Acquisition of a Leucine Zipper Motif as a Mechanism of Antimorphy for an Allele of the 
Drosophila Hox Gene Sex Combs Reduced

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ABSTRACT
In 1932, Müller first used the term "antimorphic" to describe mutant alleles that have an effect that is antagonistic to that of the wild-type allele from which they were derived. In a previous characterization of mutant alleles of the Drosophila melanogaster Hox gene, Sex combs reduced (Scr), we identified the missense, antimorphic allele Scr14, which is a Ser10-to-Leu change in the N-terminally located, bilateranspecific octapeptide motif. Here we propose that the cause of Scr14 antimorphy is the acquisition of a leucine zipper oligomerization motif spanning the octapeptide motif and adjacently located protostome-specific LASCY motif. Analysis of the primary and predicted secondary structures of the SCR N-terminus suggests that while the SCR+ encodes a short, α-helical region containing one putative heptad repeat, the same region in SCR14 encodes a longer, α-helical region containing two putative heptad repeats. In addition, in vitro cross-linking assays demonstrated strong oligomerization of SCR14 but not SCR+. For in vivo sex comb formation, we observed reciprocal inhibition of endogenous SCR+ and SCR14 activity by ectopic expression of truncated SCR14 and SCR+ peptides, respectively. The acquisition of an oligomerization domain in SCR14 presents a novel mechanism of antimorphy relative to the dominant negative mechanism, which maintains oligomerization between the wild-type and mutant protein subunits.

KEYWORDS
Homeotic selector genes body patterning Sex combs reduced evolutionarily conserved protein motifs leucine zipper interaction interface antimorph dominant negative

Body patterning in bilaterans requires the Homeotic selector genes (Hox) (Lewis 1978; McGinnis and Krumlauf 1992; Carroll 1995). In Drosophila melanogaster, Hox genes are expressed in spatially restricted domains along the anterior–posterior axis of the embryo and function as sequence-specific, DNA-binding transcription factors to establish the unique identity of each body segment. One focus in the study of Hox genes is to understand how the evolution of Hox regulation of development has contributed to the morphological variation observed in Bilateria (Carroll et al. 2005). Many previous studies have examined the contribution of cis-regulatory motifs to changes in Hox function; however, our understanding of HOX protein structure and functional contribution of highly conserved motifs and domains to HOX activity is limited. Functionally important regions are often synonymous with highly conserved proteins domains, which are the products of purifying selection; therefore, these regions offer a starting point for the dissection of protein function. In the study of Drosophila HOX protein structure, the greatest emphasis has been placed on identifying the developmental roles of two HOX protein motifs found in all Bilaterans: the homeodomain (HD) and the YPWM motif (Zhao et al. 1996; Curtis et al. 2001; Galant et al. 2002; Hittinger et al. 2005). Surprisingly, although both of these regions are important for HOX function, the only essential domain is the DNA-binding HD (Berry and Gehring 2000; Tayyab et al. 2004; Joshi et al. 2007). An explanation for this finding is the observation of differential pleiotropy: the additive, context-dependent function of HOX protein motifs and domains (Hittinger et al. 2005; Prince et al. 2008; Sivanantharajah and Percival-Smith 2009; Merabet et al. 2011; Percival-Smith et al. 2013).
Here, we are interested in identifying the regions that have been functionally conserved in the HOX protein, Sex comb reduced (SCR).

SCR has 10 protein motifs and domains conserved at different taxonomical levels: bilateran-specific (octapeptide, YPWM, HD, and KMAS); protostome-specific or arthropod-specific (LASCY, SCKY, PQDL, and NANGE); and insect-specific (DTYQL and C-terminal domain) (Percival-Smith et al. 2013). During development, SCR establishes the identity of the labial and prothoracic segments. In the labial segment, SCR function is required for development of the proboscis, which is the adult feeding tube, and the larval salivary glands; however, in the prothoracic segment, SCR is required for establishing the identity of the prothoracic legs, which are characterized by the presence of sex comb on the fifth tarsal segment (Lewis et al. 1980; Struhl 1982; Panzer et al. 1992; Percival-Smith et al. 1997). The previously identified missense, antimorphic allele, Scr14, is a Ser10 to Leu substitution of the most conserved residue in the SSYF submotif of the octapeptide motif (Tour et al. 2005; Sivanantharajah and Percival-Smith 2009). The analysis of the interaction between Scr14 and Scr4 and the interaction between Scr14 and null alleles led to the classification of this allele as an antimorph, which is a class of mutant alleles that are antagonistic to the wild-type allele (Müller 1932; Sivanantharajah and Percival-Smith 2009). The subsequent interpretation of this genetic evidence led to the proposal that the octapeptide motif of SCR may mediate dimerization between SCR N-termini (Sivanantharajah and Percival-Smith 2009).

