

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 some examples have been reported in other species,⁶ few convincing examples 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 ($p < 2.91 \times 10^{-16}$). We tested the discovery interactions for replication in two independent data sets.^{11,12} Three hundred and forty-five interactions had replication interaction p -values that were more extreme than the 2.5% confidence interval of the distribution under the null hypothesis of no epistasis, with 30 significant at a conservative $p < 0.05$ Bonferroni level. There was evidence of functional enrichment for the interacting SNPs, for instance 44 of the genetic interactions are located within 2Mb of regions of known intra-cellular 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 epistatic genetic effects emerging from natural genetic variation in humans.

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 there is little empirical exploration of the role that epistasis plays in the architecture of complex traits in humans,^{7,8} though its contribution to phenotypic variance is frequently the subject of debate.¹⁻³ Outside the prism of human association studies there is evidence for epistasis, not only at the molecular scale from artificially induced mutations⁴ but also at the evolutionary scale in fitness adaptation¹⁵ and speciation.¹⁶

Methods are now available to overcome the computational problems involved in searching for epistasis, but its detection still remains problematic due to reduced statistical power. For example increased dependence on linkage disequilibrium (LD) between causal SNPs and observed SNPs,^{17,18} increased model complexity in fitting interaction terms,¹⁹ and more extreme significance thresholds to account for increased multiple testing⁹ all make it more difficult to detect epistasis in comparison to additive effects. Thus, 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 less polygenic than high level phenotypes,²¹ thus many genetic effects are relatively large, 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 exhaustively tested every pair of SNPs for genetic interactions against each of 7339 expression traits in peripheral blood. After stringent filtering and multiple testing correction (Methods) we identified 501 putative genetic interactions influencing the expression levels of 238 genes (Supplementary Table 4). Of the 501 discovery interactions, 434 had available data and passed filtering (Methods) in two independent replication datasets, Fehrmann¹² and the Estonian Genomics Centre University of Tartu (EGCUT),¹¹ in which we saw convincing evidence for replication. We used the summary statistics from the replication datasets to perform a meta analysis to obtain an independent p -value for the putative interactions, and 30 were significant after applying a Bonferroni correction for multiple testing (Table 1). These significant interactions exhibited remarkable similarity in GP maps between all three datasets (Figure 2).

In addition, using the meta analysis from the replication samples only, we observed that 316 of the remaining 404 discovery SNPs had replication interaction p -values more extreme than the 2.5% confidence interval of the distribution under the null distribution of no epistatic effects ($p < 1.0 \times 10^{-16}$, Figure 3 and Supplementary Figure S1). The congruence of the epistatic networks in discovery and replication datasets is shown in Figure 1, 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 27 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. 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 because the dependence on LD between observed SNPs and causal variants is up to three orders of magnitude

higher than it is for independent additive effects.^{17,18} 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 are not designed to resemble biological function.²⁴

Of the discovery interactions, 47 were *cis-cis* acting (both SNPs were on the same chromosome as the expression gene), 441 were *cis-trans*-acting, and 13 were *trans-trans*-acting. We observed a wide range of significant GP maps (Figure 2) 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 S5). Each of these interactions have evidence for replication in at least one dataset and six are significant at the Bonferroni level (Supplementary Figure S2). We see similar epistatic networks involving multiple (eight or more) *trans*-acting SNPs for other gene expression levels too, for example TMEM149 (Supplementary Figure S6), NAPRT1 (Supplementary Figure S7), TRAPPC5 (Supplementary Figure S8), and CAST (Supplementary Figure S9). 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-4,3-methylation (H3K4me3) chromatin marks, in 34 cell types²⁶ (Supplementary Figure S4). 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 the *cis*-SNPs provide tissue specificity in these interactions. There is also strong enrichment for SNPs to be localised in enhancer regions,²⁷

consistent for both *cis* and *trans* SNPs ($p < 1 \times 10^{-6}$).

We also demonstrate spacial organisation of interacting loci 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 2Mb ($p < 1.8 \times 10^{-10}$), (Supplementary Figure S10). 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. How does this compare to the number of traits influenced by additive effects? The BSGS dataset has been previously analysed for additive effects at all expression traits,²² and if we take all the additive eQTLs that were significant at the epistatic threshold of $p < 2.91 \times 10^{-16}$ we find that 453 gene expression levels out of the 7339 analysed had at least one significant expression quantitative trait locus (eQTL). Therefore it can be argued that the number of instances of detectable epistasis are substantial.

However 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) at the $p < 2.91 \times 10^{-16}$ threshold, we calculate that on average they explain 1.73% of the phenotypic variance of each of the 7339 probes. By contrast, the epistatic variance from the interacting SNPs detected in this study on average explain 0.25% of phenotypic variance, approximately seven times lower than the additive variance. If we use the effect estimates taken from the Fehrmann or EGCUT datasets to perform the same comparison we obtain ratios of additive to epistatic variance of 36:1 and 34:1, respectively (Supplementary Figure S12, Methods). There are several caveats to this comparison. Firstly, the ratio of additive to epistatic variance may differ at different effect sizes, 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 rela-

tive 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 and who have gene expression levels measured in peripheral blood samples for 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 Fehrman 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.

Acknowledgements

We are grateful to the volunteers for their generous participation in these studies. We thank Bill Hill, Chris Haley and Lars Ronnegard for helpful discussions and comments.

This work could not have been completed without access to high performance GPGPU compute clusters. We acknowledge iVEC for the use of advanced computing resources located at iVEC@UWA (www.ivec.org), and the Multimodal Australian ScienceS Imaging and Visualisation Environment (MASSIVE) (www.massive.org.au). We also thank Jake Carroll and Irek Porebski from the Queensland Brain Institute Information Technology Group for HPC support.

The University of Queensland group is supported by the Australian National Health and Medical Research Council (NHMRC) grants 389892, 496667, 613601, 1010374 and 1046880, the Australian Research Council (ARC) grant (DE130100691), and by National Institutes of Health (NIH) grants GM057091 and GM099568.

The QIMR researchers acknowledge funding from the Australian National Health and Medical Research Council (grants 241944, 389875, 389891, 389892, 389938, 442915, 442981, 496739, 496688 and 552485), the and the National Institutes of Health (grants AA07535, AA10248, AA014041, AA13320, AA13321, AA13326 and DA12854). We thank Anthony Caracella and Lisa Bowdler for technical assistance with the micro-array hybridisations.

The CHDWB study funding support from the Georgia Institute of Technology Research Foundation. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript

The Fehrmann study was supported by grants from the Celiac Disease Consortium (an innovative cluster approved by the Netherlands Genomics Initiative and partly funded by the Dutch Government (grant BSIK03009), the Netherlands Organization for Scientific Research (NWO-VICI grant 918.66.620, NWO-VENI grant 916.10.135 to L.F.), the Dutch Digestive Disease Foundation (MLDS WO11-30), and a Horizon Breakthrough grant from the Netherlands Genomics Initiative (grant 92519031 to L.F.). This project was supported by the Prinses Beatrix Fonds, VSB fonds, H. Kersten and M. Kersten (Kersten Foundation), The Netherlands ALS Foundation, and J.R. van Dijk and the Adessium Foundation. The research leading to these results has received funding from the European Communitys Health Seventh Framework Programme (FP7/2007-2013) under grant agreement 259867.

The EGCUT study received targeted financing from Estonian Government SF0180142s08, Center of Excellence in Genomics (EXCEGEN) and University of Tartu (SP1GVARENG). We acknowledge EGCUT technical personnel, especially Mr V. Soo and S. Smit. Data analyzes were carried out in part in the High Performance Computing Center of University of Tartu.

1 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

2 Figures

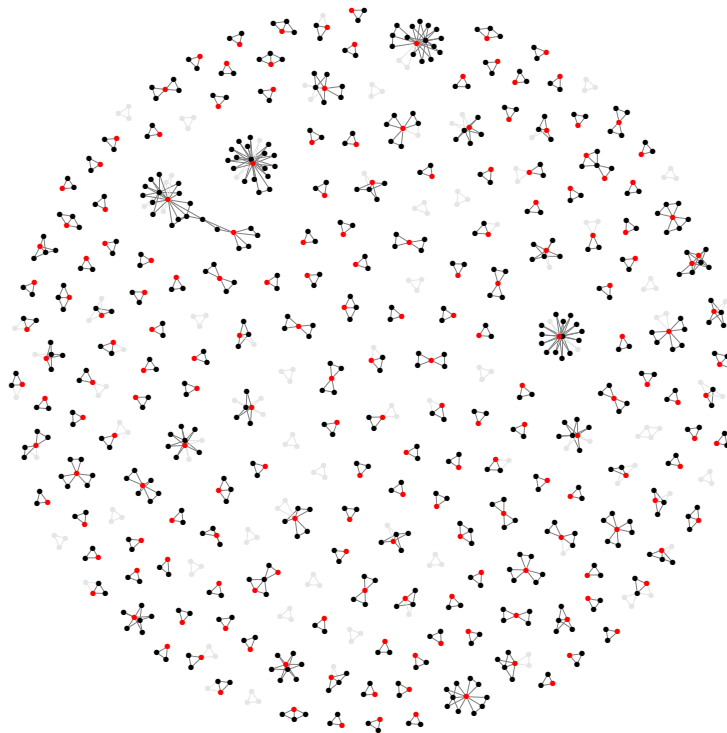


Figure 1: **Discovery and replication of epistatic networks** All 434 putative genetic interactions (edges) with data common to discovery and replication sets is shown, where black nodes represent SNPs and red nodes represent traits (gene expression probes). Three hundred and forty-five interactions had p -values exceeding the 2.5% confidence interval following meta analysis of the replication data, but 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.

