

The ontogeny of color: developmental origins of divergent pigmentation in *Drosophila americana* and *D. novamexicana*

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SUMMARY Pigmentation is a model trait for evolutionary and developmental analysis that is particularly amenable to molecular investigation in the genus *Drosophila*. To better understand how this phenotype evolves, we examined divergent pigmentation and gene expression over developmental time in the dark-bodied *D. americana* and its light-bodied sister species *D. novamexicana*. Prior genetic analysis implicated two enzyme-encoding genes, *tan* and *ebony*, in pigmentation divergence between these species, but questions remain about the underlying molecular mechanisms. Here, we describe stages of pupal development in both species and use this staging to determine when pigmentation develops and diverges between *D. americana* and *D. novamexicana*. For the developmental stages encompassing pigment divergence, we compare mRNA expression of *tan* and *ebony* over time

and between species. Finally, we use allele-specific expression assays to determine whether interspecific differences in mRNA abundance have a *cis*-regulatory basis and find evidence of *cis*-regulatory divergence for both *tan* and *ebony*. *cis*-regulatory divergence affecting *tan* had a small effect on mRNA abundance and was limited to a few developmental stages, yet previous data suggests that this divergence is likely to be biologically meaningful. Our study suggests that small and developmentally transient expression changes may contribute to phenotypic diversification more often than commonly appreciated. Recognizing the potential phenotypic impact of such changes is important for a scientific community increasingly focused on dissecting quantitative variation, but detecting these types of changes will be a major challenge to elucidating the molecular basis of complex traits.

INTRODUCTION

Pigmentation is a classic model for investigating the links between gene sequence and morphology (e.g. McClintock 1967; Levis et al. 1985; Martin and Gerats 1993; Wittkopp et al. 2002; Linnen et al. 2009; Cooley et al. 2011), in part because it is one of the most variable traits within and between species. It is also easy to visualize and provides a direct readout of the activity level of the underlying pigment biosynthetic pathways. Finally, the genes comprising pigmentation pathways in invertebrates, vertebrates, and plants are well characterized (Clegg and Durbin 2003; True 2003; Hoekstra 2006; Cazzonelli and Pogson 2010), further contributing to the emergence of pigmentation as one of the premier systems for studying the evolution of development (Kopp 2009).

The genus *Drosophila*, which combines the advantages of a well-developed model system with substantial phenotypic diversity, has been a particular focus of pigmentation studies (Waddington 1942; Levis, Hazelrigg, and Rubin 1985; Hollocher et al. 2000; Dombeck and Jaenike 2004; Gompel et al. 2005; Jeong et al. 2006; Rebeiz et al. 2009; Telonis-Scott et al. 2011). Pigmentation variation in *Drosophila* includes changes in overall body color as well as changes in melanin patterns

specific to the abdomen, wing, and thorax (reviewed in Wittkopp, Carroll et al. 2003). Here, we focus on a difference in overall body pigmentation between *D. novamexicana* and *D. americana*, members of the *virilis* species group. *Drosophila novamexicana* has a derived light-colored phenotype that is both striking and recent (Fig. 1A). It is estimated to have evolved from the dark body color seen in its sister species, *D. americana*, during the last 0.4 million years (Caletka and McAllister 2004). These two species appear to have allopatric distributions, with the dark-bodied *D. americana* found east of the Rocky Mountains and the light-bodied *D. novamexicana* found in sparse populations confined to the southwestern desert of the United States (Throckmorton 1982).

Prior genetic analysis has shown that changes linked to the pigmentation genes *tan* and *ebony* explain 87% of the body color difference between *D. americana* and *D. novamexicana* (Wittkopp et al. 2009). Interestingly, alleles contributing to this interspecific divergence also contribute to a longitudinal pigmentation cline in *D. americana* that is apparently maintained by natural selection (Wittkopp et al. 2009, 2011). Biochemically and phenotypically, *tan* and *ebony* have opposite effects on pigmentation (Fig. 1B): the Ebony protein promotes the accumulation of light yellow sclerotin by