A common interaction motif found in the bZIP and the bHLH-ZIP classes of transcription factors is the leucine zipper motif (Vinson et al. 1989; Murre et al. 1989). These proteins, when bound to DNA, are long α-helices with N-terminal DNA binding domains and C-terminal dimerization domains that form a leucine zipper coiled coil (Landschultz et al. 1988; Vinson et al. 1989; Ellenberger et al. 1992). The leucine zipper regions in these transcription factors typically comprise four or more heptad repeats; however, there is growing evidence that shorter and moderately stable leucine zipper motifs, consisting of only two heptad repeats, can function independently as oligomerization domains (Burkhard et al. 2000; Ferera et al. 2001; Nikolaev and Pervushin 2009). The leucine zipper heptad contains seven amino acids with a standard notation of (a b c d e f g)n (Krylov et al. 1994). Coiled coil interactions promote oligomerization between the leucine zipper motifs of one protein and its partner, and the partner’s heptad repeats are designated (a’ b’ c’ d’ e’ f’ g’).n. The structure and biochemical properties of the coiled coil structure formed by the interaction of two leucine zipper motifs are dependent on the intermolecular interactions that occur between the hydrophobic, stability-conferring residues in positions a-a’ and d-d’, and the interaction between charged, specificity-conferring residues in positions g-e’ (Krylov et al. 1994). Because the amino acid at position g of one heptad interacts with the amino acid occupying position e’ in the adjacent heptad of the dimerization partner, the designation (a b c d e f g)n accounts for all relevant intermolecular interactions. Here we report that the antimorphic–hypermorphic Scr4 allele may be the result of the acquisition of a short, functional leucine zipper motif spanning two evolutionarily conserved motifs in the SCR N-terminus: the bilateran-specific octapeptide motif and the adjacentally located, protostome-specific LASY motif.

**MATERIALS AND METHODS**

**Ab initio method for prediction of secondary structure of the SCR** and SCR14 N-termini

The Protein Data Bank (PDB) was searched with two queries: the wild-type sequence of the SCR N-terminus (FAMSLYQFVNSLACYCYPQMN) and the sequence of SCR14 (FAMSLYQFVNSLACYCYPQMN). All peptide sequence matches were assessed using the following criteria: peptide sequences of six or more amino acids in length were used, except if a sequence of five amino acids had perfect identity and all matches with insertions or deletions were ignored; to improve the accuracy of identifying homologous sequences without perfect identity, the BLOSUM62 substitution matrix (Henikoff and Henikoff 1992) was applied. Matches with amino acid substitution scores that equaled zero or above were used. If two substitutions were present, then homologous peptide sequences were used only if the substitutions had positive scores; matches with the same peptide sequences were used if the proteins were from different protein families. Once a match was identified, Cn3D (NCBI) was used to identify the secondary (2δ) structure encoded by a protein sequence (i.e., α-helix, β-strand, coil, turn) within a protein with solved 3D structure.

**Expression and affinity purification of SCR peptides from Esherichia coli**

For biochemical analyses, Esherichia coli expression constructs were made for the purification of triple-tagged (TT) (composed of 6xHis, Strep, and Flag tags) (Tiefenbach et al. 2010), 29-amino-acid-long SCR peptides, SCR* (SCR8), SCR14 (SCR8), SCR14, and SCR27, by inserting the relevant coding sequence into the multiple cloning site of pET-3a (Studier and Moffatt 1986; Studier et al. 1990). All constructs were transformed into BL21(DE3) cells (Studier and Moffatt 1986) and protein expression was induced using the Novagen toxic cells protocol. Tagged SCR peptides were purified from E. coli using Strep-Tactin Sepharose (IBA) according to the manufacturer’s protocol.

**Protein cross-linking and Western analysis**

Purified proteins were incubated without and with (1%) the protein cross-linker, formaldehyde, for 10 min at room temperature. Samples were heated to 65°C in SDS buffer for 8 min and run on 12%–15% SDS PAGE. Immobilon-P transfer membrane (Millipore) was used for Western blotting. The anti-FLAG mouse monoclonal antibody (Sigma) was diluted 1:40,000. Antibody–antigen complexes were detected with anti-mouse HRP-conjugated secondary antibody (Sigma) and the SuperSignal West Femto Chemiluminescent kit (Pierce). Images were collected on a Fluorchem 8900 gel documentation system (Alpha Innotech).