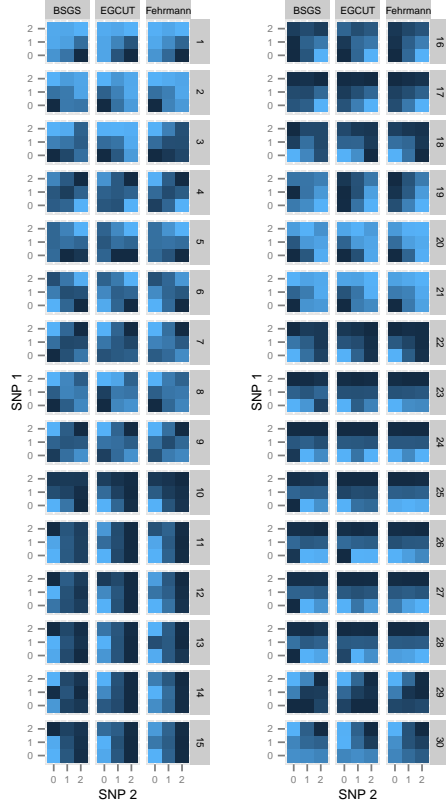


Figure 2: Replication of GP maps in two independent populations
The GP maps for each epistatic interaction that is significant at the Bonferroni level in both replication datasets are shown. Each GP map consists of nine tiles where each tile represents the expression level for that two-locus genotype 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 population. 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.

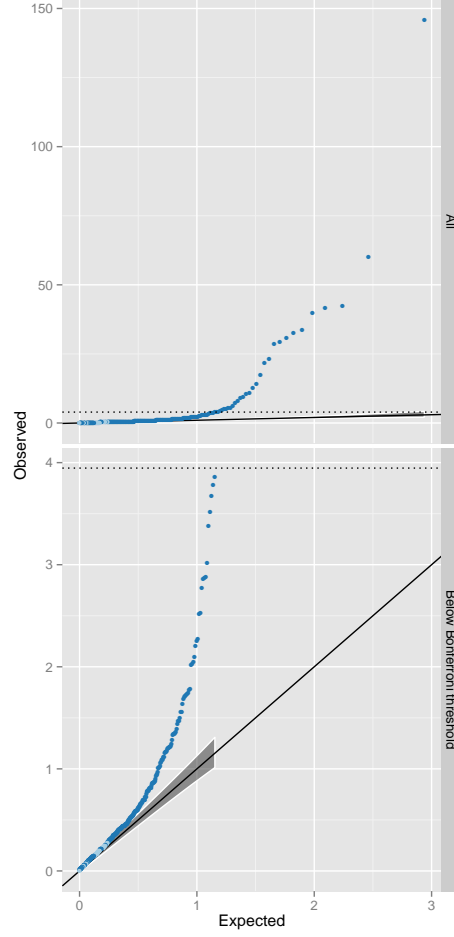


Figure 3: **Q-Q plots of interaction p -values from replication datasets** The top panel shows all 434 discovery SNPs that were tested for interactions. Observed p -values (y -axis, $-\log_{10}$ scale) are plotted against the expected p -values (x -axis, $-\log_{10}$ scale). The multiple testing correction 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.

3 Supplementary Figures

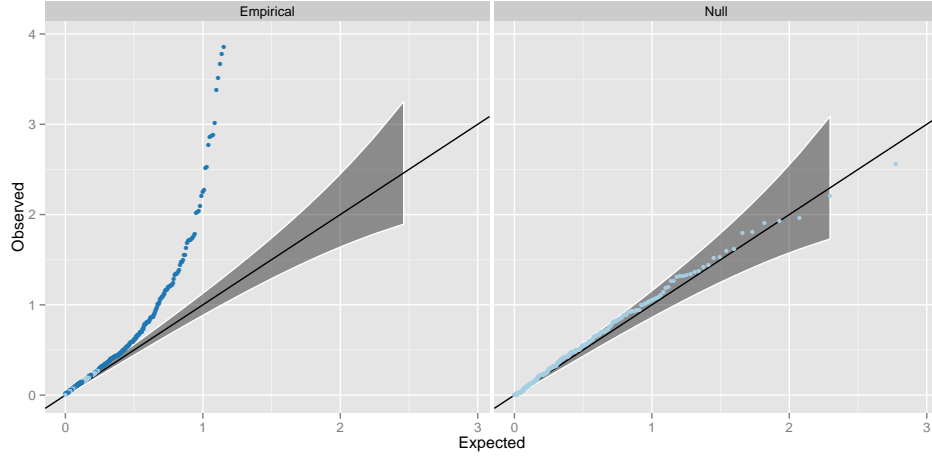


Figure S1: **Q-Q plots of interaction p -values from replication datasets, excluding the 30 points significant at the Bonferroni level** The right panel (Null) shows the interaction p -values from a meta analysis across two independent datasets on 434 randomly drawn SNP pairs. The left panel (Empirical) 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 3.

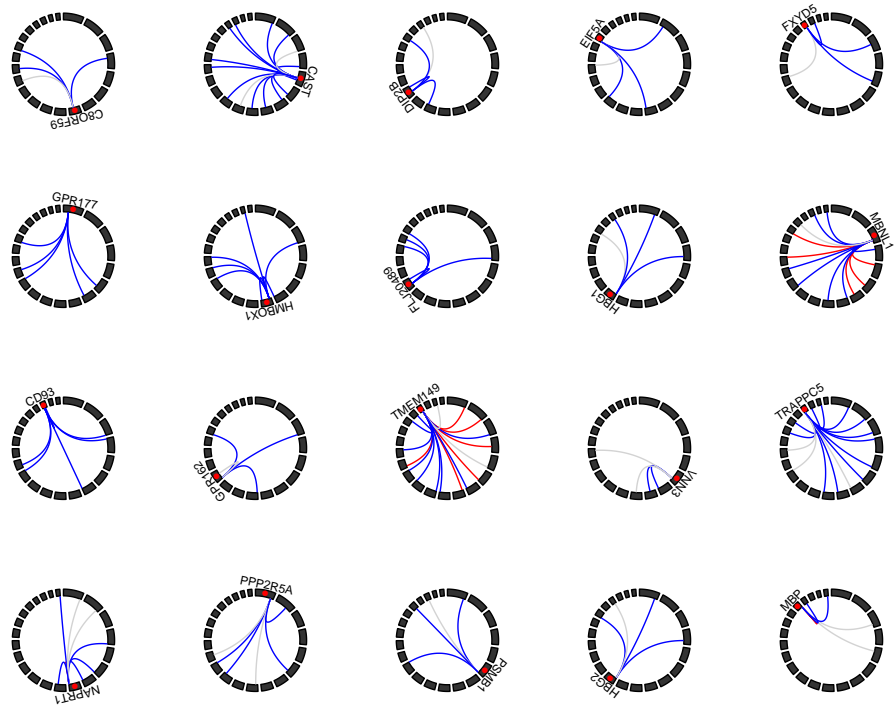


Figure S2: **Gene expression traits with four or more genetic interactions** Circle plots represent the genomic positions for SNPs (linking lines) and expression probes (red points). Chromosomes are represented by black blocks and ordered from 1 to 22 clockwise, starting from the top. Grey lines represent no evidence for replication, blue lines denote interactions that are outside the 97.5% confidence interval or the Q-Q plot (Figure 3), and red lines denote replication at the Bonferroni correction level. Most interactions are characterised as being *cis-trans* to the expression probe.

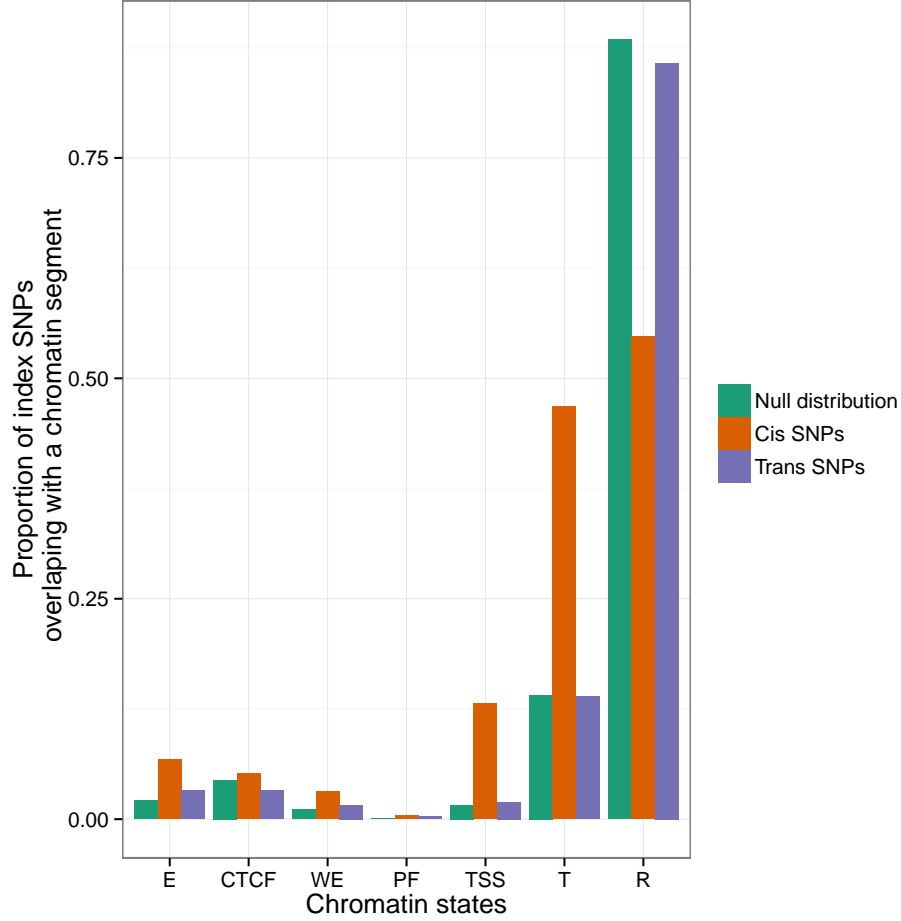


Figure S3: Location of SNPs relative to genomic features All SNPs within 1Mb and $r^2 > 0.8$ of each *cis*- and *trans*-SNP were taken to find which genomic features (x -axis) were covered by the SNPs that compose the 501 significant interactions. Green bars represent the proportion (y -axis) of the 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

