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Gene Expression Networks in the Drosophila Genetic Reference Panel

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ABSTRACT

A major challenge in modern biology is to understand how naturally occurring variation in DNA sequences affects complex organismal traits through networks of intermediate molecular phenotypes. This question is best addressed in a genetic mapping population in which all molecular polymorphisms are known and for which molecular endophenotypes and complex traits are assessed on the same genotypes. Here, we performed deep RNA sequencing of 200 Drosophila Genetic Reference Panel inbred lines with complete genome sequences and for which phenotypes of many quantitative traits have been evaluated. We mapped expression quantitative trait loci for annotated genes, novel transcribed regions, transposable elements and microbial species. We identified host variants that affect expression of transposable elements, independent of their copy number, as well as microbiome composition. We constructed sex-specific expression quantitative trait locus regulatory networks. These networks are enriched for novel transcribed regions and target genes in heterochromatin and euchromatic regions of reduced recombination, and genes regulating transposable element expression. This study provides new insights regarding the role of natural genetic variation in regulating gene expression and generates testable hypotheses for future functional analyses.

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INTRODUCTION

Understanding how naturally occurring genetic variation affects variation in organismal quantitative traits by modifying underlying molecular networks is a key challenge in modern biology. Most traits are highly polygenic (Visscher et al. 2012; Mackay et al. 2009; Mackay and huang, 2018) and associated molecular variants have small additive effects on trait variation (Manolio et al. 2009). Most of these variants are in intergenic regions, up- or down-stream of coding regions, or in introns, and presumably play a regulatory role in modulating gene expression.

Systems genetics analysis seeks to determine how naturally occurring molecular variation gives rise to genetic variation in organismal phenotypes by examining genetic variation in gene expression (expression quantitative trait loci, or eQTLs) and other intermediate molecular phenotypes (Sieberts et al. 2007; Chen et al. 2008; Emilsson et al. 2008; Rockman 2008; Cookson et al. 2009; Mackay et al. 2009; Civelek and Lusis 2014; Albert and Kruglyak 2015; Gibson et al. 2015; Ogura and Busch 2016; Schughart and Williams 2017). Polymorphic variants associated with variation in gene expression are classified as *cis*- or *trans*-eQTLs depending on whether they are proximal or distal to the gene encoding the transcript, respectively. Genetic variation in gene expression is pervasive; *cis*-eQTLs can have large effects on gene expression that are detectable in small samples; and variants associated with human diseases and quantitative traits tend to be enriched for *cis*-eQTLs (Sieberts et al. 2007; Chen et al. 2008; Emilsson et al. 2008; Rockman 2008; Cookson et al. 2009; Mackay et al. 2009; Nicolai et al. 2010; Civelek and Lusis 2014; Albert and Kruglyak 2015; Gibson et al. 2015; Ogura and Busch 2016; Boyle et al. 2017; Schughart and Williams 2017). eQTLs with both *cis*- and *trans*- effects can be assembled into directed transcriptional networks of regulator and target genes (Liu et al. 2008; Bryois et al. 2014; Fagny et al. 2017). Elucidating such regulatory transcriptional networks will facilitate understanding how the effects of individual variants

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propagate through the network, how multiple variants together regulate gene expression and affect complex traits (Liu et al. 2008; Nicolae et al. 2010; Bryois et al. 2014; Fagny et al. 2017), and improve genomic prediction (Zhou et al. 2020).

Here, we performed deep RNA sequencing of the *Drosophila melanogaster* Genetic Reference Panel (DGRP) of inbred lines with complete DNA sequences (Mackay et al. 2012; Huang et al. 2014). We mapped eQTLs for annotated genes, novel transcribed region (NTRs), transposable elements (TEs) and microbiome composition; constructed *de novo cis-trans* eQTL gene expression networks; and evaluated associations of eQTLs and expression traits with organismal phenotypes.

RESULTS

We collected and sequenced ribo(-) RNA from replicate pools of young flies from each of 200 DGRP lines, separately for males and females. In total we sequenced 1.94 Terabases of RNA, of which on average 13.4 million reads per sample uniquely aligned to the *D. melanogaster* genome (Supplemental Table S1). The sequences were processed through a pipeline (Supplemental Fig S1) that (i) removes adapter and rRNA sequences; (ii) aligns and quantifies expressed TE sequences and microbial transcripts; (iii) verifies the origin of each sample; and (iv) quantifies known and novel *D. melanogaster* transcripts and corrects for potential alignment bias due to line-specific sequence variation. We then analyzed normalized expression values for endogenous genes, TEs and microbial species.

Genetic Variation in Gene Expression

We quantified expression levels of all RNA sequences that aligned to the reference genome in each DGRP line. After elimination of sequences with low expression, we found that 12,806 of 17,097 known *D. melanogaster* genes (75%) were expressed consistently in young adult males

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and/or females (Supplemental Table S2A). In addition, we identified 4,282 novel transcribed regions (NTRs) (Supplemental Table S2B) that showed no overlap with exons on the same strand. A total of 3,846 of the NTRs were located in introns; 290 were anti-sense to known genes, and 146 were intergenic. Most (95.6%) of the NTRs are ≥ 200 bp; the majority (4,149 or 96.9%) lack protein coding potential (Kang et al. 2017) (Supplemental Table S2C). These NTRs in total represent 5.61 Mb new transcribed mature RNA sequences that eluded prior annotation efforts. This increase is likely due to the multiple genetic backgrounds profiled in this study. Although RNA-seq alignment and assembly alone are not sufficient to prove genuine transcriptional activities, our stringent expression-based filter was able to narrow down the NTRs to a subset that were similar to known genes in terms of mapping ambiguity (Supplemental Fig S2) and expression in at least one Drosophila cell line (Supplemental Fig S3).

Variation in gene expression among the DGRP lines may be confounded by variation in alignment rate to the reference strain due to variation in DNA sequences between the DGRP lines and the reference. Indeed, 2,735 genes (2,117 known genes and 618 NTRs) were affected by alignment bias (Supplemental Table S2D). We corrected for alignment bias, and partitioned variation in gene expression between males and females, DGRP lines, the sex by line interaction, and residual (environmental) terms (Supplemental Table S2D), using a false discovery rate of FDR ≤ 0.05. Similar to previous studies (Ayroles et al. 2009; Massouras et al. 2012; Huang et al. 2015), we found that gene expression is sexually dimorphic: 98% (96%) of expressed known genes (NTRs) have a significant sex effect (Fig 1A, Supplemental Table S2D). There is genetic variation in the magnitude of sex dimorphism: 69% (10%) of expressed known genes (NTRs) have a significant sex by line interaction (Supplemental Table S2D). Therefore, we assessed genetic variation in gene expression separately for males and females (Supplemental Tables S2D, S2E), and found that 12,151 genes (10,354 known genes and 1,797 NTRs) were genetically variable in females (Fig 1B) and 13,819 genes (11,393 known

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genes and 2,426 NTRs) were genetically variable in males (Fig 1C). These numbers of genes with significant genetic variation are much higher than previously reported studies, which used microarrays (4,308 in females and 5,814 in males) rather than RNA-seq (Huang et al. 2015). Relative to tiling arrays, RNA-seq has a higher dynamic range and greater precision in quantifying gene expression, although the results from both analyses are positively correlated (Supplemental Fig S4).

Broad sense heritabilities (proportion of phenotypic variance due to genotype differences) ranged from $H^2 = 0.148 – 0.986$ in females and $H^2 = 0.145 – 0.986$ in males (Figs 1B, 1C). Notably, 472 (514) of the genetically variable genes in females (number for males in parenthesis) were located in molecularly defined heterochromatin (*2LHet*, *2RHet*, *3LHet*, *3RHet*, *XHet*, and *YHet*) and Chromosome *4*. While there are 6.92× (5.52×) as many annotated genes relative to NTRs in euchromatic regions in females (males); there are 2.21 × (3.18×) as many NTRs in heterochromatin and Chromosome *4* in females (males) (Supplemental Table S2G). Thus, NTRs are highly enriched in heterochromatic regions.

We used weighted gene co-expression network analysis (WGCNA, Langfelder and Horvath 2008) to assess the extent to which gene expression levels are genetically correlated in each sex (Figs 1D, 1E, Supplemental Table S3). We found 13 (15) co-expression modules in females (males). We assessed the extent to which each module was significantly (FDR ≤ 0.05) enriched for Gene Ontology (GO) terms and pathway and protein domain annotations (Lyne et al. 2007) (Supplemental Table S3). For example, female Module 2 (149 genes) is enriched for GO terms involved in ovary function and male Module 6 (365 genes) is enriched for biological process GO terms involved in male reproduction. Female Module 12 (88 genes) and male Modules 13 (35 genes) and 14 (165 genes) are enriched for GO terms affecting small molecule metabolism. Female Modules 3 (26 genes), 6 (27 genes), and 7 (21 genes) and male Modules 9 (42 genes) and 12 (44 genes) are enriched for GO terms affecting innate immunity, and female Module 13 (560 genes) is enriched for GO terms affecting chemosensation.

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Gene Expression QTLs (eQTLs)

We performed genome wide association eQTL analyses for each of the genetically variable genes in each sex. We used ~1,932,427 common (minor allele frequency > 0.05) polymorphisms and accounted for effects of Wolbachia infection, polymorphic inversions and polygenic relatedness on gene expression (Huang et al. 2014; Huang et al. 2015). We mapped 90,634 eQTLs in females and 147,412 eQTLs in males (FDR ≤ 0.05). A total of 2,053 genes in females (1,818 known genes and 235 NTRs) and 3,178 genes in males (2,790 known genes and 388 NTRs) were associated with at least one significant eQTL. We defined potentially *cis*- and *trans*-regulatory eQTLs as ≤ 1 kb and > 1 kb of their respective gene bodies. We mapped putative *cis*-eQTLs to 1,284 (2,154) genes in females (males) and *trans*-eQTLs to 1,653 (2,521) genes in females (males), of which 902 (1,305) were *trans*-eQTLs located on different chromosomal arms (Supplemental Tables S4A, S4B).

