Evolution of phenotypic plasticity in response to ethanol between sister species with different ecological histories (*Drosophila melanogaster* and *D. simulans*)

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Running title: Evolution of plasticity to ethanol

D. melanogaster sequence data have been submitted to GenBank: accession number PRJNA482662. *D. simulans* sequence data will be made available upon acceptance.

Abstract

The role of phenotypic plasticity in evolution is contentious, in part because different types of plasticity – adaptive, neutral, or non-adaptive, are often not distinguished. Adaptive plasticity is expected to facilitate expansion into new environments, while non-adaptive plasticity will result in a mean phenotype further from the adaptive optimum and/or an increase in variance due to the expression of variation that was neutral and shielded from selection in the prior environment. We explore these patterns here by exposing *Drosophila melanogaster* and *D. simulans* to high ethanol concentrations, with the knowledge that *D. melanogaster* associates with high concentrations of ethanol in nature while *D. simulans* does not. Using changes in gene expression and splicing we find that in *D. simulans* there is a large genotype-specific response to ethanol, orders of magnitude larger than any effect that is not genotype-specific. In *D. melanogaster* the response to ethanol is limited, and it is concordant among different genotypes. This response to ethanol in *D. simulans* is enriched for non-protein coding nested genes that do not have orthologs in *D. melanogaster*, suggesting rapid evolution of transcription. Sequence variation in ethanol-implicated genes is consistent with balancing selection or selection relaxation in *D. simulans*, and they are more divergent from *D. melanogaster* than the genome average. Overall, these patterns are consistent with a maladaptive passive response to ethanol-induced stress in *D. simulans*, while in *D. melanogaster* the reduced response and lack of genotype-specific variation suggests selection for an optimal response.

Keywords: phenotypic plasticity, *Drosophila*, ethanol, alternative splicing

Introduction

Genetic variation (G) occurs when there are differences in a phenotype due to genotypic differences among individuals within a population (Figure 1A). In contrast, plasticity – i.e. environment specific (E) adjustment in phenotype - can occur without genetic variation between individuals and is a change in a given phenotype in response to an external stimulus (Figure 1A). When components of variance are estimated plasticity is the part of phenotypic variation attributable to environmental variation. Plasticity is a ubiquitous property of organisms, and it may be adaptive, neutral, or maladaptive with regard to the fitness of a plastic individual (Ghalambor *et al*. 2007; Lande 2009; Marais *et al*. 2013). Genetic variation in plasticity (G x E) describes variation between genotypes for the response to the environment (Figure 1A). Quantifying genetic variation, plasticity, and genetic variation for plasticity is important for understanding the genotype to phenotype map and adaptation. For example, the evolution of a reduced plastic response (E) can reconfigure the relationship between fitness and the environment, redrawing the genotype to phenotype map. In another example, genetic variation in plasticity can make the evolution of locally adapted populations unpredictable in response to environmental change.

It is often assumed that plasticity is adaptive, however phenotypic plasticity can be adaptive, neutral, or maladaptive. This depends on the evolutionary history of exposure to the agent causing plasticity. Were such exposure infrequent, plasticity could arise as a passive consequence of environmental stress. This is likely an important, or perhaps prevalent, form of phenotypic plasticity (Ghalambor *et al*. 2007; 2015). One potential prediction arising from stress inducing passive phenotypic plasticity is that within population variation for performance is expected to be greater, as stress breaks down buffering mechanisms that might normally reduce between individual variation (Joshi & Thompson 1997). Thus under lower stress conditions, populations will exhibit less individual and genetic variation compared to high stress, rare conditions (Ghalambor *et al*. 2007). The novel or rare high stress environment is essentially uncovering cryptic variation that was allowed to neutrally accumulate in lower stress environments where selection was unable to shape the reaction norm (Rutherford 2000).

When a population colonizes a new habitat, this could manifest as a transient increase in G x E, followed by a reduction in genetic variation for plasticity, assuming that there is selection for a single optimal phenotype in response to the environment (Via & Lande 1985; Guntrip & Sibly 1998; Lande 2009; Matzkin 2012; Huang & Agrawal 2016). The maintenance of plasticity, or loss of plasticity, would then depend upon the frequency with which the population continues to experience multiple environments, or the cost of maintaining the plastic response (Joshi & Thompson 1997; Lande 2009; Scoville & Pfrender 2010; Lee *et al*. 2011; Hodgins-Davis *et al*. 2012). Thus if phenotypic plasticity is costly to maintain, or the organism no longer experiences the previous environment, the expectation is that phenotypic plasticity will transition to local adaptation.

In this manuscript we will examine plasticity and genetic variation for plasticity in response to ethanol in *D. simulans* and *D. melanogaster*, two species with divergence adaptive histories with respect to ethanol. *D. melanogaster* and *D. simulans* are thought to have originated in or around Africa (*D. melanogaster*, southern-central Africa, *D. simulans*, Madagascar) followed by an out-of-Africa expansion ~10,000 years ago, however *D. melanogaster* may have colonized Europe and America earlier than *D. simulans* (David and Capy 1988; Lachaise et al. 1988; Baudry et al. 2004; Li and Stephan 2006; Thornton and Andolfatto 2006). *Drosophila simulans* is thought to have colonized the Americas in the last 100 years, creating a large and more recent population bottleneck than in *D. melanogaster* (Sturtevant 1920; Signor *et al.* 2017c).

D. melanogaster and *D. simulans* are cosmopolitan species and human commensals, however while *D. melanogaster* is commonly found inside houses, breweries, and wineries, *D. simulans* is more often observed in orchards and parks (though their niches overlap significantly and they are often found on the same patches) (David & Bocquet 1976; Parsons & King 1977; McKenzie & McKechnie 1979; Parsons & Spence 1981; Dickinson *et al.* 1984; Gibson & Wilks 1988; Thomson *et al.* 1991; Chakir *et al.* 1993; Mercot *et al.* 1994; Joshi & Thompson 1997). *D. melanogaster* is considerably more ethanol tolerant than *D. simulans*, and is found regularly feeding upon and ovipositing in resources with ethanol

concentrations greater than 8% (Hoffmann & McKechnie 1991; Fry 2014; Zhu & Fry 2015). At concentrations of 4% ethanol *D. simulans* shows reductions in survivorship and increased development time relative to *D. melanogaster* (Joshi & Thompson 1997). However, at low concentrations of ethanol (0.5 - 3.0%) adult *D. simulans* show increases in longevity (less so than adult *D. melanogaster*, which increases longevity from 0.5 - 9.0%) (Parsons *et al.* 1979). During multi-generation exposure to ethanol-rich substrates in the lab, *D. simulans* evolves tolerance (Joshi & Thompson 1997; Lefèvre *et al.* 2012). It has been demonstrated that ethanol tolerance comes as a trade-off that reduces the efficacy of processing other biochemical targets, thus it is possible that in *D. simulans* this trade-off is under different selective pressure (Chakraborty & Fry 2016). However, ethanol is by no means a novel resource for *D. simulans*.

In general genetic adaptation and phenotypic plasticity are considered as alternative adaptive strategies in response to the environment (Schlichting & Pigliucci, 1998). For a variety of traits *D. simulans* has been observed to be 'plastic' while *D. melanogaster* is described as being locally adapted (David & Bocquet 1975; Hyytia *et al.* 1985; Watada *et al.* 1986; Singh 1989) . For example, for temperature tolerance *D. simulans* has been shown to exhibit less difference between populations and to produce 'optimal' phenotypes under a wider range of environments (consistent with adaptive plasticity) (Austin & Moehring 2013). This includes less differentiation between populations, in that populations from different temperatures show similar levels of plasticity across temperatures. Population genetic surveys generally corroborate these phenotypic observations, where *D. melanogaster* also shows greater differences between populations, and more evidence of clinal variation than in *D. simulans* (Singh 1989; Weeks *et al.* 2002; Hoffmann & Weeks 2006; Schmidt & Paaby 2008; Machado *et al.* 2015; Sedghifar *et al.* 2016; Bergland *et al.* 2016; Signor *et al.* 2017c). However, *D. simulans* has generally been observed to harbor more within population diversity than *D. melanogaster* (Machado *et al.* 2015; Sedghifar *et al.* 2016; Signor *et al.* 2017c).

Gene expression is both affected by the environment and mediates an organisms response to the environment. It has been shown to be plastic in response to many environmental variables such as heat stress, ethanol, and the presence of predators (Hodgins-Davis & Townsend 2009; Levine *et al*. 2011; Yampolsky *et al*. 2012; Hodgins-Davis *et al*. 2012). While many studies have quantified the effect of the environment on gene expression, far fewer have performed these experiments in multiple genotypes within a common garden to obtain the interaction between genotype and the environment for gene expression (DeBiasse & Kelly 2015). Indeed, most work on gene expression plasticity has not quantified G x E, and has found inconsistent results with regard to theoretical expectations for the maintenance of plasticity (Cheviron *et al*. 2008; McCairns & Bernatchez 2009; Hodgins-Davis *et al*. 2012; DeBiasse & Kelly 2015; Heckel *et al*. 2016; Mathur & Schmidt 2016). Furthermore, many plastic responses are time dependent, and if performed in a common garden without sampling across time points may not detect early plastic responses that usher in 'maintenance' phenotypes, or dynamic responses to the environment (Aubin-Horth & Renn 2009; Lewis *et al*. 2014). In addition, alternative splicing has been shown to respond to changes in the environment, and it is thought to diverge more rapidly than gene expression between lineages (Barbosa-Morais *et al*. 2012; Merkin *et al*. 2012; Gueroussov & Gonatopoulos-Pournatzis 2015; Jakšić & Schlötterer 2016; Wang *et al*. 2017; Pajoro *et al*. 2017; Singh *et al*. 2017; Calixto *et al*. 2018). As such it may be an important component of plasticity, necessary for adaptation to rapidly changing and heterogeneous environments (Marais *et al*. 2013; Preußner *et al*. 2017; Price *et al*. 2018; Calixto *et al*. 2018). Furthermore, it has been previously implicated in the response to ethanol, and in *D. melanogaster* may be a more important component of the response than gene expression changes (Oomizu *et al*. 2003; Newton *et al*. 2004; Pietrzykowski *et al*. 2008; Sasabe & Ishiura 2010; Hemby 2012; Zaharieva *et al*. 2012; Robinson & Atkinson 2013; Signor & Nuzhdin 2018). However, alternative splicing is rarely accounted for in studies of plasticity.

