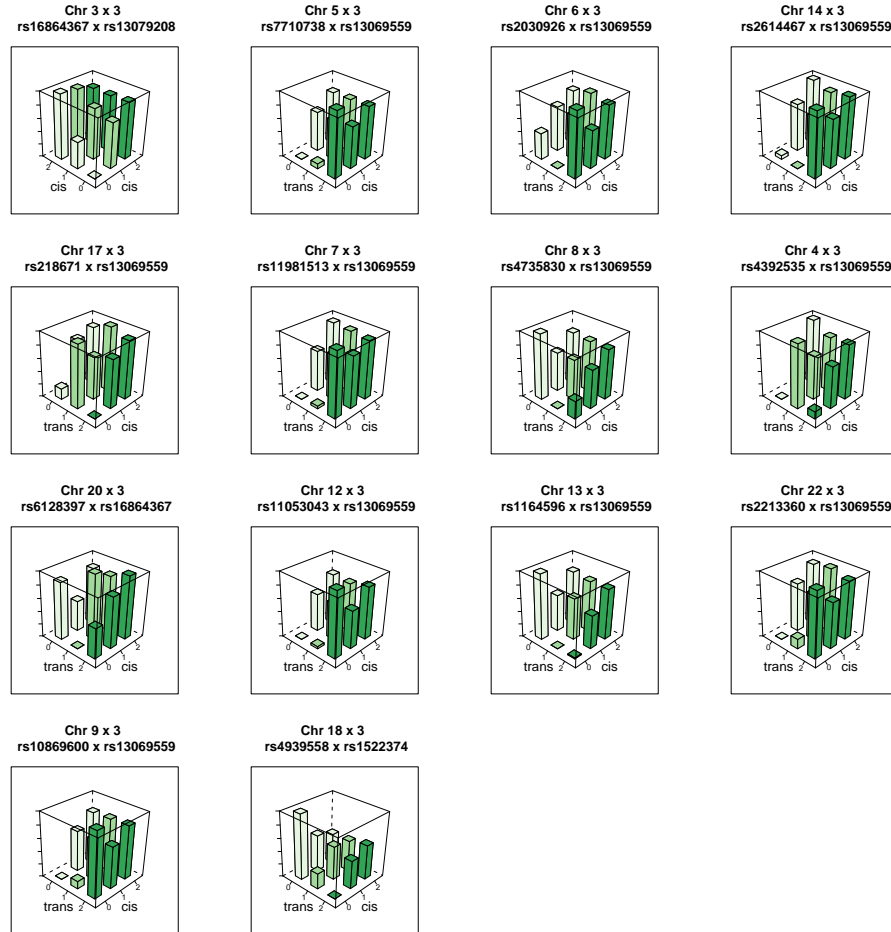


Figure S10: **Genotype-phenotype maps for 14 interactions influencing the expression of MBNL1** Each bar represents the mean phenotypic value for individuals in that genotype class. The rs13069559 SNP typically has a *cis*-additive decreasing effect on the expression of MBNL1, but in many of these interactions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

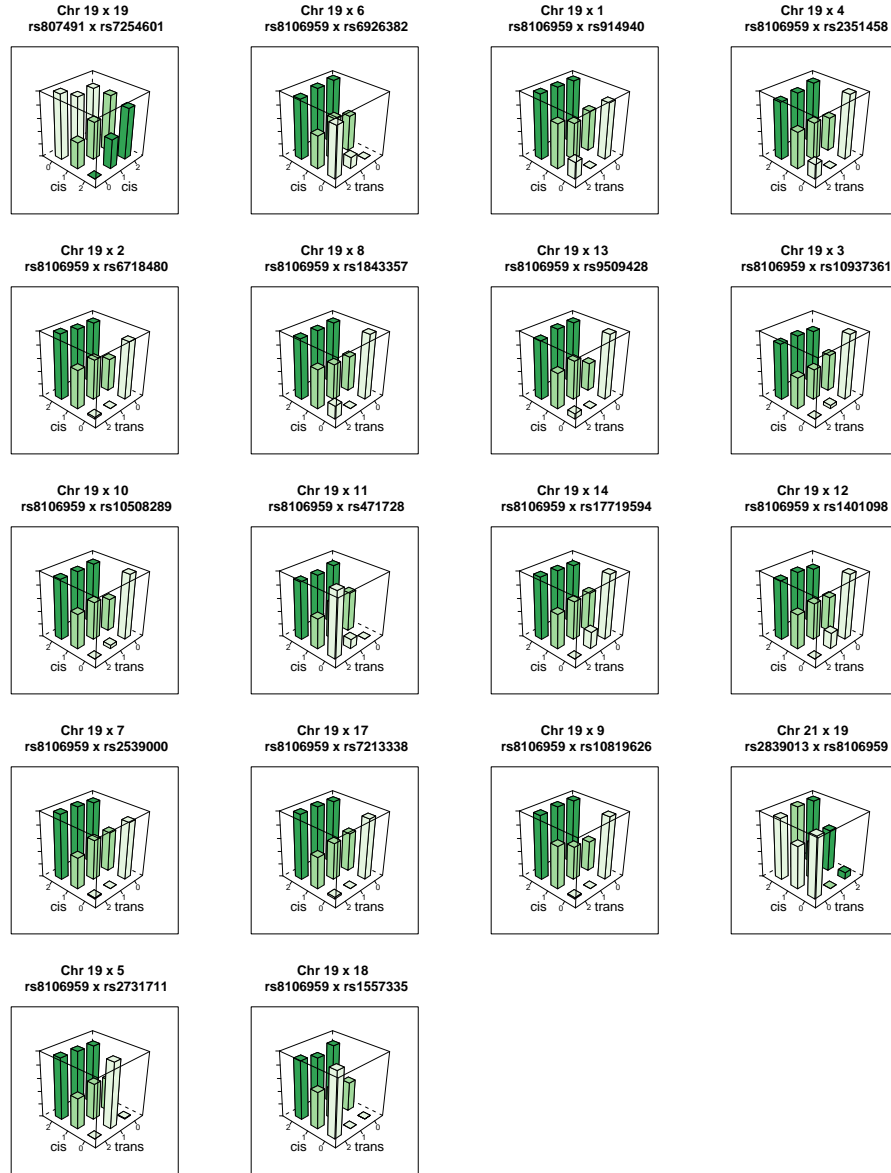


Figure S11: **Genotype-phenotype maps for 19 interactions influencing the expression of TMEM149** Each bar represents the mean phenotypic value for individuals in that genotype class. The rs13069559 SNP typically has a *cis*-additive decreasing effect on the expression of TMEM149, but in many of these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