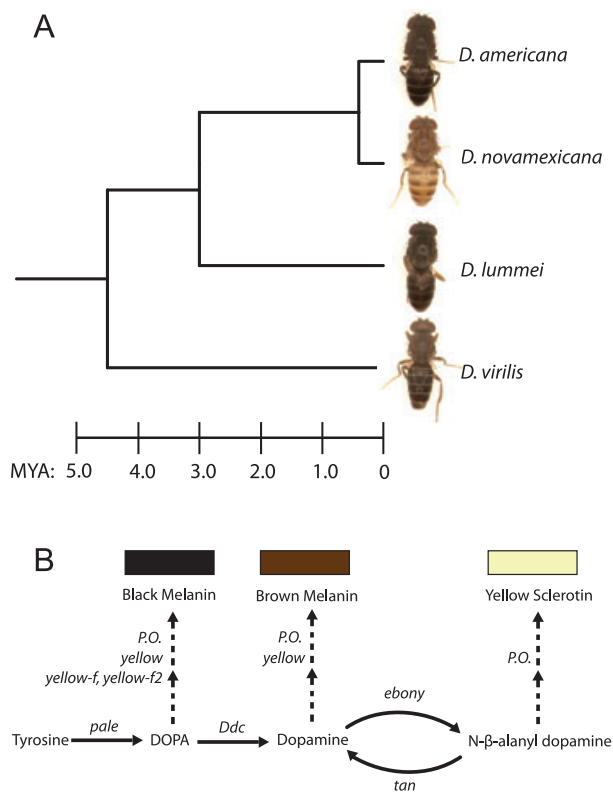


Fig. 1. Pigmentation in the *virilis* group of *Drosophila*. (A) The light body color of *D. novamexicana* is recently evolved and dramatically different from pigmentation in the other members of the *virilis* group. Estimated divergence times (MYA = millions of years ago), from Caletka and McAllister (2004), are shown on the scale at the bottom. (B) A simplified depiction of the melanin and sclerotin biosynthesis pathway. Gene(s) controlling each enzymatic step are shown in italics. *P.O.*, phenol oxidase-encoding genes. Arrows indicate chemical reactions, and branches with two consecutive dashed arrows indicate multiple enzymatic steps that are not completely known.

catalyzing the conversion of dopamine into N-β-alanyl-dopamine (NBAD), whereas Tan promotes the accumulation of dark brown melanin via the reverse biochemical reaction, converting NBAD to dopamine (Wittkopp, Carroll et al. 2003). Consistent with these functions, Ebony protein was reported to be more abundant in the lightly pigmented *D. novamexicana* (Wittkopp, Williams et al. 2003) whereas *tan* mRNA was reported to be slightly more abundant in the darkly pigmented *D. americana* (Wittkopp et al. 2009).

Expression differences between species can result from changes in either *cis*- or *trans*-acting sequences. The fact that genetic changes contributing to pigmentation divergence were found to be linked to *tan* and *ebony* suggests that the observed interspecific expression differences for these genes might be caused by divergent *cis*-regulatory sequences, although this has yet to be demonstrated. Consistent with

this hypothesis, fine-scale genetic mapping has shown that noncoding changes in the 5'UTR and/or first intron of *tan* (gene regions that often harbor *cis*-regulatory elements in *Drosophila*) contribute to pigmentation divergence between *D. americana* and *D. novamexicana* (Wittkopp et al. 2009).

Here, we take a closer look at how pigment patterns and expression of pigmentation genes develop during the late pupal stages of *D. americana* and *D. novamexicana*. Specifically, we answer the following three questions: (1) How and when does pigment development visibly diverge between *D. americana* and *D. novamexicana*? (2) How are the pigmentation genes *tan* and *ebony* expressed in each species during this developmental window of time? and, (3) at each developmental stage examined, are differences in *tan* or *ebony* expression between species due to divergent *cis*-regulation? Answers to these questions provide a more comprehensive and integrated understanding of the mechanisms underlying pigmentation divergence. They also illustrate how tests for *cis*-regulatory divergence are influenced by the changing cellular environment over developmental time and suggest that small changes in gene regulation can have significant phenotypic consequences.

RESULTS

Pupal development of *D. americana* and *D. novamexicana*

In order to compare pigmentation development between species, we generated a chronology for the sequential appearance of developmental features in *D. americana* and *D. novamexicana*. The definition of discrete developmental stages, each marked by the appearance of one or more morphological features, is an important tool within the *melanogaster* group (e.g. Ashburner 1989; Warren et al. 2006; Ninov and Martin-Blanco 2007), but has never been established for any species in the *virilis* group. We timed the appearance of stage-defining morphological features (supporting information Table S1) in *D. americana* and *D. novamexicana*, with identically aged *D. melanogaster* as a control, initiating data collection when the first pupae in the sample reached pupal stage P8 and continuing through adult stage A2.

Drosophila americana reached the A2 developmental stage 208.5 ± 5.7 hours (mean \pm 95% CI) after the start of the experiment. *Drosophila novamexicana* took a similar amount of time (212.1 ± 3.3 hours; Fig. 2A). Consistent with qualitative observations of the closely related species *D. virilis* (Markow and O'Grady 2005), these developmental times are more than twice that in *D. melanogaster* (98.9 ± 1.7 hours between start of experiment and occurrence of stage A2). Two-sample, two-sided *t*-tests showed that the time taken to reach stage A2 is significantly different between *D. melanogaster* and *D. americana* ($P < 0.0001$) and between *D. melanogaster*