Overall, we compare plasticity in response to ethanol, including genetic variation for plasticity over time and between genotypes, among *Drosophila* species with different patterns of ethanol utilization. We measure plasticity and variation in expression and splicing in response to ethanol treatment, and we observe that in *D. melanogaster*, which regularly exploits ethanol-rich substrates, there is essentially no genetic variation for plasticity. This suggests the following scenario: for *D. simulans* high ethanol

concentrations are essentially a novel stressful environment that reveals cryptic variation for phenotypic plasticity, while *D. melanogaster* does not experience high stress under 15% ethanol and cryptic variation is not revealed. Furthermore, *D. melanogaster* does not experience high stress as it is adapted to ethanol and genetic variation has been removed in favor of an optimal, and reduced, plastic response

Methods

Fly lines

Male flies for *D. melanogaster* originated from six genotypes collected from an orchard in Winters, California in 1998 and made isogenic by 40 generations of full sibling inbreeding (Yang & Nuzhdin 2003; Campo *et al*. 2013; Signor *et al*. 2017a; b). Male flies for *D. simulans* came from six genotypes collected from the Zuma Organic Orchard in the winter of 2012 and made isogenic by 15 generations of full sib inbreeding (Signor *et al*. 2017b; c). Residual heterozygosity is similar between these lines of *D. melanogaster* and *D. simulans* (Campo *et al*. 2013; Signor *et al*. 2017c). In natural conditions flies will not by homozygous at all loci, thus each inbred genotype was crossed to a white eyed 'tester' strain to create the F1 flies used in the ethanol exposure assays (*D. melanogaster*, w^{1118}, Bloomington stock number 3605; *D. simulans*, w^{501}, Cornell species stock number 14021-0251.011). This design allows us to replicate observations of gene expression because the flies are identical twins, while also maintaining heterozygosity such that they more closely resemble wild type flies. Rearing occurred on a standard medium at 25 °C with a 12-h light/12-h dark cycle. Several measures were taken to standardize offspring quality: F1 flies were produced from females of the same age, these females were held at the same density (10 individuals per sex/vial), males used for ethanol exposure assays were collected as virgins and reared in single sex vials, and males used for assays were held at a standard density (24-30 per vial).

During the ethanol exposure assay a female was included as a stimulus, but was not collected for RNA-seq (Signor *et al*. 2017a; b). The females were from y^1w^1 mutants for both *D. simulans* and *D. melanogaster* (*D. melanogaster*, Bloomington stock number 1495, *D. simulans*, Cornell species stock number 14021-0251.013). Quality of the females was controlled in the same manner as previously described for males, and both males and females were aged three to four days before being used in the ethanol exposure assay.

Experiment setup

Ethanol exposure took place in a circular arena, each of which was part of a larger chamber containing 12 arenas each with a diameter of 2.54 cm (VWR cat. no. 89093-496). Prior to the assay the flies were sedated through exposure at 4 °C for ten minutes, to avoid the confounding effects of CO_2 exposure on behavior and gene expression. Once the flies were sedated they were placed in the arenas with a paintbrush (two males and one female per arena) and the chambers were secured with two small pieces of cryogenic tape. It is standard to allow organisms to recover in a new location for ten minutes prior to beginning behavioral assays, which was a portion of the goal of this study, so the chambers were left upside down for ten minutes while the flies regained consciousness and oriented themselves (Signor *et al*. 2017a). The bottom of each chamber contained a standard amount of either grapefruit medium or medium in which 15% of the water has been replaced with ethanol. During ethanol administration the flies were recorded using VideoGrabber (http://code.google.com/p/video-grabber/), and set-up of the assays was facilitated with FlyCapture (PointGrey, Canada). In order to better standardize the transcriptional response, these videos were also used to determine if any male/female pairs mated during the assay or any flies were damaged during setup, and flies in those chambers were not collected for RNA-seq.

Assays were conducted for ten, 20, or 30 minutes for three replicates of each of the two conditions. Flies are most active during the hours following dawn, thus to standardize behavior and circadian rhythms all assays were conducted within a two-hour window after dawn. Replicates for both species were conducted randomly under standardized conditions (25 °C, 70% humidity). At the conclusion of the assay

205 the chambers were flash frozen in liquid nitrogen, allowed to freeze through, transferred to dry ice, and all
206 of the males were collected for RNA-seq. For both *D. melanogaster* and *D. simulans* the expectation is
207 that intoxication is occurring through inhalation of ethanol vapors, and evidence of the behavioral effects
208 in both species and the efficacy of this approach have been published previously (Signor *et al*. 2017a; b).

209 *Sample preparation and RNA sequencing*

210 Sample preparation has been described previously, and will be briefly summarized here (Signor & Nuzhdin
211 2018). Flash-frozen flies were freeze dried and ten to 12 heads were placed into a 96-tube plate (Axygen
212 MTS-11-C-R). mRNA purification, cDNA synthesis and library preparation were carried out by RAPiD
213 GENOMICS (http://rapid-genomics.com) using a robot. mRNA was purified using Dynabeads mRNA
214 DIRECT Micro kit (Invitrogen # 61021) with slight modifications. To fragment the RNA mRNA-beads
215 were resuspended in 10 uL 2X first strand buffer (Invitrogen # 18064-014), incubated at 80 C for two
216 minutes and placed on ice, then the supernatant was collected after five minutes on magnetic stand. First
217 strand synthesis was performed using standard protocols for Superscript II (Invitrogen #18640-014) and
218 reverse transcription (25 C 10 min, 42 C for 50 min, 70 C for 15 min, 10 C hold). Second strand synthesis
219 was carried out using standard protocols with DNA Pol I and incubated at 16 C for 2.5 hours. cDNA was
220 purified with 1.8 volume of AMPure XP following manufactures instructions (Beckman Coulter A63880).
221 Illumina RNAseq libraries were prepared by Rapid Genomics (http://rapid-genomics.com/home/) using
222 dual barcodes. Sequencing was performed using the Illumina HiSeq 2500 as both 2×150 bp or 2×50 bp
223 reads. The two run lengths (and runs) were intended to provide extra coverage, and all replicates were
224 sequenced in both runs.

225 *Gene expression analysis*

226 It is common in organisms with alternative splicing for exons from different isoforms of a single gene to
227 overlap with one another, or be shared between all or most isoforms (Figure 1B). Short read data
228 fundamentally cannot resolve these exons to individual isoforms, however, one approach is to quantify each
229 exon separately and decompose exoms overlapping between isoforms into those which are shared and
230 unique. When the differences between overlapping exons are less than 10 bp, there is no appreciable
231 amount of information loss in not decomposing overlapping exons, and this approach has been taken in the
232 past (Dalton *et al*. 2013; Graze *et al*. 2014; Newell *et al*. 2016; Fear *et al*. 2016). However, in many cases
233 the differences in exon overlap are much larger than this, so to address this issue we use a classification
234 scheme where reads may be assigned to exons, exonic regions, or exon fragments (Signor & Nuzhdin 2018)
235 (Figure 1B). Exon boundaries were determined using the *D. melanogaster* FlyBase 6.17 genome features
236 file and the *D. simulans* 2.02 genome features file. A single exon is one that does not overlap any other
237 exons, and it may be unique to a single isoform, shared between several (common), or shared between all
238 isoforms (constitutive). If a gene has only a single transcript then every exons it contains will be both unique
239 and constitutive. When exons overlap other exons from different isoforms, they are grouped together into
240 an exonic region, and they may be common to some isoforms or constitutive to all, but they are not unique
241 given that by definition they require overlap between exons. When overlapping exons differ measurably,
242 we used the 5' and 3' positions of the exons within the region to create exon fragments. An exon fragment
243 may exclusive to a single isoform (unique) or common/constitutive (Figure 1B). For exon fragments there
244 is only one unique situation in which it may be both constitutive and unique – when the exons of two genes
245 overlap, and one of those genes has a single transcript, such that the non-overlapping portion of the exon
246 belonging to the gene with one transcript will become a unique and constitutive exon fragment. Alignment
247 was performed using BWA-MEM version 0.7.15 and BED files were used to count reads in each region
248 and obtain the length adjusted read count (reads in region divided by the length of region), and the APN
249 (average per nucleotide) (Li 2015).

250 The APN was summed for technical replicates of the same read length then averaged between read
251 lengths to handle the mixture of read lengths for each sample (2×150 bp and 2×50 bp). If the APN was
252 greater than zero in at least half of all samples per condition it was considered detected. While we

considered several approaches to normalize coverage counts upper-quartile normalization with log-transformation and median centering within time × treatment × genotype were selected due to better performance of the residuals (Bullard *et al*. 2010; Dillies *et al*. 2013).

To test the significance of components of expression variation, the log APN for each exonic region was modeled as

$$\Upsilon_{ijkl} = \mu + g_i + t_j + (gt)_{ik} + m_k + (gm)_{ik} + (tm)_{ijk} + \epsilon_{ijkl}$$

for the ith genotype (g_i), j^{th} treatment (t_j; j = ethanol or no ethanol), k^{th} time point (m_k; k = 10 min, 20 min, 30 min), and l^{th} replicate. For the interaction between treatment and time point, the log APN for each exonic region was modeled as

$$\Upsilon_{ij} = \mu + t_j + \epsilon_{ij}$$

or the ith condition (time × treatment) and j^{th} replicate. Contrasts to compare treatments within time point (ethanol versus no ethanol, for 10 min, 20 min and 30 min) were conducted. Residuals were evaluated for conformation with normality assumptions, and assumptions were met in excess of 95% of the models.