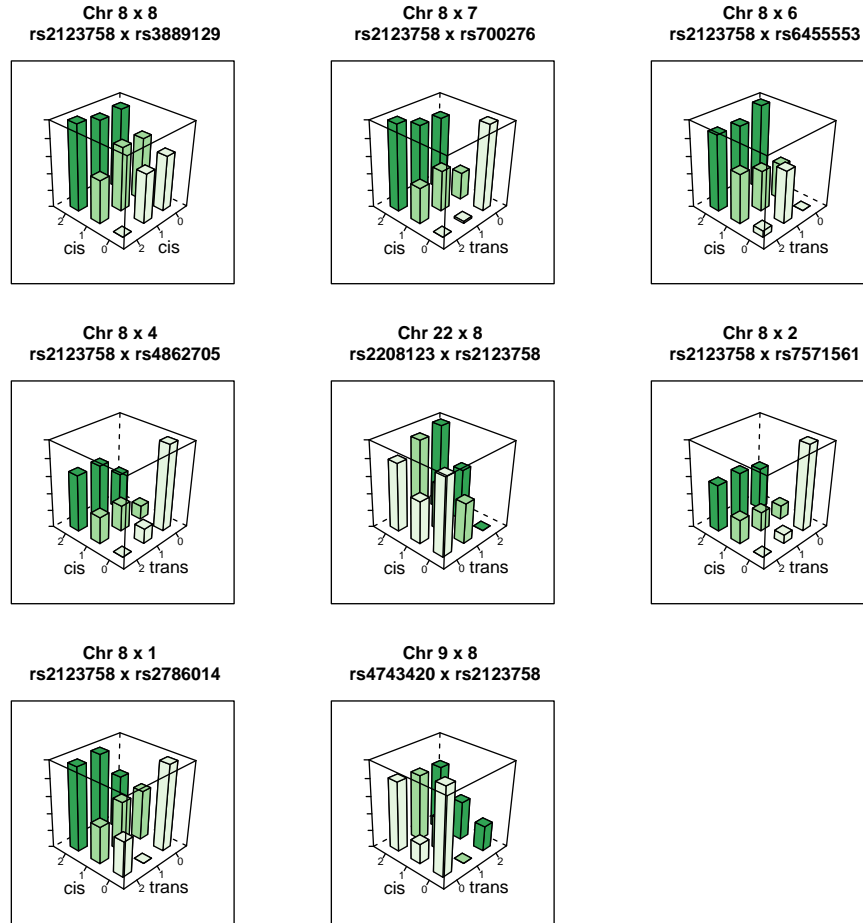


Figure S12: **Genotype-phenotype maps for 8 interactions influencing the expression of NAPRT1** Each bar represents the mean phenotypic value for individuals in that genotype class.

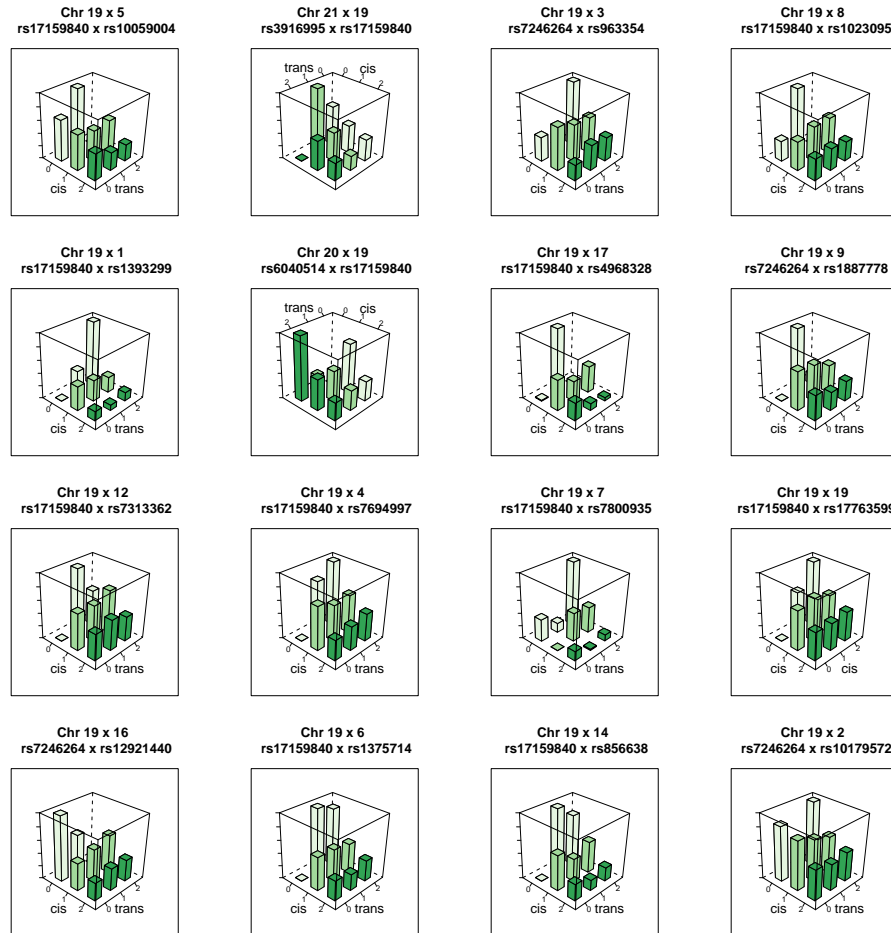


Figure S13: **Genotype-phenotype maps for 16 interactions influencing the expression of TRAPPC5** Each bar represents the mean phenotypic value for individuals in that genotype class.

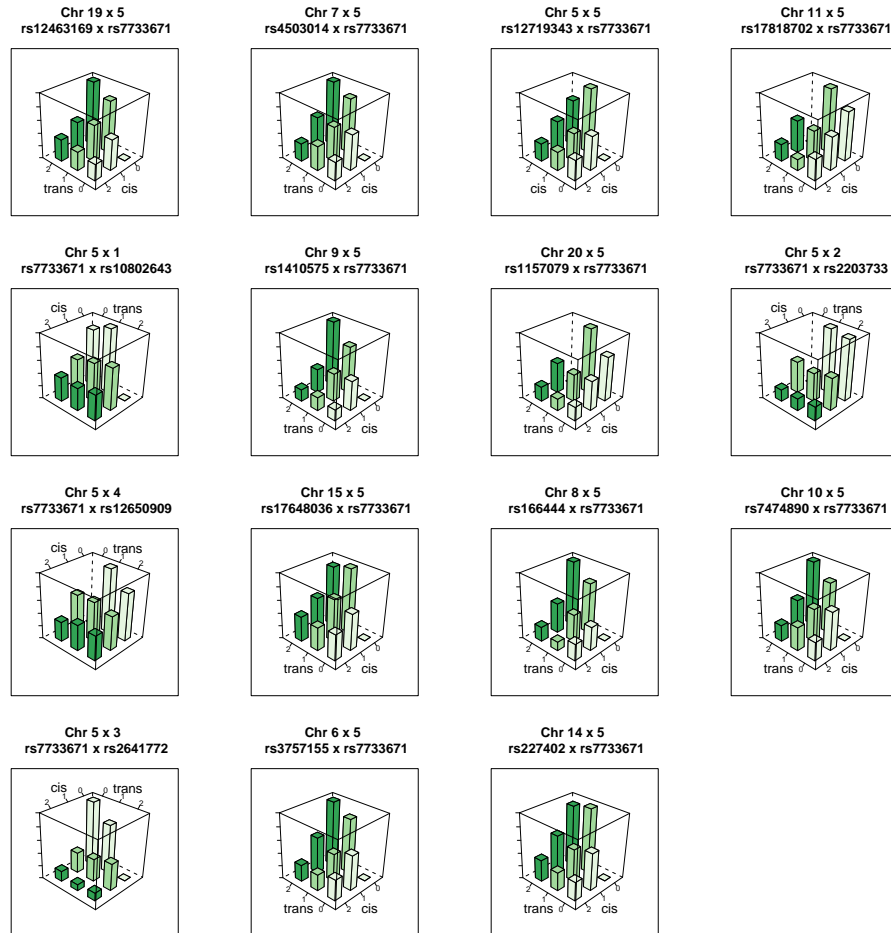
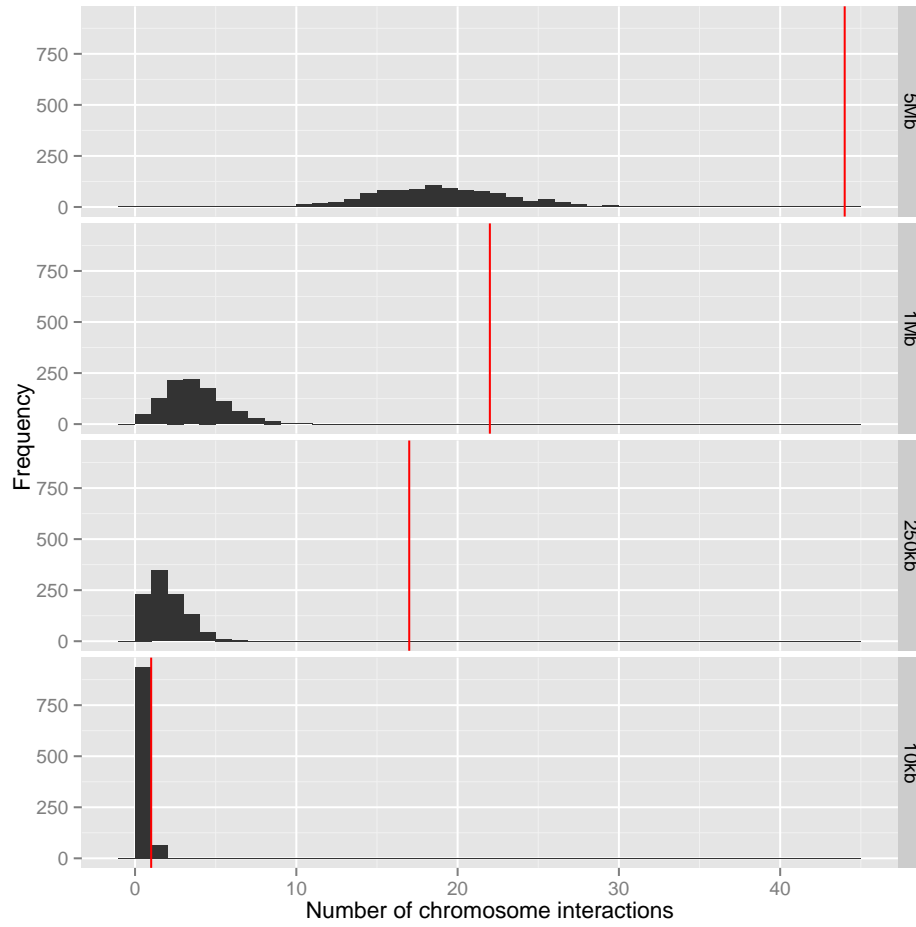