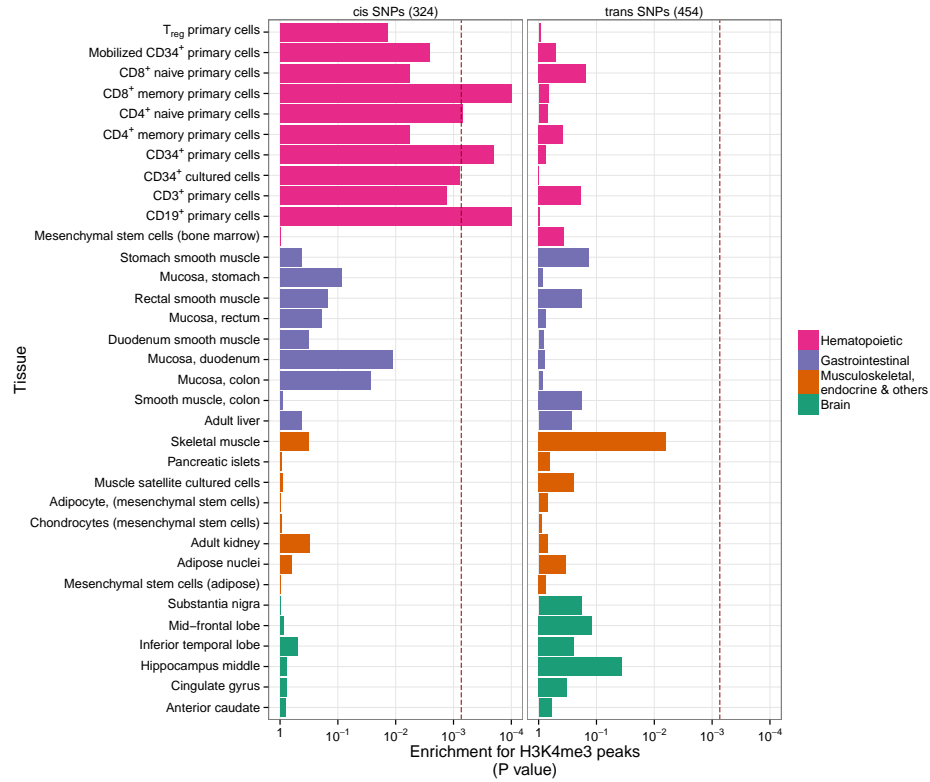


Figure S4: Tissue specific enrichment of SNPs in transcriptionally active regions The locations of transcriptional activity can be predicted by chromatin marks, assayed by H3K4me3.²⁶ Enrichment *p*-values are calculated using permutation analysis for 34 different cell types (*y*-axis) in four tissue types (Rows of boxes). There is enrichment for *cis*-acting SNPs in Haematopoietic tissue types only. *Trans*-acting SNPs have no tissue specificity.

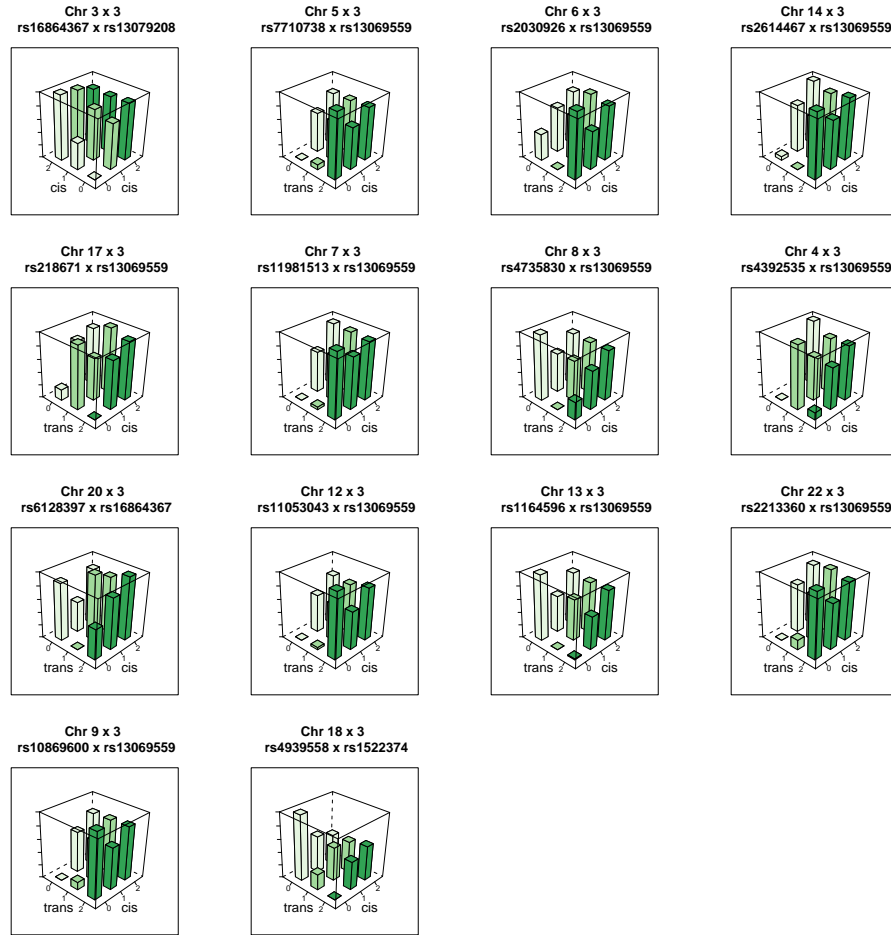


Figure S5: **Genotype-phenotype maps for 14 interactions influencing the expression of MBNL1** Each bar represents the mean phenotypic value for individuals in that genotype class. The rs13069559 SNP typically has a *cis*-additive decreasing effect on the expression of MBNL1, but in many of these interactions the *cis* effect is masked when the *trans* SNP is homozygous.

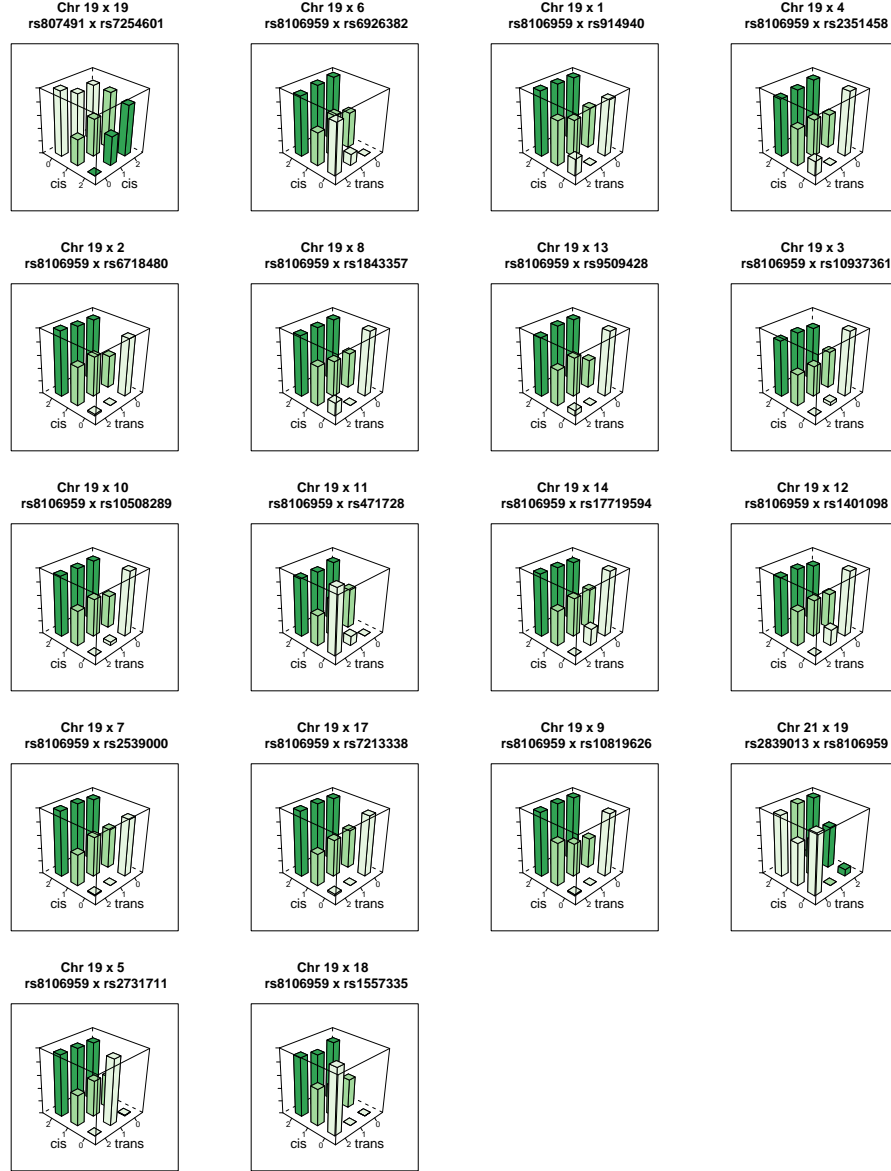


Figure S6: **Genotype-phenotype maps for 19 interactions influencing the expression of TMEM149** Each bar represents the mean phenotypic value for individuals in that genotype class. The rs13069559 SNP typically has a *cis*-additive decreasing effect on the expression of TMEM149, but in many of these interactions the *cis* effect is masked when the *trans* SNP is homozygous.