To evaluate whether there was evidence for splicing differences among times or treatments, exonic regions for each gene and for each sample were ranked and the most expressed region ranked as one, the least expressed exonic region as three and all others as two. Exon ranks for each gene were modeled as

$$\Upsilon_{ijk} = \mu + r_i + t_j + (rt)_{ij} + \epsilon_{ijkl}$$

where Υ_{ijk} is the exon rank (1,2,3) of the ith exonic region of the gene, j^{th} condition (time × treatment), and the kth replicate; r_i is the exonic region of the gene; t_j is condition; and $(rt)_{ij}$ is the interaction between exonic region and condition. A more traditional GLM test could not be used due to a lack of normality in the distribution of model residuals. Accordingly, a non-parametric test must be relied upon to look for changes in exon or exonic region representation between exons of a gene and we used of a rank test to summarize changes in exon representation (Supplemental File 3). F-tests for the significance of the mean square attributed to the effect tested versus the mean square attributed to error, or the appropriate interaction term, were used. The false discovery rate was controlled using the Benjamini-Hochberg procedure, with a significance cutoff of $\alpha = 0.05$ (Benjamini & Hochberg 1995).

GO Analysis

When a gene had more than one ortholog in *D. melanogaster* only one ortholog was included for the GO analysis, so as not to inflate the number of genes involved in a given process. This does presume that orthologs will be annotated with the same GO terms, and this is generally the case. For example, in *D. simulans* there is only *AOX4*, while in *D. melanogaster* there is *AOX3* and *AOX4*, but the GO terms for each are the same. However, as *D. simulans* genes are generally not independently annotated, especially those without *D. melanogaster* orthologs, if there was no *D. melanogaster* ortholog the gene was not included in the enrichment analysis. This is a significant fraction of the overall genes that were involved in the response to treatment, treatment by genotype, and genotype by treatment by time, however there is no viable alternative. Lists of significant genes were tested for GO enrichment using the PANTHER classification system (Mi *et al.* 2017). They were corrected for multiple testing and a *p*-value of .01 was required for significance.

Functional class enrichment

To test for functional class enrichment multigene exonic regions were not included. Every test of functional class enrichment compared the frequency of a given subcategory among all exons and exonic regions detected in the dataset compared to the frequency within a significant list of exons and exonic

295 regions. A χ^2 test was performed in R to test the significance of the enrichment of each of these
296 categories.

297 *D. melanogaster polymorphism and divergence*

298 To compare polymorphism between the significant sets of genes and genome-wide averages we
299 calculated Tajima's *D* genome-wide for the source populations of *D. melanogaster* and *D. simulans*. We
300 obtained the VCF files from the Winters population sequenced by (Campo *et al.* 2013), which includes all
301 six of the genotypes assayed here as well as 29 other inbred lines sampled from the same population at the
302 same time. The coordinates of the genes implicated for exons, exonic regions, and exon fragments were
303 converted from the current assembly coordinates (v6) to those used in the previous study (v5). For *D.*
304 *simulans* we used data previously obtained from 170 individuals from this population (Signor *et al.*
305 2017c). To obtain estimates that were consistent with our dataset (which is gene regions) we extracted
306 regions from genome-wide VCFs that corresponded to genes as annotated in the latest assembly. Note
307 that we are considering gene regions here rather than trying to include regulatory regions for a number of
308 reasons: 1) Gene regions includes introns, which will include some regulatory regions and splice sites. 2)
309 The location of regulatory regions for these genes is not well established, much less so between species
310 where we do not know if they may have shifted locations. This approach is more conservative than
311 including arbitrary amounts of putative upstream regulatory regions. 3) If there has been recent selection
312 on a regulatory region it may still be linked to the gene region and show the same differences in
313 polymorphism frequency. We excluded regions of reduced recombination near the centromeres, either 1
314 MB or more if there were significant reductions in diversity for a broader region, calculated as extended
315 negative Tajima's *D* or values of $\pi < ½$ the chromosomal average (Sedghifar *et al.* 2016; Signor *et al.*
316 2017c). SNPs within these coordinates were separated using bedtools intersect (v2.26.0). We excluded
317 regions from the fourth chromosome and unassembled scaffolds for the analysis of population genetic
318 differences. In both species this is a trivial number of significant regions. Alternate SNPs that were
319 present in > 99% of the mapping population were excluded, as were SNPs with more than 10% missing
320 data. Tajima's *D* was calculated in windows of 1 KB using VCFtools v1.12a. Windows of Tajima's *D*
321 that overhung the ends of genes were included in the analysis.
322 We calculated per-gene D_{XY} for the significant exons, exonic regions, and genes with significant
323 changes in rank abundance, to determine if there were any unusual patterns of divergence for these
324 subsets. To calculate divergence comparable regions of the genome need to be identified, and there has
325 been considerable evolution of transcription start sites between *D. melanogaster* and *D. simulans* (Main *et*
326 *al.* 2013). Comparing regions annotated as the gene regions in each species introduces large and
327 unexpected gaps in the start of the alignments. While the evolution of transcription start sites is of
328 interest, if unaccounted for it will artificially inflate estimates of D_{XY}. Thus for genes implicated in
329 differences in *D. melanogaster* the regions annotated as genes were blasted using ncbi blastn (v2.4.0) to
330 the *D. simulans* assembly and then back to the *D. melanogaster* assembly and used for divergence
331 statistics, and vice versa for *D. simulans* genes. This means that for different genes small non-coding
332 regions may be included depending upon the direction of evolution of transcription start sites, or small
333 portions of coding regions may be excluded. Furthermore, for genes from *D. simulans* with more than one
334 ortholog in *D. melanogaster* the top blast hit was used, and while there may have been orthologous
335 sequence in *D. melanogaster* (or *D. simulans*) if there was no annotated ortholog it was excluded from the
336 analysis. For example, some of the non-protein coding genes from *D. simulans* have very conserved
337 BLAST hits in *D. melanogaster* that are not annotated as containing genes. The sequences were aligned
338 using the R package DECIPHER (Wright 2016) and if necessary they were reverse complimented using
339 Biostrings (Pages *et al.* 2017, for example, in the region on 3R in *D. melanogaster* that is an inversion).
340 Following alignment the *D. simulans* and *D. melanogaster* reads were output separately in aligned fasta
341 format and D_{XY} was calculated using an R script courtesy of Dr. Emily Delaney, which incorporates
342 commands from the R packages ape and pegas (Supplemetnal file 4, Paradis *et al.* 2004; Paradis 2010)) .

343 This was done for every orthologous gene in the genome from *D. melanogaster* and *D. simulans*, and
344 each significant subset from each component of variance, to compare distributions of divergence.
345
346 Results

347 *Gene expression and isoform usage*

348 It is difficult to decouple alternative isoform usage from gene expression, given that many exons are
349 shared between isoforms or overlap other exons. To infer isoforms from short read data, one must rely
350 upon unique junctions or regions of individual isoforms and extrapolate to shared regions. This requires
351 accurate isoform annotation (knowing that any given exon/junction is found in combination with other
352 exons/junctions) and in general can be very noisy. Accordingly, we subscribe to a simpler but more robust
353 approach and summarize the abundance of different exons and exonic regions separately. We detail
354 overall abundance of exons, exonic regions, and exon fragments, and changes in the rank abundance of
355 exons within a gene (Figure 1 C&D). As illustrated in Figure 1 C&D, these approaches summarize
356 different features of expression and splicing – for example differences in the expression level of exon
357 fragments from *cabut* suggest both that there are overall expression differences for the gene between
358 environments, and that in both *D. simulans* and *D. melanogaster* one of the alternative isoforms is more
359 abundant with ethanol (Figure 1C). While exons within *Drat* change their expression in response to
360 ethanol by time, and between environments at 30 minutes, that doesn't capture the fact that the third exon
361 is most abundant in ethanol while the first is most abundant without it, suggesting differences in isoform
362 abundance that may also belong to unannotated isoforms (Figure 1D). In the following sections we will
363 first summarize changes in exon, exonic region, and exon fragment abundance followed by differences in
364 rank abundance between environments.

	G	ETOH	ETOH x G	ETOH x T	ETOH x G x T	10	20	30
Exons								
Unique	96	6	16	2	75	1	0	6
Unique/Constitutive	1547	58	324	19	943	5	1	16
Common	52	2	3	0	19	0	0	1
Constitutive	299	10	44	3	121	3	2	6
Exonic Regions								
Common	119	9	22	1	76	1	0	2
Constitutive	287	14	76	6	223	2	1	2
Fragments								
Unique	101	2	5	2	26	1	1	2
Unique/Constitutive	169	7	45	3	93	0	0	1
Common	147	6	6	1	32	1	0	2
Constitutive	191	3	25	0	74	2	1	1

365

366 *Table 1: A summary of the exons, exonic regions, and exon fragments that are signifigantly differently*
367 *expressed for each component of variance. Abbreviations are as follows: (G) Genotype, (ETOH)*
368 *Ethanol, and (T) Time.*

369 *Exons, exonic regions, and exon fragments*

370 1994 exons, 406 exonic regions, and 608 exon fragments changed their expression in response to
371 genotype in *D. simulans*, while in *D. melanogaster* 1445 exons, 631 exonic regions, and 1135 exon
372 fragments altered their representation . Note that the results for *D. melanogaster* are summarized in
373 (Signor & Nuzhdin 2018), but are included here for comparison. 76 exons, 23 exonic regions, and 18
374 exon fragments changed expression in response to ethanol in *D. simulans*, compared to 15 exons, 13
375 exonic regions, and 21 exon fragments in *D. melanogaster*. Seven exons and exonic regions were shared
376 between species for treatment, *Drat*, *cabut*, *CG11741*, *CG32512*, *CG4607*, *Pino*, and *sugarbabe.* For all
377 discussion of shared genes, the particular exon or exonic region may or may not be the same, as well as
378 the direction or nature of the change in expression. The complexity of this comparison is shown in Figure

379 1C, where in *D. melanogaster cabut* has three annotated transcripts and four exon fragments (one exonic
380 region), while in *D. simulans cabut* there are two transcripts, one exon, and two exon fragments (one
381 exonic region). In addition, in *D. melanogaster cabut* is nested within the gene *ush*, while in *D. simulans*
382 it is not nested. In *D. melanogaster* an area annotated as an intron in *D. simulans* is unique to its third
383 isoform, and this is more frequent with ethanol than without; in *D. simulans* there is one unique exon
384 fragment belonging to one isoform which is more frequent with ethanol than without, suggesting that
385 isoform is more common in ethanol environments (Figure 1C). In *D. melanogaster* the exon fragment
386 which is unique in *D. simulans* is not unique, and only increased frequency of its unique third isoform can
387 be inferred (Figure 1C). 15 exons and exonic regions and three fragments had no ortholog in *D.*
388 *melanogaster*, 12 exons and exonic regions and one exon fragment of which were non-protein coding
389 genes, and one exon fragment and one exon/exonic region of which are labeled as pseudogenes. The
390 interaction between genotype and treatment was significant for 387 exons, 98 exonic regions, and 81 exon
391 fragments in *D. simulans* and three exons, no exonic regions, and no fragments in *D. melanogaster*
392 (Figure 2 A&B). Of these, 82 exons and exonic regions and 9 exon fragments originate from genes that
393 do not have an ortholog in *D. melanogaster,* and 58 exons and exonic regions and seven exon fragments
394 of these were non-protein coding, while ten exons and exonic regions and one exon fragment were
395 pseudogenes. No fusions or fragments were shared between *D. simulans* and *D. melanogaster*.