**Transgenic flies**

Transgenic lines were created to express full-length and truncated SCR proteins fused with a C-terminal triple-tag using the GAL4-UAS system (Brand and Perrimon 1993). All constructs were made and independently incorporated into the Drosophila genome using P-element–mediated transformation (Rubin and Spradling 1982). Full-length (SCR14) and wild-type (SCR*) and the SCR14 sequence and were expressed in all three leg discs using the rotund (rn)–Gal4 (P{GawB}) driver. Truncated SCR peptides encoded the first 121 amino acids of the SCR N-terminus with wild-type sequence, specific deletions, singular or in combination, of the first eight amino acids of SCR (8aa), the octapeptide (octa), and the LASY motif (LASY), and point mutations of Ser10 to Ala. Fly lines of the genotype, y w; P(UAS-Scr14;TT, w+); p[14]TM6B, were established and crossed to either P{Armidillo (Arm)-Gal4, w+}; Scr14;TM6B or P{Arm-Gal4, w+}; Scr14;TM6B driver lines to produce progeny in which UAS constructs were constitutively
expressed in Scrx or Scr14 hemizygotes; these flies have one endogenous copy of Scrx because pb34 is a deficiency encompassing the Scrx locus. Flies of genotype P[Arm-Gal4, w1]; Scrx/pb34 and P[Arm-Gal4, w1]; Scr14/pb34 were used as controls for GAL4 expression. Three genotypes were quantified: the number of sex comb bristles on the T1 leg; the number of pseudotrachea that developed on the proboscis; and the number of nuclei in the salivary glands. A description of the genetic markers and balancer chromosomes used can be found elsewhere (Lindsley and Zimm 1992).

Expression of UAS constructs in vivo
Protein was extracted from third instar larvae of the genotype y w; P[УΑS-Scr14FL121TT, w1]; P[Arm-Gal4, w1]; Scrx/pb34. The FLAG epitope tag was detected with the anti-FLAG antibody. Tubulin was used as a loading control and was detected using anti-β-tubulin E7 mouse monoclonal antibody (Klymkowsky et al. 1987). Scrx and tubulin expression levels were quantified from three independent Western blots using the AlphaEase Fluorchem software (v.4.0.1).

Statistical analyses
All data were analyzed in SPSS v.20.0 (SPSS Inc. 2011). Comparisons of phenotypic data collected in 2013 and 2009 (Sivanantharajah and Percival-Smith 2009) for the mean number of sex comb bristles, pseudotracheal rows, and nuclei in the salivary gland were performed using Student t tests. Analyses of multiple means were performed using ANOVA and Kruskal-Wallace tests and, if significant differences were detected, then multiple pair-wise comparisons were performed using Tukey tests and Dunnett T3 tests, respectively. In characterizing Scrx antimorph in vivo using full-length proteins, data for number of sex comb bristles on the T1 leg were log10-transformed to meet the requirements of homoscedasticity and normality and then analyzed using a one-way ANOVA. The data for number of sex comb bristles on the T2 and T3 legs were analyzed using Student t tests. In characterizing Scrx antimorph in vivo using truncated peptides, all salivary gland data were analyzed using one-way ANOVA. All data for number of sex comb bristles in Scrx hemizygotes and pseudotrachea in Scrx hemizygotes were analyzed using one-way ANOVA; the latter data set was log10-transformed first. All data for number of sex comb bristles in Scrx hemizygotes and pseudotrachea in Scrx hemizygotes were analyzed using Kruskal-Wallace tests. In mapping the region of Scrx required for antimorph, sex comb bristle and salivary gland data were analyzed using one-way ANOVA. UAS-Scrx peptide expression data were square root-transformed and analyzed using a one-way ANOVA.