Figure S14: **Genotype-phenotype maps for 15 interactions influencing the expression of CAST** Each bar represents the mean phenotypic value for individuals in that genotype class.



**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.

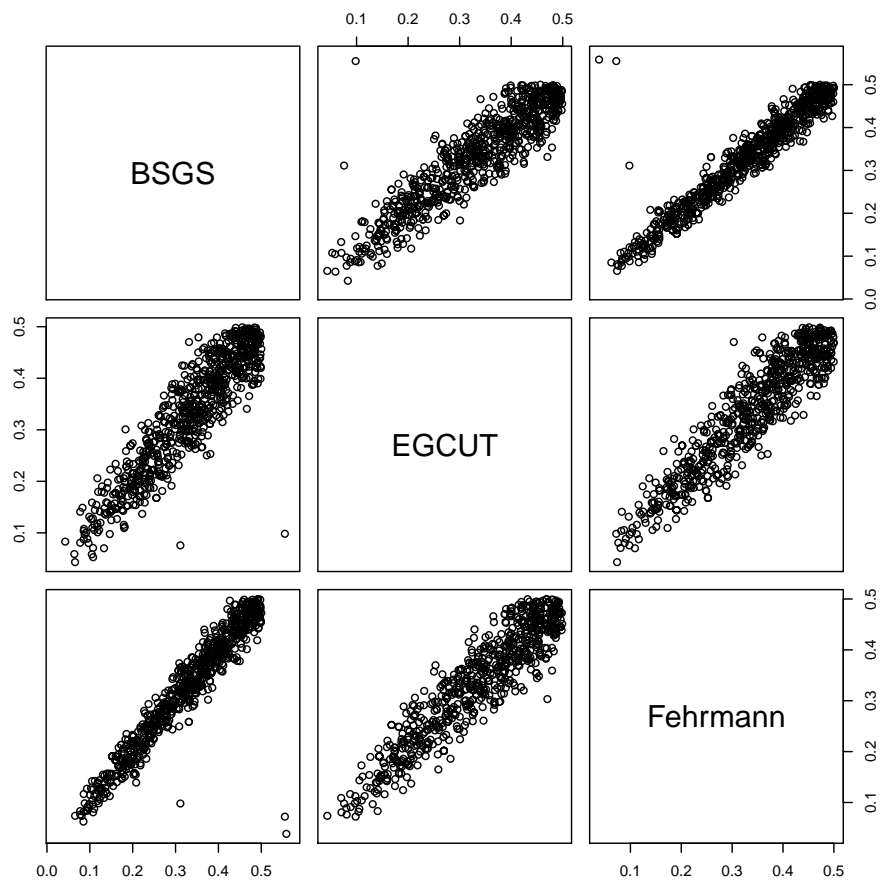
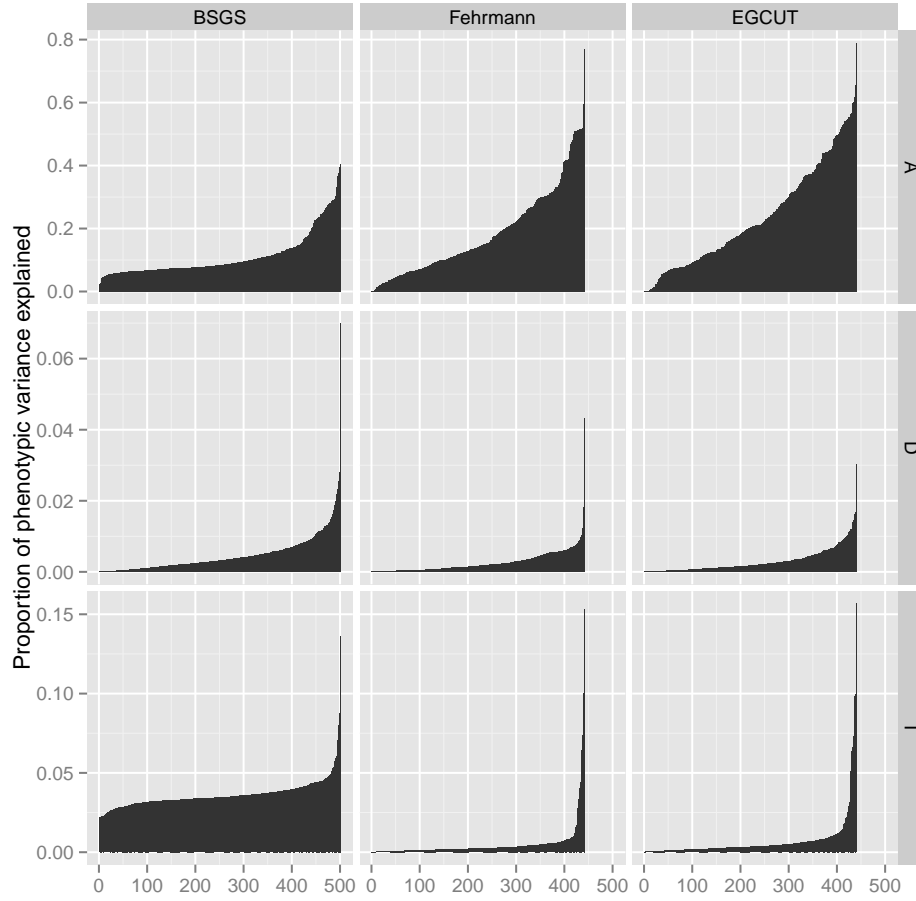


Figure S16: **Comparison of allele frequencies for 781 SNPs involved in genetic interactions across independent populations** Outliers were removed from the analysis as part of the filtering stage during replication.