396 24 exons, seven exonic regions, and six exon fragments were significantly different in response to
397 the interaction between ethanol and time in *D. simulans*, compared to 22 exons, eight exonic regions, and
398 12 fragments in *D. melanogaster*. Three genes were shared between exons and exonic regions for these
399 species, *Drat*, *cabut*, and *CG43366*. *CG43366* is the *Drosophila* homolog of human *Serpina2*, which has
400 previously been implicated in susceptibility to chemical dependence (Agrawal *et al.* 2008). In *D. simulans*
401 1158 exons, 299 exonic regions, and 225 exon fragments were significantly different for the interaction
402 between genotype, ethanol, and time, while in *D. melanogaster* no exons, two exonic regions, and four
403 exon fragments were significantly different (Figure 2 A&B). No exon fusions or fragments were shared
404 between species for this component of variance. Of the exons and exonic regions 24 were pseudogenes,
405 172 were non-protein coding genes, and 265 had no ortholog in *D. melanogaster*. Among exon fragments
406 39 had no ortholog in *D. melanogaster*, of which 29 were non-protein coding genes and four were
407 pseudogenes. None of those with orthologs were non-protein coding genes. At ten minutes nine exons,
408 three exonic regions, and four exon fragments are significantly different between treatments, compared to
409 two exons, one exonic region, and three exon fragments in *D. melanogaster* (Figure 2A). *cabut* is shared
410 between exons and exonic regions in *D. melanogaster* and *D. simulans*. At 20 minutes three exons, one
411 exonic region, and two exon fragments were different in *D. simulans*, two of which belong to *cabut* in
412 both exons/exonic regions and exon fragments. In *D. melanogaster* one exon, four exonic regions, and
413 five exonic fragments are different at this time point, and the gene *cabut* is shared between species for
414 both categories. At 30 minutes 29 exons, four exonic regions, and six fragments in *D. simulans* were
415 different between treatments, all of which have *D. melanogaster* orthologs. In *D. melanogaster* 46 exons,
416 24 exonic regions, and 20 exon fragments were significantly different between treatments at 30 minutes.
417 Among exons and exonic regions *Drat*, *CG32103*, *sugarbabe*, *cabut*, *Pinocchio*, *CG32512*, and *CG4607*
418 are shared between species. Among exon fragments *cabut* and *Pinocchio* are shared.

419 <u>*Shared and unique exons, exonic regions, and fragments*</u>

420 Comparisons between *D. simulans* and *D. melanogaster* are shown in Figure 2C for the three components
421 of variance in which enough genes are implicated in *D. melanogaster* to make the frequency of different
422 categories meaningful. When an exon is unique/constitutive it belongs to the only annotated transcript for
423 that gene, and as such no exonic regions are unique/constitutive. However, the combined counts are
424 shown for the common and constitutive categories. However, in *D. melanogaster* constitutive exons and
425 exonic regions are much more common than in *D. simulans*, where unique/constitutive exons are
426 overwhelmingly implicated. This could potentially be explained by differences in annotation, for example

427 if fewer genes have multiple transcripts annotated in *D. simulans*. While this does appear to be the case
428 (Supplemental Figure 1), it is unclear if it is enough to explain the discrepancy between species. It is also
429 possible that in response to the environment genes with alternative splicing are more important in *D.*
430 *melanogaster* compared to *D. simulans*. The proportion of genes in each category in *D. simulans* is
431 consistent between components of variance, including ethanol by genotype and ethanol by genotype by
432 time, again suggesting the possibility that differences in annotation are responsible. If annotation
433 differences are responsible this does not affect the overall results, for example the number of exons
434 implicated, but may make comparison between species for the number of unique versus constitutive
435 differences not meaningful. It also underlines the importance of not relying upon isoform annotation
436 when trying to understand differences in expression and alternative splicing.

437

		Biological Process		Cellular Component		Molecular Function	
D. simulans	Ethanol x genotype						
				axonemal dynein complex	3×10^{-3}	dynein light chain binding	3.3×10^{-3}
	Ethanol x genotype x time						
		sensory perception of taste detection of chemical stimulus	6.6×10^{-3} 8.2×10^{-3}	nucleplasm cytoskeletal part ciliary part	6.5×10^{-3} 8.4×10^{-3} 4.6×10^{-3}	RNA binding	7.8×10^{-3}

438

439 *Table 2: A summary of the significant GO enrichment terms for D. simulans. Two components of variance*
440 *(genotype x time; genotype x ethanol x time) have significant terms, the cutoff is p < .01.*
441
442 <u>*GO enrichment analysis*</u>
443 We report here only the results of enrichment for exons and exonic regions (Table 2). In *D. simulans* the
444 response to treatment, treatment by time, and the three time points were not enriched for any GO terms.
445 Much of the lack of enrichment is likely due to annotation issues – for example 9% of the exons
446 implicated in treatment over time could not be resolved to a single gene, and of the remaining genes 14%
447 do not have a *D. melanogaster* ortholog. Of those with annotated orthologs, 20% do not have any gene
448 ontology terms associated with them. A summary of significant GO terms for the remaining components
449 of variance is shown in Table 2. In *D. melanogaster* exons and exonic regions were not significantly
450 enriched for any category of genes. In general, the small number of genes implicated for many categories
451 precludes any conclusion of enrichment.
452

453 <u>*Changes in rank abundance*</u>

454 In response to treatment 54 genes showed changes in rank abundance, two of which are without a *D.*
455 *melanogaster* ortholog and one of which is non-protein coding. 94 genes change the rank abundance of
456 their exons for the interaction between treatment and time, including seven genes with no ortholog in *D.*
457 *melanogaster*, five of which are non-protein coding (three are pseudogenes). In *D. melanogaster* 71 genes
458 changed the rank abundance of their exons in response to ethanol and 145 changed for treatment by time
459 (Signor & Nuzhdin 2018). No genes were shared between species. For the interaction between treatment
460 and time *lola*, *Mhc*, and *Prm* were shared between species. *lola* is well established as being involved in
461 the response to ethanol and is the *Drosophila* ortholog of *ZBTB20*, hypermethylation of which has been
462 associated with major depressive disorder (Davies *et al.* 2014), as well as alcohol related cancer (Shi *et al.*
463 2018), and the development of fatty liver disease (Liu *et al.* 2017). Figure 1D illustrates an example of a
464 change in rank abundance for the gene *Drat* in *D. simulans*, where with ethanol the third exon is most
465 abundant, and without ethanol the first exon is most abundant. While differences in exon abundance

466 summarized above capture some of this variation, as *Drat* is significant for some exons for ethanol by
467 time and 30 minutes, the change in rank abundance highlights potential differences in exon inclusion
468 between environments.

469 <u>*GO enrichment analysis*</u>
470 Changes in rank abundance were not enriched for any GO terms in *D. simulans*, with the exception of
471 peroxiredoxin activity in response to ethanol, albeit slightly above the more conservative cut-off used in
472 our other tests (p = .017). Peroxiredoxin activity has been associated with protection against alcohol
473 induced liver damage (Bae *et al.* 2011; Chattopadhyay *et al.* 2015). In *D. melanogaster* genes implicated
474 in changes in rank abundance in response to ethanol by time were enriched for cellular components actin
475 cytoskeleton, again somewhat above our more conservative cutoff (p = 0.018) (Signor & Nuzhdin 2018).
476
477 *In D. simulans genotype-specific reactions to the environment are abundant*
478 In *D. melanogaster* components of variance for interaction terms have very few significant genes, with
479 the largest category being exons and exonic regions that respond to ethanol and that are different at 30
480 minutes. *D. simulans* is roughly comparable for the majority of these categories. However, many more
481 exons and exonic regions are significant for components of variance that interact with genotype: 1457 for
482 the interaction between genotype, treatment and time, and 486 for the interaction between genotype and
483 treatment, compared to two and three exons and exonic regions respectively in *D. melanogaster* (Figure 2
484 A&B). This suggests that in *D. simulans* interactions with genotype are a far more important component
485 of the response to ethanol than in *D. melanogaster*. It is also worth noting that in *D. simulans* the
486 response to ethanol (plasticity) is three times as large (28 exons and exonic regions in *D. melanogaster*
487 compared to 99), though this is many order of magnitudes less of a difference than for genetic variation
488 for plasticity.
489
490 *Genotype-specific responses are enriched for non-coding genes in D. simulans*
491 A large number of non-protein coding genes were implicated in gene expression changes in *D. simulans*
492 in these analyses, so we applied a χ^2 test to understand if our gene lists were enriched for this functional
493 category. The number of non-protein coding genes that were significant for ethanol, ethanol by genotype,
494 and ethanol by genotype by time, were more than would be expected by chance in *D. simulans* (13, $\chi =$
495 $49.598, p < 0.0005$; 68, $\chi = 235.21, p < 2.2\ x\ 10^{-16}$; 196, $\chi = 727.17, p < 2.2\ x\ 10^{-16}$). Other than
496 the pseudogenes, these are long non-coding RNAs (lncRNAs) as the shortest is annotated at 713 bp, and
497 the majority are over 4,000 bp (long non-coding RNAs being any non-protein coding genes over 200 nt).
498 The interaction between treatment and time, and the differences in expression at ten, 20, and 30 minutes
499 were not enriched for non-protein coding genes, nor were changes in rank abundance.
500 *D. melanogaster* is not enriched for non-protein coding genes in response to ethanol, ethanol by
501 time, ethanol by genotype, ethanol by genotype by time, ten or 20 minutes, exons or exonic regions
502 expressed only in one environment, or genes implicated in changes in rank abundance. However, at 30
503 minutes *D. melanogaster* is enriched for non-protein coding genes, though this concerns far fewer genes
504 than in *D. simulans* (5, $\chi = 12.831, p = 0.005$).
505
506 *Non-coding genes that respond to the environment in D. simulans do not have D. melanogaster orthologs*
507 Given that a considerable number of non-protein coding genes were implicated in this analysis for *D.*
508 *simulans*, we were suspicious that this may be an annotation issue in *D. melanogaster*. Using the exons
509 associated with every transcript from each of these non-protein coding genes as a reference we did not
510 find that *D. melanogaster* reads mapped to these exons across a range of relaxed mapping parameters
511 allowing for mismatches and gaps (using bwa mem, from defaults to -B 2 -O 3; using both the *D.*
512 *simulans* exons or existing homologous regions in *D. melanogaster*). In several cases, such as *CG30377*,
513 an annotated protein coding gene in *D. melanogaster* is annotated as a pseudogene in *D. simulans*, with
514 no noted relationship between them. Therefore there may be a small number of cases in which an ortholog
515 exists but is not recognized, but in general these non-protein coding genes appear to be unique to *D.*