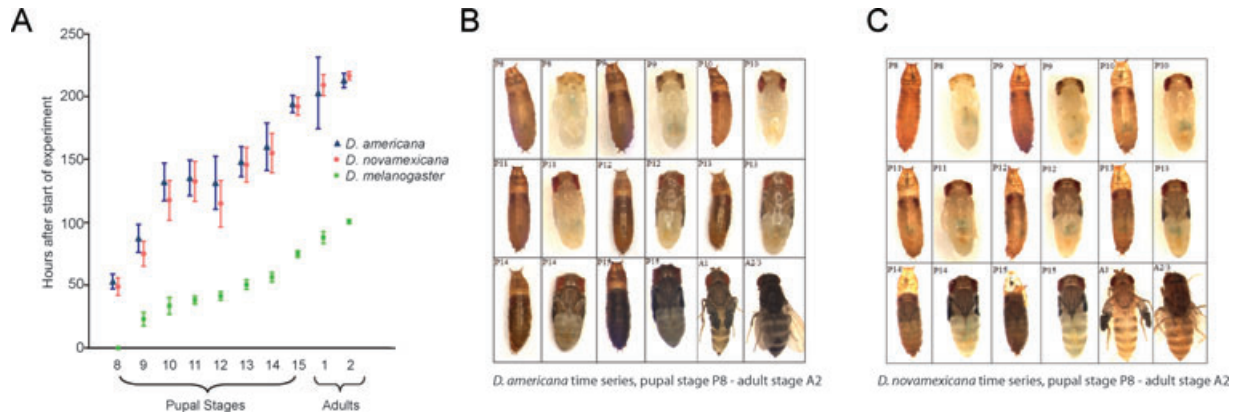


Fig. 2. Developmental stages P13–A1 encompass most of the visible accumulation of abdominal pigmentation in *D. americana* and *D. novamexicana*. (A) *Drosophila americana* and *D. novamexicana* develop more slowly than *D. melanogaster*. Beginning at stage P8 of *D. melanogaster*, samples of five developing flies per species were collected at regular time points, and their developmental stages were determined using morphological markers described by Bainbridge and Bownes (1981) (supporting information Table S1). The data were used to calculate the mean value (hours after initiation of P8 in *D. melanogaster*) of each developmental stage. Error bars indicate 95% confidence intervals. (B, C) Pigment accumulation differs between species in the late pupal and early adult stages. Photographs of *D. americana* (B) and *D. novamexicana* (C) were taken at pupal stages P8–P15 and adult stages A1–A2. For the eight pupal stages, flies were photographed both inside and outside of the pupal case (left and right photo, respectively, of each photo pair). Because the initial photos (taken for an earlier, unpublished project; L. Shefner, undergraduate senior thesis) showed an unusually dark stage P13 *D. americana* fly, we re-took the photos for that as well as the two surrounding stages (P12 and P14).

and *D. novamexicana* ($P < 0.0001$), but not between *D. americana* and *D. novamexicana* ($P = 0.287$).

Drosophila melanogaster accumulates melanic pigment during the latter half of pupal development (Walter et al. 1991), followed by yellow-pigmented sclerotization (Wright 1987). Similarly, visible abdominal pigmentation differences between *D. americana* and *D. novamexicana* accumulated between developmental stages P13 and A1 (Fig. 2B and C). The developing flies became darker in stages P13–P15, presumably reflecting melanization. They became more yellow at stage A1, implying the deposition of yellow sclerotin. This two-step process suggested that *tan*, which is necessary for melanization, might be transcriptionally active earlier in development than *ebony*, which is required for the production of yellow sclerotin. We tested this hypothesis by measuring changes in transcript abundance over developmental time for both genes.

Developmental profiles of *tan* and *ebony* differ over time and between species

To better understand the molecular changes underlying pigmentation divergence, we compared mRNA abundance of the *tan* and *ebony* genes between *D. americana* and *D. novamexicana* during the late pupal stages when the adult pigmentation differences are developing. mRNA levels were quantified using pyrosequencing, which can reliably detect expression differences as small as 10% between genotypes (Wittkopp et al. 2008). Because pyrosequencing directly mea-

sures the relative frequency of two alleles, we combined A1 stage flies from one species with flies from each pupal stage of interest from the other species (Fig. 3A). These A1 flies served as an internal reference that allowed us to compare expression within and between each species across developmental time, as described in the Materials and Methods.

For *tan*, the P13–P15 stages in which melanin visibly accumulated were marked by maximal gene expression in *D. americana* and relatively high gene expression in *D. novamexicana* (Fig. 4A and C). For *ebony*, expression increased dramatically at A1, coincident with the visible appearance of yellow sclerotin (Fig. 4B and C). Expression differences between the species were observed for both *tan* and *ebony*, in directions consistent with each gene's biological function and previously published comparisons at single developmental time points (Wittkopp, Williams et al. 2003; Wittkopp et al. 2009). *ebony* expression was approximately 2-fold higher in *D. novamexicana* than *D. americana* at each developmental time point (Fig. 4B; supporting information Table S2), whereas *tan* expression was generally higher in the dark-colored *D. americana* than in the light-colored *D. novamexicana*. The expression difference for *tan* was significant only at pupal stages P14 and P15, with a marginally significant difference at stage P13 (Fig. 4A; supporting information Table S2).