Due to correlation between genotypes at putative eQTL positions, some genes contained a large number of eQTLs that were not independent from each other. To develop a more parsimonious model, we used forward stepwise model selection to select putative eQTLs from the significant candidates, conservatively requiring that the last eQTL entering the model had a conditional *P* value < 1×10^{-5}. The models contained between one and seven eQTLs, with more than 60% of genes containing only one eQTL (Supplemental Tables S4A, S4B). After model selection, we visualized the significant eQTLs by plotting the polymorphism positions on the *X*- axis and the gene positions on the *Y*-axis such that the diagonal corresponds to *cis*-eQTLs and the off-diagonal to *trans*-eQTLs (Fig 2). We found the majority of eQTLs retained by model selection to be in *cis* with the genes they controlled, though *trans*-eQTLs were not uncommon (Fig 2).

eQTL Regulatory Networks

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The existence of eQTLs that are *cis*-eQTL for gene X and also *trans*-eQTL for gene Y (Supplemental Tables S5A, S5B) enables us to construct gene regulatory networks based on multifactorial variation in a natural population. Although significant putative eQTLs may not remain in the selected models, we still considered them when constructing regulatory networks because we could not genetically distinguish them and their associations with gene expression when all *P*-values were highly significant. We identified 408 (794) such regulatory interactions supported by at least one *cis-trans* eQTL connecting 257 (471) regulatory genes (*cis* end) to 251 (447) target genes (*trans* end) in females (males) (Supplemental Tables S5C, S5D). There are two or three large regulatory networks in each sex, and many smaller networks (Supplemental Figs S5, S6, S7). The regulatory genes are largely distinct between the two sexes, although many target genes are in common between males and females (Fig 3, Supplemental Fig S5, Supplemental Table S5E). Genes from the sex-specific regulatory networks or from the common networks are not enriched for any GO terms. It is not clear from their anatomical gene expression patterns how the sex-specificity could arise, since the majority of these genes are expressed in multiple tissues, including the reproductive tissues of both sexes (Gramates et al. 2017).

Examination of *cis* and *trans* eQTLs (Supplemental Table S5) showed that there are more NTRs than expected among genes with *cis-trans* eQTLs based on the total number of NTRs with eQTLs among the target genes ($\chi^2_1 = 29.74$, $P = 4.95 \times 10^{-8}$ in females; $\chi^2_1 = 60.54$, $P = 7.20 \times 10^{-15}$ in males) but not the regulatory genes ($\chi^2_1 = 1.54$, $P = 0.21$ in females; $\chi^2_1 = 1.49$, $P = 0.22$ in males). The regulatory genes tend to be located in pericentromeric regions of reduced recombination (Fiston-Lavier et al. 2010) ($\chi^2_1 = 17.28$, $P = 3.23 \times 10^{-5}$ in females; $\chi^2_1 = 120.28$, $P < 2.2 \times 10^{-16}$ in males) and target gene locations are enriched for heterochromatin and pericentromeric regions of reduced recombination ($\chi^2_1 = 28.53$, $P = 9.21 \times 10^{-8}$ in females; $\chi^2_1 = 147.78$, $P < 2.2 \times 10^{-16}$ in males). Regulatory genes with many target genes thus tend to have

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multiple *cis*-eQTLs in linkage disequilibrium (LD) near the centromere, and regulate other NTRs both in heterochromatic regions across the genome and euchromatic regions on other chromosomes (Fig 3, Supplemental Figs S5, S6, S7). The smaller networks with fewer regulators and targets tend to consist of genes in euchromatin in regions of normal recombination (Fig 3, Supplemental Figs S5, S6, S7; Supplemental Tables S5C, S5D). Regulatory genes often have many *cis*-eQTLs; a single *cis*-eQTL can regulate multiple target genes; and multiple *cis*-eQTLs (which may or may not be in LD) within a gene can regulate different target genes. It is possible that multiple *cis*-eQTLs in LD can be classified as a *trans*-eQTL for different target genes due to differences in thresholding and ranking of eQTLs among the target genes. Each gene with at least one *cis*-eQTL may itself be regulated in *trans* by *cis*-eQTLs in one or more upstream genes, and the genes regulated by a focal *cis*-eQTL may themselves have *cis*-eQTLs regulating other genes.

Genetic Variation in TE Expression

A total of 9% of the *D. melanogaster* genome contains TEs spanning multiple families (Spradling and Rubin 1981). Active retrotransposon sequences are present in our RNA-seq libraries. We aligned reads to the RepBase database of known repetitive elements (Jurka et al. 2005), and quantified TE RNA levels based on normalized read counts. Overall, 1.3% of the RNA-seq reads align to RepBase. The most abundant families of TE sequences were *gypsy*, *copia*, *BEL*, *jockey* and *Mariner*/*Tc1* elements, but all TE families represented in RepBase were detected (Fig 4A, Supplemental Table S6A).

Line-specific differences in TE RNA levels can be driven by both differences in underlying copy number (Lee and Langley 2010) and differences in the rate of transcription per genomic copy. We quantified DNA copy variation for each TE sequence (Supplemental Table S6B) and used linear models to estimate the percentage of variation in TE expression that arises from differences in copy number (Supplemental Table S6C). We then partitioned the

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remaining copy number-independent variation in TE expression between sexes, DGRP lines, the line by sex interaction and residual terms (Supplemental Table S6C), using FDR ≤ 0.05 as the significance threshold for each term in the analysis. Since the majority (153, 79%) of TEs had a significant sex by line interaction effect, we assessed genetic variation in TE expression for each transposon sequence separately for each sex (Supplemental Tables S6D, S6E). We observed significant genetic variation in expression for 187 (97%) TE sequences in females (Fig 4B) and 186 (96%) TE sequences in males (Fig 4C). Broad sense heritabilities of TE expression ranged from H^2 = 0.15 – 0.99 in females and H^2 = 0.15 – 0.98 in males (Figs 4B, 4C). Thus, there is host genetic control of expression for most *D. melanogaster* TEs.

We assessed whether different TE sequences had similar patterns of expression across the DGRP lines (Langfelder and Horvath 2008), separately for males and females (Figs 4D, 4E, Supplemental Tables S6F, S6G). We found minimal correlation structure in the activity scores of different TEs (Supplemental Table S6H), with the strongest correlations between pairs of TE sequences from the same family. This suggests that host genetic factors independently affect variation in expression of each TE family.

TE eQTLs

We mapped eQTLs for each of the TEs with genetically variable expression in females and males (Supplemental Table S7). We found 54 TEs with significant eQTLs (FDR ≤ 0.05), 36 in females and 39 in males. A total of 20 TE sequences were expressed in both males and females; 16 (18) TE sequences were expressed only in females (males). The number of eQTLs per TE sequence ranged from 1-1,020, with on average more eQTL associations for TEs in males than females (Supplemental Tables S7A-C). However, forward model selection retained between one and four eQTLs associated with TE activity, suggesting substantial LD among the eQTLs. Indeed, the large numbers of eQTLs associated with some TEs were located in LD

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blocks in pericentromeric regions and on the *4*th chromosome (Supplemental Fig S8, Supplemental Tables S7D, S7E). Many eQTLs for TEs expressed in both males and females overlapped between the sexes, but typically additional eQTLs were present in males. Although there was little clustering of expression patterns of different TE sequences, 202 (1,032) eQTLs were associated with two or more sequences in females (males) (Supplemental Tables S7F, S7G).

Many eQTLs associated with TE expression were within 1 kb of annotated genes and NTRs. Indeed, 19.8% (17.7%) of TE eQTLs were within 1 kb of NTRs in females (males). Known genes near TE eQTLs were enriched (FDR < 0.05) for GO categories related to regulation of gene expression and protein binding (Supplemental Table S7H). We next asked to what extent eQTLs associated with gene expression were also associated with expression of TE sequences. We found 1,206 eQTLs associated with 85 genes (37 known genes and 48 NTRs) and 23 TEs in females; and 3,656 eQTLs associated with 166 genes (79 known genes and 87 NTRs) and 30 TEs in males (Supplemental Fig S9, Supplemental Table S8). We could thus incorporate variation in TE expression into the *cis-trans* gene regulatory network via shared eQTLs (Fig 5). These eQTLs are predominantly located in pericentromeric regions, and the genes they regulate are in pericentromeric regions as well as heterochromatin.

Genetic Variation in Microbiome Composition

RNA samples extracted from pools of whole flies contain RNA from gut microbial communities, and from microbes on their exoskeleton. We assessed the contribution of microbial sequences to the RNA-seq libraries by aligning reads to a database of candidate microbial genomes (Supplemental Table S9). *Wolbachia pipientis*, a bacterial endosymbiont that infects ~50% of the DGRP lines (Mackay et al. 2012), is the most abundant source of expressed sequence, followed by multiple Acetobacter species and genome assemblies (Fig 6A, Supplemental Table S9). We estimated the total gene expression from each microbial species in all samples

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(Supplemental Table S10A) and partitioned variation in microbial gene expression between sexes, DGRP lines, the sex by line interaction and residual terms, using FDR ≤ 0.05 as the significance threshold (Supplemental Table S10B). The H^2 of *Wolbachia pipientis* abundance is extremely high (H^2 = 0.972), as expected. We next assessed whether the sum of all non-Wolbachia microbial species is genetically variable after accounting for any Wolbachia effects, and estimated H^2 = 0.595 (Fig 6B, Supplemental Table S10B). The sex by line interaction for total microbial gene expression was not significant, indicating that total microbial RNA is highly correlated between males and females. We estimated the heritability of gene expression for the 122 non-Wolbachia microbial species, and found that 84 microbial species had significant genetic variation in RNA abundance, with broad sense heritabilities ranging from H^2 = 0.07 – 0.90 (Fig 6C, Supplemental Table S10B). Microbial species that are likely to colonize the Drosophila gut (Acetobacter and Lactobacillus species) were among those with the highest H^2.