516 *simulans*. It has been noted previously that lncRNAs can share high sequence similarity between related
517 species, but be expressed in only one (Ulitsky 2016). Determining the orthology of these genes and the
518 dynamics of non-protein coding gene evolution on the *Drosophila* phylogeny is an interesting question
519 that will require additional future research.
520
521 *The response to the environment is enriched for nested non-protein coding genes in D. simulans*
522 In analyzing the non-protein coding genes identified in *D. simulans* we noted that many of them were
523 nested in the introns of other genes. We investigated the frequency with which nested genes were
524 implicated in our analysis, and nested non-protein coding genes, to understand the possibility that
525 differences in expression could also be changes in intron retention in the parental genes (as 'nestedness'
526 generally refers to exons or entire genes found within the introns of other genes). Using the criteria that an
527 exon nested in an intron must overlap the intron by at least 80 bp or 10%, we found that in *D.*
528 *melanogaster* 9.2% of exons were nested within introns, while in *D. simulans* 9.7% were nested, similar
529 to what has been previously reported (Lee & Chang 2013). This was reflected in our data, where for both
530 *D. melanogaster* and *D. simulans* 6.8% of exons and exonic regions were nested within introns (lower
531 because multi-gene exons were excluded). However, among significant exons and exonic regions in *D.*
532 *melanogaster* 18.6% were nested, while in *D. simulans* 33.5% were nested. This is a significant
533 enrichment of exons that are nested within other introns, for both *D. melanogaster* ($\chi = 12.344, p <$
534 $.0004$) and *D. simulans* ($\chi = 1721.1, p < 2.2\ x\ 10^{-16}$).
535 The remaining question then is whether nested genes are more likely to be non-protein coding, or
536 whether the dataset is enriched for both. Indeed, compared to the total number of nested genes that are
537 noncoding within the dataset, the number that are significant for the response to ethanol is enriched for
538 nested, non-protein coding genes in *D. simulans* ($\chi = 237.49, p < 2.2\ x\ 10^{-16}$), but not in *D.*
539 *melanogaster*. Overall *D. melanogaster* has more annotated non-protein coding genes (2963) than *D.*
540 *simulans* (1675), and similarly more exons from noncoding genes are nested within introns (1772 *D.*
541 *melanogaster*, 1066 *D. simulans*), suggesting that this pattern is not reflective of annotation issues
542 between the two species. However, the question remains as to why nested genes are so much more likely
543 to be non-protein coding than non-nested genes – for example in the total dataset of *D. simulans* 9.7% of
544 exons are nested, while of the annotated non-coding genes in *D. simulans* 63% are nested. Furthermore
545 the importance of nested non-protein coding genes for the response to the environment, or potentially
546 delays in splicing that are specific to certain components of variance, is unclear.
547
548 *Nested non-protein coding genes involved in the response to ethanol are regulated independently of their*
549 *parental gene*
550 In *D. simulans* 328 significant nested genes (375 nested exons and exonic regions) are on the opposite
551 strand as their parental gene, 75% of the total significant nested genes. Of non-protein coding genes, this
552 bias is stronger, with 83% having the opposite strandedness as their parental gene. In *D. melanogaster* 13
553 significant genes are nested (16 total exons and exonic regions) and of these nine are on the opposite
554 strand as their parent gene (69%). Only one non-protein coding gene shares strandedness with its parental
555 gene, *CR44660*/*drp1*. It has been observed that many nested non-protein coding genes require expression
556 of the parental gene, share strandedness, and depend on splicing out of the parental intron for activation
557 (miRNAs, (Boivin *et al.* 2018). In both *D. simulans* and *D. melanogaster*, however, we do not observe
558 non-protein coding genes as being more likely to share strandedness with their parental gene among our
559 significant nested exons and exonic regions. This suggests that whatever the reason for the observed
560 enrichment in nested non-protein coding genes, it is likely not because the parental genes are being
561 expressed and the nested non-protein coding genes are being spliced out of the introns. We could not test
562 the *D. melanogaster* dataset for correlation between the expression of parental and nested genes as it is
563 too small. However, in *D. simulans* genes that were both non-coding and shared strandedness with their
564 parental gene had highly correlated expression differences between treatments (0.83, 14 genes). All other
565 categories (i.e. nested non-protein coding, opposite strand) were essentially uncorrelated (0.09-0.27). We
566 note that of the parental genes only three are also represented as significant in the main dataset, *sima*,

mira, and *bru-3* (significant in response to ethanol or genotype by ethanol). Thus it is unlikely that overall the observed differences in expression are due to changes in the expression of the parental gene, as this would likely result in differences in expression detected at both loci.

Polymorphism and divergence in D. melanogaster and D. simulans

We were interested to determine if the patterns of polymorphism and divergence in the genes implicated in the response to ethanol suggested that they were evolving under a particular selection regime or were unusually diverged compared to background levels of polymorphism or divergence. If the genes implicated in expression differences in *D. melanogaster* had unusually low Tajima's *D*, for example, this could suggest that directional selection for adaptation to ethanol is responsible for the observed expression differences. This might help to clarify whether the observed expression differences are adaptive or show evidence of adaptation. For both polymorphism and divergence in the following sections we will compare the genome-wide background distribution of polymorphism and divergence to differences in expression and rank abundance as a whole, rather than by components of variance, for two reasons. Firstly, for the majority of categories other than genotype interaction terms in *D. simulans*, the number of independent genes on each list is ~50% confounding any attempt to separate out components of variance. Second, in *D. melanogaster* there are too few genes that are significant for many components of variance and they cannot be considered separately.

	Threshold	Abundance: # of outliers	Rank abundance: # of outliers		Threshold	Abundance: # of outliers	Rank abundance: # of outliers
***D. melanogaster* candidate gene regions**				***D. simulans* candidate gene regions**			
D. melanogaster				***D. melanogaster***			
X-linked	-1.49	5	10	X-linked	-1.49	**13**	11
	2.29	4	9		2.29	**35**	19
Autosomal	-1.5	18	68	Autosomal	-1.5	191	1
	1.9	13	49		1.9	225	4
D. simulans				***D. simulans***			
X-linked	-1.89	**0**	10	X-linked	-1.89	77	77
	3.25	**15**	6		3.25	**46**	**4**
Autosomal	-1.09	**0**	37	Autosomal	-1.09	**318**	72
	3.06	**35**	67		3.06	**267**	**15**

Table 4: A summary of the threshold values for Tajima's D and the number of intervals in each species which fall beyond this cutoff. X-linked and autosomal genes are shown separately, outlier categories which are significantly enriched are shown in bold. The same numbers are shown as percentages of the total number of intervals in Figure 4.

<u>*Polymorphism*</u>

We calculated Tajima's *D* in windows of 1 kb for both all annotated gene regions in the genome and the subset of genes implicated in differences in ethanol response in both species, and consider anything in the top ±2.5% to be an outlier. Outliers for *D. simulans* and *D. melanogaster*, separated between the X and the autosome, are shown in Table 4, and the percentage of outliers for each category are graphed in Figure 3A.

D. melanogaster

For changes in abundance of exons and exonic regions on the X and autosome, *D. melanogaster* is not enriched for outliers of Tajima's *D* but there are more outliers than expected in *D. simulans* (Autosomes: $\chi = 5.26, p < 0.02$, X: $\chi = 29.361, p < 0.0005$). In *D. melanogaster* these intervals correspond to 27 genes, while in *D. simulans* they cover a total of 25, including both the X and