To obtain more precise estimates of the relative mRNA abundance between species, we performed direct comparisons by combining *D. americana* and *D. novamexicana* flies at the same developmental stage, for the stages at which *tan* and *ebony* are highly expressed (Fig. 3B). This meant

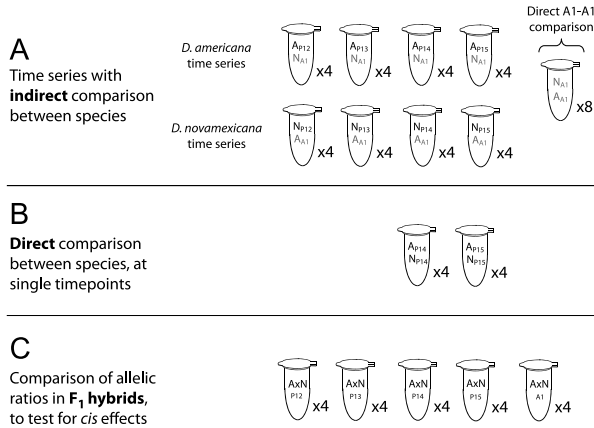


Fig. 3. Experimental designs for pyrosequencing assays of mRNA expression. Each tube represents a collection of six flies (three *D. americana* and three *D. novamexicana*, or six F_1 hybrids). Four biological replicates were collected for each stage, with technical replicates obtained for each biological replicate as described in the Materials and Methods. DNA and RNA were extracted from each tube; RNA was converted to cDNA; and the relative abundance of *D. americana* and *D. novamexicana* alleles of *tan* and *ebony* in each tube was measured using pyrosequencing. (A) To measure changes in gene expression across time, each stage of *D. americana* (black font) was combined with *D. novamexicana* stage A1 reference flies (gray font). Conversely, each stage of *D. novamexicana* was compared with *D. americana* stage A1 reference flies. To enable an indirect comparison between the species, the ratio of the two references (*D. novamexicana* stage A1 and *D. americana* stage A1) was used as a correction factor for the *D. americana* time series, as described in the Materials and Methods. (B) Expression differences at stages P14 and P15 were further investigated using a direct comparison of *D. americana* and *D. novamexicana* flies from the same stage. (C) *D. americana* \times *D. novamexicana* F_1 hybrids, collected at each of the five focal developmental stages, were used to determine whether the two alleles of each gene are still differentially expressed when they share a common *trans*-regulatory environment.

creating new pools of flies for stages P14 and P15, which showed the highest expression for *tan*. The A1 stage, which had the highest expression for *ebony*, was already directly measured as part of our initial experimental design (Fig. 3A). Relative to the indirect estimates obtained from the temporal experiment described above, the direct comparison revealed somewhat larger expression differences between the species (supporting information Fig. S1). Again, *ebony* was more highly expressed in *D. novamexicana* whereas *tan* was more highly expressed in *D. americana*. The direct-comparison data (stages P14, P15, and A1) revealed significant interspecific expression differences for *ebony* at all three stages, and for *tan* at stages P14 and P15, but not A1 (supporting information Table S2).

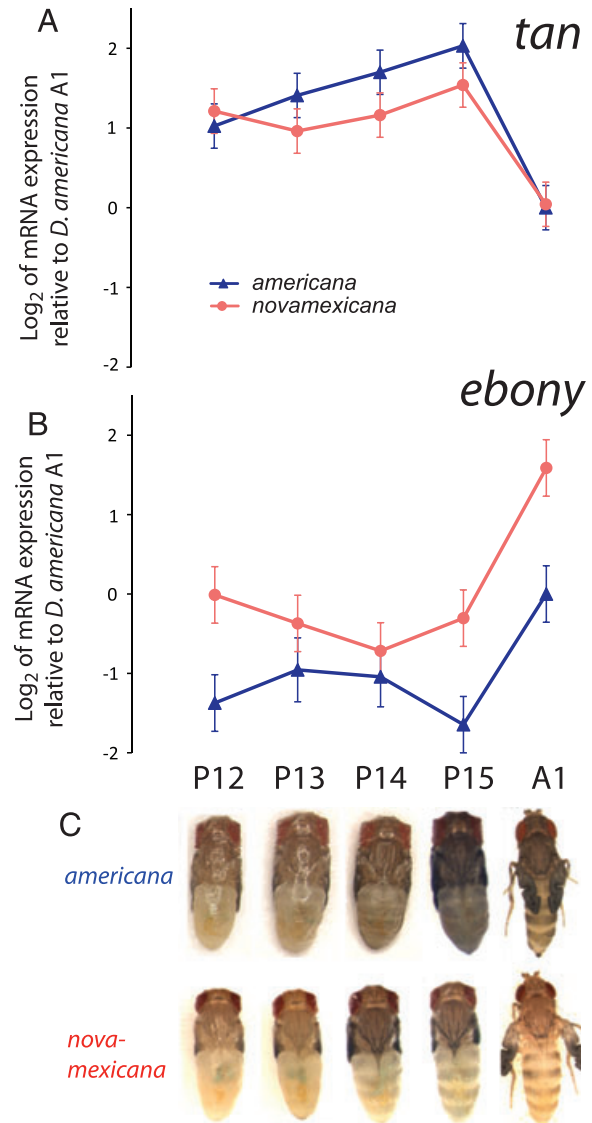


Fig. 4. Total mRNA expression of *tan* and *ebony* differs over developmental time and between species. The \log_2 of mRNA abundance at *tan* (A) and *ebony* (B) are shown for developmental stages P12–A1 (C). Each point represents the mean of four \log_2 -transformed biological replicates of either *D. americana* (blue) or *D. novamexicana* (red) mRNA expression, relative to *D. americana* stage A1. *Drosophila americana* stage A1, the internal reference point, was arbitrarily assigned a \log_2 value of 0 for both genes. Error bars are the experiment-wise 95% confidence intervals, calculated separately for *tan* and *ebony*.