RESULTS

Scr14 is an antimorphic–hypomorphic allele
Scrx14 was initially classified as an antimorphic allele based on the observation that Scr14/Scrx14 had the same phenotype as Scrx14/pb34 or Scrx14/pb34. In addition, the observation that Scrx14/pb34 and Scrx14/pb34 had the same phenotype was the major observation for the proposal that Scrx and Scrx14 formed an inactive locked complex in vivo (Sivanantharajah and Percival-Smith 2009). Reexamination of the Scrx14 allele on Scrx-dependent phenotypes in this study has revealed a change in allelic behavior. When the phenotypes observed now were compared to the phenotypes observed in 2009, the mean number of sex comb bristles in Scrx14 hemizygotes decreased from 7 to 2.5 (t(64)=14.6; P < 0.001), the mean number of pseudotrachea decreased from 5.4 to 3.0 [F(1,12)=6.9; P < 0.001], and the mean number of nuclei in the salivary glands decreased from 114.8 to 100.5 [F(1,30)=3.2; P = 0.030] (Table 1). We sequenced the Scrx14 allele over the deficiency pb34 at three different times independently: in 2007 for initial sequenc- ing analysis, in 2009 to confirm the genotype of Scrx14/pb34 flies that had a wild-type phenotype, and in 2013 (Figure 1). The missense change, causing Ser10 to Leu in the octapeptide motif, and all polymorphisms in exon 2 that were reported in 2007 are still present in the 2013 sequence. Despite the observed differences in Scrx14 phenotype, the Scrx14/Scrx14 hemizygote still had the lowest number of sex comb bristles of all other viable Scrx hypomorphic alleles with Scrx14 (Table 1) (Sivanantharajah and Percival-Smith 2009). If no interaction occurred between these two alleles, then one would expect the sum total of their individual activity to produce 8.7 sex comb bristles and not 7.0 (Table 1). This indicates that Scrx14 inhibits Scrx activity and is therefore antimorphic. Second, we observed an inhibitory interaction between Scrx14 and Scrx13A. Scrx13A is a null allele encoding a truncated protein product of 260 amino acids, where the HD and CTD have been deleted but the N-terminal region, encoding the octapeptide and LASCY motifs, is still present. If no interaction occurred between Scrx13A and Scrx14, then one would expect the sum total of their individual activity to be the same as Scrx14 alone (Scr14/pb34) because Scrx13A is a null allele that should be functionally equivalent to pb34; however, Scrx14/Scrx13A heterozygotes showed a significant decrease in sex comb bristles when compared to Scrx14/pb34 hemizygotes of 2.5 to 1.9 [F(2)=3.6; P = 0.001] (Table 1 in italics). Finally, overexpression of Scr14FL1TT and Scr14FL1TT proteins using the rn-Gal4 driver and UAS fusion genes in the T1 segment had a significant effect [F(2)=278; P < 0.001] (Figure 1G). In control flies, when GAL4 protein is expressed alone in the leg discs, a mean number of 11.3 ± 0.3 sex comb bristles develop on the first thoracic leg (T1). When SCR14FL1TT is expressed, there is a significant increase in the mean number of sex comb bristles on T1 from 11.3 ± 0.3 to 28.5 ± 1.1 (P < 0.001). In contrast, expression of SCR14FL1TT results in a significant decrease in the number of sex comb bristles on the T1 leg from 11.3 ± 0.3 to 7.9 ± 0.4 (P < 0.001). This indicates that SCR14FL1TT expression in T1 inhibits endogenous SCR activity, supporting the classification of Scrx14 as an antimorphic allele.

Table 1: Comparison of Scrx phenotypes observed in this study and in a previous study (+SEM)

| Genotype     | Mean Sex Comb Bristles, No. | Mean Pseudotrachea, No. | Mean Cells in Salivary Gland, No. |
|--------------|------------------------------|--------------------------|-----------------------------------|
|              | 2013                         | 20091                    | 2013                              | 2009 |
| Scrx14/pb34  | 6.2 ± 0.1                    | 6.3 ± 0.2                | 5.3 ± 0.1                         | 5.5 ± 0.1* |
| Scrx14/pb34  | 2.5 ± 0.1                    | 7.0 ± 0.3*               | 3.0 ± 0.1                         | 5.4 ± 0.3* |
| Scrx14/Scrx13A| 7.0 ± 0.1                    | 6.9 ± 0.2                | 6.0 ± 0.1                         | 5.5 ± 0.1* |
| Scrx14/Scrx13A| 1.9 ± 0.1                    | 2.6 ± 0.1*               | 4.8 ± 0.2                         | 5.2 ± 0.1* |
| Scrx14/Scrx14 | 7.1 ± 0.2                    | 6.3 ± 0.2*               | 6.0 ± 0.0                         | 5.9 ± 0.1* |
|              |                              |                          | 108.4 ± 3.1                       | 126.0 ± 5.9* |