We used WGCNA (Langfelder and Horwath 2008) to group species with similar abundance patterns based on the average of male and female line means (Fig 6D, Supplemental Tables S10C, S10D). We found three groups of strongly correlated species, consisting primarily of the gut-related microbes (Acetobacter and Lactobacillus species), and two additional clusters of microbes primarily consisting of viral and fungal species that are strongly anti-correlated with the abundances of species in the first three clusters. Thus, there is line-specific variation in the microbial communities living in and on DGRP flies. Species which most plausibly colonize the Drosophila gut are largely correlated across lines, with some fluctuation in the relative abundance of Acetobacter versus Lactobacillus species.

eQTLs for Microbiome Composition

There was little genetic variation in sexual dimorphism for microbial gene expression; therefore, we performed eQTL mapping using the average expression of males and females for each microbial species. Four microbial species and total microbial sequence expression were

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associated with significant eQTLs (FDR ≤ 0.05) (Supplemental Table S11A). The sum of all microbial species is associated with one eQTL that maps to an NTR; the expression of *Borrelia coriaceae*, *Acidovorax temperans* and *Podospora anserine* map, respectively, to single eQTLs in *CG2616, CG46301,* and to *cic* and an NTR; and *Leuconostoc pseudomesenteroides* expression maps to 39 variants in or near *GC* and *nSyb* (Supplemental Table S11A).

We lowered the significance threshold to $P < 10^{-5}$ to explore the extent to which common eQTLs may control the expression of multiple microbial species that cluster together based on the WGCNA analysis (Fig 6D). At this threshold, 1,455 eQTLs are associated with 88 microbial species and the sum of all species (Supplemental Table S11B); 268 variants were associated with expression of more than one microbial species, and five eQTLs were associated with expression of 10 or more microbial species (Supplemental Table S11C). These data suggest that there is genetic variation in host control of microbial gene expression and that some variants have pleiotropic effects on multiple microbial species.

We assessed whether the genes to which the eQTLs associated with variation in microbial gene expression were enriched for GO categories (FDR ≤ 0.05). The most highly enriched Biological Process GO terms were related to development and morphogenesis, including development and function of the nervous system (Supplemental Table S11D).

Gene Expression and Complex Traits

To examine the relationship between variation in gene expression and variation in organismal quantitative trait phenotypes, we chose 11 quantitative traits with published phenotypic data (chill coma recovery time and startle response (Mackay et al. 2012); starvation resistance (Huang et al. 2014); day and night sleep bout number, day and night total sleep duration, and total waking activity (Harbison et al. 2013); food consumption (Garlapow et al. 2015); male aggression (Shoprter et al. 2015); phototaxis (Carbone et al. 2016)); and additionally measured

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five metabolic traits (levels of free glucose, glycogen, free glycerol, triglyceride and protein) and three metrics of body size (body weight, thorax length, thorax width). All traits were quantified in the same laboratory under the same culture conditions used in this study. The line means for all traits are given in Supplemental Table S12; quantitative genetic analyses of the metabolic and body size traits are given in Supplemental Table S13; and the most significant associations ($P < 10^{-5}$) from GWA analyses (separately for males and females) for these quantitative traits based on the 200 lines for which we have gene expression data are in Supplemental Table S14.

We first assessed whether variants associated with all organismal traits were enriched for eQTLs, as found in human studies (Chen et al. 2008; Emilsson et al. 2008; Cookson et al. 2009Nicolae et al. 2010; Boyle et al. 2017). Of all the eQTLs (prior to model selection) and GWAS hits, only 26 in males and 8 in females were common between eQTLs and GWAS hits, with clear patterns of clustering. We found no enrichment of *cis*-eQTLs ($P = 0.13$ in females and $P = 0.71$ in males), *trans*-eQTLs ($P = 0.98$ in females and $P = 0.28$ in males) or all eQTLs ($P = 0.94$ in females and $P = 0.23$ in males) among top GWA hits in either sex. Many top GWA hits as well as eQTLs map to regions greater than 1kb from any gene, and may indicate novel regulatory regions. To exclude the possibility that the lack of overlap was due to using different mapping procedures, we performed QTL mapping for the organismal traits using the same procedure as the eQTL mapping. At an empirical FDR = 0.05, we found four SNPs associated with three traits (chill coma recovery in females, day sleep duration and free glucose level in males) and none was an eQTL.

We next performed transcriptome wide association studies (TWAS) for individual genetically variable transcripts for gene expression, TE sequences and microbial species, for each of the 18 (19) genetically variable organismal phenotypes in females (males). We found several significant (Benjamini-Hochberg FDR < 0.05) associations of transcripts with organismal phenotypes (Supplemental Table S15). These associations include a known noncoding RNA (*CR46032*) with male aggression, two NTRs with male waking activity, *Gbs-70E* with free

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glucose in both sexes, *AkhR* with starvation resistance in males and females, and *Acidovorax temperans* with male aggression (Supplemental Table S15).

DISCUSSION

Deep RNA sequencing gives accurate estimates of gene expression of annotated genes and can implicate novel non-coding RNAs and their regulatory interactions with annotated genes. We have identified 4,282 novel transcribed regions, which are unlikely to be artifacts since the majority are genetically variable, and they are not randomly distributed in the genome but are preferentially located in heterochromatic regions and in pericentromeric euchromatin bordering heterochromatin. Thus, there is genetic variation in heterochromatic gene expression, thought to be largely transcriptionally silent (Riddle et al. 2011). These heterochromatic and pericentromeric NTRs are regulated by pericentromeric *cis*-eQTLs as well as *trans*-eQTLs dispersed throughout the euchromatic genome. Genes associated with eQTLs with both *cis*- and *trans*- effects form sex-specific networks of regulator and target genes, the largest of which is enriched for NTR target genes in heterochromatin and regulator and target genes in pericentromeric euchromatin. The considerable overlap between eQTLs associated with NTRs in the large networks and eQTLs associated with TE expression recruits TEs to the network. We do not know where the TE sequences with genetically variable expression are integrated in the genome; however, heterochromatin is composed of largely silenced TE repeats (Riddle et al. 2011), raising the possibility that TEs in heterochromatin are subject to the same regulation as other heterochromatic genes. Further work is needed to confirm the regulatory networks derived from naturally occurring genetic variation and determine the regulatory mechanism(s) through which the NTRs act. We speculate that many of the NTRs may be long noncoding RNAs, operationally defined as encoding transcripts > 200 bp with no significant protein-coding

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potential, but further work is needed to establish whether this is true (Khalil et al. 2009; Wang et al. 2011; Hacisuleyyman et al. 2014; Rogoyski et al. 2017; Ransohoff et al. 2018).

The first step in systems genetic analysis is to identify eQTLs associated with both gene expression and organismal quantitative traits, for which variation in gene expression is correlated with variation in the organismal phenotypes (Sieberts and Schadt 2007; Rockman 2008; Mackay et al. 2009). We did not find any such trios, although we did find interesting transcript-trait associations. This may be because our sample size is adequate to detect eQTLs but not QTLs affecting organismal traits, which have smaller effects; because eQTLs need to be mapped in tissues relevant to the organismal trait; and because there are non-linear (epistatic) relationships between QTLs for both transcripts and organismal phenotypes. The complex and highly connected *cis-trans* regulatory networks suggest that higher order interactions need to be accommodated in systems genetic modeling, at least at the level of gene expression.

METHODS

Drosophila lines

We used 200 inbred, sequenced DGRP lines (Mackay et al. 2012; Huang et al. 2014), established by 20 generations of full sib inbreeding from gravid females collected at the Raleigh, NC USA Farmer's Market. Genome sequences of the lines were obtained previously using the Illumina platform with an average of coverage of 27×. A total of 4,565,215 molecular variants (3,976,011 single/multiple nucleotide polymorphisms (SNPs/MNPs), 169,053 polymorphic insertions (relative to the reference genome), 293,363 polymorphic deletions and 125,788 polymorphic microsatellites) segregate in the DGRP.

Sample collection

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All lines were reared on cornmeal-molasses-agar medium at 25°C, 60–75% relative humidity and a 12-hr light-dark cycle at equal larval densities. We collected two replicates of 25 females and 30 males per line, for a total of 800 samples. We used a strict randomized experimental design for sample collection. We collected mated 3-5 day old flies between 1-3 pm. We transferred the flies into empty culture vials and froze them over ice supplemented with liquid nitrogen, and sexed the frozen flies. The samples were transferred to 2.0 ml nuclease-free microcentrifuge tubes (Ambion) and stored at -80°C until ready to process.