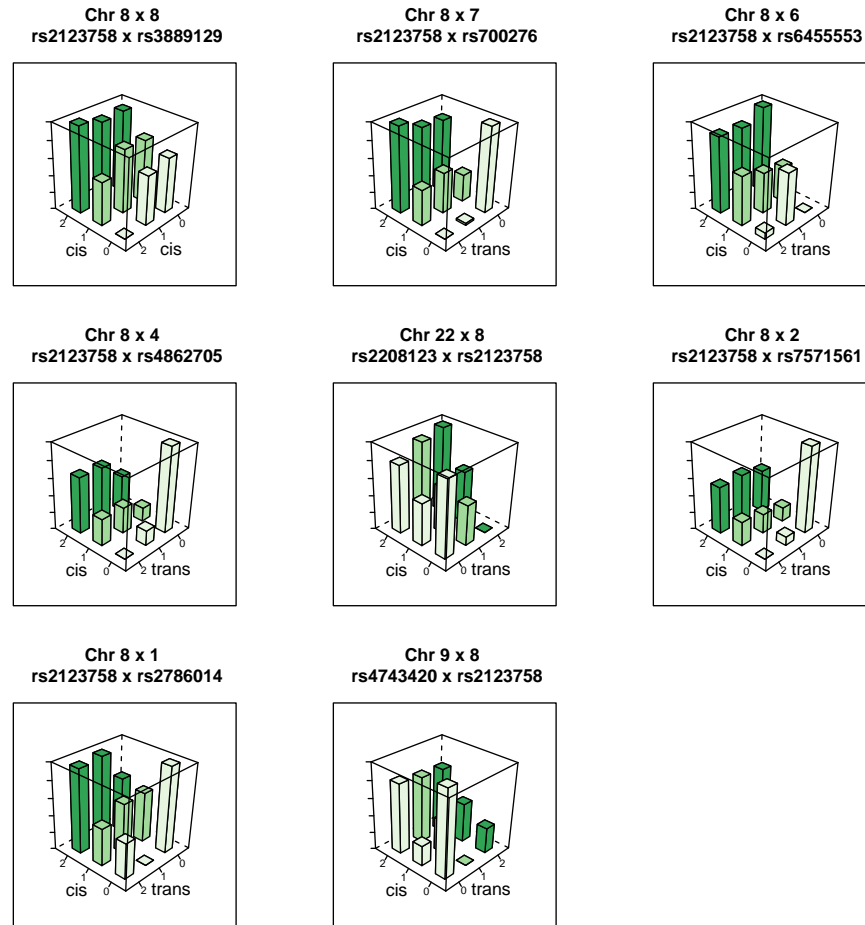


Figure S7: **Genotype-phenotype maps for 8 interactions influencing the expression of NAPRT1** Each bar represents the mean phenotypic value for individuals in that genotype class.

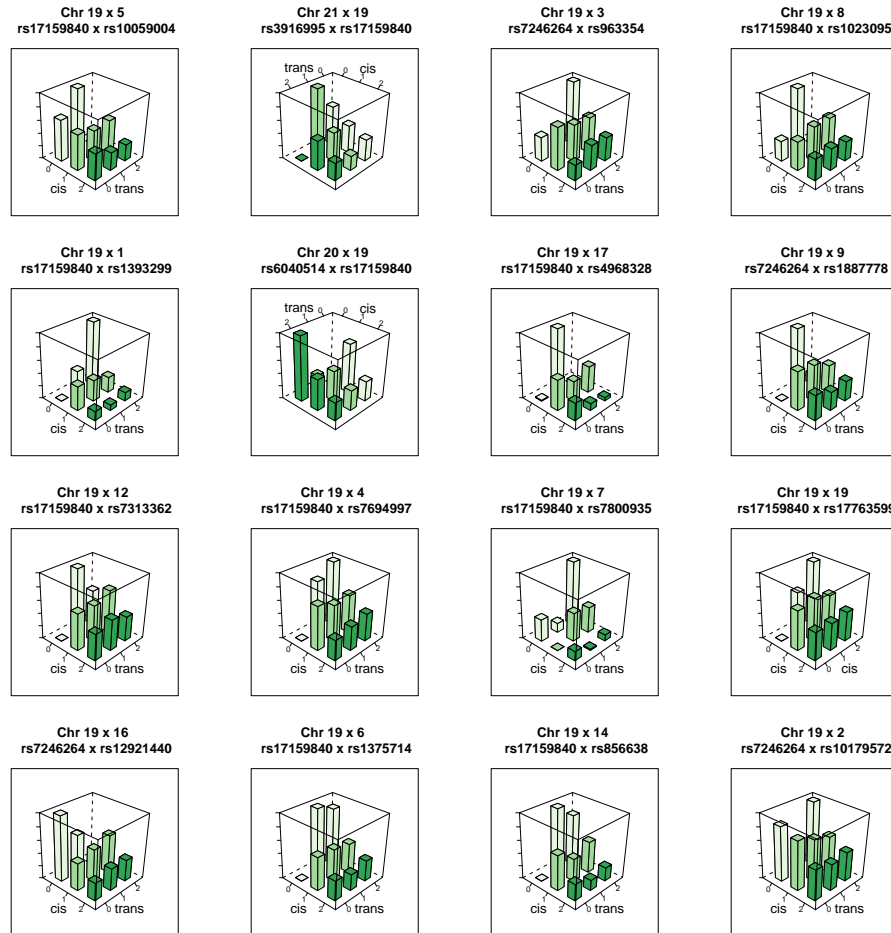


Figure S8: **Genotype-phenotype maps for 16 interactions influencing the expression of TRAPPC5** Each bar represents the mean phenotypic value for individuals in that genotype class.

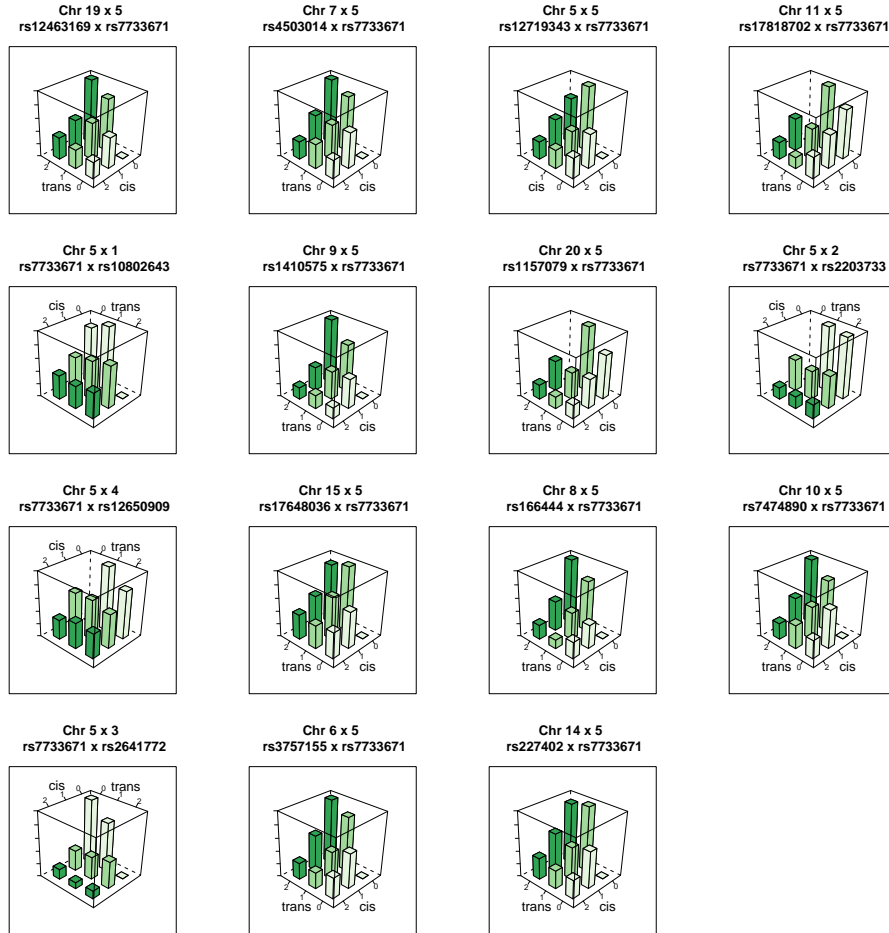


Figure S9: **Genotype-phenotype maps for 15 interactions influencing the expression of CAST** Each bar represents the mean phenotypic value for individuals in that genotype class.

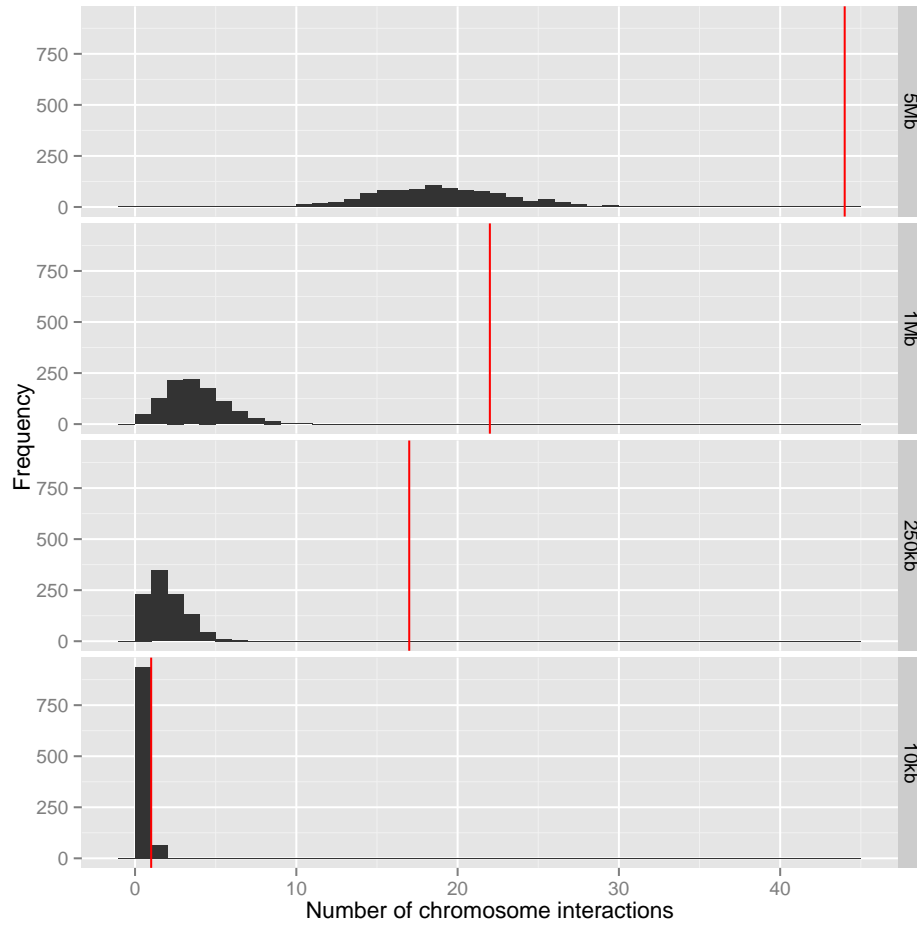


Figure S10: Number of overlaps between chromosome interactions and epistatic interactions Interacting chromosome regions may be a possible mechanism underlying epistatic interactions. The number of epistatic interactions within 20kb, 500kb, 2Mb and 10Mb of known chromosome interacting regions are shown by red vertical lines. The histograms represent the null distribution based on random sampling of 10,000 datasets for each window size.