604 autosomes. Between the two lists there is considerable overlap in the location of outlier intervals, at
605 *CG43366*, *AOX4*, *dsb*, *fumble*, *CG31875*, *Mical, Ada2b, forked, CG1986,* and *CG42749.*
606 For genes implicated in changes in rank abundance in *D. melanogaster*, there is also no
607 significant enrichment for outliers of Tajima's *D* relative to the genome-wide frequency in *D.*
608 *melanogaster* or *D. simulans.* In *D. melanogaster* on the autosome these intervals cover 59 genes, while
609 in *D. simulans* they cover 56. *14-3-3ε*, *AcCoAs*, *boi*, *CG31522*, *CG34398*, *eff*, *elF4EHP*, *fru*, *grh*, *kay*,
610 *lds*, *lola*, *mRpL21*, *osp*, *siz*, *Syp*, *Tep2*, and *tweek* are shared in containing outlier intervals for Tajima's *D*
611 in both *D. simulans* and *D. melanogaster*. In *D. simulans* on the X chromosome these intervals cover
612 four genes, while in *D. melanogaster* they cover nine, and two are in common between the species
613 (*CG34417*, *Ten-a*).
614
615 ***D. simulans***
616 Many more genes were implicated in the response to ethanol in *D. simulans*, and for exons and exonic
617 regions that changed in abundance there was an excess of outliers for Tajima's *D* for the autosomes and
618 the X chromosome (Table 4; Figure 3; Autosomes: $\chi = 9.41, p = 0.0022$, X: $X = 14.03, p = 0.00018$).
619 These intervals correspond to 428 unique genes, and many outlier intervals overlap the same gene region.
620 For example, every 1 kb interval that overlaps the gene *Cubilin* is an outlier, with an average of -2.41.
621 Polymorphisms in the human ortholog of *Cubilin*, *CUBN*, have been associated with lifetime rates of
622 heavy drinking (Hamidovic *et al.* 2013). It is worth noting here that the outliers that were significantly
623 enriched in the *D. melanogaster* dataset for *D. simulans* were all in the positive direction, while these are
624 both positive and negative. Given the tendency towards positive Tajima's *D* in this population of *D.*
625 *simulans*, negative values of Tajima's *D* may be a more suggestive measure of selection, either purifying
626 or directional (Signor *et al.* 2017c). In *D. melanogaster* the genes that were implicated in changes in the
627 expression of exons and exonic in response to ethanol in *D. simulans* were less enriched for outliers on
628 the X chromosome than expected, and not enriched on the autosomes (X: $\chi = 6.38, p = .0012$).
629 For genes implicated in changes in rank abundance in *D. simulans* there was a significant
630 enrichment compared to genome-wide frequency for both the autosomes and the X (Autosomes: $\chi =$
631 $19.78, p < 8.7\ x\ 10^{-6}$, X: $\chi = 458.73, p < 2.2\ x\ 10^{-16}$). This includes intervals in 44 unique genes
632 including 16 intervals in the gene *Mondo*. In *D. melanogaster* this did not constitute an enrichment
633 relative to genome-wide. Outlier intervals that occur within shared genes between the two species include
634 *CG7881*, *lola*, *mbf1* (a stress response gene), *Mhc*, *PI31*, and *Pka-1* (which has been implicated in the
635 response to ethanol previously (Chen *et al.* 2008)). Note that for changes in the abundance of exons and
636 exonic regions in *D. simulans*, if regions of low recombination are excluded from the candidate regions
637 the enrichment is no longer significant, but there is no change among the other tests.
638
639 <u>*Divergence*</u>
640 We calculated D_{XY} for orthologous genes on the autosomes (10,435 genes) and the X (1,943 genes)
641 between *D. melanogaster* and *D. simulans* (Figure 3B). D_{XY} was similar to that previously reported, with
642 an average of 0.52 on the autosomes and 0.55 on the X (compared to 0.048 on the autosomes, 0.054 on
643 the X (Nolte *et al.* 2013)). For the subsets of genes implicated in expression differences the mean was
644 0.047 on the autosomes and 0.057 on the X for expression differences in *D. melanogaster* and 0.053 on
645 the autosomes and 0.06 on the X in *D. simulans*. The genes implicated in changes in rank abundance in *D.*
646 *melanogaster* had a mean D_{XY} of 0.047 on the autosomes and 0.05 on the X. In *D. simulans* this was
647 0.049 on the autosomes and 0.059 on the X. To determine if the distribution of D_{XY} for any of the
648 significant subsets of genes differed compared to the genome-wide distribution we used a Kolmogorov-
649 Smirnov two-sample test, separately for genes on the X and the autosomes. In *D. melanogaster* exons and
650 exonic regions that changed abundance in response to ethanol were not significantly different on the
651 autosomes or the X. However, exons and exonic regions with significant differences in rank abundance
652 did differ in distribution on the autosomes, but not the X (Autosomes: D = 0.14, $p = 0.002$). The genes
653 implicated in expression differences on the autosomes in *D. simulans* differed significantly from the

genome-wide distribution, but not on the X (Autosomes: D = 0.061, $p = 0.005$). However, those implicated in changes in rank abundance did not, on the autosomes or the X. A difference in distribution does not indicate the direction of effect, and in fact in *D. melanogaster* the mean D_{XY} for changes in rank abundance is lower than genome-wide, while for expression differences in *D. simulans* it is slightly higher. This suggests that the genes implicated in changes in rank abundance in response to ethanol in *D. melanogaster* are somewhat more conserved than expected based on genome-wide distributions, while those implicated in expression differences in *D. simulans* are slightly less conserved. Note that because we can only include genes with orthologs here this does not include any of the abundant non-protein coding genes implicated in *D. simulans*, likely to be among the more rapidly evolving genes.

Population genetic patterns – or a lack thereof- among genes that respond to ethanol are difficult to interpret, in part because plastic responses may be due to a single regulatory change at an upstream gene. As the number of regulatory (or coding) changes involved in the plastic response are not known, it is difficult to interpret a lack of enrichment, as it is possible, for example, that only one of the genes whose expression changed could be under selection. Furthermore, it is not known whether the response to ethanol is adaptive – while it is presumed to be so in *D. melanogaster* due to its relative fitness on ethanol compared to *D. simulans*, functional links between gene expression differences and the phenotype were not established.

Discussion

In this study both *D. melanogaster* and *D. simulans* have a plastic response to the environment, but *D. melanogaster* is lacking in genetic variation for plasticity. Given the observed patterns, it appears most likely that for *D. simulans* 15% ethanol is a novel environment, and that the marked G x E is a passive stress response. This maladaptive plasticity would allow for the expression of previously cryptic variation that had accumulated in the absence of selection. In *D. melanogaster* this environment may not be stressful or novel, resulting in past selection removing genetic variation for plasticity that in *D. simulans* was allowed to neutrally accumulate. In this sense *D. melanogaster* may be more 'locally adapted', as it has evolved a reduced plastic response compared to *D. simulans*. However, it could also be that because *D. melanogaster* colonized the Americas a few thousand years prior to *D. simulans*, *D. simulans* has not yet reached an optimal level of plasticity (a nonequilibrium situation). Also, the frequency with which they encounter ethanol rich environments may vary for *D. simulans* and *D. melanogaster*, resulting in different selection pressures for developing the optimal level of plasticity – particularly if there is a trade-off with traits that are beneficial to the semi-domestic habitat of *D. simulans*. Trade-offs among genetically correlated traits may mean that a genotype with optimal phenotypic plasticity in one environment is constrained from evolving the optimal phenotype in another environment, resulting in the maintenance of variation for phenotypic plasticity.

Joshi and Thompson (1997) found that in *D. simulans* there was greater between family variation for the phenotypic response to ethanol substrate prior to selection for tolerance to ethanol. Compared to *D. melanogaster* and control populations, after selection in an ethanol environment the greatest change was a reduction in variation for plasticity between *D. simulans* families. Joshi and Thompson (1997) quantified phenotypes such as development time, which are not easily generalizable to expression patterns, but it is suggestive. However, in studies of other phenotypes in presumably adapted and non-adapted populations of *D. melanogaster*, no increase in genotype by environment response were observed in non-adapted populations (Heckel *et al.* 2016). This could be due to differences in the degree of stress or novelty of different environments, or it is possible that the response in *D. simulans* is due to something other than stress induced maladaptive plasticity.

The abundance of lncRNAs which are involved in genotype by environment interactions in *D. simulans* is perhaps not surprising, given that they have been implicated previously in the response to stress (Valadkhan & Valencia-Hipólito 2016). It is also possible they are more frequently involved because lncRNAs are often less conserved, as there is no requirement for the maintenance of ORFs and codon synonymy (Chodroff *et al.* 2010; Ulitsky *et al.* 2011; Quinn *et al.* 2016; Ulitsky 2016). It has been

705 observed previously that transcription evolves more quickly than sequences, and lncRNA are commonly
706 homologous to non-transcribed sequences in other species, however these species are typically more
707 diverged than *D. melanogaster* and *D. simulans* (Main *et al.* 2013; Ulitsky 2016). If neutral variation was
708 allowed to accumulate without selection, and then uncovered in a stressful environment, preferential
709 accumulation within less constrained sequences would be expected. *D. melanogaster* is also enriched for
710 lncRNAs for the response to ethanol at 30 minutes, which could be explained by increasing ethanol stress
711 over time. While it is possible that lncRNAs are indicative of stress-induced maladaptive plasticity this
712 cannot be separated from a more general involvement in the stress response, which would not necessarily
713 occur as a result of maladaptive plasticity.
714 Why the lncRNAs are preferentially nested in *D. simulans* is less easily envisioned, though four
715 potential scenarios for an increase in nested lncRNAs are depicted in Figure 4. The simplest explanation
716 is only that there has been a change in the expression of nested lncRNAs, and perhaps nested lncRNAs
717 are more commonly involved in less essential processes than other lncRNAs and are therefore less
718 constrained. It is also possible that ethanol causes a change in the intron stability of the spliced transcript,
719 causing increased (or decreased) detection of the nested lncRNA (Figure 4). It is also possible that a
720 change in the processing of the parental gene occurred, causing a change in the number of reads mapping
721 to unspliced introns, or that the parental gene simply changed expression. These latter two explanations
722 would predict correlation between the parental gene expression and nested gene expression, which was
723 only observed for the very small fraction of lncRNAs that shared strandedness with their parental gene.
724 Thus it is more likely that either nested lncRNAs are less constrained, or ethanol alters the stability of
725 introns during or after the process of splicing.
726 The population genetic patterns observed in these two species are not easily interpreted. *D.*
727 *melanogaster* does not show any increase in outliers for Tajima's *D* in any direction for either changes in
728 rank abundance or expression (with the exception of fewer than expected on the X chromosome for *D.*
729 *simulans* genes implicated in expression differences). *D. simulans* is an outlier in the positive direction for
730 *D. melanogaster* genes implicated in differential expression, which could indicate that the genes involved
731 in expression differences in *D. melanogaster* are subject to relaxed purifying or balancing selection in *D.*
732 *simulans*. As these are all in the positive direction, it is worth noting that genome-wide *D. simulans* is
733 biased towards positive values of Tajima's *D*, likely due to recent population contraction (Signor *et al.*
734 2017c). While they are still more positive than expected due to background levels of Tajima's *D*, caution
735 is also warranted in interpreting these patterns as due to selection. Among genes with expression
736 differences and changes in rank abundance in *D. simulans*, there is an enrichment of outliers for
737 expression differences on the autosome and X. This is due in large part to negative Tajima's *D* outliers,
738 which given genome-wide patterns is overall more suggestive of selection, in this case directional
739 selection. It may be that *D. simulans* phenotypic plasticity is currently not in equilibrium, and there is
740 selection for an optimal phenotypic response. This can be true whether or not the observed response is
741 due to maladaptive plasticity – diversity of the passive stress response to ethanol implies that some
742 responses will be more beneficial than others, and there may be selection against the less adaptive stress
743 responses.
744 Between species divergence (D_{XY}) suggests that in *D. melanogaster* the genes involved in
745 changes in rank abundance are less diverged than expected compared to background levels of divergence.
746 In *D. simulans* the genes implicated in expression differences for exons and exonic regions are more
747 diverged than expected based on background levels of divergence, which combined with being outliers
748 for largely negative Tajima's *D* could indicate that they have been important for adaptation in *D.*
749 *simulans*. However, given that the number of nucleotide differences involved in the response to the
750 environment is unknown – for example all the observed patterns could be due to a single *trans* variant, it
751 is difficult to interpret the results of Tajima's *D* and D_{XY} in terms of selection.
752 Inferring that gene expression differences are adaptive or non-adaptive remains a major challenge
753 in the study of gene expression reaction norms, given the lack of direct correlation between gene
754 expression phenotypes and organismal phenotypes. However, the patterns observed in *D. simulans* do
755 suggest maladaptive plasticity in response to ethanol exposure. In this scenario abundant genotype by