RNA sequencing

Total RNA was extracted with QIAzol lysis reagent (Qiagen) and the Quick-RNA MiniPrep Zymo Research Kit (Zymo Research). Ribosomal RNA (rRNA) was depleted from 5 ug of total RNA using the Ribo-Zero^{TM} Gold Kit (Illumina, Inc). Depleted mRNA was fragmented and converted to first-strand cDNA using SuperScript III reverse transcriptase (Invitrogen). During the synthesis of second strand cDNA, dUTP instead of dTTP was incorporated to label the second strand cDNA. cDNA from each RNA sample was used to produce barcoded cDNA libraries using NEXTflex™ DNA Barcodes (Bioo Scientific, Inc.) with an Illumina TruSeq compatible protocol. Libraries were size-selected for 250 bp (insert size ~130 bp) using Agencourt Ampure XP Beads (Beckman Coulter, Inc.). Second strand DNA was digested with Uracil-DNA Glycosylase before amplification to produce directional cDNA libraries. Libraries were quantified using Qubit dsDNA HS Kits (Life Technologies, Inc.) and Bioanalyzer (Agilent Technologies, Inc.) to calculate molarity. Libraries were then diluted to equal molarity and re-quantified. A total of 50 pools of 16 libraries were made, again randomly assigning samples to each pool. Pooled library samples were quantified again to calculate final molarity and then denatured and diluted to 14pM. Pooled library samples were clustered on an Illumina cBot; each pool was sequenced on one lane of Illumina HiSeq 2500 using 125 bp single-read v4 chemistry.

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RNA sequence analysis

Barcoded sequence reads were demultiplexed using the Illumina pipeline v1.9. Adapter sequences were trimmed using cutadapt v1.6 (Martin 2011) and trimmed sequences shorter than 50bp were discarded from further analysis. Trimmed sequences were then aligned to multiple target sequence databases in the following order, using BWA v0.7.10 (MEM algorithm with parameters '-v 2 –t 4') (Li and Durbin 2010): (1) all trimmed sequences were aligned against a database containing the complete 5S, 18S-5p8S-2S-28S, mt:lrRNA, and mt:srRNA sequences to filter out residual rRNA that escaped depletion during library preparation; (2) remaining sequences were then aligned against a custom database of potential micriobiome component species (see below) using BWA; (3) sequences that did not align to either the rRNA or microbiome databases were aligned to all *D. melanogaster* sequences in Repbase (Jurka et al. 2005). The remaining sequences that did not align to any of the databases above were then aligned to the *D. melanogaster* genome (BDGP5) and known transcriptome (FlyBase v5.57) using STAR v2.4.0e (Dobin et al. 2013). Libraries with fewer than 5 million reads uniquely aligned to the *D. melanogaster* reference genome were re-sequenced to achieve sufficient read depth.

Generation of microbiome database

We first performed a preliminary alignment of RNA-seq reads by filtering only rRNA sequences, and then aligning directly to the *D. melanogaster* genome using the tools and parameters described above. Sequences that did not align to the rRNA database or *D. melanogaster* reference genome were then analyzed with Trinity v2.1.1 (Garbherr et al. 2011) to perform *de novo* assembly of longer sequences from the short reads. Assembled sequences > 1kb in length were then searched against the RefSeq_genomic database (downloaded from NCBI on 1/27/16) using BLAST. We then compiled a list of all RefSeq genomes that were found as a top

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BLAST hit for at least two assembled sequences. We compiled all FASTA files for each of these RefSeq genomes into a single database for alignment with BWA.

Genotype validation

To validate the DGRP line assigned to each RNA-seq sample, we identified single nucleotide polymorphisms (SNPs) from the RNA-seq reads that aligned to the *D. melanogaster* reference genome using STAR as described above. We retained only those SNP calls covered by at least 3 reads and at least 75% of all reads supporting the major genotype (note that DGRP lines are inbred and therefore the majority of SNPs are homozygous). This filtering process produced >400k usable SNPs per sample, primarily located in transcribed regions of the genome. We then performed two validation tests of the DGRP line assigned to each sample *X* by comparing to the previously published genotype calls for each DGRP line (http://dgrp2.gnets.ncsu.edu/data/website/dgrp2.tgeno; Huang et al. 2014). First, we computed the "line mean error" (LME) for each line as follows: given the set of homozygous SNPs from line *Y* that have sufficient coverage (described above) in sample *X*, LME(*X*,*Y*) = # of mismatching SNPs / total # of comparable SNPs. We confirmed that for each sample *X*, the DGRP line Y_{lab} labeled for that samples produced the minimum value of LME(*X*,*Y*) as compared to all other possible line assignments Y_{alt}, and further confirmed that LME(X,Y_{lab}) was below 1%. Second, we performed competitive tests between the labeled line Y_{lab} and each possible alternate line Y_{alt}. Under this test, we considered only the SNPs that are homozygous for different genotypes in Y_{lab} and Y_{alt} (*i.e.*, only the segregating SNPs for the two lines) and which have sufficient coverage in sample *X*. We then computed the "line error ratio" (LER) = # of SNPs matching Y_{lab} / # of SNPs matching Y_{alt}. We confirmed that for each sample *X*, the lowest LER for any Y_{alt} was > 1 (*i.e.*, the majority of SNP calls always supported the labeled line compared to any alternative line).

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Inference of novel transcripts

We constructed a *de novo* transcriptome for each individual sample by inputting the RNA-seq reads aligned to the *D. melanogaster* reference genome into Cufflinks v2.2.1 (Trapnell et al. 2012). We also considered the novel transcribed regions (NTRs) identified in a previous study based on unstranded pooled RNA sequencing of the DGRP lines (Huang et al. 2015). However, the previously published data do not provide strand-specific signal, while our current RNA-seq data uses a strand-specific library preparation. Therefore, we reassigned the strand for each of the previously published NTRs that was supported by the greater number of total aligned reads across all samples. We then merged all *de novo* sample transcriptomes and the previously published NTRs using the cuffmerge tool included with Cufflinks v2.2.1, then removed all merged transcript models with any exon overlapping on the same strand any exon in the known *D. melanogaster* transcriptome. We defined the known transcriptome here as all gene models in FlyBase v5.57 plus all subsequently added gene models in FlyBase v6.11 to account for recently discovered lncRNA sequences. Thus, the final output of this analysis was a set of NTRs constructed from both our current RNA-seq data and previously published pooled RNA-seq data that do not overlap any known gene exons on the same strand.

Gene expression estimation

Read counts for individual microbial species were computed as all reads aligning to any sequence in any genome for any strain of that species. Reads aligning to multiple species were ignored for individual species read counts. We also aligned microbiome-aligning reads to the *D. melanogaster* genome, and removed all reads that aligned to both microbial and *D. melanogaster* sequences before computing read counts, to account for several domains which are highly conserved between microbial and metazoan species. Read counts were computed for transposon sequences by computing the number of reads uniquely aligned to each transposon sequence in Repbase. Highly homologous sequences were grouped together for

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computing transposon read counts. Read counts were computed for known and novel gene models using HTSeq-count (Anders et al. 2015) with the 'intersection-nonempty' assignment method for exonic reads only. Tabulated read counts for each expression feature type (microbiome, transposon, endogenous genes) were then normalized across all samples using edgeR (Robinson et al. 2010) as follows. First, genes with low expression overall (<10 aligned reads in >75% of the libraries) were excluded from the analysis. Library sizes were re-computed as the sum of reads assigned to the remaining genes, and further normalized using the Trimmed Mean of M-values (TMM) method (Robinson and Oshlack 2010). At this point, we retained only genes (known or novel) whose expression in both biological replicates was above an empirical threshold in more than 200 line-sex combinations (400 samples total). This criterion retains genes expressed in only one sex. The threshold was determined by fitting all $\log_2$ transformed FPKM expression data points using a 2-component Gaussian mixture model and finding the expression value (FPKM = 0.280263) where the posterior probability of being in the lower expression component is 0.95. Genes on Chr U and Chr Uextra were also removed. We further adjusted transposon expression estimates to account for differences in transposon copy number across lines by fitting a linear model: RNA ~ DNA + ε, where RNA = the normalized $\log_2$ (RNA-seq read count); and DNA = normalized $\log_2$ (DNA read count) derived from the previously published DNA-seq data for each DGRP line (Huang et al. 2014). After fitting the linear model for each transposon sequence, ε estimates the relative transcription rate in each line independent of copy number, and was used as the adjusted transposon expression for all subsequent analysis. We further adjusted endogenous gene expression values by estimating and removing the effect of alignment bias resulting from higher rates of non-reference variants clustering in some lines. We computed the alignment bias score $A(g,L)$ defined as the number of non-reference nucleotides per kb in all exons of gene g in DGRP line L, based on the previous map of genomic variation in the DGRP (Huang et al. 2014). We then fit a linear model for each endogenous gene: $Y = A + \varepsilon$, where Y is the normalized expression profile for gene g

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after the read counting and edgeR normalization described above. After fitting these linear models, ε represents the alignment bias-corrected expression, and was used as the normalized gene expression in all subsequent analysis.Read mapping ambiguity could affect the confidence in defining NTRs. We assessed this by using blat (https://genome.ucsc.edu/FAQ/FAQblat.html) to map all RNA transcripts to the fly genome and identified all possible alignments. We used a metric (ΔBitscore) to characterize the mapping ambiguity of the full length RNA transcripts for known genes, NTRs filtering for low expression across the DGRP, and NTRs retained after filtering. ΔBitscore is defined as the difference between the bit score for the best alignment and the second best one. The greater the ΔBitscore, the less ambiguous is the alignment. In addition, we assessed whether the NTRs identified in the DGRP were present in an independent data set of 41 Drosophila cell lines that were either untreated or treated with the hormone ecdysone (Stoiber et al. 2016). We computed the median and maximum expression across all cell lines for each transcript using kallisto (Bray et al. 2016), an alignment-free abundance estimator, and calculated the median and maximum expression RNA transcripts for known genes, NTRs filtered for low expression across the DGRP, and NTRs retained after filtering.