cis*-regulatory variation contributes to divergent expression of both *tan* and *ebony

Variation in gene expression can arise from differences in *cis*- or *trans*-regulation. In order to test whether *cis*-regulatory divergence contributes to the expression differences observed between *D. americana* and *D. novamexicana*, we quantified the relative mRNA abundance of the two alleles for each gene

in a *D. americana* × *D. novamexicana* F₁ hybrid at developmental stages P12–A1 (Fig. 3C). Regulatory differences that act in *cis* cause allele-specific expression, whereas regulatory differences that act in *trans* affect expression of both alleles in a diploid cell. These two types of changes can therefore be differentiated by comparing allele-specific expression differences in an interspecific hybrid (Cowles et al. 2002; Wittkopp et al. 2004). Because both alleles in the hybrid are exposed to the same set of transcription factors, expression differences between the alleles are inferred to result from differences in *cis*-regulatory sequences.

In the F₁ hybrids, the *D. americana* allele of *tan* was expressed more highly than the *D. novamexicana* allele (Fig. 5A). This difference was nominally significant ($P < 0.05$) for stages P13–P15, suggesting differences in *cis*-regulatory activity, although only stage P14 survived a (conservative) Bonferroni correction for multiple testing (supporting information Table S2).

For *ebony*, the *D. americana* allele was expressed approximately 2-fold lower than the *D. novamexicana* allele, at each developmental stage (Fig. 5B). The *cis*-regulatory contribution to the expression differences between the two species was significant at each developmental stage (supporting information Table S2). Divergent *cis*-regulation explained the majority of the interspecific expression difference for *ebony*, but less than half of the interspecific expression difference at *tan* (Fig. 5). This suggests that *trans*-regulatory changes also contribute to expression divergence between *D. americana* and *D. novamexicana*, especially for *tan*.

DISCUSSION

The recent and dramatic difference between the dark-bodied *D. americana* and the light-bodied *D. novamexicana* presents an ideal opportunity for better understanding the mechanisms of trait divergence. The visibility of the phenotype, the lack of strong mating barriers between the two species, and the extensive history of *Drosophila* pigmentation research all make this system especially useful for genetic, molecular, and developmental investigation.

Developmental genetic changes underlying pigmentation divergence

Most of the body color evolution between *D. americana* and *D. novamexicana* is explained by genetic variants linked to two biosynthetic enzyme-coding genes, *tan* and *ebony* (Wittkopp, Williams et al. 2003; Wittkopp et al. 2009). The causal variant(s) at each locus could be amino acid changes that affect protein function, or noncoding changes that affect mRNA or protein abundance in either a *cis*-regulatory (allele specific) or *trans*-regulatory (nonallele-specific) fashion.

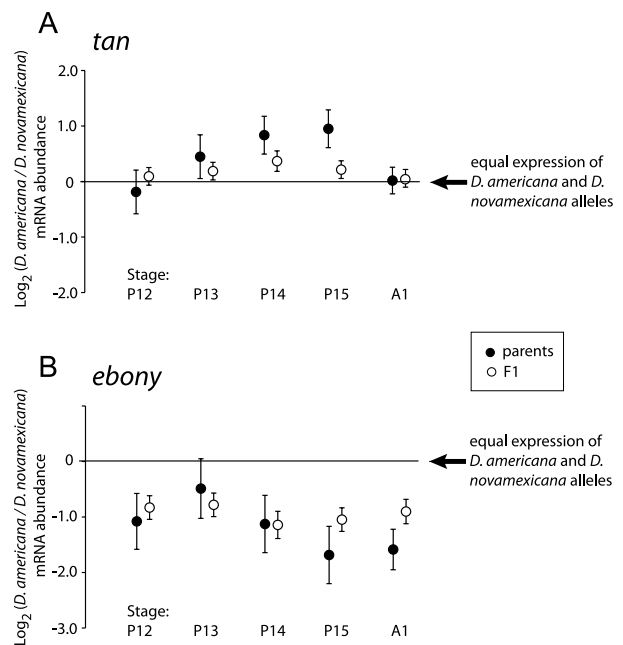


Fig. 5. Species- and allele-specific differences in *tan* and *ebony* mRNA expression over developmental time. The log₂ of mRNA abundance at *tan* (A) and *ebony* (B) are shown for developmental stages P12–A1 (x-axis). Filled circles represent the ratio of *D. americana* to *D. novamexicana* gene expression between the species (parental data). Open circles represent the ratio of expression of the two alleles within F₁ hybrids. A value of 0 corresponds to no expression difference between the two alleles; alleles are significantly differentially expressed ($P < 0.05$) at all except stages P12 and A1 (both parental and F₁ data) for *tan*, and stage P13 (parental data only) for *ebony*. Parental data were obtained using indirect comparisons for stages P12 and P13, and direct comparisons for stages P14–A1. Each point shows the mean of four biological replicates. Error bars are experiment-wise 95% confidence intervals, calculated separately for each dataset (parental indirect; parental direct; and F₁ hybrid).

Neither Tan nor Ebony protein sequences contain fixed differences between *D. americana* and *D. novamexicana* (Wittkopp et al. 2009), suggesting that nonsynonymous mutations are unlikely to be responsible for the fixed difference in pigmentation.