1 Values taken from Sivanantharajah and Percival-Smith (2009).

* Significant differences for a particular phenotype examined in 2009 from values collected in 2013 (P < 0.05).
Two observations show that Scr14 is also hypomorphic. When compared to the Scr+ hemizygotes, Scr14 flies show a decreased number of rows of pseudotrachea and sex comb bristles (Figure 1, A–D). Second, relative to full-length SCR+FLTT, expression of full-length SCR14FLTT in the T2 and T3 segments with the rn-Gal4 driver had a decreased capacity to induce ectopic sex comb bristles (Figure 1G). In control flies, when GAL4 protein is expressed alone, no sex comb bristles develop on the T2 and T3 legs. When SCR14FLTT is expressed, there is an increase to 23.1 ± 2.7 sex comb bristles on T2 and to 16.5 ± 3.1 bristles on T3. Compared to SCR+FLTT, there is an increase to 23.1 ± 2.7 sex comb bristles on T2 and to 16.5 ± 3.1 bristles on T3. Compared to SCR+FLTT, there is an increase to 23.1 ± 2.7 sex comb bristles on T2 and to 16.5 ± 3.1 bristles on T3. Compared to SCR14FLTT, there is a significant decrease in ectopic sex comb bristles to 2.1 ± 0.3 on T2 [t(13)=7.9; P < 0.001] and to 2.23 ± 0.4 on T3 [t(16)=4.6; P < 0.001]. Relative to SCR+, SCR14 has reduced function, which is the definition of hypomorphy. All evidence together suggests that Scr14 in 2013 is better classified as an antimorphic–hypomorphic allele. The locked complex model previously proposed is no longer applicable because it was dependent on SCR+ and SCR14 having wild-type activity when alone but forming inactive heterodimers when together (Sivanantharajah and Percival-Smith 2009).

The octapeptide and LASCY motifs may comprise a putative leucine zipper motif in SCR14

An examination of the SCR14 N-terminus, encoding the octapeptide motif and LASCY motifs, revealed the presence of a putative leucine zipper motif based on three observations. First, we identified two putative heptad repeats in the SCR14 N-terminus; the same region in SCR+ encodes a single putative heptad repeat. A sequence logo compiled from leucine zipper motifs annotated in 27 Drosophila bZIP

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**Figure 1** Proboscis and sex comb phenotypes of the antimorphic–hypomorphic allele, Scr14. Scr14 is a missense allele in the highly conserved, octapeptide motif. (A) Representations of the SCR proteins encoded by Scr+ and Scr14 alleles are shown. Scanning electron micrographs of the adult labial palps and of the fifth tarsal segment of the adult prothoracic leg of flies hemizygous for Scr+ (B, C) and Scr14 (D, E). When compared to the Scr+ hemizygotes, Scr14 flies show a decreased number of rows of pseudotrachea and sex comb bristles, indicating that Scr14 is a hypomorphic allele. The arrows indicate the position of the sex comb. Images of Scr+ flies were taken from Sivanantharajah and Percival-Smith (2009). (F) Electropherogram sections illustrating the missense (C29→T) and five silent mutations (A171→G, C190→T, A345→G, C747→T, and A933→G) initially identified in exon 2 of Scr14. (G) Expression of full-length SCR+FLTT and SCR14FLTT proteins in the three legs (T1–T3) demonstrates that SCR14 is both antimorphic and hypomorphic. The numbers of sex comb bristles that form on the three legs are shown for expression from the m-Gal4 driver of GAL4 alone (control), SCR+FLTT, and SCR14FLTT. P values listed above black lines indicate the results of specific pair-wise comparisons.

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**Table**

| Mean number of sex comb bristles + SEM | T1 | T2 | T3 | T1 | T2 | T3 |
|--------------------------------------|----|----|----|----|----|----|
| Control                              |    |    |    |    |    |    |
| SCR+FLTT                             |    |    |    |    |    |    |
| SCR14FLTT                            |    |    |    |    |    |    |

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proteins identified by Fassler et al. (2002) revealed a general consensus sequence of E\_gL\_a\_Le\_b\_A\_c\_L\_d\_R\_e\_f\_g for a Drosophila heptad (Figure 2A) (Schneider and Stephens 1990; Fassler et al. 2002; Crooks et al. 2004). The two heptad repeats found in SCR\^14 both contain hydrophobic residues at positions a and d; however, charged residues are not present at residues e and g. When compared to SCR\^14, the sequence of the same region in SCR\(^{+}\) encodes a single putative heptad repeat spanning part of the octapeptide and the LASCY motifs (Figure 2A).

A second property of leucine zipper motifs is their amphipathic \(\alpha\)-helical structure. The analysis of the SCR N-terminus using \textit{ab initio} protein \textit{2} prediction software identified a \(\alpha\)-helix in the

![Figure 2](image-url)