Genetics of gene expression

For each class of expression features (endogenous genes, transposons, microbiome), we fit mixed-effect models to the gene expression data corresponding to: $Y = S + W + W{\times}S + L + L{\times}S + \varepsilon$, where Y is the observed $\log_2$ (normalized read count), S is sex, W is Wobachia infection status, $W{\times}S$ is Wolbachia by sex interaction, L is DGRP line, $L{\times}S$ is the line by sex interaction and ε is the residual error. We also performed reduced analyses ($Y = W + L + \varepsilon$) independently for males and females. We identified genetically variable transcripts as those that passed a 5%

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FDR threshold (based on Benjamini-Hochberg (1995) corrected *P*-values) for the *L* and/or *L*×*S* terms. We computed the broad sense heritabilities (H^2) for each gene expression trait separately for males and females as $H^2 = \sigma_L^2/(\sigma_L^2 + \sigma_\varepsilon^2)$, where σ_L^2 and σ_ε^2 are, respectively, the among line and within line variance components.

Clustering by genetic correlation

For each feature type (microbiome, transposons, endogenous genes), we clustered line means using the WGCNA R package v1.51 (Langfelder and Horvath 2008) as follows. Only genes with sufficient average expression ($\log_2$ FPKM > 0) and genetic variance (line mean variance > 0.05) were considered in these analyses. First, the Pearson correlation coefficient for every pair of line means, the soft-power threshold was computed using the pickSoftThreshold function, and used to convert the correlation matrix to an adjacency matrix with approximately scale-free connectivity. The adjacency matrix was then converted to a dissimilarity matrix based on the topological overlap map (Langfelder and Horvath 2008). Expression features were then clustered using hierarchical clustering (hclust function) based on the dissimilarity matrix, and split into distinct modules using the cutreeDynamic with deepSplit=4 and minClusterSize=20 (for endogenous gene expression, minClusterSize=4 was used for microbiome and transposon clustering). Module eigengenes were computed for each cluster, and highly similar clusters were combined using the mergeCloseModules function with cutHeight = 0.25. Expression features assigned to module 0 (insufficient similarity) were discarded. Modules consisting of >1,000 features were also discarded as insufficiently split into distinct modules. For each expression feature, the degree was computed as the overage topological overlap with all other features assigned to the same module. The average degree of each module was computed as the mean degree across all features in the module. Modules were sorted by average degree, such that module 1 has the highest average degree in each analysis.

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Gene set enrichment analyses

Lists of known gene IDs (FlyBase FBgn accessions) were uploaded to FlyMine (Lyne et al. 2007) or Panther (Mi et al. 2017) for functional enrichment. For analysis of gene lists from WGCNA clusters, the list of known genes input to WGCNA was used as the background set, to correct for any biases inherent to highly heritable expression patterns in general.

Expression QTL (eQTL) mapping

For each gene expression feature, we performed eQTL analysis as previously described (Huang et al. 2015). Briefly, we adjusted mean expression values in each sex for fixed effects of Wolbachia infection status, five major polymorphic inversions (*In2L(t)*, *In2R(NS)*, *In3R(P)*, *In3R(K)*, *In3R(Mo)*), and the first 10 principal components of the genetic relatedness matrix of all DGRP lines using a linear model. We mapped QTLs for the adjusted line means using PLINK (Purcell et al. 2007) against 1,932,427 SNPs with major allele frequency > 0.05 and missing genotypes in fewer than 25% of the 200 DGRP lines profiled by RNA-seq. Instead of controlling for experiment-wise type I error rate, which can be overly conservative, we controlled for the false discovery rate (FDR, Benjamani and Hochberg 1995). We computed FDR of eQTL calls by comparing observed eQTL *P*-value distributions to those obtained from running PLINK on 100 permutations of the observed line means for each expression feature. At any given *P*-value cut-off *X*, the estimated false positive rate of eQTLs for a specific gene expression feature is the average number of eQTLs with *P*-value < *X* across all permutations. The FDR at the same *P*-value is then computed as the estimated false positive rate divided by the number of eQTLs with *P*-value < *X* in the observed data. Using this formulation of FDR, we identified the unadjusted *P*-value cut-off corresponding to 5% FDR for each gene expression feature. No further model selection was performed; however, we classified eQTLs as being either *cis*-eQTLs (within 1kb of the gene body for the associated expression feature) or *trans*-eQTLs (> 1 kb of the gene body). To eliminate eQTLs whose genotypes are correlated with each other and cannot be genetically

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distinguished, we used forward model selection to iteratively add eQTLs to the model in the order of their conditional association (Huang et al. 2015). The model selection was stopped when none of the remaining putative eQTLs can enter the model with $P < 0.00001$. When two putative eQTLs had equal *P*-values, the one closer to the transcription start site was added.

Construction of eQTL networks

We then constructed regulatory eQTL networks based on individual SNPs which were called as both *cis*- and *trans*-eQTLs for multiple expression features. Specifically, we assign a directed edge $X \rightarrow Y$ if there is at least one variant that is both a *cis*-eQTL for gene *X* (defined as within 1 kb of gene *X*) and a *trans*-eQTL for gene *Y* at 5% FDR. We then broke all loops in the regulatory network for each sex by dropping the edge in each loop with the highest minimum *P*-value from all associated SNPs to create a directed, acyclic network.

Quantitative traits

We retrieved phenotypic data documented from previous publications on the same fly lines for male aggression (Shorter et al. 2015); chill coma recovery time and startle response (Mackay et al. 2012); food consumption (Garlapow et al. 2015); phototaxis (Carbone et al. 2016); sleep traits (Harbison et al. 2013) (day and night bout number, day and night total sleep duration, total waking activity); and starvation resistance (Huang et al. 2014).

To measure body weight and size, we collected 10 replicates of 10 flies per line and sex into pre-weighed 1.7 ml tubes, and weighed and flash froze them for downstream analyses. Virgin flies were used to avoid body weight variation due to variation in egg production. In addition we measured thorax length and thorax width as metrics for body size.

Frozen flies were homogenized in 250 µL Dulbecco's phosphate-buffered saline, and after gentle centrifugation supernatants were collected for measurements of free glucose, glycogen, free glycerol, triglyceride and total protein (further diluted 10 fold). For free glucose

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and glycogen, samples were denatured at 95°C for 25 minutes to prevent glycogenolysis. Measurements were performed following protocols provided by the Glycogen Colorimetric/Fluorometric Assay Kit (BioVision Inc.). For free glycerol and triglyceride, we used the Serum Triglyceride Determination Kit (Sigma-Aldrich Inc.), and incubated samples with the Triglyceride Reagent for 1 hour at 37°C. For total protein measurement, we used the Qubit Protein Assay Kit (Thermo Fisher Scientific Inc.).

Quantitative trait genetic parameters

We used mixed model, factorial ANOVAs ($Y = S + L + L\times S + Rep(L) + S\times Rep(L) + \varepsilon$, to partition variation of the quantitative traits between sexes (*S*), DGRP lines (*L*) and replicate vials within lines (*Rep*). Broad sense heritabilities were estimated as $H^2 = (\sigma_L^2 + \sigma_{SL}^2)/(\sigma_L^2 + \sigma_{SL}^2 + \sigma_\varepsilon^2)$, where σ_L^2, σ_{SL}^2 and σ_ε^2 are, respectively, the among line, sex by line and within line variance components.

eQTL-GWA enrichment analysis

We performed GWA analyses for all quantitative traits, separately for females and males. All phenotypes (line means) were first adjusted for the effect of Wolbachia infection and major polymorphic inversions using a linear model. The residuals (plus the intercept) from this analysis were then used as phenotype in a linear mixed model to test for the effect of each common variant individually, while adjusting for sample structure using a genomic relationship matrix (GRM), as implemented in GCTA-MLMA (Yang et al. 2011). The GRM was built as $\frac{WW'}{p}$ where ***W*** is a matrix of centered and scaled genotypes for the 200 lines and p is the total number of genetic variants. Similarly, we have also mapped trait QTLs using the same procedure as the eQTL mapping described above, by deriving empirical FDR based on 100 permutations of phenotypes.

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For each trait and sex, variants with $P < 10^{-5}$ were retained for downstream analysis. We then combined the lists of variants associated with each trait, separately for females and males, to obtain a single list of unique variants (*i.e.*, no duplicates) associated with any of the traits of interest. The enrichment analysis proceeded as described in Nicolae et al. (2010), within each sex. Briefly, GWAS hits were divided into minor allele frequency bins of width equal to 0.05. Then, an equal number of common variants (which may or may not have included actual GWAS hits) per bin were sampled at random and the overlap with eQTLs was calculated. This procedure was repeated 10,000 times and an empirical *P*-value for the enrichment was calculated as the number of replicates where the overlap between randomly sampled variants and eQTLs was greater than or equal to the observed overlap between GWAS hits and eQTLs over the total number of replicates.

Association of expression and quantitative traits

A transcriptome-wide association study (TWAS), *i.e.,* regressing the phenotype on each gene's expression level, was performed for each sex separately for each quantitative trait. We developed a method that accounts for structure present in the transcriptome due correlations between transcripts. This was achieved by fitting a linear mixed model of the type: $\mathbf{y} = \mathbf{1}\mu + \boldsymbol{w}\beta + \mathbf{t} + \mathbf{e}$, where **y** = n-vector of mean phenotypic values for n lines, μ = fixed population mean effect, **w** = n-vector of the tested gene's centered and scaled expression level, β = fixed effect of the gene, **t** = n-vector of random transcriptomic line effect (**t** ~$N(0, \boldsymbol{T}\sigma^2_t)$), and **e** = n-vector of random error (**e** ~$N(0, \boldsymbol{I}\sigma^2_e)$).

The key term in the model that accounts for sample structure is **T**, the transcriptomic relationship matrix (TRM). The TRM was computed as $\frac{W^{-}W^{-\prime}}{p}$, where $\boldsymbol{W}^{-}$ is a matrix of centered and scaled gene expression levels for the 200 lines, excluding the gene tested to maximize the

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power to find an association (Yang et al. 2014), and *p* is the total number of genes. The TRM in TWAS has similar role to the GRM in GWAS.