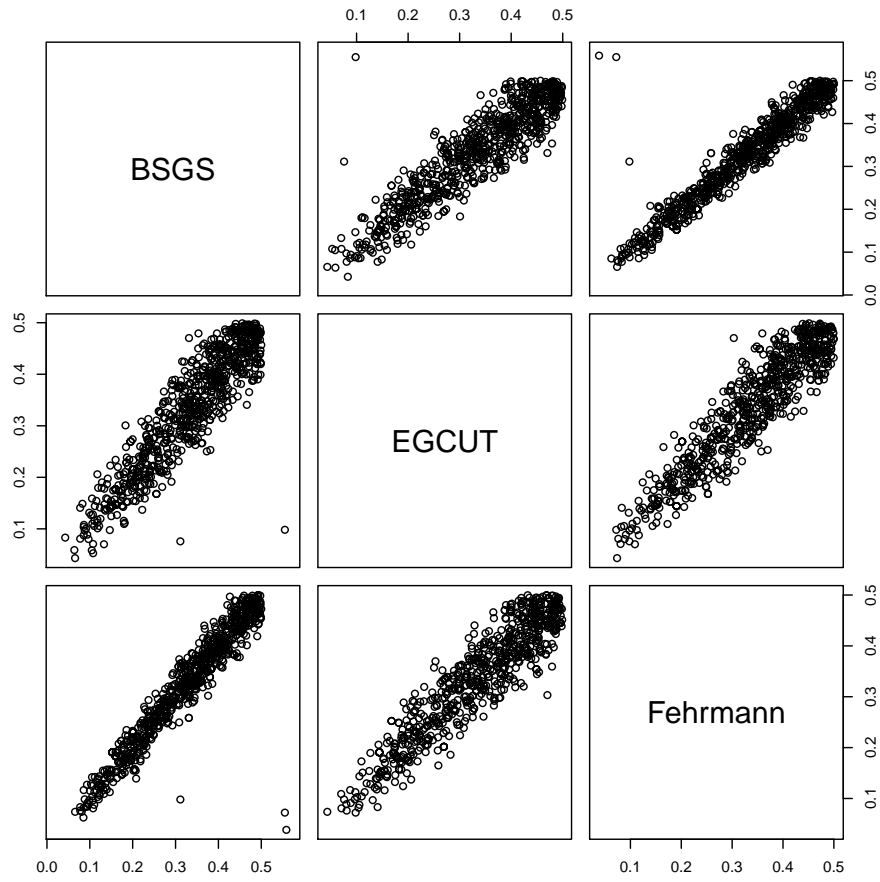


Figure S11: **Comparison of allele frequencies for 781 SNPs involved in genetic interactions across independent populations** Outliers were removed from the analysis as part of the filtering stage during replication.

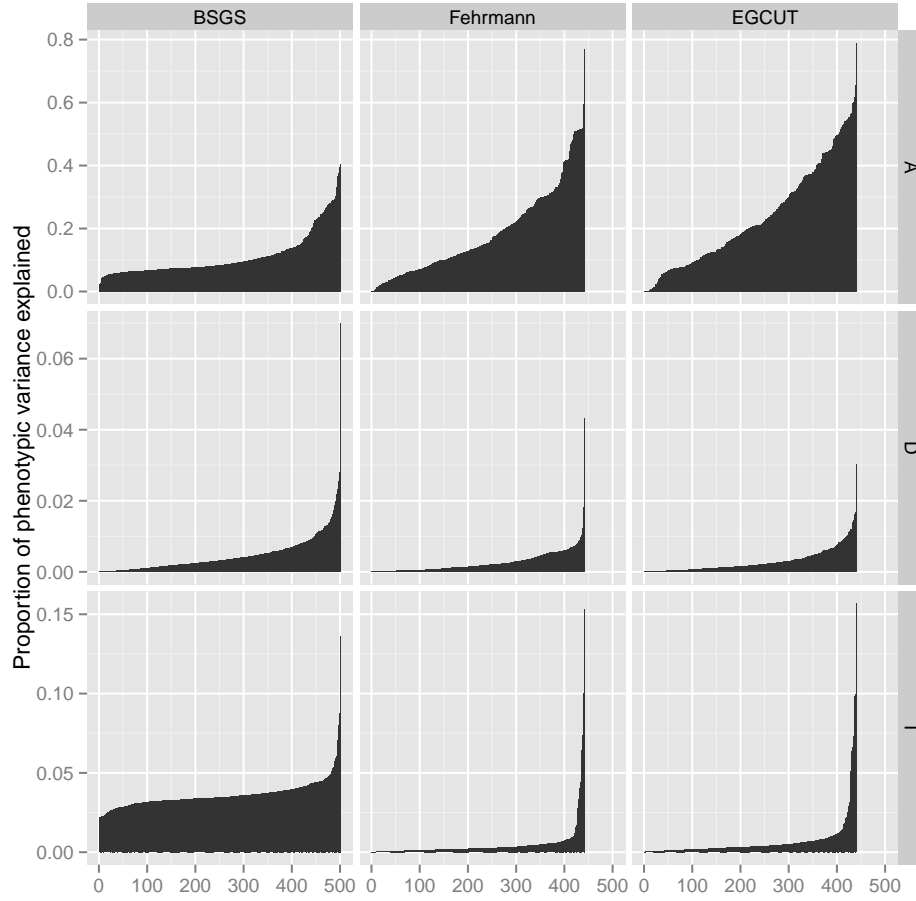


Figure S12: Comparison of variance explained by additive, dominant and epistatic effects from different cohorts How does the estimated variance decomposition change in different cohorts? The proportion of the phenotypic variance that is additive (A), dominant (D), or epistatic (I) for each putative interaction 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.