**Figure 2** The N-terminus of SCR\(^{+}\) satisfies the criteria for a leucine zipper motif. (A) General consensus of a heptad repeat of a leucine zipper motif in Drosophila. The sequence logo was made from the sequences of 130 heptad sequences from 27 Drosophila bZIP proteins that were identified and aligned in the study by Fassler et al. (2002). Amino acids have been color-coded according to their biochemical properties. Basic residues are green (K, R, H), acidic residues are red (D, E), hydrophobic residues are blue (G, A, V, L, I, M, P, F, W), and hydrophilic residues are orange (S, T, N, Q, Y, C). The x-axis displays the general consensus for a heptad repeat (g a b c d e f g) of a Drosophila bZIP protein, which is ELAEQLRQ. The amino acids of the heptad are displayed from N-terminal (N) to C-terminal (C). The y-axis is a measure of uncertainty (bits). The height of a stack indicates the degree of sequence conservation at that position, and the height of an amino acid within a stack indicates the relative frequency of that residue at that position. Below the logo are the sequences of the two heptad repeats found in SCR\(^{14}\). The sequence logo was created using WebLogo 3 (v.2.8.2). (B) The N-termini of SCR\(^{+}\) and SCR\(^{14}\) are predicted to have distinct secondary structures. SCR\(^{14}\) is predicted to encode a longer \(\alpha\)-helical region than SCR\(^{+}\). (C) Helical wheel plot of the SCR\(^{14}\) N-terminus demonstrating the capacity for formation of an amphipathic helix. The circles of the wheel are labeled with letters a through g, designating the standard nomenclature of a heptad repeat of a coiled coil. The diagram illustrates hydrophobic amino acids in positions a and d (gray) and a general occupancy of positions b, c, and f (white) by uncharged, polar amino acids. Charged amino acids at g (blue) and e (red) are not present. Figure adapted from Nikolaev and Pervushin (2009). (D) Testing the capacity of SCR peptides to oligomerize. The sequence of the N-terminus of wild-type SCR is shown, highlighting the two putative heptad repeats spanning the octapeptide and LASCY motifs. The sequences are given for SCR peptides with substitutions in position d of one or both heptads, SCR\^L (SCR\(^{14}\) (i), SCR\^L (SCR\(^{+}\) (ii), SCR\^L (iii), and SCR\^L (iv). (E) Oligomerization is strongest with SCR\(^{14}\) peptides. SCR\(^{14}\) (SCR\(^{14}\)) peptides oligomerize strongly on addition of 1% (+) protein cross-linker, whereas SCR\(^{+}\) (SCR\(^{+}\)) peptides do not. (F) The ability of SCR peptides to oligomerize is dependent on the presence of two leucine residues. SCR peptides with point mutations in two putative heptad repeats were incubated with 1% (+) protein cross-linker. Removal of either or both leucine residues significantly reduced oligomerization. Proteins were detected on a Western blot using anti-FLAG antibody. Relative sizes of SCR\(^{14}\) oligomers, when run on 15% (E) and 12% (F) SDS-PAGE, are indicated in brackets.
region encoded by the octapeptide in both SCR+ and SCR14 using the Garnier-Robson method; however, this α-helix is slightly longer in SCR14 (data not shown; Protein v.4.0.3, DNA Star 1990–1999). The same software predicted α-helical regions in the octapeptide of SCR14, but not SCR+, using the Chou-Fasman method. An alternative ab initio method for predicting the potential 2° protein structure is to search for the primary (1°) amino acid sequence of interest in the PDB of proteins with known 3D structure. This method predicted that the N-termini of SCR+ and SCR14 have distinct biochemical structures; where the SCR+ N-terminal encodes a short, α-helical region spanning part of the octapeptide and LASYC motif, the same region in SCR14 encodes a longer, α-helical region spanning both motifs entirely (Figure 2B; Supporting Information, Figure S1). A helical wheel plot of the SCR14 N-terminus predicts the formation of an amphipathic α-helix with a hydrophobic core formed by leucines at position d and hydrophilic residues at position a, and a hydrophilic surface formed primarily by noncharged, polar residues at positions b, c, and f (Figure 2C).

In vitro analysis of SCR14

A third property of a leucine zipper motif is the capacity for oligomerization. To test the oligomerization capacity of SCR+ and SCR14 peptides, these peptides were expressed and purified from bacteria and incubated with and without protein cross-linker. First, it was noted that SCR+ and SCR14 ran at different relative molecular weights (Mr). Second, the addition of protein cross-linker detected strong oligomerization of SCR+ peptides but not SCR+ peptides (Figure 2D). To identify the amino acids required for SCR14 oligomerization, modified SCR peptides were constructed with substitutions at the leucine residue present in each of the two putative heptad repeats of SCR+ (Figure 2C). Oligomerization of SCR+ peptides was dependent on the presence of two leucine residues, because removal of either or both leucines of the heptad repeats reduced oligomerization (Figure 2E). In addition, the Mr of modified SCR peptides was dependent on the number of serine residues present; there was an increase in Mr as leucines were substituted with serine residues.