The effect of each gene's expression level on the phenotype was tested using a Wald test of the form $\frac{\beta^2}{(SE(\beta))^2} \sim \chi_1^2$. Raw *P*-values and Benjamini-Hochberg (1995) FDR-corrected *P*-values were computed.

The phenotypes were adjusted for the effects of Wolbachia and major polymorphic inversions as described in the previous section. Because the phenotypes were adjusted, we did not adjust gene expression in this analysis to avoid spurious associations due to adjustment on both sides of the equation.

We also performed similar associations of quantitative traits with TEs and microbial gene expression, using the same models as for TWAS but substituting TE and microbial expression for gene expression levels. Quantitative trait phenotypes were adjusted for the effects of Wolbachia and major polymorphic inversions but the TE and microbial expression data were not. The TE analysis was performed for males and females separately, while sex-pooled microbe expression data was used with female or male quantitative trait phenotypes since microbial gene expression was not sex-specific.

DATA ACCESS

All raw and processed sequencing data generated in this study have been submitted to the NCBI Gene Expression Omnibus (GEO; https://www.ncbi.nlm.nih.gov/geo/) under accession number GSE117850. The DGRP lines are available from the Bloomington Drosophila Stock Center (Bloomington, IN). All analysis codes are available in Supplemental Codes and on GitHub (https://github.com/qgg-lab/dgrp-rna-seq/).

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AUTHOR CONTRIBUTIONS

L. E., W. H. S. Z., F. M., R. R. H. A. and T. F. C. M. wrote the manuscript. L. E., W. H. S. Z., F. M. and T. F. C. M. analyzed the data. M. A. C., R. L., G. A., M. S. G., J. M., G. S. A. and L. T performed the research.

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FIGURE LEGENDS

Figure 1. Genetic variation of gene expression in the DGRP. (**A**) Sexual dimorphism of gene expression. Red (blue) indicates significant up-regulation in females (males). (**B**) Distribution of H^2 estimates for annotated genes and NTRs in females. (**C**) Distribution of H^2 estimates for annotated genes and NTRs in males. (**D**) WGCNA modules for annotated genes and NTRs in females. (**E**) WGCNA modules for annotated genes and NTRs in males. Heatmaps show the pairwise correlation of all genes in each module, sorted by average connectivity, with the most tightly connected module at the top left.

Figure 2. Genomic location of eQTLs for gene expression and genes they regulate. eQTL chromosome positions (bp) are given on the *X*-axis, and the genes with which they are associated on the *Y*-axis. Red points denote female-specific eQTLs, blue indicates male-specific eQTLs, and black shows eQTLs shared by males and females.

Figure 3. Large *cis-trans* eQTL genetic network in females and males. Node interior colors indicate genomic location of genes (yellow: euchromatic regions with normal recombination; gray: euchromatic regions with reduced recombination; blue: heterochomatin). Node border colors denote annotated gene (gray) or NTR (red). Node shape indicates whether a gene is a regulator and/or target (triangles: regulator only; squares: target only; circles: both regulator and target). The node size indicates the number of node connections. Arrows on the edges point to the target. Edges are color coded to show female-specific regulation (red), male-specific regulation (blue) and regulation common to both sexes (black).

Figure 4. Genetic variation of TE expression in the DGRP. (**A**) Total signal for each TE family, summed over all individual transposon sequences and averaged across all DGRP lines,

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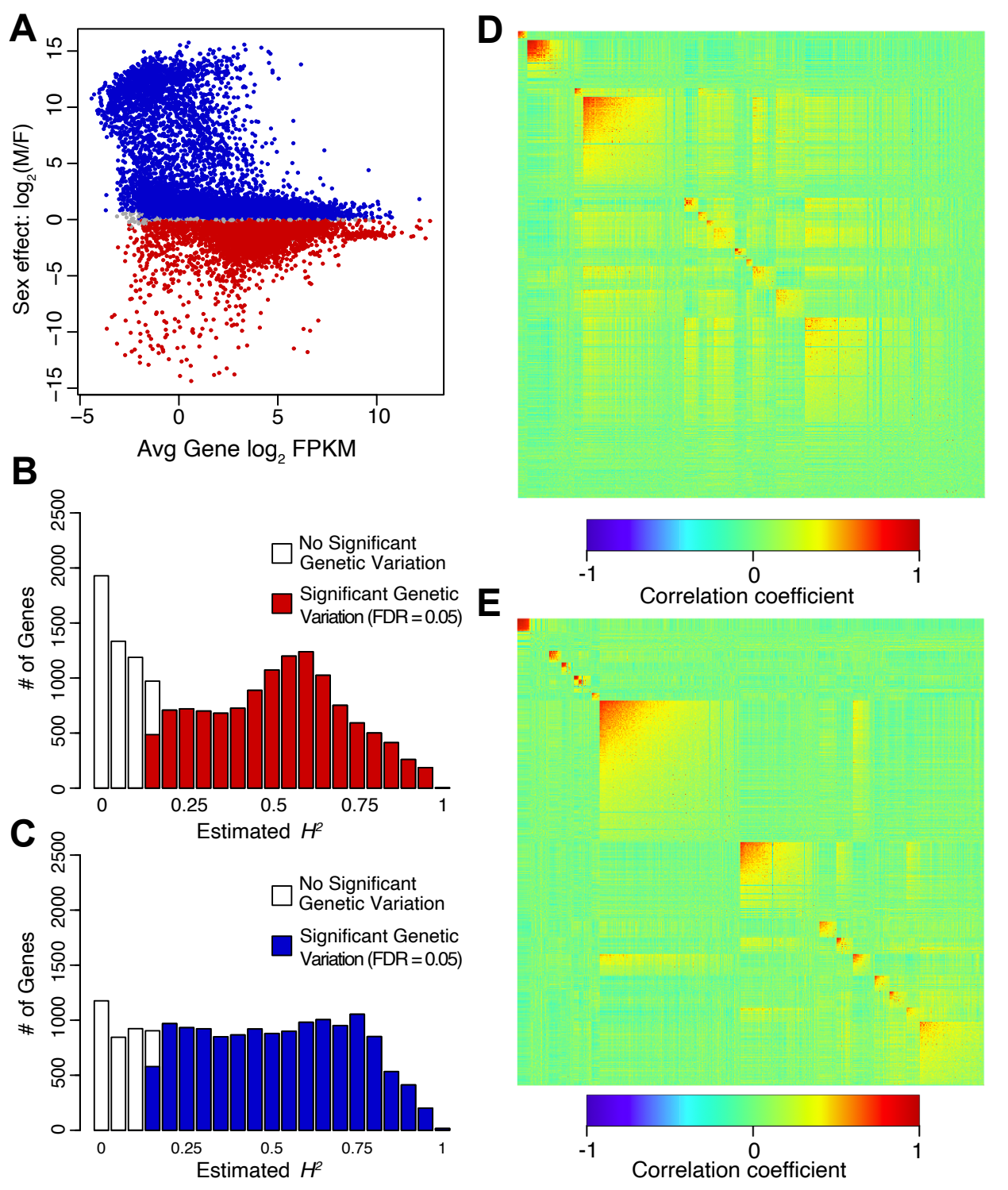
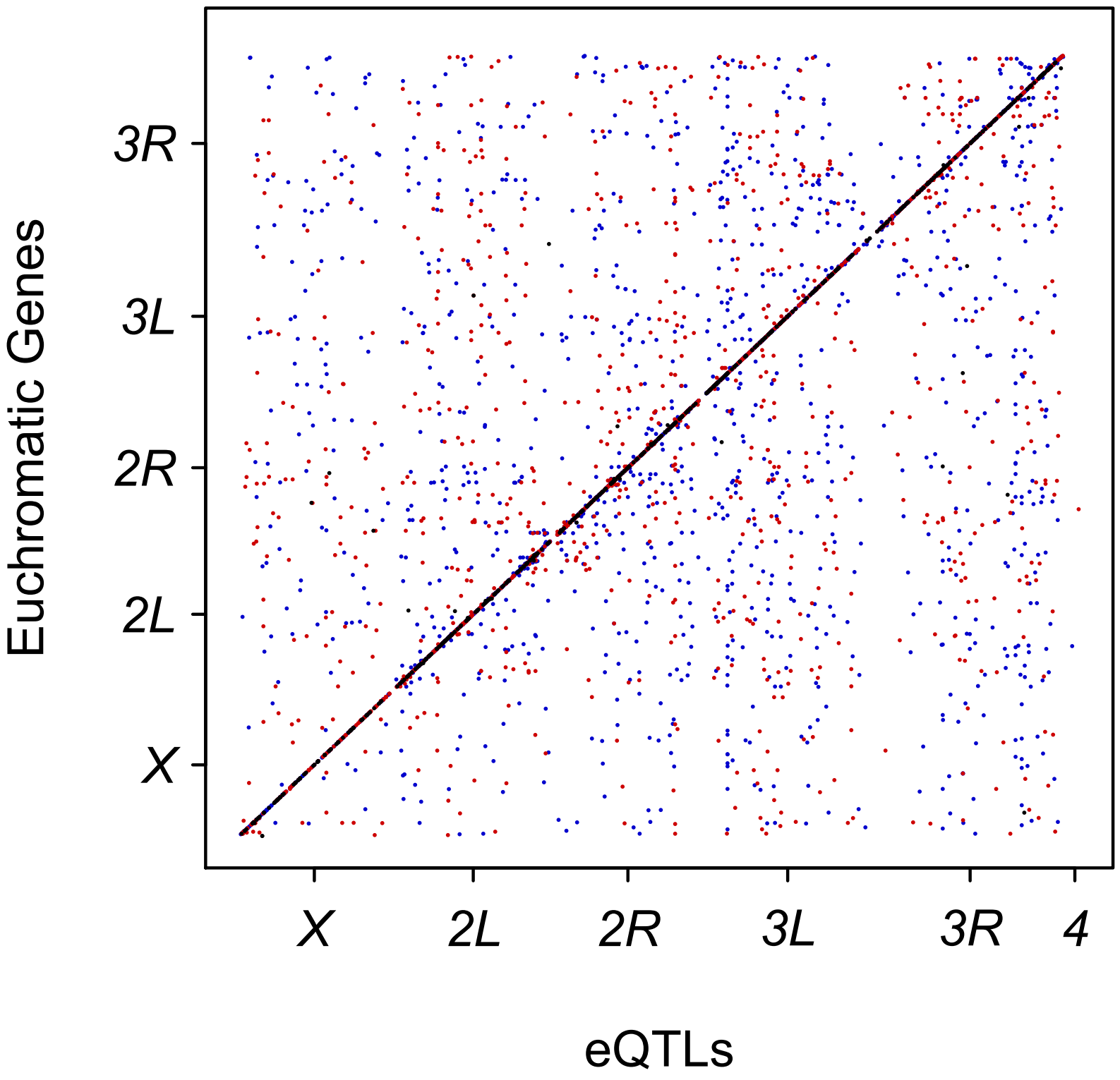
sex, and replicates. (**B**) Distribution of copy number independent H^2 estimates for TE sequences in females. (**C**) Distribution of copy number independent H^2 estimates for TE sequences in males. (**D**) WGCNA modules of TEs for females. (**E**) WGCNA modules of TEs for males. Heatmaps are depicted as in Figure 1. TE sequences not assigned to any module are included at the bottom right.