At *tan*, fine mapping has identified a functionally divergent region that contains no coding differences. The fine mapped region spans 2.7 kb, from the 5'UTR to the end of the first intron (Wittkopp et al. 2009). It contains 56 single nucleotide polymorphisms (SNPs) and 19 insertions or deletions (indels) in the first intron (which is a common location for regulatory elements in *Drosophila*) and one SNP just upstream of the first exon. These data show that the pigmentation phenotype must be controlled at least in part by noncoding changes in *tan*. The location of these divergent sites—5'UTR and first intron—suggests, but does not prove, that an allele-specific (*cis*-regulatory) mechanism

such as a mutation in a transcription factor binding site is responsible for divergent *tan* activity. A *cis*-regulatory difference was previously looked for but, surprisingly, not found (Wittkopp et al. 2009). In the present study, however, we sampled developmental stages more broadly and precisely and found evidence of significant *cis*-regulatory differences between the *D. americana* and *D. novamexicana* *tan* alleles, although the difference observed was small and limited to a narrow window of developmental time.

Qualitative differences in the abundance of the Ebony protein have been reported between *D. americana* and *D. novamexicana* (Wittkopp, Williams et al. 2003); *ebony* mRNA abundance has not previously been examined in these species. The difference in Ebony expression could be due to genetic changes at *ebony* itself or at other (*trans*-acting) loci. Genetic mapping has shown that variation linked to *ebony* contributes to pigmentation divergence; however, in the population used for mapping, *ebony* was linked to approximately 20% of the second chromosome because of a chromosomal inversion between species (Wittkopp, Williams et al. 2003; Wittkopp et al. 2009). Consequently, genetic mapping between *D. americana* and *D. novamexicana* cannot be used to discriminate genetic changes at *ebony* from genetic changes at other loci within the inversion. The *cis*-regulatory difference affecting *ebony* mRNA expression we report here therefore provides the most direct evidence to date that genetic changes at *ebony* itself contribute to pigmentation divergence.

Small changes in mRNA abundance may have functional consequences

With many studies now using gene expression (i.e. mRNA and/or protein abundance) as a way to understand how genotypes are converted into phenotypes, it is important to understand the relationship between changes in gene expression and changes (or lack thereof) in phenotype. We found that *cis*-regulatory divergence for *tan* is modest, increasing expression of the *D. americana* allele by only 29% compared with the *D. novamexicana* allele, yet multiple lines of evidence suggest that these *cis*-acting changes contribute to pigmentation divergence (Wittkopp et al. 2009). This suggests that even small and hard-to-detect differences in expression can be biologically meaningful, although we cannot definitively rule out a precise combination of *cis*- and *trans*-regulatory changes between species that cause a smaller difference in relative allelic expression in hybrids than between species (Takahasi et al. 2011). Small yet highly repeatable expression changes have also been observed in other studies, for example in response to experimental diet manipulations in mice and rats (de Boer et al. 2006; Patsouris et al. 2006; Rodenburg et al. 2008) and among wild-caught *Fundulus* fish (Oleksiak et al. 2005). In the latter case, small differences in gene expression were found to have just as much predictive power for individual metabolic trait variation as larger differences.

These small differences in RNA abundance might translate into small differences in protein abundance that nevertheless have large effects on cellular function, but it is also possible that small differences in RNA abundance might be amplified to produce larger differences in protein abundance, perhaps by producing stable proteins that accumulate overtime. Regardless of the mechanism, the focus of molecular geneticists on increasingly on complex, quantitative traits suggests that the ability to accurately detect small as well as temporally limited changes in expression is likely to be ever more critical.

cis-regulatory mutations can have temporally restricted effects

Although genomic sequence remains constant throughout an individual's life, the effects of *cis*-regulatory sequence variants may only be seen in certain transcriptional contexts, such as a particular sex (Bhasin et al. 2008), tissue type (Pretetto et al. 2006), or environmental condition (Li et al. 2006; Tirosch et al. 2009). Ontogeny creates, from the perspective of a single cell, a dynamic environment within a single individual. For any cell, the surrounding tissue types, extracellular molecules, and/or intracellular regulatory factors are subject to change over developmental time, and these changes cause temporally dynamic gene expression. This is illustrated by the temporal specificity of *cis*-regulatory divergence at *tan*. The allele-specific expression difference is visible at developmental stages P13–P15, but is absent at P12 and A1 (Fig. 5)—despite the fact that the gene is still expressed at those times. The ephemeral nature of the allele-specific expression difference may result from a *cis*-regulatory mutation(s) that affects a binding site(s) for a transcription factor(s) that is important primarily at stages P13–P15 (Fig. 6). Other *cis*-regulatory sequences, conserved between species, presumably control expression at other times during development (Fig. 6). This illustrates that although *cis*-regulatory mutations with phenotypic effects can be mapped genetically, assessing the impact of these mutations on gene expression is dependent on the environmental and developmental context.