In vivo analysis of Scr+14 antimorphy testing reciprocity

In sex comb formation, SCR+14 inhibited SCR+ activity and the truncated SCR+13A protein inhibited SCR14 activity (Table 1). To determine if the inhibition of SCR+ by SCR+14 was reciprocal, truncated SCR+121TT and SCR+14121TT peptides were expressed in SCR+ and SCR14 hemizygotes using the GAL4-UAS system. If the inhibition is reciprocal, then SCR+14121TT is expected to inhibit endogenous SCR+ activity and SCR+121TT is expected to inhibit endogenous SCR14 activity, as observed with SCR+13A. In the prothoracic segment, there was a significant effect of expressing SCR+121TT peptides in SCR+ hemizygotes [F(2)=8.4; P < 0.001] (Table 2). As expected, expression of SCR+121TT peptides significantly reduced the mean number of sex comb bristles from 6.4 to 5.5 (P < 0.001). A significant effect was also noted with expression of SCR+ peptides in the prothoracic segment of SCR14 hemizygotes [F(2)=5.7; P < 0.001]. As expected, expression of SCR+121TT peptides caused a significant decrease in the number of sex comb bristles from 3.0 to 1.3 (P < 0.001), and expression of SCR14 peptides also caused a significant decrease from 3.0 to 1.9 (P < 0.001).

We also examined two other SCR-dependent phenotypes. In the proboscis, no effect was observed with expression of SCR121TT peptides in SCR+ hemizygotes [F(2)=5.7; P = 0.06]; however, there was a significant effect with expression of SCR peptides in SCR14 hemizygotes [F(2)=11.9; P < 0.001]. Expression of SCR14121TT peptides caused a significant increase in the number of rows of pseudotrachea from 3.1 to 3.6 (P = 0.01). In the salivary glands, there was a significant effect of expressing SCR peptides in SCR+ hemizygotes [F(2)=35.11; P < 0.001]. Expression of SCR+121TT peptides caused a significant decrease in the mean number of nuclei in the salivary gland from 113.3 to 91.9 (P < 0.001). A significant effect was also noted on expression of SCR peptides in the salivary glands of SCR14 hemizygotes [F(2)=19.6; P < 0.001]. Expression of SCR+121TT peptides caused a significant decrease in the mean number of nuclei from 100.7 to 88.8 (P < 0.001).

### Table 2 Tissue-specific inhibition of endogenous Scr activity by ectopic expression of SCR+121TT or SCR14121TT polypeptides in flies hemizygous for Scr+ or Scr14 (±SEM)

| UAS-X          | Mean Sex Comb Bristles, No. | Mean Pseudotrachea, No. | Mean Cells in Salivary Gland, No. |
|----------------|-----------------------------|-------------------------|----------------------------------|
|                | Scr+                        | Scr14                   | Scr+                             | Scr14                        |
| None           | 6.4 ± 0.1a                  | 3.0 ± 0.1a              | 113.3 ± 1.7a                     | 100.7 ± 1.8ab                |
| Scr+121TT      | 6.0 ± 0.2a                  | 1.3 ± 0.2b              | 114.5 ± 2.5a                     | 105.0 ± 2.0ab                |
| Scr14121TT     | 5.5 ± 0.1b                  | 1.9 ± 0.1c              | 91.9 ± 2.2bc                     | 88.8 ± 1.9b                 |

Data in the same column with the same letters are not significantly different (P < 0.05).
SCR14 activity was affected by expression of SCR14121TT peptides indicated by a weak yet significant decrease in the mean number of nuclei in a salivary gland from 96.4 to 84.3 (P = 0.04).

**DISCUSSION**

**Molecular mechanism of Scr14 antimorphy**

The term "antimorphic" was coined by Müller to describe mutant genes that have an effect that is antagonistic to that of the wild-type gene from which they were derived by mutation (Müller 1932). Here we test a possible mechanism to explain the antimorphic nature of an allele of Scr. An ab initio survey of the N-terminal region of SCR14 identified a putative leucine zipper motif spanning the octapeptide and LASCY motifs presenting a potential explanation for the observed antimorphy. To test this idea, we analyzed the 1° and predicted 2° structures of SCR+ and SCR14 N-termini and found that SCR14 is predicted to form an amphipathic α-helix. Next, in vitro cross-linking assays demonstrated that SCR14 peptides have a strong propensity to oligomerize that is lacking in SCR+ peptides and are dependent on the presence of leucine residues at position d of the heptads. The formation of higher-order oligomers by SCR14 peptides, rather than dimers, was not surprising given the absence of charged residues at positions e and g of each heptad, which are required for dimer specificity (Fassler et al. 2002).