Figure 5. TE genetic regulatory network. Symbols and color-coding are as for Figure 3. Black squares denote TE sequences.

Figure 6. Genetic variation of microbiome composition. (**A**) The proportion of microbiome signal in RNA-seq libraries aligned to species in each genus or viral group. (**B**) Line means of total microbial signal (excluding Wolbachia). (**C**) Distribution of H^2 estimates for individual microbe species. (**D**) WGCNA modules for microbial species. Heatmaps are depicted as in Figure 1. Species not assigned to any module are included at the bottom right.


A
Sex effect: log2(M/F)
15
10
5
0
−5
−10
−15
−5
0
5
10
Avg Gene log2 FPKM
B
# of Genes
2500
2000
1500
1000
500
0
No Significant Genetic Variation
Significant Genetic Variation (FDR = 0.05)
0
0.25
0.5
0.75
1
Estimated H2
C
# of Genes
2500
2000
1500
1000
500
0
No Significant Genetic Variation
Significant Genetic Variation (FDR = 0.05)
0
0.25
0.5
0.75
1
Estimated H2
D
-1
0
1
Correlation coefficient
E
-1
0
1
Correlation coefficient


Euchromatic Genes
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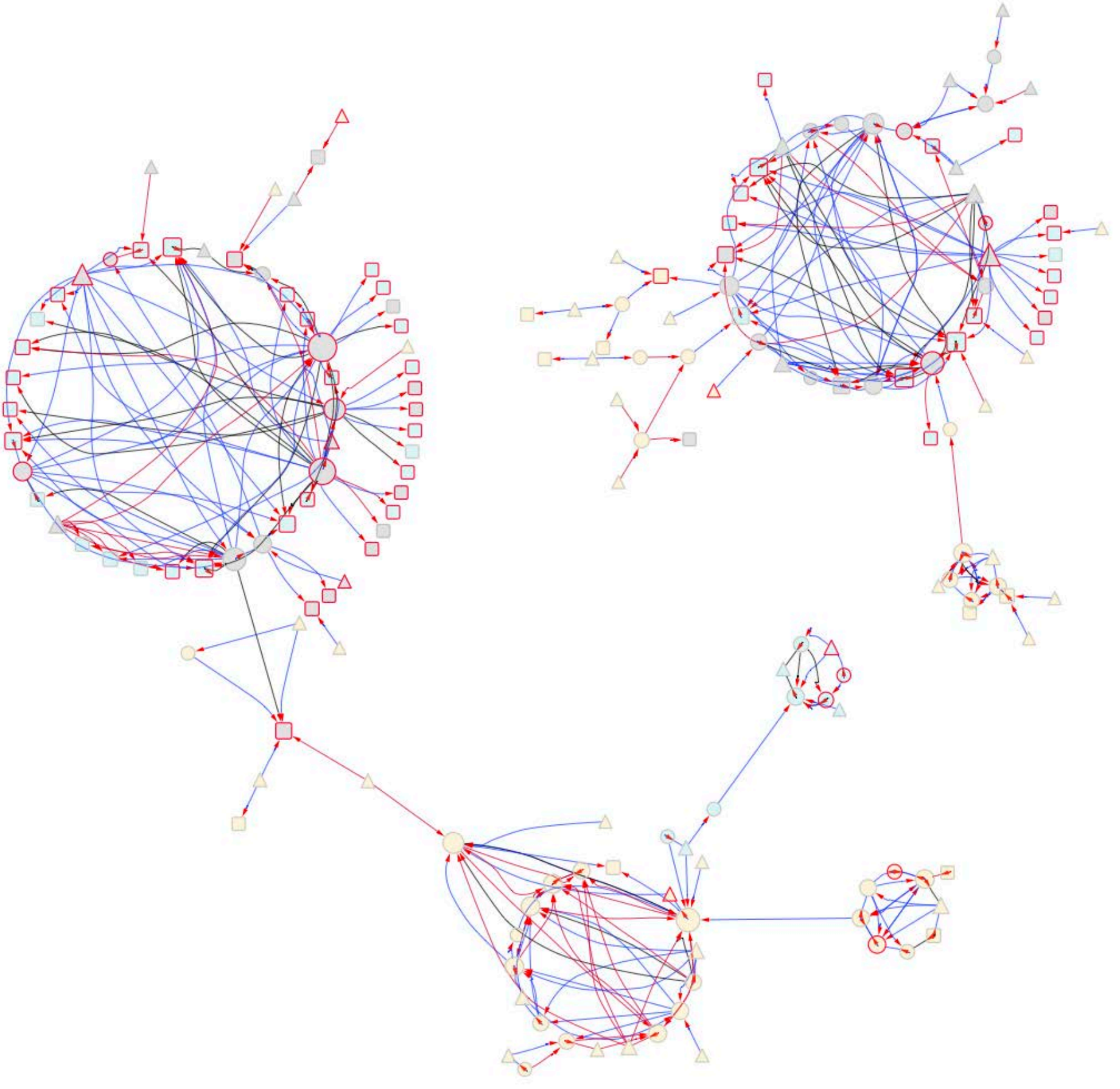