4 Supplementary Tables

Table S1 – continued from previous page

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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			
	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
CPVL	ILMN-1682928	7	rs2835998	21	39202070		rs245884	7	29185475	CPVL	5.55	0.19	0.03	0.04
CRPT	ILMN-1813256	2	rs2131290	4	188559908		rs1531133	7	46843631	CRPT	5.47	0.28	0.10	0.12
CRUS1	ILMN-1737685	20	rs6139887	20	5986234	CRUS1	rs1473927	5	42460408		6.18	0.10	0.36	0.15
CS1B	ILMN-1761797	21	rs9979356	21	43230974		rs3761385	21	43198355		11.99	25.20	16.72	42.27
CTNNA1	ILMN-1804854	5	rs924943	18	69000505		rs176382	5	138226707	CTNNA1	5.74	0.02	0.41	0.11
CTSC	ILMN-1696347	11	rs2457684	11	88139983	CTSC	rs7079264	10	10867982		5.67	0.92	0.74	0.03
CTSC	ILMN-1696347	11	rs7532236	22	26250645		rs1728352	11	88077479	CTSC	5.84	0.49	0.73	0.73
CTSC	ILMN-2242463	11	rs7930237	11	88117962		rs1556895	11	88077479		7.16	18.76	15.06	33.53
CWFI9L1	ILMN-1651886	10	rs7108734	11	11456027		rs12784396	10	102027407	CWFI9L1	5.42	0.21	0.01	0.03
CYBRD1	ILMN-1712305	4	rs2592948	4	129994690		rs888427	2	172366120	CYBRD1	5.89	0.23	0.53	0.34
CYBRD1	ILMN-1712305	2	rs7852475	9	140698856		rs888427	2	172366120	CYBRD1	5.68	0.20	0.02	0.04
CYBRD1	ILMN-2087692	2	rs11257679	10	12318284		rs888427	2	172366120	CYBRD1	5.81	0.39	1.87	1.47
CYBRD1	ILMN-2087692	2	rs6137908	20	23344590		rs888427	2	172366120	CYBRD1	5.53	0.05	0.83	0.36
CYBRD1	ILMN-2087692	2	rs888427	2	172366120	CYBRD1	rs7591849	2	160112881		5.85	0.87	0.10	0.44
CYP27A1	ILMN-1704985	2	rs6021982	20	36571928		rs933994	2	219650616	CYP27A1	5.42	0.29	0.86	0.60
CYP27A1	ILMN-2128428	5	rs7778910	17	110451383		rs835223	5	39381357	DAB2	5.44	0.48	0.41	0.44
DAB2	ILMN-1816458	17	rs9900173	17	43111688		rs1343244	9	82076988		9.12	0.00	0.58	0.14
DDT	ILMN-1690982	22	rs9760102	22	24248761	DDT	rs275341	3	187475208		5.62	0.64	0.25	0.42
DDX58	ILMN-1797001	9	rs4537097	11	125962645		rs7042042	7	32451144		5.31	0.61	0.29	0.44
DEM1	ILMN-1783996	1	rs10120023	9	137810259	COQ10A	rs10120023	9	137810259	COQ10A	5.37	0.08	0.41	0.16
DEM1	ILMN-1783996	1	rs12363827	13	106703727		rs7566044	2	169960422	DHRS9	6.39	0.77	0.02	0.29
DHRS9	ILMN-1733998	2	rs1511956	12	89468283		rs7566044	2	169960422	DHRS9	6.00	0.06	1.17	0.58
DHRS9	ILMN-1733998	2	rs1328529	12	39468285		rs2161037	2	169893419	DHRS9	6.48	0.37	0.34	0.32
DHRS9	ILMN-2384181	2	rs2831914	21	29959453		rs2161037	2	169893419	DHRS9	5.51	0.88	0.04	0.37
DHRS9	ILMN-2384181	2	rs7661304	4	187776431		rs11669332	12	50610976	LASS5	7.64	0.05	0.11	0.03
DP2B	ILMN-1755589	12	rs11080134	17	29161503	LASS5	rs11669332	12	50610976	LASS5	4.65	0.05	0.10	0.10
DP2B	ILMN-1755589	12	rs1166935	12	50636364		rs2872008	7	153134888	LASS5	4.87	0.30	0.58	0.19
DP2B	ILMN-1755589	12	rs3383855	19	41711815	LASS5	rs7134595	12	50730458	LASS5	5.31	0.38	0.22	0.19
DP2B	ILMN-1755589	12	rs7312552	12	50730458	LASS5	rs1808634	8	61971140	LASS5	4.40	0.37	0.09	0.02
DP2B	ILMN-1755589	12	rs7312552	12	50730458	LASS5	rs4532958	10	115214154	LASS5	5.03	0.09	0.02	0.01
DP2B	ILMN-1755589	12	rs7312552	12	50730458	LASS5	rs2427378	12	51074199	DNAJB6	5.92	0.48	0.00	0.11
DNAJB6	ILMN-1793770	7	rs2288842	15	171994348		rs3775539	7	157163614	DNAJB6	5.79	0.23	1.45	0.97
DPH3	ILMN-2140610	3	rs12252308	15	93409054		rs1566972	3	16320360	DPH3	6.17	1.58	0.27	1.12
ECGF1	ILMN-2232308	22	rs140522	22	50971266	ECGF1	rs4891884	18	64004670	DPH3	4.81	0.15	1.18	0.70
ECGF1	ILMN-2109708	22	rs4234091	22	241911027		rs11206043	1	53402552	ECGHC2	6.19	0.22	0.35	0.22
ECGHC2	ILMN-1671568	1	rs5092637	22	17675900		rs1206043	1	53402552	ECGHC2	5.58	0.64	0.16	0.35
ECGHC2	ILMN-1671568	1	rs5092637	22	17675900		rs1048166	15	42192040	ECGHC2	6.98	0.90	0.47	0.79
EHD4	ILMN-1720083	15	rs0403312	19	53244938		rs1754556	14	75503430	EIF2B2	5.56	0.23	0.11	0.10
EIF2B2	ILMN-1719380	14	rs6567288	18	60218334		rs1269096	14	99603119	EIF2B2	5.44	0.56	0.08	0.24
EIF5A	ILMN-1794522	17	rs7216490	17	7221707	EIF5A	rs1269096	14	99603119	EIF2B2	5.55	0.28	0.59	0.41
EIF5A	ILMN-1794522	17	rs7216490	17	7221707	EIF5A	rs1553474	2	49359676		6.36	0.08	0.05	0.02
EIF5A	ILMN-1794522	17	rs7216490	17	7221707	EIF5A	rs1553474	2	49359676		5.52	0.05	1.12	0.53
EIF5A	ILMN-1794522	17	rs7216490	17	7221707	EIF5A	rs4471434	11	126387391	EMR2	6.51	0.36	0.04	0.11
EMR2	ILMN-2353633	19	rs2827076	21	23196249		rs9305048	19	14879034	EMR2	5.56	0.45	0.40	0.41
EMR2	ILMN-2353633	19	rs6132112	20	18761714		rs3007765	13	102480759	EMR2	6.03	0.20	0.58	0.35
EMR2	ILMN-2353633	19	rs9305048	19	14879034	EMR2	rs3007765	13	102480759	EMR2	5.70	0.20	0.58	0.35
EPHX2	ILMN-1709237	8	rs1109764	11	12790396		rs13269963	8	27400604	EPHX2	5.70	0.20	0.58	0.35
EPHX2	ILMN-1731001	8	rs10894861	11	13461176		rs12115088	8	5787462	EPHX2	5.43	0.25	1.20	0.81
EPHX2	ILMN-1731001	8	rs5766218	22	45337329		rs12115088	8	5787462	EPHX2	6.11	0.20	0.11	0.09
EPHX2	ILMN-1731001	8	rs726145	18	31187910		rs12115088	8	5787462	EPHX2	5.63	0.29	0.04	0.08
ERICH1	ILMN-2104696	5	rs4735895	8	600729	ERICH1	rs1517297	4	182786760	ERICH1	5.63	0.67	1.03	1.06
ERICH1	ILMN-2104696	5	rs187076	10	55228462		rs12188164	4	4228236	EXOC3	5.63	0.67	1.03	1.06
EXOC3	ILMN-1789419	5	rs187076	10	55228462		rs12188164	4	4228236	EXOC3	5.63	0.67	1.03	1.06
FAHD1	ILMN-2246661	16	rs1560104	16	12708208		rs344363	16	1972548	FAHD1	6.33	0.27	0.30	0.23
FCN1	ILMN-1668063	9	rs12580388	12	129591144		rs10120023	9	137810259	COQ10A	6.33	0.27	0.30	0.23

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Table S1 – continued from previous page

Gene ID ^a	Expression trait	Chr.	rs ID	Chr.	SNP 1	Pos/Mb ^c	Association ^d	rs ID	Chr.	SNP 2	Pos/Mb ^c	Association ^d	BSGS ^e	Interaction statistic ^f	-log ₁₀ p-value	Distance / Mb ^g
FE22	ILMN_1739586	2	rs2356400	19	44321776	2	rs13406184	2	36791226	FE22			5.78	0.14	0.33	0.16
FE22	ILMN_1739586	2	rs969010	4	159963132	2	rs11691600	2	36810133	FE22			6.59	0.14	0.28	0.14
FGD2	ILMN_2115005	6	rs4803848	19	46203050	6	rs31486	6	37001267	FGD2			5.69	0.12	0.25	0.11
FGD2	ILMN_2115005	6	rs902634	12	133943531	12	rs17036706	12	36999652	FGD2			5.49	0.06	0.10	0.06
FLJ20489	ILMN_1761044	16	rs17015703	12	177036706	16	rs3824908	12	48169526	FLJ20489			5.81	0.06	0.10	0.29
FLJ20489	ILMN_1761044	16	rs403198	12	177036706	16	rs3824908	12	48169526	FLJ20489			5.79	0.06	0.10	0.06
FLJ20489	ILMN_1778144	12	rs7021190	15	70929121	12	rs3782908	12	48169526	FLJ20489			5.79	0.10	0.02	0.04
FLJ20489	ILMN_1778144	12	rs4984400	15	97033126	12	rs3782908	12	48169526	FLJ20489			6.90	0.38	0.17	0.21
FLJ20489	ILMN_1763663	16	rs9325634	21	43818790	16	rs7204135	16	50106594	FLJ20489			6.04	0.14	0.95	0.53
FLJ43093	ILMN_2123450	6	rs71712712	14	107276627	6	rs6906101	6	36667610	FLJ43093			5.48	0.39	0.06	0.13
FN3KRP	ILMN_1652333	17	rs898095	17	80890638	17	rs9892064	17	808927903	FUCA1			16.16	28.24	29.39	59.95
FUCA1	ILMN_1752728	17	rs4971478	17	3460963	17	rs881878	17	24168019	FUCA1			6.41	0.01	0.30	0.06
FXD5	ILMN_2309848	19	rs1633921	19	35695290	19	rs2285515	13	98328559	FXD5			3.70	0.09	0.41	0.17
FXD5	ILMN_2309848	19	rs17398183	20	55609148	19	rs2285515	19	35660450	FXD5			6.58	0.03	0.48	0.15
FXD5	ILMN_2309848	19	rs2285515	19	35660450	19	rs11739594	5	141709163	FXD5			5.70	0.07	0.17	0.05
FXD5	ILMN_2309848	19	rs2285515	19	35660450	19	rs13067700	3	95331048	FXD5			6.00	0.09	0.09	0.51
G3BP2	ILMN_23081758	4	rs10230232	7	29390239	4	rs17036504	4	7567329	G3BP2			6.10	0.28	0.08	0.08
GAA	ILMN_2410783	17	rs11150847	17	78153130	17	rs1553985	4	7654604	GAA			5.19	0.08	0.37	0.14
GAA	ILMN_2410783	17	rs8068856	17	78100731	17	rs12602462	17	78146016	GAA			13.91	19.98	12.99	32.60
GAPT	ILMN_1675191	5	rs10070522	5	57786110	5	rs10902506	12	132678089	GAPT			5.65	0.11	0.39	0.17
GAPT	ILMN_1675191	5	rs7082031	10	128038717	5	rs7605821	2	235695528	GAPT			5.85	0.01	0.78	0.28
GATS	ILMN_1699631	7	rs1147447	14	66460742	7	rs2950520	7	99827148	GATS			5.72	0.26	0.11	0.11
GATS	ILMN_1699631	7	rs2425256	20	35056572	7	rs2950520	7	99827148	GATS			5.47	0.83	0.63	0.87
GDPD3	ILMN_1774901	16	rs8090624	16	30102802	14	rs2197465	14	48572632	GDPD3			6.22	0.42	0.35	0.33
GDPD3	ILMN_1774901	16	rs7204270	16	30156963	4	rs1015111	4	128972357	GDPD3			5.86	0.55	0.09	0.24
GDPD3	ILMN_1790692	2	rs145072	13	110899955	4	rs7577293	4	128972357	GDPD3			5.78	0.02	0.45	0.13
GNLY	ILMN_3239426	12	rs198646	16	26084476	6	rs7960552	12	111164237	GNLY			5.72			
GNLY	ILMN_3239426	12	rs1860563	16	6478898	12	rs7960552	12	111164237	GNLY			5.72			
GPR162	ILMN_1730816	12	rs2272500	12	79685913	12	rs2707210	12	6902002	GPR162			5.07	0.26	0.46	0.39
GPR162	ILMN_1730816	12	rs2272500	12	79685913	12	rs2707210	12	6902002	GPR162			5.47	0.25	0.06	0.07
GPR162	ILMN_1730816	12	rs2707210	12	6902002	12	rs4740848	9	6554558	GPR162			5.47	0.25	0.06	0.44
GPR177	ILMN_1660549	1	rs11057383	12	124369421	3	rs9827054	3	188880113	GPR177			6.21	0.96	0.06	0.44
GPR177	ILMN_1660549	1	rs12527241	6	120468039	1	rs12065581	1	68732819	GPR177			5.76	0.72	0.67	0.81
GPR177	ILMN_1660549	1	rs12532999	7	127939793	7	rs12065581	1	68732819	GPR177			5.40	0.79	0.17	0.40
GPR177	ILMN_1660549	1	rs25613	16	11169683	13	rs12065581	1	68732819	GPR177			6.50	0.71	1.43	1.50
GPR177	ILMN_1660549	1	rs9575097	13	82986268	13	rs12065581	1	68732819	GPR177			5.43	0.79	0.11	0.13
GPR177	ILMN_1660549	1	rs656669	18	70506011	1	rs12065581	1	68732819	GPR177			6.04	0.95	0.21	0.60
GPR177	ILMN_2283325	1	rs656669	18	70506011	1	rs12065581	1	68732819	GPR177			5.86	0.24	0.34	0.23
GPR177	ILMN_2283325	1	rs9290426	3	171399321	1	rs12065581	1	68732819	GPR177			6.50	0.01	0.24	0.04
GSDMB	ILMN_2347193	17	rs11557467	17	38028634	15	rs4965745	15	101508261	GSDMB			5.88	0.68	0.20	0.41
GSTM1	ILMN_2391861	1	rs12244673	10	53192833	1	rs11101992	1	110266754	GSTM1			6.11	0.27	1.14	0.79
GSTM1	ILMN_2391861	1	rs1547574	13	83344527	1	rs11101992	1	110266754	GSTM1			5.91	0.27	1.14	0.79
GSTM2	ILMN_2201580	1	rs6432807	13	96139560	1	rs3754446	1	110253241	GSTM1			6.77			
H1FO	ILMN_1757467	22	rs139898	22	38399979	15	rs4533353	15	77919015	H1FO			6.36	0.52	0.66	0.65
H1FO	ILMN_1757467	22	rs139898	22	38399979	21	rs6497007	17	85877017	H1FO			6.52	0.27	0.31	0.23
H1FO	ILMN_1757467	22	rs139898	22	38399979	21	rs9885949	21	19532546	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	38399979	17	rs39167	17	85877017	H1FO			5.40	0.25	0.48	0.32
H1FO	ILMN_1757467	22	rs139898	22	383999											