MATERIALS AND METHODS

Timing of developmental stages

Three fly strains—*D. melanogaster* Canton S; *D. americana* A00 (strain 15010–0951.01); and *D. novamexicana* N14 (strain 15010–1031.14)—were raised on standard cornmeal media at room temperature. Sixty wandering third instar larvae of each strain were removed from their vials and distributed into new vials, with larval density standardized to 20 per vial. Vials were incubated at 20°C until pupal cases had formed and the pupae had hardened. Pupae were affixed to pieces of tape placed inside clear boxes, which were returned

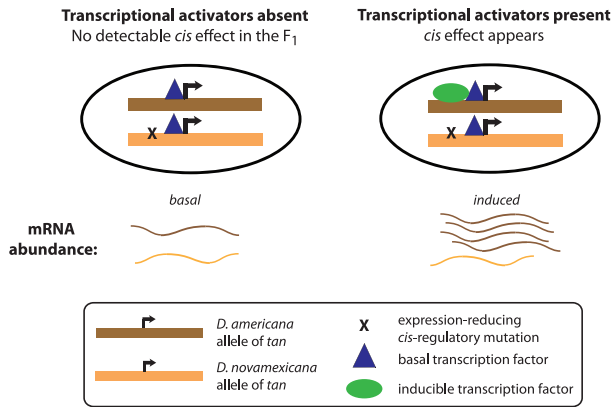


Fig. 6. *cis*-regulatory mutations are constant, but their effects can be ephemeral. Here we illustrate a hypothetical example in which two alleles of the *tan* gene are functionally differentiated by a single SNP in the *cis*-regulatory region of the gene. The derived allele, in *D. novamexicana*, contains a mutation that reduces the binding affinity of a stage-specific transcription factor (green oval), leading to reduced gene expression. The blue triangle represents a transcription factor controlling basal levels of expression that binds similarly to each species' allele. When this type of scenario occurs, the effects of the *cis*-regulatory mutation may only be noticed in environmentally or developmentally specific environments, when a sufficient amount of the relevant transcription factor is active.

to the 20°C incubator. Boxes were checked every 2–6 hours, and each time a sample of five pupae from each species was removed for analysis. The stage of each fly was determined according to the traits listed in supporting information Table S1 by examination under a dissecting microscope, removing the pupa from its case when necessary for unambiguous staging. Data were recorded from stage P8 onward; time 0 was defined as the time at which the first fly reached stage P8. One individual from *D. americana* and one individual from *D. novamexicana* was photographed for each pupal stage using a Leica MZ6 microscope and a Scion color digital camera, model CFW-1308C. Both males and females were used in this experiment, as pigment patterning is not sexually dimorphic in *D. americana* or *D. novamexicana* (Wittkopp et al. 2011). Because the initial photos (taken for an earlier, unpublished project; L. Shefner, undergraduate senior thesis) showed an unusually dark stage P13 *D. americana* fly, we retook the photos for that as well as the two surrounding stages (P12 and P14) in *D. americana*. The P12 and P14 stages were used as references to verify that lighting conditions were consistent between the two photo sets.

Experimental design for three types of gene expression analyses

We used three experimental designs to investigate different aspects of *tan* and *ebony* mRNA abundance: over time, be-

tween species, and in F₁ hybrids (Fig. 3). In all three cases, ratios of *D. americana*:*D. novamexicana* gene expression were determined using pyrosequencing.

Gene expression over time. To track changes in expression over time, we measured mRNA abundance of each *D. americana* developmental stage relative to *D. novamexicana* A1-stage flies, and mRNA abundance of each *D. novamexicana* developmental stage relative to *D. americana* A1-stage flies. In order to compare the time courses between the species, we wished to plot all of the time points relative to *D. americana* A1. Because the raw pyrosequencing data provided the ratio of *D. americana* alleles at each stage relative to *D. novamexicana* alleles at stage A1, a transformation was necessary to make *D. americana* A1 the common denominator:

$$\log_2 \left(\frac{A_{stageX}}{N_{A1}} \right) + \log_2 \left(\frac{N_{A1}}{A_{A1}} \right) = \log_2 \left(\frac{A_{stageX}}{A_{A1}} \right)$$

where A_{stageX} = the mean *D. americana* value for stage X and N_{A1} = the mean *D. novamexicana* value for stage A1. This is simply the log equivalent of:

$$\frac{A_{stageX}}{N_{A1}} \times \frac{N_{A1}}{A_{A1}} = \frac{A_{stageX}}{A_{A1}}$$

Gene expression differences between species. To more directly quantify differences in mRNA abundance between the species, at developmental time points of particular interest, we combined *D. americana* and *D. novamexicana* flies from the same stage and measured their relative mRNA abundance. This assay was performed for stages P14 and P15. A direct comparison of mRNA abundance at the A1 stage was obtained from the previous experiment (Fig. 3A).

Gene expression in F₁ hybrids. To test whether *cis*-regulatory divergence contributes to the total expression difference observed between the species, the relative mRNA abundance of *D. americana*:*D. novamexicana* alleles was measured in F₁ hybrids (Fig. 3B). Although the total expression difference between the species can reflect both *cis*- and *trans*-regulatory divergence, allelic expression differences in hybrids represent only the *cis*-regulatory component, because both alleles in the hybrid share a common *trans*-regulatory environment.