**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.

292 **Supplementary Tables**

**Table S1: Details on 501 interactions discovered in BSGS dataset**

[illegible]

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Table S1 – continued from previous page

Gene ID <sup>a</sup>	Expression trait	Chr.	rs ID	Chr.	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	rs ID	Chr.	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	BSGS <sup>e</sup>	Interaction statistic <sup>f</sup>	EGCUT <sup>g</sup>	Meta <sup>g</sup>	Distance / Mb
ILMN_1653205	ILMN_1653205	8	rs8051751	16	7188323	C9ORF72	rs2896452	8	86102223	CSORF59	5.79	1.39	0.18	0.87	
ILMN_1741881	ILMN_1741881	9	rs10122902	9	27556780	C9ORF72	rs2526698	1	242029101		6.36	0.96	0.01	0.37	
ILMN_1731064	ILMN_1731064	10	rs12765847	4	4353908	CABP1	rs7338782	1	227174210	CABP1	6.36	0.94	0.00	0.34	
ILMN_1712532	ILMN_1712532	9	rs4266763	9	139289825	INPP5E	rs684040	1	82128660		5.81				
ILMN_1712532	ILMN_1712532	9	rs4576361	11	6026661		rs139266496	9	139266496	INPP5E	6.61	0.09	0.86	0.42	
ILMN_1717234	ILMN_1717234	5	rs11157079	20	6778978		rs4077515	9	96000269	CAST	7.07	0.23	0.96	0.62	
ILMN_1717234	ILMN_1717234	5	rs12463169	19	17321669		rs7733671	5	96000269	CAST	7.07	0.23	0.96	0.62	
ILMN_1599264	ILMN_1599264	16	rs12599264	16	81840122		rs7733671	15	96000269	CAST	5.73	0.02	2.85	1.75	
ILMN_1717234	ILMN_1717234	5	rs127119343	5	125369113		rs7733671	5	96000269	CAST	7.00				29.369
ILMN_1717234	ILMN_1717234	5	rs141010575	8	78255630		rs7733671	5	96000269	CAST	7.08	0.36	1.57	1.20	
ILMN_1717234	ILMN_1717234	5	rs166444	8	78392770		rs7733671	5	96000269	CAST	6.55	0.13	1.34	0.78	
ILMN_1717234	ILMN_1717234	5	rs17648036	15	27311111		rs7733671	5	96000269	CAST	7.01	0.27	0.52	0.37	
ILMN_1717234	ILMN_1717234	5	rs17818702	11	86107920		rs7733671	5	96000269	CAST	7.81	0.97	0.03	0.41	
ILMN_1717234	ILMN_1717234	5	rs227402	14	70496867		rs7733671	5	96000269	CAST	6.62	1.15	0.59	1.09	
ILMN_1717234	ILMN_1717234	5	rs2822124	21	15166804		rs7733671	5	96000269	CAST	6.12	0.11	0.01	0.01	
ILMN_1717234	ILMN_1717234	5	rs3757155	6	136458593		rs7733671	5	96000269	CAST	6.87				
ILMN_1717234	ILMN_1717234	5	rs4503014	7	31149140		rs7733671	5	96000269	CAST	7.24	0.07	0.33	0.12	
ILMN_1717234	ILMN_1717234	5	rs4744890	10	59590078		rs7733671	5	96000269	CAST	5.88	1.72	1.56	1.72	
ILMN_1717234	ILMN_1717234	5	rs7733671	5	96000269	CAST	rs10802643	1	238120177		6.74	0.49	0.12	0.23	
ILMN_1717234	ILMN_1717234	5	rs7733671	5	96000269	CAST	rs12650909	4	170192890		7.42	0.75	0.78	0.75	
ILMN_1717234	ILMN_1717234	5	rs7733671	5	96000269	CAST	rs2203733	2	224093101		7.42	0.23	0.78	0.54	
ILMN_1717234	ILMN_1717234	5	rs7733671	5	96000269	CAST	rs2641772	3	195531841		6.93	0.22	0.87	0.54	
ILMN_1651705	ILMN_1651705	11	rs872311	18	66175386		rs11032695	11	34447586	CAT	6.41	0.26	0.30	0.22	
ILMN_1722208	ILMN_1722208	11	rs3253203	19	17099980		rs541207	11	64125142	CDC88B	5.68	0.33	0.37	0.31	
ILMN_1722208	ILMN_1722208	11	rs694739	11	70997233	CDC88B	rs12771349	10	96998193		5.62	0.23	0.18	0.14	

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Gene ID <sup>a</sup>	Expression trait	Chr.	rs ID	Chr.	SNP 1	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	rs ID	Chr.	SNP 2	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	BSGS <sup>e</sup>	Interaction statistic <sup>f</sup> – Fehrmann <sup>g</sup>	– log <sub>10</sub> p-values	Metag <sup>h</sup>	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	rs6139887	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	rs6137908	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.204985	3	rs6021982	20	33651928	20	33651928	rs7591849	2	219650616	CYBRD1	5.42	0.29	0.86	0.60		
DAB2	ILMN.2128428	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		
DRH1	ILMN.1733998	12	rs1519956	12	89468283	12	89468283	rs7566044	2	169960422	DRH1	6.48	0.37	0.			

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Table S1 – continued from previous page