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Table S1 – continued from previous page

Gene ID ^a	Expression trait	Chr.	rs ID	Chr.	SNP 1	Pos/Mb ^c	Association ^d	rs ID	Chr.	SNP 2	Pos/Mb ^c	Association ^d	BSGS ^e	Interaction statistic ^f	-log ₁₀ p-value	Metas ^g	Distance / Mb ^h
HBG2	ILMN_2084825	11	rs12975066	19	35723301	19	HBG2	rs2855039	11	5271671	21	HBG2	5.77	0.08	0.13	0.05	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs12042181	1	213088494	1	LQN1	6.84	0.06	0.34	0.21	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs12033379	4	141533832	4	LQN1	5.98	0.00	0.46	0.10	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs47060636	12	48173352	12	HBG2	5.75	0.35	0.46	0.10	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671	11	HBG2	rs17086655	17	133220622	17	HBG2	5.98	0.15	0.59	0.32	
HBG2	ILMN_2084825	11	rs2855039	11	5271671												

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Table S1 – continued from previous page

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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.	Pos / Mb ^c	Association ^d	BSGS ^e	Fehrmann ^f	EGCUT ^g	Meta ^g
REBE	ILMN-1802380	1	rs4982958	14	24987865		rs301819	1	8501786	REBE	5.66	0.61	1.23	1.17
REBE	ILMN-1802380	1	rs7697290	4	135248366		rs301819	1	8501786	REBE	5.74	0.14	0.10	0.06
REBE	ILMN-2327795	1	rs11085829	19	13174312		rs301819	1	8501786	REBE	5.12	0.21	0.33	0.21
REBE	ILMN-2327795	1	rs3852011	3	112844086	RNASE6	rs301819	1	8501786	REBE	5.71	0.08	0.60	0.26
RNASE6	ILMN-1780533	14	rs11628398	14	8106521		rs7324365	13	100601327	RNASE6	5.48	0.42	0.21	0.26
RNASE6	ILMN-1780533	14	rs6603134	19	8106521		rs1628398	14	21182800	RNASE6	5.11	0.09	0.22	0.08
RNF167	ILMN-1794726	17	rs2382330	17	4875566		rs4884857	13	54668512		4.37			
RNF167	ILMN-1794726	17	rs400688	17	4839930		rs11706900	3	36348968		5.59	0.71	0.46	0.64
RNFEP	ILMN-1738347	1	rs1107121	21	46127549		rs2819365	1	201983242		6.27	0.11	0.30	0.13
RNFEP	ILMN-1738347	1	rs8071611	17	67153386		rs2819365	1	201983242		4.32	1.48	0.52	1.28
RPL13	ILMN-2413278	16	rs352935	16	89648580		rs2965817	16	89513234		4.98	3.79	14.41	17.24
RPL23AP7	ILMN-2222750	2	rs1401202	16	80320056		rs4849261	2	114450030	RPL23AP7	5.55	0.13	0.73	0.38
RPL36AL	ILMN-2186933	14	rs3007033	14	50310816		rs17495030	9	138038093		5.46	0.09	0.06	0.02
RPL36AL	ILMN-2186936	14	rs4009028	14	50020817		rs1502991	6	66137260		5.86	0.32	0.20	0.19
RPL8	ILMN-1764721	8	rs2958482	8	145984615		rs1619856	1	234585790		4.59	0.10	0.37	0.15
RPL8	ILMN-1764721	8	rs4143674	20	4741304		rs2958482	8	145984615	RPL8	4.33	0.13	0.45	0.22
SEC13	ILMN-3297880	3	rs4889214	16	80913946		rs696221	3	10342876	SEC13	6.48			
SEC13	ILMN-3297880	3	rs17085428	5	95388015		rs7695	1	156147326	SEC13	5.70	0.22	1.73	1.17
SESN3	ILMN-1694027	11	rs12147460	14	104412137		rs684856	11	94906111	SESN3	5.50	0.02	0.51	0.15
SESN3	ILMN-1694027	11	rs355391	15	46391793		rs684856	11	94906111	SESN3	5.67	0.31	0.06	0.10
SESN3	ILMN-1694027	11	rs684856	15	46391793	SESN3	rs7004947	8	134606425	SESN3	5.60	0.21	0.51	0.31
SH3BGL2	ILMN-1762764	6	rs10838191	11	43893658		rs1354034	3	56849749	PPBP	5.52	0.70	0.12	0.35
SH3BGL2	ILMN-1762764	6	rs2545385	5	66833979		rs1354034	3	56849749	PPBP	5.97	0.20	0.51	0.30
SH3BGL2	ILMN-1762764	6	rs6845394	4	88280592		rs1354034	3	56849749	PPBP	5.23	0.32	0.71	0.53
SH3BGL2	ILMN-2158336	9	rs1034120	21	18196922		rs17455517	9	131785369	SH3BGL2	7.40	0.22	0.18	0.13
SIRPG	ILMN-1771801	20	rs1535883	20	5128119	SIRPG	rs6842739	4	60489510		5.74	0.29	0.18	0.17
SIRPG	ILMN-2382505	11	rs11673260	19	56218798		rs367035	11	2923826	SLC22A18	5.47	0.09	0.24	0.09
SLC22A18	ILMN-2382505	11	rs367035	19	2923826	SLC22A18	rs3110874	7	153224179		5.70	0.15	0.10	0.06
SLC22A18	ILMN-2382505	11	rs367035	11	2923826	SLC22A18	rs3772054	2	241678528		6.15	0.39	0.13	0.19
SLC41A3	ILMN-236111	3	rs1912136	11	24616743		rs6771703	3	125801067	SLC41A3	5.88	1.10	0.82	1.24
SLC45A4	ILMN-1745778	8	rs2337734	8	142337734	SLC45A4	rs7981190	5	174598073	SLC45A4	5.95	0.86	0.07	0.40
SLC46A3	ILMN-1658639	13	rs949805	17	55602091		rs7981190	5	174598073	SLC46A3	5.52	0.09	0.58	0.26
SMG7	ILMN-1706553	1	rs8035259	15	97403923		rs10911353	1	183489203	SMG7	6.52	0.17	0.09	0.06
SMOX	ILMN-1775380	20	rs11677215	20	4161500	SMOX	rs11677215	2	65800982	SMOX	5.68	0.39	0.62	0.52
SMOX	ILMN-1775380	20	rs11677215	20	4161500		rs705837	4	119225940	SNHG8	6.11			
SNHG8	ILMN-3309349	4	rs1505621	9	133050233		rs214097	11	17291499	SNHG8	6.11	0.29	1.03	0.72
SNORD14A	ILMN-1799381	11	rs15204429	15	46259108		rs6486334	11	17015557	SNORD14A	6.60	0.29	1.03	0.72
SNORD14A	ILMN-1799381	11	rs2634462	11	17339127		rs6486334	11	17015557	SNORD14A	7.31	13.11	10.96	23.22
SNORD89	ILMN-3238662	2	rs10445863	2	115929241		rs750783	2	101889306	SNORD89	5.96			
SNORD89	ILMN-3238662	2	rs10445863	2	115929241		rs750783	2	101889306	SNORD89	5.96			
SNORD89	ILMN-3238662	2	rs10445863	2	115929241		rs750783	2	101889306	SNORD89	5.96			
SNORD89	ILMN-3238662	2	rs2135064	5	26778066		rs750783	2	101889306	SNORD89	6.33			
SNUPN	ILMN-1733932	15	rs8134646	21	46376528	SNUPN	rs17185362	16	81889005		6.45	0.13	1.41	0.83
SNUPN	ILMN-2364535	15	rs1346466	21	46376528	SNUPN	rs1472075	3	193706323		5.59	0.34	0.00	0.06
SPATA5L1	ILMN-1729179	15	rs1131620	19	41117869		rs4774580	15	45652086	SPATA5L1	5.44			
STARD10	ILMN-2210752	11	rs2221406	13	90174526		rs1006620	15	45652086	STARD10	5.65	0.67	0.12	0.33
STARD10	ILMN-2210752	11	rs4073164	14	104947517		rs1006620	15	45652086	STARD10	5.44	0.34	0.00	0.06
STYXL1	ILMN-2345142	20	rs4073164	14	104947517		rs17685	7	75616105	STYXL1	5.51	0.57	0.17	0.31
SULT2	ILMN-2345142	20	rs1463965	18	74332954	SULF2	rs17685	7	75616105	SULT2	5.85	0.26	0.16	0.14
SULT2	ILMN-2345142	20	rs1463965	18	74332954		rs392994	4	180439236	SULT2	5.51	0.57	0.17	0.31
SULTIA4	ILMN-2336133	16	rs2436657	20	40119768		rs392994	4	180439236	SULTIA4	7.05	0.01	0.05	0.00
SURF6	ILMN-1778032	9	rs2636657	21	56013994		rs3785354	16	28550667	TUFM	5.83			
SURF6	ILMN-1778032	9	rs6099626	20	40119768		rs3785354	16	28550667	TUFM	5.83			
SYTL2	ILMN-2336609	11	rs1375719	13	10341078		rs3118663	9	136281753	SURF6	6.14	0.26	0.16	0.14
THBS3	ILMN-1804663	1	rs1939875	14	95422867		rs485485	11	85495269	SYTL2	5.47	0.28	0.31	0.24
THBS3	ILMN-1804663	1	rs1939875	14	95422867		rs485485	11	85495269	THBS3	5.55	0.03	0.15	0.03
THBS3	ILMN-1804663	1	rs8014956	14	20687978		rs4072037	1	155194980	THBS3	5.65	0.31	0.76	0.55
TIPRL	ILMN-1781457	1	rs2823245	21	16745523		rs1320993	1	168154599	TIPRL	5.22	0.07	0.40	0.15