In vivo experiments to characterize the properties of the Scr14 leucine zipper motif found that inhibition is reciprocal, because SCR14121TT peptides inhibit SCR+ activity and SCR+121TT peptides inhibit SCR14 activity. However, the pattern of reciprocal inhibition was observed only in sex comb formation and not in the other tissues tested, the salivary glands, and proboscis. In the salivary glands only inhibition of SCR+ by ectopic SCR14121TT expression is observed. This result may be a consequence of haploinsufficiency of the Scr locus for sex comb but not salivary gland development (Sivanantharajah and Percival-Smith 2009). In the proboscis, ectopic expression of SCR14121TT resulted in an increase in the number of rows of pseudotrachea. A recent study suggests that SCR has two distinct activities that determine the labial and prothoracic segmental identities; therefore, this positive interaction may reflect a distinct effect of SCR14 on SCR labial activity compared with SCR prothoracic activity (Percival-Smith et al. 2013).

Closer examination of the SCR N-terminus to pinpoint the minimal region required for inhibition of SCR14 found that inhibition requires the presence of heptad 2, which spans part of the octapeptide and the LASCY motif. There were two results that were not consistent with experimental expectations. First, we did not expect to see inhibition of SCR14 activity by SCR ASCYP121TT. However, this is likely because the five-amino-acid deletion in SCR ASCYP121TT fails to remove the second heptad motif; rather, the original heptad is replaced with another, which is also predicted to be α-helical in structure (Figure S2). Second, we did not see inhibition of SCR14 activity by SCRSer10→Ala121TT, though this peptide encodes a hydrophobic residue at position d of heptad 1. Is it possible that the formation of asymmetric d-d pairs between Leu and Ala residues affects dimerization stability in some way.

**Scr14 presents a novel mechanism of antimorphy**

A classic mechanistic explanation of antimorphy is the dominant negative, where inactive mutant protein subunits retain the capacity to form complexes with wild-type proteins, resulting in inactive protein complexes (Herskowitz 1987). There are a number of examples of leucine zipper encoding proteins behaving in a dominant negative manner. For instance, truncated versions of the developmentally important Arabidopsis protein ERECTA, a leucine-repeat–rich receptor-
like Ser/Thr kinase (LRR-RTK), encoding the LRR were able to inhibit normal ERECTA function in a dominant negative fashion (Shpak et al. 2003). The introduction of a point mutation into the LRR abolished this dominant negative effect, indicating that a functional leucine zipper motif is essential for the antimorphic activity of truncated ERECTA proteins. In another example, truncated forms of the Drosophila transcription factor Fos, encoding a leucine zipper motif, were able to inhibit wild-type Fos activity (Pierre et al. 2008). Truncated peptides also demonstrated the ability to form novel heterodimers with other leucine zipper encoding proteins. With both ERECTA and Fos, it is presumed that inhibition occurs by forming inactive heterodimeric complexes with normal, and sometimes novel, binding partners (Shpak et al. 2003; Pierre et al. 2008).

When compared to the above examples, Scr14 antimorphy is unique because the mode of antimorphy is not dominant negative. There is no evidence that wild-type SCR proteins have an active leucine zipper motif and form oligomers via the octapeptide motif in vivo. Rather, a missense mutation introduces a leucine zipper motif into the SCR N-terminus, resulting in an acquired capacity for inhibiting endogenous SCR activity. The ability of a single amino acid change in SCR14 to alter SCR activity is supported by the observation that deletions of either the octapeptide motif or the LASCY motif do not decrease SCR activity for inducing ectopic sex comb bristles on the T2 leg as drastically as the SCR14 substitution (Percival-Smith et al. 2013). We have been unsuccessful in a number of assays in detecting hetero-oligomerization between SCR and SCR14 peptides; therefore, there are two possible explanations for the inhibition noted in this study. First, SCR peptides may form interactions with endogenously produced SCR molecules to directly inhibit their activity. Second, SCR14 peptides may interact with a protein other than SCR to inhibit SCR activity.

**SCR N-terminus as a transcriptional activation domain**

Our inability to capture wild-type homodimers of SCR in vitro indicates that the capacity to form strong, detectable dimers is a specific function of SCR14. A recent study found that SCR dimerizes on DNA through the HD (Papadopoulos et al. 2012). It is possible that in a wild-type cell, dimerization on DNA via the HD facilitates interaction between two wild-type SCR N-termini through the shared heptad 2. Such an interaction could be highly dependent on DNA binding for bringing the SCR N-termini into close proximity and for stability.
Given the necessity for protein cross-linker in our in vitro experiments, it is likely that the SCR–SCR interactions observed in this study are transient and unstable. This lack of stability may have hampered our attempts to determine the structure of the SCR+ and SCR14 termini using NMR spectroscopy, which found these regions to be unstructured in vitro (Anthony Percival-Smith, unpublished observations). However, the SCR N-terminus may be structured using NMR spectroscopy, which found these regions to be ordered until bound to a target molecule (Wright and Dyson 1999; