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Tinker where the tinkering's good

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Abstract

Do general principles govern the genetic causes of phenotypic evolution? One promising idea is that mutations in *cis*-regulatory regions play a predominant role in phenotypic evolution because they can alter gene activity without causing pleiotropic effects. Recent evidence that revealed the genetic basis of pigmentation pattern evolution in *Drosophila santomea* supports this notion. Multiple mutations that disrupt an abdominal enhancer of the pleiotropic gene *tan* partly explain the reduced pigmentation observed in this species.

Which mutations underlie phenotypic evolution?

Do developmental mechanisms influence genetic evolution? That is, can we predict what types of mutations underlie phenotypic evolution? The simplest model predicts that mutations with the least pleiotropic effects are most likely to contribute to phenotypic variation and divergence. The rationale for this model comes from a synthesis of Fisher's geometrical model of phenotypic evolution [1] with modern developmental biology. Fisher conceived of mutational effects as randomly oriented with respect to phenotypes. Given that phenotypes are complex and multidimensional, he assumed that mutations affect many traits at the same time. Fisher's model thus implies that most mutations generate pleiotropic effects. Such pleiotropic mutations will typically degrade fitness, because it is difficult to improve fitness if each mutation simultaneously affects many aspects of a phenotype. Modern developmental biology has provided an escape from this conundrum by demonstrating that *cis*-regulatory DNA is often functionally modular: genetically separable elements are responsible for discrete phases and patterns of expression [2–5]. Thus, although nearly all genes perform pleiotropic roles in development, *cis*-regulatory regions ensure that not all mutations have pleiotropic effects [3]. Accordingly, evolution is expected to favor mutations that cause fewer pleiotropic effects [6, 7], such as mutations in *cis*-regulatory regions. Is this model correct? The decisive data—the mutations themselves—have been hard to identify. Although mapping natural variation to chromosomal regions is now easy, the final step of identifying the causal genes and mutations remains a massive challenge. Now Jeong *et al.* [8] have added an important piece to the puzzle by demonstrating that the evolution of *Drosophila santomea* pigmentation patterning has been driven by mutations at a *cis*-regulatory element of the pleiotropic enzyme *tan*, not once, but three separate times (Figure 1).

Pale abdomens lack *tan*

Drosophila abdominal pigmentation patterns vary among species. *D. santomea*, endemic to São Tomé, an island off the west coast of Africa, lost pigmentation in the posterior abdominal segments since divergence from their darkly pigmented sister species *D. yakuba*. Linkage mapping studies implicated a locus of large effect on the X chromosome as one of four loci contributing to the pigmentation difference between these species [9]. The linkage peak falls

near the candidate gene *tan*, an enzyme required for pigmentation as well as for vision [10]. In *D. yakuba*, *tan* mRNA is expressed in the abdominal epidermal cells that produce darkly pigmented cuticle. In *D. santomea*, *tan* mRNA is undetectable in the epidermal cells. In both species, however, *tan* is expressed in the cells that produce pigmented bristle cells. Thus, the absence of *tan* expression in epidermal cells likely resulted from altered regulation of *tan* and not a complete loss-of-function *tan* allele.

Jeong *et al.* [8] performed simple crosses to demonstrate that the absence of *tan* expression in *D. santomea* results from X-linked factors and not from the evolution of *transacting* factors located on other chromosomes. Together with an earlier study [9] showing tight linkage of pigmentation pattern with *tan*, these results suggested that the change in *tan* expression patterns might have resulted from a change in the *tan cis*-regulatory region. As this *cis*-regulatory region was previously uncharacterized, Jeong *et al.* [8] made a series of reporter constructs using *D. melanogaster* DNA and found a *cis*-regulatory region 3–4 kb upstream from the transcription start site, in the intergenic region between two other genes, that drives expression in the same abdominal pattern observed for endogenous *D. melanogaster tan*.

Jeong *et al.* [8] found that this *cis*-regulatory region could drive *tan* cDNA expression and rescue abdominal pigmentation in *D. melanogaster tan* mutants. This transgene also partially restored pigmentation in *D. santomea*, implying that altered *tan* activity is indeed a major cause of the reduced pigmentation in *D. santomea* abdomens.

Additional observations indicate that changes in this *cis*-regulatory region have caused pigmentation evolution. First, the *D. yakuba* and *D. santomea tan* genes encode identical proteins, ruling out any effects caused by coding changes. Second, the *D. yakuba* abdominal *cis*-regulatory element drives *tan* expression in the posterior abdomen, whereas the same region from *D. santomea* fails to do so. Jeong *et al.* [8] dissected this regulatory element further and identified two single nucleotide substitutions that together cause the loss of enhancer activity.

Thus, mutations that alter *tan* abdominal expression, without affecting the gene's other roles, underlie a portion of the pigmentation pattern divergence observed in *D. santomea*. Surprisingly, a survey of the *tan* locus from multiple *D. santomea* natural isolates uncovered three separate alleles that independently eliminate *tan* abdominal enhancer activity [8]. A second allele carries a 30-bp deletion in the *tan* abdominal *cis*-regulatory element, and a third harbors a 212-bp deletion in the same region. No *tan* alleles with a functional abdominal *cis*-regulatory element were found in the *D. santomea* isolates. Remarkably, the three nonfunctional regulatory sequences arose independently from a functional allele, based on the sequences inferred for their ancestors.

The mutations that cause these three evolutionarily independent alleles all occur within a few bases of one another in the *tan* abdominal *cis*-regulatory element. These results are consistent with a model wherein the pleiotropic roles of *tan* bias evolutionarily relevant mutations toward the abdominal *cis*-regulatory module and away from gene regions that might alter other *tan* functions.

Concluding remarks

These new results provide the latest example of phenotypic evolution via parallel mutations at a single *cis*-regulatory element. In humans, for example, a change in temporal regulation of *lactase* expression has arisen multiple times through mutations in a small *cis*-regulatory element [11]. In this case, expression later in development has been added without altering the structure of the enzyme or its early expression.

The *cis*-regulatory hypothesis for evolutionary change does not require that all genetic changes occur at *cis*-regulatory sites. Rather, it predicts that natural selection will often favor the least pleiotropic route to phenotypic evolution. When evolution requires spatially or temporally regulated gene expression, the least pleiotropic mutations are likely to be found in *cis*-regulatory regions. At other times, natural selection will favor a change in protein sequence. By example, when host immunity gene products physically interact with pathogens we expect that protein sequence evolution will occur in response to changes in pathogen populations [12]. However, even in such cases, pathogen attack can sometimes be foiled by specifically altering the regulation of a target protein. In human populations from West Africa, a *cis*-regulatory mutation eliminates *DARC* (*Duffy blood group, chemokine receptor*) expression in red blood cells, and blocks malaria infection, without altering *DARC* expression in other cells types [13]. Of course, evolution will always use the available molecular variation. However, given the choice, natural selection will favor mutations that cause the fewest pleiotropic effects.

The new work from Jeong *et al.* [8] on *tan* evolution provides yet one more clearly documented case for the importance of *cis*-regulatory evolution. Such examples have been slow in coming, largely because it is more technically demanding to identify functional *cis*-regulatory changes than functional changes in proteins. As the field develops new approaches to fine mapping natural variation, we expect to gain deeper understanding of the rules that influence when evolution favors *cis*-regulatory mutations. For now, the basic conceptual model remains healthy. Evolution tinkers where the tinkering is good. For traits that require regulated gene expression, *cis*-regulatory DNA is the place.

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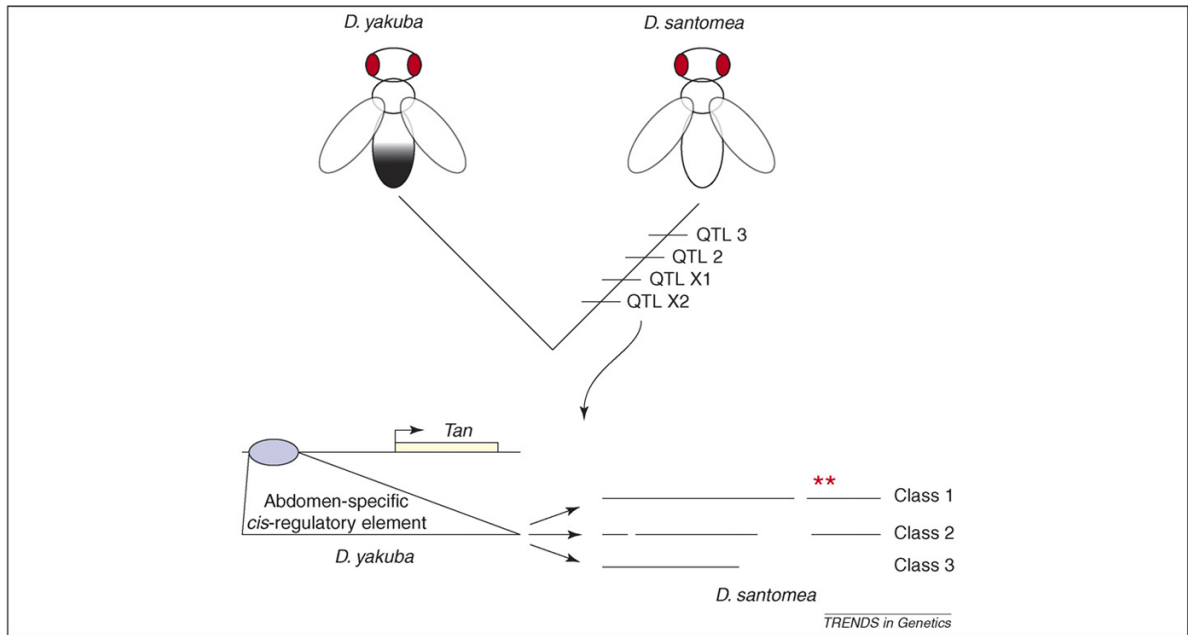


Figure 1.

Genetic causes of abdominal pigmentation evolution. The two closely related species, *Drosophila yakuba* and *Drosophila santomea*, shown schematically at the top, display different levels of abdominal pigmentation. This difference results from changes in at least four quantitative trait loci (QTLs) that all occurred on the lineage leading to *D. santomea* [9]. Two QTLs reside on the X chromosome (QTL X1 and QTL X2), one on chromosome II (QTL 2) and one on chromosome III (QTL 3). The order of evolved QTLs is shown for illustration purposes only. The true order of QTL evolution is unknown. Jeong *et al.* [8] investigated the genetic changes underlying QTL X2. They found that changes in an abdomen-specific cis-regulatory element of the *tan* gene, shown below, had evolved in *D. santomea*. They identified three classes of alleles carrying different mutations that all incapacitate the cis-regulatory element. The cis-regulatory enhancer sequences are represented as horizontal lines at the bottom. Point mutations that alter function are shown as asterisks above the *D. santomea* sequences. Deletions are shown as gaps in the sequence.