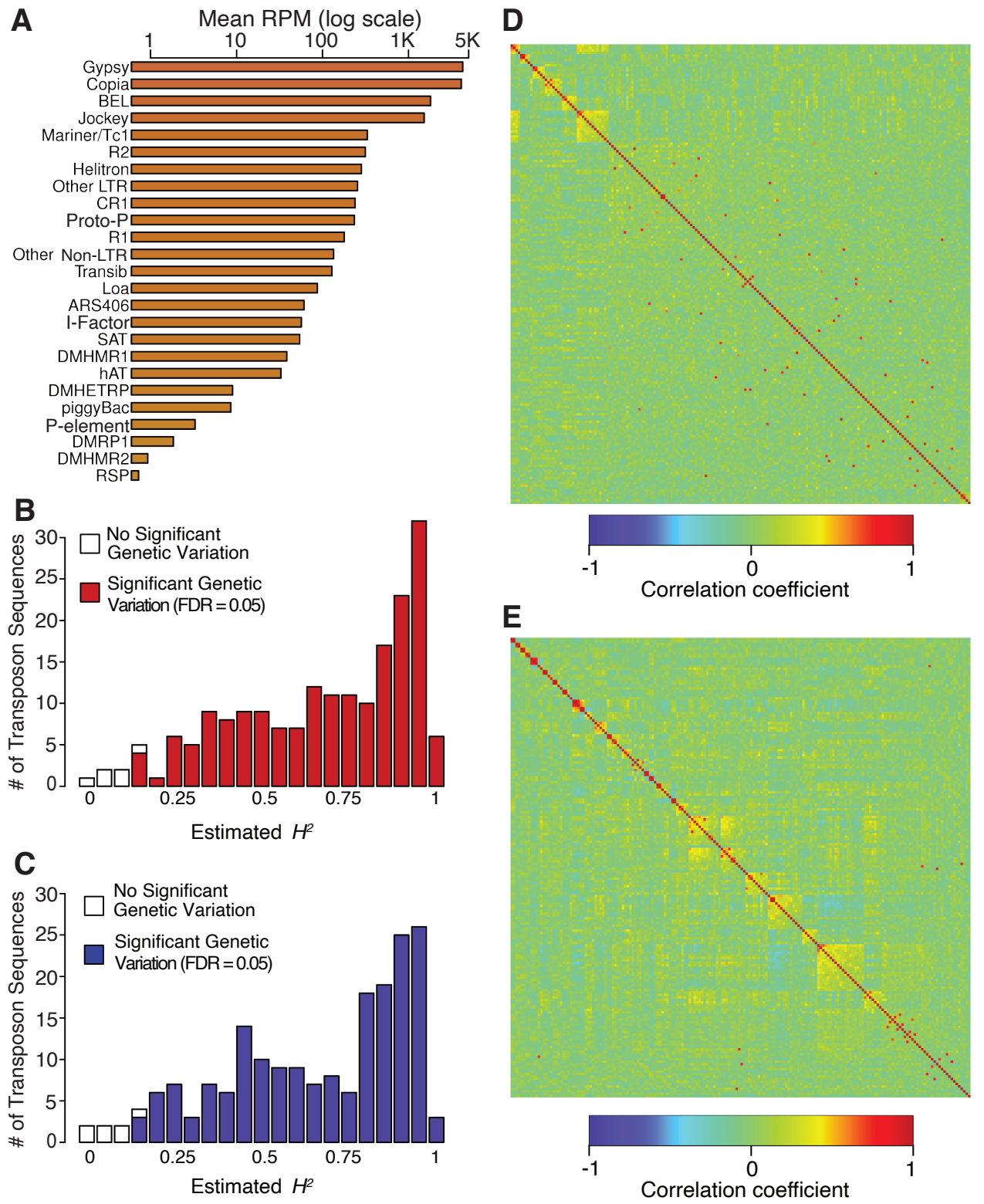
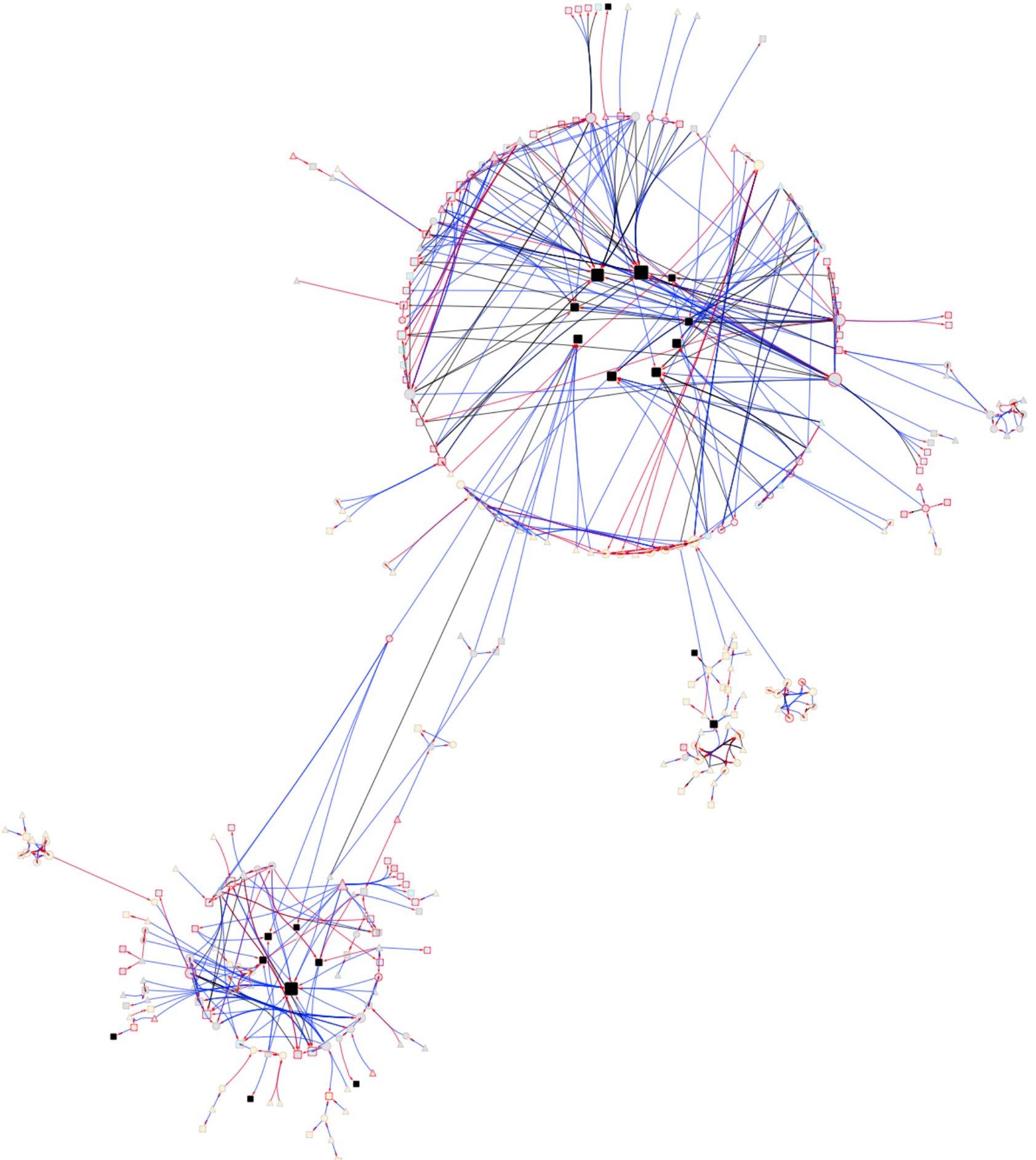
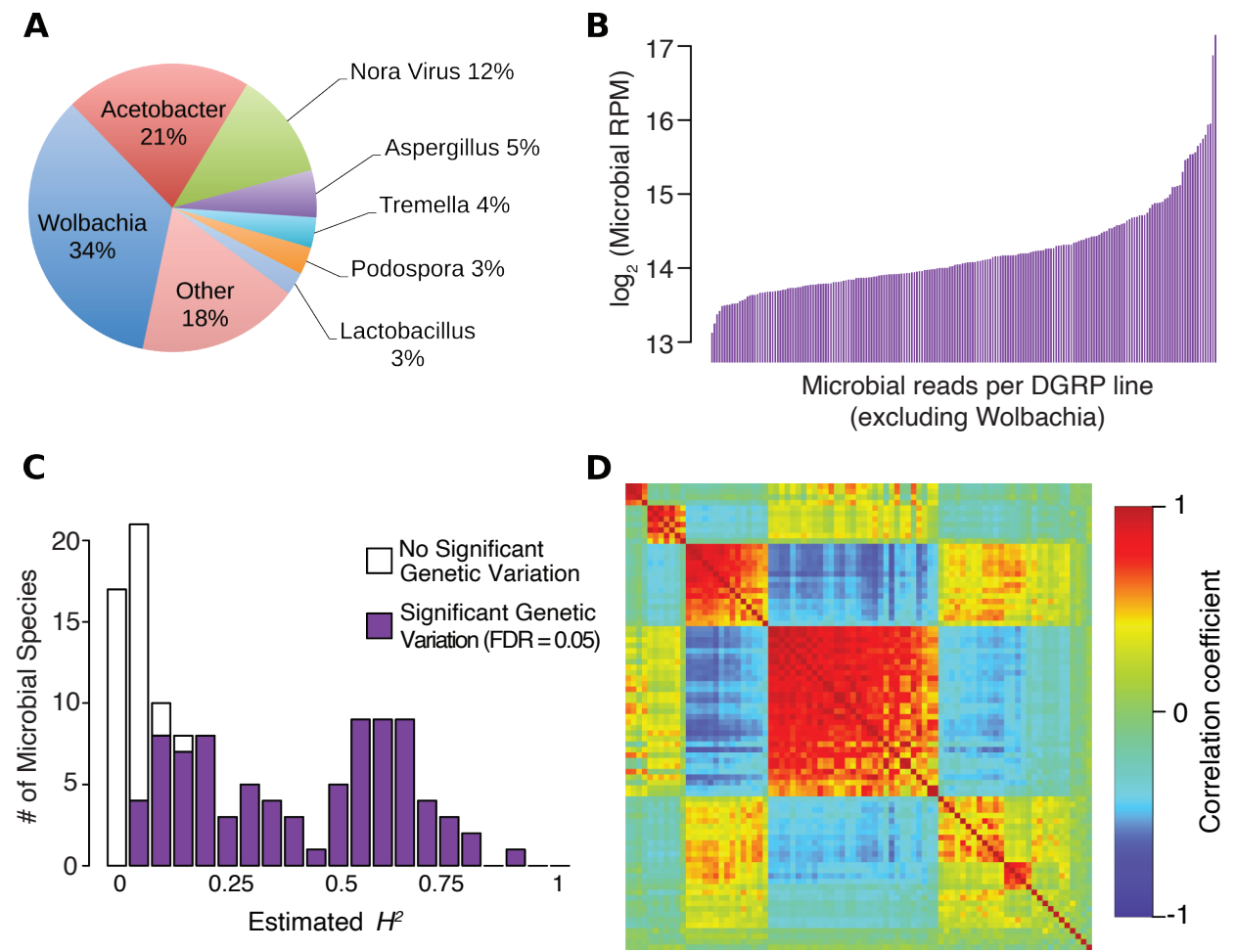
3L
2R
2L
X
X
2L
2R
3L
3R
4
eQTLs




A
Mean RPM (log scale)
1
10
100
1K
5K
Gypsy
Copia
BEL
Jockey
Mariner/Tc1
R2
Helitron
Other LTR
CR1
Proto-P
R1
Other Non-LTR
Transib
Loa
ARS406
I-Factor
SAT
DMHMR1
hAT
DMHETRP
piggyBac
P-element
DMRP1
DMHMR2
RSP
B
No Significant Genetic Variation
Significant Genetic Variation (FDR = 0.05)
# of Transposon Sequences
Estimated H^2
C
No Significant Genetic Variation
Significant Genetic Variation (FDR = 0.05)
# of Transposon Sequences
Estimated H^2
D
-1
0
1
Correlation coefficient
E
-1
0
1
Correlation coefficient




A
Nora Virus 12%
Aspergillus 5%
Tremella 4%
Podospora 3%
Lactobacillus 3%
Other 18%
Wolbachia 34%
Acetobacter 21%
B
log2 (Microbial RPM)
17
16
15
14
13
Microbial reads per DGRP line (excluding Wolbachia)
C
No Significant Genetic Variation
Significant Genetic Variation (FDR = 0.05)
# of Microbial Species
20
15
10
5
0
0
0.25
0.5
0.75
1
Estimated H²
D
1
0
-1
Correlation coefficient