Gene ID <sup>a</sup>	Expression trait <sup>b</sup>	Chr.	SNP 1			SNP 2			Interaction statistic / -log <sub>10</sub> p-values			Distance / Mb <sup>h</sup>		
			rs ID	Chr.	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	rs ID	Chr.	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	BSGS <sup>e</sup>		Fehrmann <sup>f</sup>	EGCUT <sup>g</sup>
FEZ2	ILMN-1739586	2	rs2356400	19	44321776		rs13406184	2	36791226	FEZ2	5.78	0.14	0.33	0.16
FEZ2	ILMN-1739586	2	rs969010	4	159963132		rs3810133	2	36810133	FEZ2	6.59	0.14	0.28	0.14
FGD2	ILMN-2115005	6	rs4803848	19	46205050		rs831486	6	37001267	FGD2	5.69	0.12	0.25	0.11
FGD2	ILMN-2115005	6	rs902634	10	133943951		rs831489	6	36999682	FGD2	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	rs3782908	12	48169526		rs875111	4	167695661	FLJ20489	5.53	0.03	0.11	0.02
FLJ20489	ILMN-1778144	12	rs4792199	17	7992118		rs3782908	12	48169526	FLJ20489	5.74	0.19	0.02	0.04
FLJ20489	ILMN-1778144	12	rs4984440	15	97033129		rs3782908	12	48169526	FLJ20489	6.49	0.31	0.47	0.36
FLJ20489	ILMN-1778144	12	rs7204135	16	50626195		rs3782908	12	48169526	FLJ20489	6.90	0.38	0.17	0.21
FLJ20718	ILMN-1763663	16	rs9325634	21	43818790		rs2827197	16	50106594	FLJ20718	6.04	0.14	0.95	0.53
FLJ43093	ILMN-2123450	6	rs17112712	14	107276627	FLJ43093	rs6906101	6	36667610	FLJ43093	5.48	0.39	0.06	0.13
FN3KRP	ILMN-2123450	6	rs6906101	6	36667610		rs13214069	6	32705248	FLJ43093	5.44	0.00	0.64	0.18
FN3KRP	ILMN-1652333	17	rs898095	17	80890638		rs9892064	17	80827903		16.16	28.24	29.39	59.95
FUCA1	ILMN-1732728	1	rs4971478	2	1346063		rs12744386	1	24168019	FUCA1	6.41	0.01	0.30	0.06
FUCA1	ILMN-2309848	19	rs1633921	19	35695200		rs788178	13	98328559		3.70	0.09	0.41	0.17
FXYD5	ILMN-2309848	19	rs17398183	20	35609148		rs2285515	19	35660450	FXYD5	6.58	0.03	0.48	0.15
FXYD5	ILMN-2309848	19	rs2285515	19	35660450	FXYD5	rs11739594	5	141709563		5.70	0.07	0.17	0.05
FXYD5	ILMN-2309848	19	rs2285515	19	35660450	FXYD5	rs13067700	3	95331048		6.00	0.09	0.51	0.22
FXYD5	ILMN-2309848	19	rs2285515	19	35660450	FXYD5	rs17036504	2	47567329		6.10	0.28		
G3BP2	ILMN-231758	4	rs10230232	17	29390239		rs1553985	4	76554604		5.19	0.08	0.37	0.14
GAA	ILMN-2410783	17	rs11190847	17	78153130		rs12602462	17	78146016		13.91	19.98	12.99	32.60
GAA	ILMN-2410783	17	rs8068856	17	78100731	GAA	rs10902506	12	132678089		5.65	0.11	0.39	0.17
GAPT	ILMN-1675191	5	rs10070522	5	57786110	GAPT	rs7605821	2	235695228		5.85	0.01	0.78	0.28
GAPT	ILMN-1696311	7	rs7082031	10	128038717		rs10070522	5	57786110	GAPT	5.72	0.26	0.11	0.11
GATS	ILMN-1696311	7	rs1147447	14	66460742		rs2950520	7	99827148	GATS	5.47	0.83	0.63	0.87
GATS	ILMN-1696311	7	rs2423256	16	30565672		rs2950520	7	99827148	GATS	6.37	0.38	0.35	0.33
GDPD3	ILMN-174901	16	rs3809624	16	30102802	GDPD3	rs2197465	14	48572632		5.86	0.55	0.09	0.24
GDPD3	ILMN-174901	16	rs7204270	16	31056963	GDPD3	rs1015111	4	128972357		5.78	0.02	0.45	0.13
GNLY	ILMN-1790692	13	rs4145072	13	11089955		rs7577293	2	85935282	GNLY	5.72	0.11	1.16	4.237
GNLY	ILMN-3239426	12	rs7198646	16	26084476		rs7060552	12	111164237	GNLY	5.72	0.36	0.46	0.39
GPR162	ILMN-1730816	12	rs1860563	16	6478898		rs2707210	12	6902002	GPR162	5.49	0.25	0.03	0.06
GPR162	ILMN-1730816	12	rs2272500	12	79685913		rs2707210	12	6902002	GPR162	5.07	0.25	0.06	0.07
GPR162	ILMN-1730816	12	rs2707210	12	6902002		rs4740848	9	6554558		5.47	0.96	0.06	0.44
GPR177	ILMN-1730816	12	rs2707210	12	6902002	GPR162	rs9827054	3	188880113		5.45	0.72	0.67	0.81
GPR177	ILMN-1660549	1	rs11057383	12	124369421		rs12065581	1	68732819	GPR177	5.76	0.17	0.40	0.22
GPR177	ILMN-1660549	1	rs12527241	6	120408039		rs12065581	1	68732819	GPR177	5.76	0.17	0.40	0.22
GPR177	ILMN-1660549	1	rs12532999	7	127939793		rs12065581	1	68732819	GPR177	6.50	0.79	1.43	1.50
GPR177	ILMN-1660549	1	rs725613	16	11169683		rs12065581	1	68732819	GPR177	5.43	0.31	0.11	0.13
GPR177	ILMN-1660549	1	rs9575097	13	82986268		rs12065581	1	68732819	GPR177	6.04	0.95	0.21	0.60
GPR177	ILMN-2283325	1	rs6566669	18	70506011		rs12065581	1	68732819	GPR177	5.86	0.24	0.34	0.23
GPR177	ILMN-2283325	1	rs9290426	3	171399321		rs12065581	1	68732819	GPR177	6.50	0.24	0.24	0.04
GPR177	ILMN-2347193	17	rs11557467	17	38028634		rs4965745	15	101508261		5.88	0.68	0.20	0.41
GSDMB	ILMN-2391861	1	rs12248673	10	83192833		rs11101992	1	110266754	GSTM1	6.11	0.27	0.19	0.16
GSTM1	ILMN-2391861	1	rs1547574	13	85344527		rs3754446	1	110253241	GSTM1	5.91	0.27	1.14	0.79
GSTM2	ILMN-2201580	1	rs6492807	13	96150560		rs453333	2	77919015		6.77	0.52	0.66	0.65
H1FO	ILMN-1757467	22	rs139898	22	38399979		rs6497007	15	85877017		6.36	0.27	0.31	0.23
H1FO	ILMN-1757467	22	rs139898	22	38399979		rs9983949	21	19532546		6.52	0.25	0.48	0.32
H1FO	ILMN-1757467	22	rs139898	22	38399979		rs2855039	11	5271671	HBC2	5.47	0.00	0.66	0.19
HBC1	ILMN-1796678	11	rs11078523	11	4523167		rs2855039	11	5271671	HBC2	5.98	0.15	0.24	0.12
HBC1	ILMN-1796678	11	rs12975066	19	35723501		rs2855039	11	5271671	HBC2	6.78	0.08	0.52	0.21
HBC1	ILMN-1796678	11	rs2855039	11	5271671	HBC2	rs12042181	1	213088494	LQK1	6.42	0.01	0.46	0.10
HBC1	ILMN-1796678	11	rs2855039	11	5271671	HBC2	rs12503379	4	141533832		6.06	0.01	0.41	0.10
HBC2	ILMN-2084825	11	rs11078523	17	4523167		rs16912979	11	5309695	HBC2	6.06	0.01	0.41	0.10

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Table S1 – continued from previous page

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Table S1 – continued from previous page