environment interactions are expected to have accumulated neutrally and become uncovered in response to environment stress. In contrast, in *D. melanogaster* this ethanol environment is not novel and maladaptive plasticity has been selected out in favor of an adaptive phenotypic response. lncRNAs are preferentially differentially expressed in *D. simulans* in response to ethanol either because they are less constrained and can accumulate more neutral variation, or because they are involved in the general stress response. It is also possible that environmental heterogeneity has caused *D. simulans* to maintain balanced polymorphisms for plasticity in a way that has not occurred in *D. melanogaster*, though we believe there is less evidence in favor of this interpretation. In the future comparing African, non-ethanol adapted populations of *D. melanogaster* to cosmopolitan populations may be a way of discerning between these hypotheses.

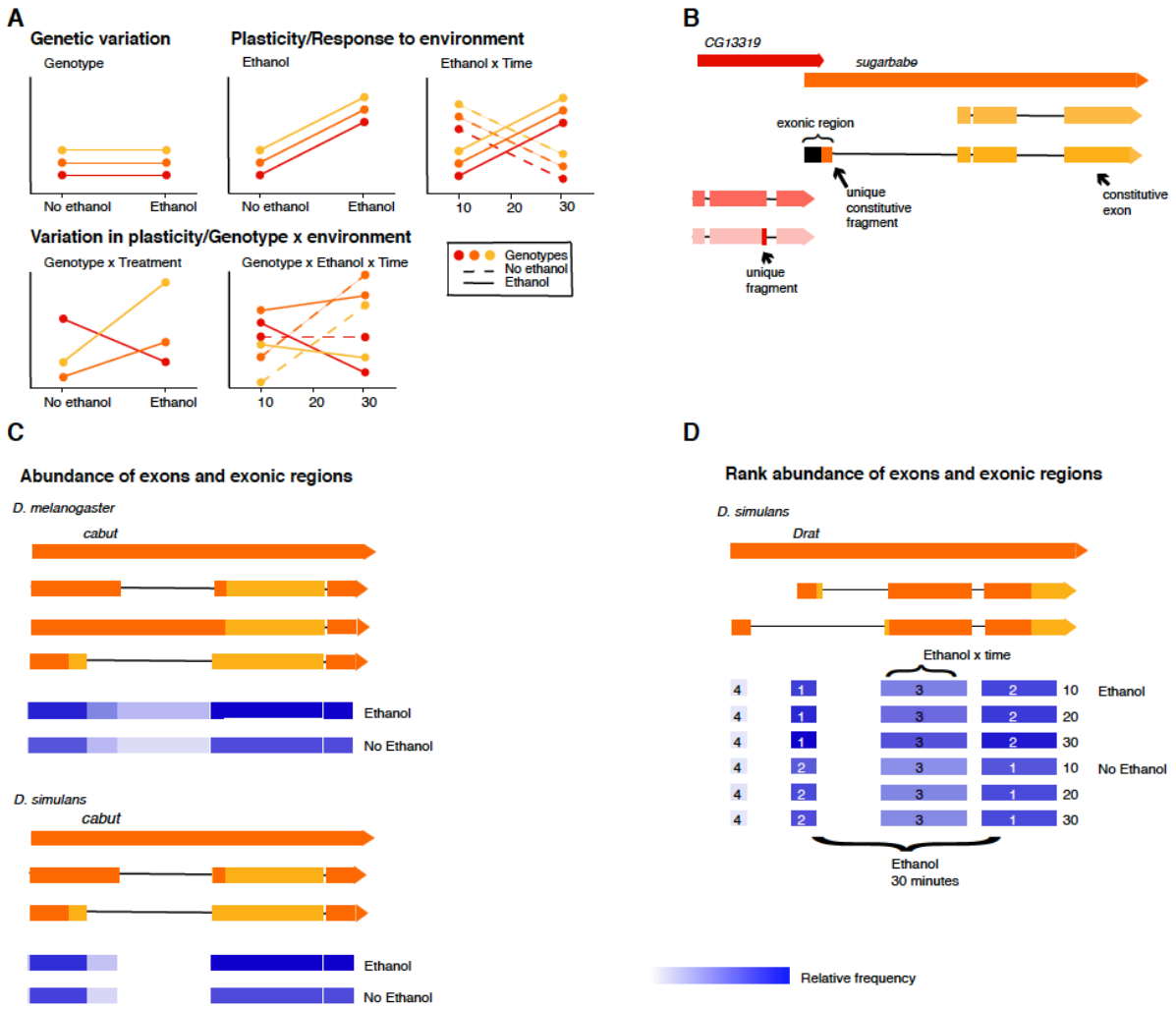

Figure 1: **A**. An illustration, for a single gene, of hypothetical gene expression changes for different genotypes with different main and interaction effects. For main effect genotype, no difference is seen in expression between environments but different genotypes have different expression levels. In response to the environment (ethanol), gene expression level changes between environments but for every genotype it

775 changes in the same way. Ethanol by time is complicated by the fact that the x-axis is now time, and there
776 are separate lines for ethanol (solid) and non-ethanol (dashed) environment. Here again there is a
777 component of time for gene expression changes, but every genotype changes in the same manner. For
778 genotype by environment interactions in the first panel different genotypes respond to ethanol differently,
779 and in the second panel different genotypes respond to ethanol differently over time. **B**. An illustration of
780 the classification scheme for exons, exonic regions, and exon fragments using the gene *sugarbabe*, which
781 overlaps by an exon with is neighboring gene *CG13319*. Exons either do not overlap other exons in
782 different isoforms (exons), or are fused regions consisting of a set of overlapping exons (exonic regions).
783 Exonic regions can be decomposed into exon fragments, depending upon their overlap between different
784 isoforms. Exons may be unique (found in a single isoform), common (found in some isoforms but not
785 all), or constitutive (present in all isoforms), and if they are unique to a single isoform they may also be
786 unique and constitutive if that is the only transcript annotated for that particular gene. As exonic regions
787 require overlap between exons they may not be unique, but they may be common or constitutive. Exon
788 fragments can be unique to a single isoform, common to several isoforms, or constitutive. Exon fragments
789 may be unique and constitutive only in the situation of a multi-gene exonic region where one gene has a
790 single isoform. **C.** An example of the information about gene expression and isoform abundance that can
791 be inferred from measures of differential exon abundance. *cabut* is a relatively simple example, having
792 few isoforms and exons, and yet it is still a complicated inter-species comparison given that *D.*
793 *melanogaster* has more annotated isoforms that *D. simulans*. In *D. simulans* the first of two isoforms
794 shown has a unique portion (fragment) that is more abundant in the presence of ethanol, suggesting that
795 that isoform is more abundant. In *D. melanogaster* only the second of the three isoforms has a unique
796 fragment, which is also more abundant in ethanol and suggests differential isoform usage. For both
797 environments and species the most frequent exon is assigned 100% pigment, and all other frequencies are
798 represented relative to that. **D.** An illustration of a change in the rank abundance of a gene, using the gene
799 *Drat*. Only the example of *D. simulans* is shown, as *D. melanogaster* and *D. simulans* only overlapped by
800 three genes with very complicated patterns of expression, such as *lola* with 24 annotated isoforms. *Drat*
801 has two annotated isoforms in *D. simulans*, with two constitutive exons and two exons unique to each of
802 the two isoforms. With ethanol, the exon unique to the first annotated transcript is the most abundant,
803 while without ethanol the first constitutive exon is most abundant. In addition, the second constitutive
804 exon is significant for changes in abundance (**A**) for the interaction between treatment and time, and the
805 two constitutive exons and exon unique to the first transcript have significant differences in expression for
806 ethanol, and between environments at 30 minutes. The dynamic changes in the expression of different
807 exons are captured in different ways by these three approaches to quantifying changes in expression and
808 splicing, where even within one gene there can be four complicated effects on expression characterized by
809 two approaches.