Fly collection and processing

Drosophila americana A00 and *D. novamexicana* N14 lines for gene expression analysis were raised on standard cornmeal media at 20°C. F₁ hybrid females were obtained by mating virgin female *D. americana* flies with male *D. novamexicana* flies (three of each gender per vial). Parents were discarded prior to hybrid eclosion, in order to prevent backcrossing. Prior work shows that female hybrids from reciprocal crosses have comparable pigmentation (Wittkopp et al. 2003).

Drosophila americana, *D. novamexicana*, and F₁ hybrid females were collected at stages P12–A1. Stages P12–P15 were first removed from pupal cases. A1 flies were collected within 15 min of eclosion. Because we wished to enrich for abdominal expression of *tan* and *ebony*, and wings are visibly pigmented from stage P12 on, wings were removed and discarded. Heads were also removed because *ebony* is expressed strongly in the eyes (Hovemann et al. 1998; Richardt et al. 2002) and both *tan* and *ebony* are expressed in the brain (True et al. 2005). The remaining tissue (thorax, abdomen, and legs) was frozen on dry ice and stored at -70°C until used for DNA and RNA extraction.

Each sample, containing either three flies of each species or six F₁ hybrids, was homogenized and used for sequential RNA and DNA extractions followed by cDNA synthesis, as in Wittkopp et al. (2004). The success of each extraction and cDNA synthesis was verified by visualizing 5 μl of final product on an agarose gel.

Pyrosequencing analyses of gene expression

In order to quantify species differences in *tan* and *ebony* mRNA expression, we compared the abundance of *D. americana* and *D. novamexicana* alleles using pyrosequencing (Ahmadian 2000). Polymerase chain reactions (PCR) and pyrosequencing primers were developed for both *tan* and *ebony*, in each case surrounding a site that differs between *D. americana* and *D. novamexicana* (supporting information Table S3). PCR and pyrosequencing reactions were performed as described in Wittkopp et al. (2008), using four biological replicates per developmental stage. For each biological replicate, cDNA was measured in triplicate. Genomic DNA, used to correct for allelic differences in extraction or PCR efficiency, was measured in duplicate for the indirect (Fig. 3A) and direct (Fig. 3B) parental comparisons. Genomic DNA from the F₁ hybrids (Fig. 3C) was measured once per biological replicate, and measurements were pooled across all four biological replicates for analysis purposes. We took this approach because there is no concern about disproportional representation of the two alleles within F₁ hybrids—genomic DNA from each hybrid is expected to contain the same number of *D. novamexicana* and *D. americana* alleles. Pyrograms were examined by eye to remove failed or poor-quality reactions. Ratios of allelic abundance were calculated as described in Methods in supporting information.

Statistical analyses

Pyrosequencing expression data were analyzed using PROC MIXED in SAS v9.2 (Cary, NC, USA), with models fitted using restricted maximum likelihood. The “indirect” dataset (Fig. 3A) was fitted with the following model to test for differences between species and over time:

$$Y_{ijkl} = S_i + T_j + ST_{ij} + PST_{ijk} + e_{ijkl}$$

where Y_{ijkl} is the \log_2 of the ratio of *D. americana* to *D. novamexicana* mRNA abundance, corrected using genomic DNA controls. S_i and T_j are fixed effects of species and time point (developmental stages P12–A1), respectively. ST_{ij} is a fixed-effect interaction of species and time; PST_{ijk} is the random effect of biological replicate (P) nested within species and developmental time point; and e_{ijkl} is the residual error.

The “direct” and “F₁” datasets (Fig. 3B and C) were fitted to the same set of variables, minus the effect of species. (For these two datasets a single ratio comparing the two species was obtained, whereas in the “indirect” comparison, the two species were measured separately.) The model used was:

$$Y_{ijk} = T_i + PT_{ij} + e_{ijk}$$

with variable names as above. Least-squares means and 95% confidence intervals of the log-transformed gene expression ratios were calculated and used to plot Figs. 4, 5, and S1. For the “direct” and “F₁” datasets, one-sample t -tests were performed within PROC MIXED to test the least-squares means for a significant difference from zero; that is, for significant difference in *D. americana* versus *D. novamexicana* allele abundance. For the “indirect” dataset, the *D. americana* and *D. novamexicana* data were compared at each stage using two-sample t -tests, again within PROC MIXED.

Acknowledgments

The authors thank U. Rosas for developing pyrosequencing primers to *tan* and *ebony*, E. Walker for assistance with fly collection, and members of the Wittkopp lab for feedback on the manuscript. This work was supported by grants from the National Science Foundation [DEB-0640485] and National Institutes of Health [5-R01-GM-089736-03] to P.J.W. and an NIH Ruth Kirschstein Award [1 F32 GM087928-01] to A.C. P.J.W. is an Alfred P. Sloan Research Fellow.

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SUPPORTING INFORMATION

Additional Supporting Information may be found in the online version of this article:

Table S1. Traits used to mark developmental stages P8–A2.

Table S2. Significance tests for differential expression of *D. americana* versus *D. novamexicana* alleles of *tan* and *ebony*.

Table S3. Primers and sequences used for pyrosequencing estimates of mRNA expression.

Fig. S1. Direct measurements confirm indirect estimates of species expression differences, but provide more precise estimates.

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