Gene ID <sup>a</sup>	Expression trait	SNP 1			SNP 2			Interaction statistic / -log <sub>10</sub> p-values				Distance / Mb <sup>b</sup>	
		rs ID	Chr.	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	rs ID	Chr.	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	BSCS <sup>e</sup>	Fehrmann <sup>f</sup>		EGCUT <sup>g</sup>
NRBF2	ILMN-3237385	rs6025645	20	56157341		rs7923609	10	65133822	NRBF2	5.45			
NRBF2	ILMN-3237385	rs6017815	21	19819016		rs7923609	10	65133822	NRBF2	6.11	0.47	0.05	0.17
NRD1	ILMN-1800897	rs4852124	2	240080022		rs6588415	1	52334047	NRD1	6.13	0.03	0.46	0.15
NUDT18	ILMN-1787885	rs5017351	11	25353442		rs1005901	8	21964378	NUDT18	5.44	4.27	1.35	9.03
OAS1	ILMN-1658247	rs11613438	12	113480510		rs1047944	6	163994767		8.59	0.43	0.81	3.86
OAS1	ILMN-1658247	rs113311	12	113480510		rs2072133	12	113409260		4.13	4.12	0.81	3.86
OAS1	ILMN-1675640	rs2892233	19	49160255		rs3741981	12	13169066	OAS1	5.64	0.87	0.46	0.14
OPIN	ILMN-2381899	rs7192613	16	74286646		rs17312962	10	13169066	OPIN	5.00	0.36	0.00	0.07
OSBPL5	ILMN-2307032	rs2829679	21	26625433		rs998639	11	3149249	OSBPL5	5.42	0.16	0.87	0.49
OSTF1	ILMN-1742456	rs17780195	9	77755469	OSTF1	rs2273770	9	77755469	OSTF1	5.42	1.20	0.08	0.62
OSTF1	ILMN-1742456	rs2273770	9	77755469		rs1264898	5	119928923	OVGP1	5.43	0.13	1.48	0.88
OVGP1	ILMN-1734542	rs10802822	1	240132968		rs1264894	1	111969719	OVGP1	6.04	0.25	1.21	0.82
OVGP1	ILMN-1734542	rs347351	3	104148107	PAM	rs784600	1	40139553	HPCAL4	5.59	0.66	0.44	0.59
PAM	ILMN-2313901	rs28092	5	102149795	PCYOX1L	rs2731939	3	21395989		6.20	0.19	0.26	0.16
PCYOX1L	ILMN-1815951	rs24358490	5	148726162		rs4328748	12	7364442	PEX5	5.85	0.09	0.71	0.32
PRX5	ILMN-1660232	rs10444467	12	128052636		rs4328748	12	7364442	PEX5	5.74	0.34	0.09	0.13
PRX5	ILMN-1660232	rs7495797	15	27246462		rs4329748	12	7364442	PFAAP5	5.64	0.87	0.36	0.67
PFAAP5	ILMN-1797893	rs131969	22	49151303	PGLYRP1	rs1263806	14	21982957	PFAAP5	5.51	0.03	0.65	0.24
PGLYRP1	ILMN-1704870	rs14298233	19	46529456		rs10736812	11	76708086	PHCA	5.90	0.36	0.90	0.70
PHCA	ILMN-1812552	rs495642	11	123097386	PIK3P1	rs2065841	1	61728597		7.11	0.02	0.87	0.33
PIK3P1	ILMN-1719986	rs4141404	22	31675185	PISD	rs10498313	14	30398876		5.23	0.20	0.81	0.03
PISD	ILMN-1739934	rs615752	22	332334931	PISD	rs1465754	22	32097775	PISD	4.12	0.00	1.19	0.48
PISD	ILMN-1739934	rs715572	22	332334931		rs6518754	22	32097775	PISD	7.11	0.05	0.42	0.15
PNKD	ILMN-1774604	rs6869411	9	458781604		rs928284	2	219182481	PNKD	6.35	0.16	0.04	0.04
PNPLA7	ILMN-1662587	rs11639998	16	15271109		rs928284	9	140487108	PNPLA7	5.15	0.31	0.78	0.56
PPF1BP2	ILMN-1662587	rs910119	20	49668255		rs4758001	11	7559930	PPF1BP2	4.44	0.29	0.33	0.26
PP2P2R3C	ILMN-1662617	rs12914603	15	5830896		rs11156875	14	35619816	PP2P2R3C	5.81	0.12	0.42	0.19
PP2P2R5A	ILMN-1738784	rs10930170	2	163999467		rs12120009	1	212447167	PP2P2R5A	5.63	0.72	0.48	0.66
PP2P2R5A	ILMN-1738784	rs124243255	12	123595064		rs12120009	1	212447167	PP2P2R5A	5.72	0.08	0.95	0.46
PP2P2R5A	ILMN-1738784	rs1889083	13	66222691		rs12120009	1	212447167	PP2P2R5A	5.61	0.36	0.13	0.17
PP2P2R5A	ILMN-1738784	rs6822334	11	107417238		rs12120009	1	212447167	PP2P2R5A	5.65	1.69	0.28	1.21
PP2P2R5A	ILMN-1738784	rs757871	6	135030045		rs12120009	1	212447167	PP2P2R5A	5.95	0.37	0.06	0.16
PP2P2R5A	ILMN-1738784	rs7871178	9	27148475		rs12120009	1	212447167	PP2P2R5A	5.72	0.16	0.30	0.16
PRDX5	ILMN-1716069	rs8019823	14	95040482		rs11600909	11	64082807	PRDX5	6.43	0.81	0.14	0.44
PRKCB1	ILMN-1713603	rs2188355	16	23867776		rs10492793	16	12639800		7.34	0.53	0.11	0.25
PRMT2	ILMN-1675038	rs1029231	21	47931653	C21ORF57	rs958127	18	31497346	C21ORF57	4.81	0.19	0.03	0.04
PRMT2	ILMN-1675038	rs2382372	21	48063862		rs11701058	21	47776382	PSMB1	5.79	0.44	0.26	0.04
PSMB1	ILMN-1789176	rs38962607	18	43983954		rs13207114	6	170877444	PSMB1	5.14	0.44	0.21	0.27
PSMB1	ILMN-1789176	rs6060930	20	30347832		rs6928843	6	170890384	PSMB1	5.44	1.95	0.64	1.78
PSMB1	ILMN-1789176	rs6928843	6	170890384	PSMB1	rs2965415	6	225797957	PSMB1	5.42	1.18	0.32	0.86
PSMB1	ILMN-1789176	rs7298749	12	131727816		rs13207114	6	170877444	PSMB1	5.00	0.03	0.48	0.15
PWPI	ILMN-1743049	rs4969205	17	76598123		rs11036212	11	5221825	PTDSS1	5.42	1.03	0.32	0.86
PWPI	ILMN-1743049	rs631562	11	126852438		rs11036212	11	5221825	PTDSS1	5.90	0.80	0.08	0.38
QDPR	ILMN-1672443	rs4946705	6	106348246		rs10020773	4	17526682	QDPR	5.70	0.02	0.40	0.11
RAB3P	ILMN-1803197	rs1075728	22	33375704		rs7305037	12	70235726		5.75	1.03	1.25	1.55
RAB3P	ILMN-2207363	rs9931702	19	42462788	RAB3P	rs79051628	11	120161117		6.55	0.25	0.08	0.09
RBL2	ILMN-1756999	rs9931702	16	53526551	AKTIP	rs1863464	15	26938488		6.42	0.28	0.84	0.59
RCN1	ILMN-1800276	rs10879131	12	41147155		rs4922579	11	32136436		6.38	0.03	0.31	0.08
RCN1	ILMN-1800276	rs4922579	11	32136436	RCN1	rs11166957	8	141177468		5.23	0.58	0.37	0.43
RCN1	ILMN-1800276	rs4922579	11	32136436	RCN1	rs1341899	1	102740645		4.32	0.41	0.09	0.17

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Table S1 – continued from previous page