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Table S1 – continued from previous page

Expression trait			SNP 1			SNP 2			Interaction statistic / -log ₁₀ p-values						
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	Meta ^g	Distance / Mb ^h
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	rs7512594	1	214514361		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	1.643
VASP	LMN-1743646	19	rs1264226	19	40063167		rs2276470	19	45974668	VNN2	5.09	0.94	5.14	4.95	0.088
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		rs2267932	6	133067782	VNN3	6.44	0.16	0.74	0.41	
VNN3	LMN-1804935	6	rs1443946	8	73006453		rs2267932	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	rs4552100	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	rs10500316	19	54553697	VSTM1	5.88	0.81	1.38	1.47	
WDR48	LMN-1763455	3	rs9628570	22	30261219		rs6778963	3	39091812	WDR48	5.79	0.09	0.33	0.09	
WDR48	LMN-1762103	3	rs1388935	4	18827822	RAPGEF1	rs853349	3	39067325	WDR48	5.34	0.57	1.35	1.22	
WDR48	LMN-1762103	3	rs1887778	9	134635088		rs7619193	3	39044116	WDR48	5.85	0.18	0.61	0.35	
WDR6	LMN-1762103	3	rs9554833	13	102624790		rs7619193	3	39044116	WDR6	5.86	1.64	1.43	2.25	
WDR6	LMN-1669484	3	rs12362253	11	123571708		rs17715581	15	49194351		5.79	2.38	0.17	1.63	
XAF1	LMN-2330573	17	rs1535031	21	9673170	XAF1	rs12591171	15	93119799		5.79	0.67	0.36	0.15	
ZFP90	LMN-1684628	16	rs909446	16	37040648		rs182968	16	68573945	ZFP90	5.79	0.09	0.36	0.15	
ZNF500	LMN-1700238	16	rs4823723	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

References

- ¹ Carlborg, O. & Haley, C. S. Epistasis: too often neglected in complex trait studies? *Nature Reviews Genetics* **5**, 618–25 (2004).
- ² Hill, W. G., Goddard, M. E. & Visscher, P. M. Data and Theory Point to Mainly Additive Genetic Variance for Complex Traits. *PLoS Genetics* **4** (2008).
- ³ Crow, J. F. On epistasis: why it is unimportant in polygenic directional selection. *Philosophical transactions of the Royal Society of London. Series B, Biological sciences* **365**, 1241–4 (2010).
- ⁴ Costanzo, M. *et al.* The genetic landscape of a cell. *Science (New York, N.Y.)* **327**, 425–31 (2010).
- ⁵ Bloom, J. S., Ehrenreich, I. M., Loo, W. T., Lite, T.-L. V. o. & Kruglyak, L. Finding the sources of missing heritability in a yeast cross. *Nature* 1–6 (2013).
- ⁶ Carlborg, O., Jacobsson, L., Ahgren, P., Siegel, P. & Andersson, L. Epistasis and the release of genetic variation during long-term selection. *Nature Genetics* **38**, 418–420 (2006).
- ⁷ Strange, A. *et al.* A genome-wide association study identifies new psoriasis susceptibility loci and an interaction between HLA-C and ERAP1. *Nature Genetics* **42**, 985–90 (2010).
- ⁸ Evans, D. M. *et al.* Interaction between ERAP1 and HLA-B27 in ankylosing spondylitis implicates peptide handling in the mechanism for HLA-B27 in disease susceptibility. *Nature Genetics* **43** (2011).
- ⁹ Cordell, H. J. Detecting gene-gene interactions that underlie human diseases. *Nature Reviews Genetics* **10**, 392–404 (2009).
- ¹⁰ Hemani, G., Theocharidis, A., Wei, W. & Haley, C. EpiGPU: exhaustive pairwise epistasis scans parallelized on consumer level graphics cards. *Bioinformatics (Oxford, England)* **27**, 1462–5 (2011).
- ¹¹ Metspalu, A. The Estonian Genome Project. *Drug Development Research* **62**, 97–101 (2004).
- ¹² Fehrmann, R. S. N. *et al.* Trans-eQTLs reveal that independent genetic variants associated with a complex phenotype converge on intermediate genes, with a major role for the HLA. *PLoS genetics* **7**, e1002197 (2011).
- ¹³ Lieberman-Aiden, E. *et al.* Comprehensive mapping of long-range interactions reveals folding principles of the human genome. *Science (New York, N.Y.)* **326**, 289–93 (2009).

- ¹⁴ Visscher, P. M., Brown, M. a., McCarthy, M. I. & Yang, J. Five years of GWAS discovery. *American journal of human genetics* **90**, 7–24 (2012).
- ¹⁵ Weinreich, D. M., Delaney, N. F., Depristo, M. a. & Hartl, D. L. Darwinian evolution can follow only very few mutational paths to fitter proteins. *Science (New York, N.Y.)* **312**, 111–4 (2006).
- ¹⁶ Breen, M. S., Kemena, C., Vlasov, P. K., Notredame, C. & Kondrashov, F. a. Epistasis as the primary factor in molecular evolution. *Nature* **490**, 535–538 (2012).
- ¹⁷ Weir, B. S. Linkage disequilibrium and association mapping. *Annual review of genomics and human genetics* **9**, 129–42 (2008).
- ¹⁸ Hemani, G., Knott, S. & Haley, C. An Evolutionary Perspective on Epistasis and the Missing Heritability. *PLoS Genetics* **9**, e1003295 (2013).
- ¹⁹ Marchini, J., Donnelly, P. & Cardon, L. R. Genome-wide strategies for detecting multiple loci that influence complex diseases. *Nature Genetics* **37**, 413–417 (2005).
- ²⁰ Lango Allen, H. *et al.* Hundreds of variants clustered in genomic loci and biological pathways affect human height. *Nature* **467**, 832–8 (2010).
- ²¹ Powell, J. E. *et al.* Congruence of Additive and Non-Additive Effects on Gene Expression Estimated from Pedigree and SNP Data. *PLoS Genetics* **9**, e1003502 (2013).
- ²² Powell, J. E. *et al.* The Brisbane Systems Genetics Study: genetical genomics meets complex trait genetics. *PloS one* **7**, e35430 (2012).
- ²³ Preiner, M. *et al.* Blood-informative transcripts define nine common axes of peripheral blood gene expression. *PLoS genetics* **9**, e1003362 (2013).
- ²⁴ Cockerham, C. C. An extension of the concept of partitioning hereditary variance for analysis of covariances among relatives when epistasis is present. *Genetics* **39**, 859–882 (1954).
- ²⁵ Ho, T. H. *et al.* Muscleblind proteins regulate alternative splicing. *The EMBO journal* **23**, 3103–12 (2004).
- ²⁶ Trynka, G. *et al.* Chromatin marks identify critical cell types for fine mapping complex trait variants. *Nature genetics* **45**, 124–30 (2013).
- ²⁷ Ward, L. D. & Kellis, M. HaploReg: a resource for exploring chromatin states, conservation, and regulatory motif alterations within sets of genetically linked variants. *Nucleic acids research* **40**, D930–4 (2012).
- ²⁸ Lan, X. *et al.* Integration of Hi-C and ChIP-seq data reveals distinct types of chromatin linkages. *Nucleic acids research* **40**, 7690–704 (2012).

- ²⁹ Osborne, C. S. *et al.* Active genes dynamically colocalize to shared sites of ongoing transcription. *Nature genetics* **36**, 1065–71 (2004).
- ³⁰ Rieder, D., Trajanoski, Z. & McNally, J. G. Transcription factories. *Frontiers in genetics* **3**, 221 (2012).
- ³¹ Visscher, P. M., Hill, W. G. & Wray, N. R. Heritability in the genomics era—concepts and misconceptions. *Nature Reviews Genetics* **9**, 255–66 (2008).
- ³² Churchill, G. A. & Doerge, R. W. Empirical threshold values for quantitative trait mapping. *Genetics* **138**, 963–71 (1994).