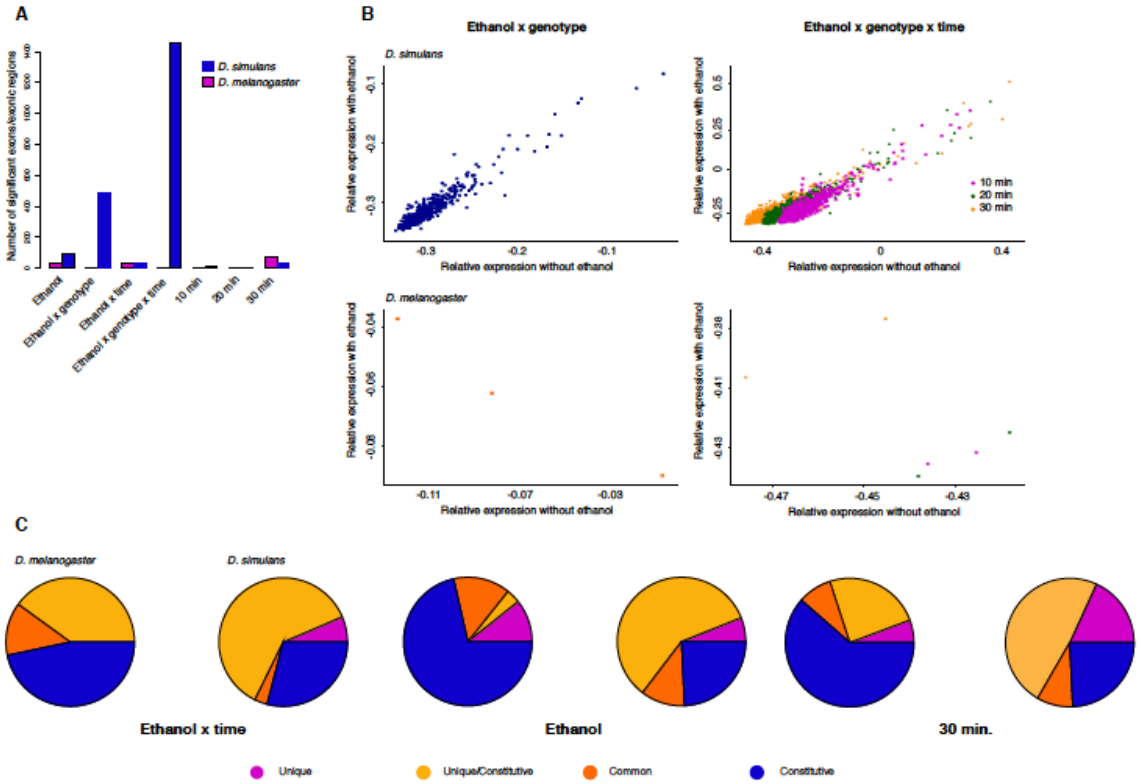

Figure 2: A. Components of variance for *D. melanogaster* and *D. simulans*. The number of significant exons and exonic regions for each component of variance are shown. Please note that in cases where so few exons and exonic regions are significant that no color can be seen, *D. melanogaster* is always on the left. Many more exons and exonic regions are significant for *D. simulans* than *D. melanogaster*, but only for components of variance that contain an interaction with genotype. **B.** *D. melanogaster* and *D. simulans* differ markedly in their genotype-specific reactions to ethanol, thus shown here is the relative expression in ethanol (*y*-axis) against the relative expression without ethanol (*x*-axis) for all genes with an interaction term with genotype. Too few genes are significant in *D. melanogaster* to make anything but a qualitative comparison, but the graphs serve to illustrate the magnitude and nature of this contrast. **C**. The proportion of exons and exonic regions that are either unique, unique/constitutive, common, or constitutive for the three components of variance for which a reasonable number of genes are significant in *D. melanogaster*. Note that while species names are only indicated for the first component of variance, *D. melanogaster* is on the left for every comparison. *D. melanogaster* has far more 'constitutive' exons and exonic regions compared to *D. simulans*, where unique/constitutive is more common. This is likely due to differences in the annotation of isoforms between the species, where *D. melanogaster* has more isoforms per gene annotated. It is also possible that alternatively spliced genes are more important for expression differences in *D. melanogaster*.

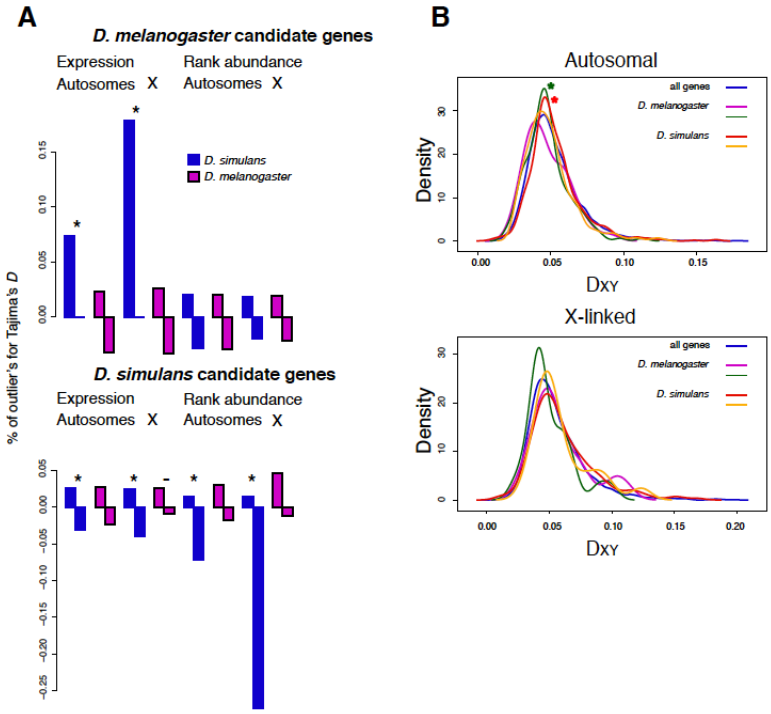

Figure 3: **A.** The percentage of intervals that were considered outliers in *D. melanogaster* and *D. simulans*, split between the autosomes and the X. The percentage of intervals that are outliers are shown separately for positive and negative, with the percent negative shown as a negative number. Note that the percentage is not actually negative but rather serves to illustrate the distinction between positive and negative. Enrichment for outlier intervals was determined as a composite of positive and negative intervals. Significant enrichment is indicated by a star, significant under-enrichment is illustrated with a minus sign. *D. melanogaster* is always on the left. The genome-wide cutoff for outliers was ±2.5%, thus enrichment is anything significantly more than or less than 5% for the positive and negative intervals together. **B**. The distribution of D_{xy} for *D. simulans* and *D. melanogaster*, compared to the genome-wide background distribution of D_{xy}. The genes are split between the autosomes (top) and X-linked (bottom), and the genome-wide background distribution is shown in blue. For *D. melanogaster* and *D. simulans* the color corresponding to changes in the abundance of exons or exonic regions is shown first, while that corresponding to changes in rank abundance is shown second , i.e. *D. melanogaster* pink – changes in overall abundance; green – changes in rank abundance.

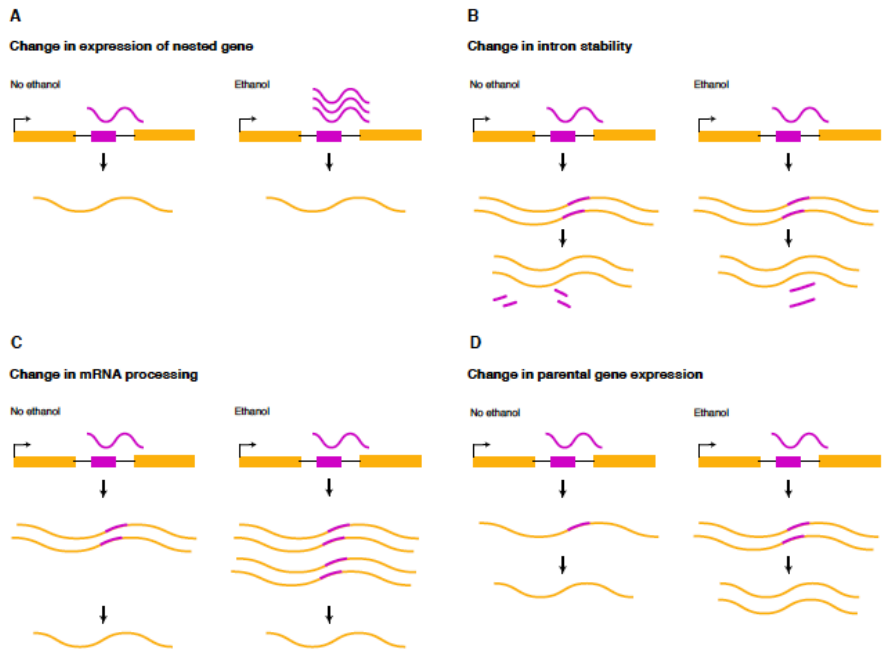

Figure 4: An illustration of the possible explanations for an enrichment in nested genes, and nested non-protein coding genes, in *D. simulans*. Note that in each illustration there are many possible details for each scenario – for example gene expression could increase or decrease in the presence of ethanol, but we illustrate here an increase in scenario **A.** This does not assume that this particular change is more likely, only that illustrating both an increase and a decrease will not aid in the readers comprehension. A. In the first scenario the nested gene changes its expression without any relationship with the expression of its parental gene, as shown by the number of squiggly lines representing RNA. Generic exons are represented by colored squares, introns by lines. The parental gene is shown in mustard yellow, and the nested gene in magenta, with RNA belonging to either in matching colors **B.** It is possible that ethanol alters the stability of introns after they are spliced out of the maturing transcript from the parental gene, potentially altering the expression level of the non-protein coding gene within the intron. Shown here and in the remaining scenarios is an extra step, with an unspliced RNA transcript from the parental gene preceding a spliced transcript. We do not attempt to accurately depict the process or machinery of splicing, only to illustrate that while expression of the nested gene may occur independently, some expression may also occur from introns as they are spliced out of the parental gene. **C.** Expression of the nested gene could also be affected if there were changes in the number of unspliced transcripts, for example through ethanol-induced alterations in the rate of splicing. **D.** The expression of the nested gene could also appear to change, or actually change through changes in the expression of the parental gene. Note that **C-D** would all predict some amount of correlation between parental gene expression and nested gene expression – such that if a change in nested gene expression is detected a change in parental gene expression would generally be expected as well.

Competing interests

The authors declare that they have no competing interests.

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Author's contributions

S.S. performed the experiments, analyzed the dataset, and wrote the manuscript. S.V.N. conceived of the experiment and assisted in writing the manuscript.

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