Gene ID <sup>a</sup>			Expression trait			SNP 1			SNP 2			Interaction statistic <sup>f</sup>			BSCS <sup>e</sup>			-log <sub>10</sub> p-values			Distance / Mb <sup>b</sup>		
Gene	ID <sup>a</sup>	Chr.	rs ID	Chr.	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	rs ID	Chr.	Pos/Mb <sup>c</sup>	Association <sup>d</sup>	rs ID	Chr.	Pos/Mb <sup>c</sup>	F <sub>max</sub>	F <sub>max</sub> p-value	SNP1	SNP2	SNP1	SNP2	SNP1	SNP2	SNP1	SNP2
RENE	ILMN_1802830	1	rs4982958	14	24987865		rs301819	1	8501786	RENE	rs301819	1	8501786	5.66	0.61	1.23	1.17						
RENE	ILMN_1802838	1	rs7697290	4	135248366		rs301819	1	8501786	RENE	rs301819	1	8501786	5.74	0.14	0.10	0.06						
RENE	ILMN_1802840	1	rs11085629	19	13174312		rs301819	1	8501786	RENE	rs301819	1	8501786	5.12	0.21	0.33	0.21						
RENE	ILMN_1802842	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802844	1	rs10218598	14	21182850	RNASE6	rs10218598	13	100601327	RNASE6	rs10218598	13	100601327	5.48	0.42	0.21	0.26						
RENE	ILMN_1802846	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802848	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802850	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802852	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802854	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802856	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802858	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802860	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802862	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802864	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802866	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802868	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802870	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802872	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802874	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802876	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802878	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802880	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802882	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802884	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802886	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802888	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802890	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802892	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802894	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802896	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802898	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802900	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802902	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802904	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802906	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802908	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802910	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802912	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802914	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802916	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802918	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802920	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802922	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802924	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802926	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802928	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802930	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802932	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802934	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802936	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802938	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802940	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802942	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802944	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802946	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802948	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802950	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802952	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802954	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802956	1	rs38262011	3	12844086		rs301819	1	8501786	RENE	rs301819	1	8501786	5.71	0.08	0.60	0.26						
RENE	ILMN_1802958	1	rs38262011	3	12844086		rs3018																

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Table S1 – continued from previous page

Gene ID <sup>a</sup>		Expression trait		SNP 1		SNP 2		Interaction statistic / -log <sub>10</sub> p-values		Distance / Mb <sup>h</sup>						
TMEMD4	Probe ID <sup>b</sup>	Chr.	rs ID	Chr.	Pos / Mb <sup>c</sup>	Association <sup>d</sup>	rs ID	Chr.	Pos / Mb <sup>c</sup>		Association <sup>d</sup>	BSGS <sup>e</sup>	Fehrmann <sup>f</sup>	EGCUT <sup>g</sup>	Meta <sup>g</sup>	
TMEMD4	ILMN-17804148	7	rs19340400	11	132389627	SNX26	rs17725246	7	44581986	TMEMD4	3.70	0.06	1.34	0.70	0.122	
TMEM149	ILMN-1780426	19	rs28390113	21	47248981		rs81069359	19	36219525	TMEM149	8.11	0.16	0.48	0.26		
TMEM149	ILMN-1780426	19	rs5762235	22	27925288		rs10609359	19	36219525	TMEM149	6.79					
TMEM149	ILMN-1780426	19	rs6090518	20	43207005		rs81069359	19	36219525	TMEM149	11.09	0.76	45.78	145.78		
TMEM149	ILMN-1780426	19	rs807491	19	36268923		rs10609359	19	36219525	TMEM149	12.16	81.55	3.09	3.07		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525		rs10609359	19	36219525	TMEM149	8.12	1.55	0.99	0.80		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525		rs10609359	19	36219525	TMEM149	8.02	0.40	0.99	0.80		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525		rs10609359	19	36219525	TMEM149	8.39	3.61	1.18	3.78		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525		rs10609359	19	36219525	TMEM149	7.37	2.41	1.00	2.52		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525		rs10609359	19	36219525	TMEM149	6.93	0.08	0.07	0.03		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs17719594	14	90932398	TMEM149	6.95	3.06	0.77	2.87		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs1843357	8	13822381	TMEM149	6.21	3.72	3.33	6.00		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs2351458	4	113317583	TMEM149	7.30	0.04	9.61	8.00		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs2359000	7	147619772	TMEM149	6.70	1.57	1.52	2.27		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs2731711	5	171792273	TMEM149	5.92	0.19	0.33	0.19		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs471728	11	12959460	TMEM149	8.89	0.90	3.62	3.51		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs6718480	2	23387906	TMEM149	8.55	3.31	5.15	7.36		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs6926382	6	161683974	TMEM149	5.80	3.06	8.80	10.72		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs7213338	17	80357420	TMEM149	5.49	0.07	3.14	2.10		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs914940	1	242889492	TMEM149	6.22	3.36	6.96	9.20		
TMEM149	ILMN-1780426	19	rs81069359	19	36219525	TMEM149	rs9509428	13	21473952	TMEM149	9.44	0.10	5.75	4.47		
TMEM63A	ILMN-1719649	1	rs1254086	13	72890603	SNX26	rs1419226	1	226027323	TMEM63A	5.60				0.031	
TMEM80	ILMN-1708482	11	rs1548475	9	58058246		rs4963126	11	65845	TMEM80	5.79	0.64	0.12	0.32		
TMEM80	ILMN-17083811	7	rs1537146	9	4859303		rs10488630	7	128593948	IRF5	5.61	0.11	0.15	0.07		
TMEM80	ILMN-17083811	7	rs199793	20	22287303		rs10488630	7	128593948	IRF5	5.52	1.03	0.17	0.62		
TMEM80	ILMN-17083811	7	rs199793	20	22287303		rs11770192	7	23498358	IRF5	8.23	3.19	1.89	4.09		
TRAPPC4	ILMN-17130443	7	rs7776572	13	133531675		rs3916581	11	118887887	TRAPPC4	5.61	0.28	0.40	0.29		
TRAPPC4	ILMN-1814650	11	rs1278760	13	131018917		rs3916581	11	118887887	TRAPPC4	5.52	0.93	0.01	0.36		
TRAPPC5	ILMN-1814650	11	rs17159840	19	7758194		rs10059004	5	166970604	TRAPPC5	5.97	0.21	1.60	1.07		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194		rs1023095	8	132022957	TRAPPC5	6.92	0.37	0.87	0.68		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194		rs1375714	6	156404902	TRAPPC5	7.79	0.12	0.18	0.09		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs1393299	1	242329791	TRAPPC5	6.43	0.63	0.47	0.58		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs17763599	19	2369415	TRAPPC5	6.38	0.21	0.24	0.16		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs4968328	17	57495457	TRAPPC5	6.51	0.50	0.38	0.44		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs7313362	12	129644342	TRAPPC5	7.08	0.04	0.65	0.25		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs7313362	4	9947811	TRAPPC5	5.86	0.20	0.36	0.22		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs7809035	7	146690926	TRAPPC5	6.27	0.15	0.33	0.16		
TRAPPC5	ILMN-2372639	19	rs17159840	19	7758194	TRAPPC5	rs856638	14	85439550	TRAPPC5	6.73	0.24	0.07	0.08		
TRAPPC5	ILMN-2372639	19	rs380708	22	22740855	SNX26	rs17159840	19	7758194	TRAPPC5	7.58				5.389	
TRAPPC5	ILMN-2372639	19	rs3916995	19	45128455		rs17159840	19	7758194	TRAPPC5	7.73	0.85	0.78	1.01		
TRAPPC5	ILMN-2372639	19	rs6040514	20	11272861		rs17159840	19	7758194	TRAPPC5	8.10	0.51	0.55	0.56		
TRAPPC5	ILMN-2372639	19	rs7246264	19	7762978		rs10179572	2	228504503	TRAPPC5	6.71	0.02	0.02	0.02		
TRAPPC5	ILMN-2372639	19	rs7246264	19	7762978		rs12921440	16	30408765	TRAPPC5	7.34	0.14	0.26	0.13		
TRAPPC5	ILMN-2372639	19	rs7246264	19	7762978		rs1887778	3	154635088	TRAPPC5	7.05	0.08	0.86	0.40		
TRAPPC5	ILMN-2372639	19	rs7246264	19	7762978		rs963354	3	157393770	TRAPPC5	7.41	0.36	0.90	0.69		
TRAPPC5	ILMN-2372639	19	rs7246264	19	7762978		rs2395771	6	41264577	TRAPPC5	5.42	0.11	0.25	0.11		
TRAPPC5	ILMN-2372639	19	rs10862975	12	85749398		rs2032447	6	26044369	TRAPPC5	5.92	1.20	1.23	1.69		
TRAPPC5	ILMN-2372639	19	rs12412964	7	158808412		rs10748536	10	82273079	TRAPPC5	6.46	0.04	0.91	0.39		
TRAPPC5	ILMN-1697971	6	rs2527180	17	17194634	rs12800935	11	2317951	TRAPPC5	5.01	0.07	0.18	0.06			
TRAPPC5	ILMN-1780606	17	rs968726	17	27635049	SNX26	rs12800935	11	2317951	TRAPPC5	5.01				45.345	
TRAPPC5	ILMN-1718621	11	rs10838738	11	47605049		rs12800935	11	2317951	TRAPPC5	5.01					
TRAPPC5	ILMN-1718621	11	rs10838738	11	47605049		rs12800935	11	2317951	TRAPPC5	5.01					
TRAPPC5	ILMN-2329970	22	rs12800698	22	50971266		rs12800935	11	2317951	TRAPPC5	5.01					
TRAPPC5	ILMN-2329970	22	rs12800698	22	50971266		rs12800935	11	2317951	TRAPPC5	5.01					
TRAPPC5	ILMN-2329970	22	rs12800698	22	50971266		rs12800935	11	2317951	TRAPPC5	5.01					
TRAPPC5	ILMN-2329970	22	rs12800698	22	50971266		rs12800935	11	2317951	TRAPPC5	5.01					
TRAPPC5	ILMN-2329970	22	rs12800698	22	50971266		rs12800935	11	2317951	TRAPPC5	5.01					
TRAPPC5	ILMN-2329970	22	rs12800698	22	50971266		rs12800935	11	2317951	TRAPPC5	5.01					
TRAPPC5	ILMN-2329970	22	rs12800698	22	50971266		rs12800935	11	2317951	TRAPPC5	5.01					

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Table S1 – continued from previous page

Expression trait			SNP 1			SNP 2			Interaction statistic <sup>f</sup> / -log <sub>10</sub> p-values			Distance / Mb <sup>h</sup>		
Gene ID <sup>a</sup>	Probe ID <sup>b</sup>	Chr.	rs ID	Chr.	Pos / Mb <sup>c</sup>	Association <sup>d</sup>	rs ID	Chr.	SNP-2	Association <sup>d</sup>	BSGS <sup>e</sup>	Fehrmann <sup>f</sup>	EGCUT <sup>g</sup>	Meta <sup>g</sup>
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	5.04	0.84	0.15	0.46
VNN2	LMN-1678939	6	rs13044386	20	9116155		rs1883617	6	133072650	VNN2	5.44	0.39	0.69	0.57
VNN2	LMN-1678939	6	rs134447	22	49927332		rs1883617	6	133072650	VNN2	5.72			
VNN3	LMN-1678939	6	rs216495	11	16834510		rs1883617	6	133072650	VNN2	5.77	0.33	0.19	0.19
VNN3	LMN-1678939	6	rs10278073	7	151662184		rs2267952	6	133067782	VNN3	6.44	0.16	0.74	0.41
VNN3	LMN-1804935	6	rs1443946	8	73006453		rs2267952	6	133067782	VNN3	5.74	0.23	0.48	0.31
VNN3	LMN-1804935	6	rs348462	9	75547169		rs2267952	6	133067782	VNN3	6.44	0.31	0.17	0.17
VNN3	LMN-1804935	6	rs7157055	14	83262064		rs2267952	6	133067782	VNN3	5.82	0.03	0.19	0.04
VNN3	LMN-2387680	6	rs2823165	21	5694253		rs2267952	6	133067782	VNN3	6.12	0.73	1.15	1.21
VNN3	LMN-2387680	6	rs9596457	13	51692548		rs2267952	6	133067782	VNN3	4.83	0.46	0.05	0.16
VSTM1	LMN-1763455	19	rs9596457	19	54553697	VSTM1	rs4532100	18	71024750		5.60	0.53	0.54	0.57
VSTM1	LMN-1763455	19	rs10500316	19	54553697	VSTM1	rs7895870	10	123095249		5.71	0.48	0.17	0.26
VSTM1	LMN-1763455	19	rs10500316	19	54553697	VSTM1	rs7895870	10	123095249		5.71	0.48	0.17	0.26
VSTM1	LMN-1763455	19	rs9628570	22	30261219		rs10500316	19	54553697	VSTM1	5.88	0.81	1.38	1.47
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		rs11715581	3	49194331	WDR6	4.86	1.64	1.43	2.25
XAF1	LMN-2330573	17	rs1535031	21	36773170	XAF1	rs12591171	15	68119799		5.48	2.38	0.17	1.63
ZEP00	LMN-1684628	16	rs906446	21	37040648		rs12591171	15	68119799	ZEP00	5.79	0.09	0.36	0.15
ZNF500	LMN-1700258	16	rs4282793	22	48283177		rs12591171	15	68119799	ZEP00	5.79	0.09	0.36	0.15
ZNF500	LMN-1700258	16	rs4282793	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

<sup>a</sup> Phenotypes are expression levels of RefSeq Genes<sup>b</sup> Illumina probe ID used to measure gene expression<sup>c</sup> Physical SNP position in base pairs (HG19)<sup>d</sup> RefSeq Gene ID of gene expression level that is influenced by the SNP (BSGS discovery dataset, significance threshold = 1.29 × 10<sup>-11</sup>)<sup>e</sup> Interaction - log<sub>10</sub> p-value from discovery dataset<sup>f</sup> Interaction - log<sub>10</sub> p-value from replication dataset<sup>g</sup> Interaction - log<sub>10</sub> p-value from meta analysis of replication datasets only<sup>h</sup> Distance in Mb between interacting SNPs for *cis-cis* acting SNP pairs<sup>i</sup> p-values are absent if the interaction did not pass the QC filtering in the replication dataset<sup>j</sup> 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 2013<sup>22</sup>

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

<sup>a</sup> “All” denotes 434 discovery interactions and “Significant” denotes 30 interactions with significant replication  $p$ -values

<sup>b</sup> Number of tests for concordance

<sup>c</sup> Expected number of concordant cases under the null hypothesis of no interactions

<sup>d</sup> Observed number of concordant cases

<sup>e</sup> The sign of the most significant epistatic variance component in discovery is the same as the corresponding variance component in the replication data.

<sup>f</sup> The largest epistatic variance component in the discovery is the same as in the replication with the same sign in both.

<sup>g</sup> 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 <sup>a</sup>	Dataset	$n^b$	0 <sup>c</sup>	1 <sup>c</sup>	2 <sup>c</sup>	3 <sup>c</sup>	4 <sup>c</sup>	$p$
Expected <sup>d</sup>	-	-	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	$4.72 \times 10^{-4}$
Significant	Fehrman	30	0.03	0.07	0.33	0.27	0.30	$6.69 \times 10^{-4}$
Significant	Combined	60	0.05	0.05	0.32	0.30	0.28	$5.49 \times 10^{-8}$

<sup>a</sup> “All” denotes 434 discovery interactions and “Significant” denotes 30 interactions with significant replication  $p$ -values.

<sup>b</sup> Number of tests for concordance.

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

<sup>d</sup> 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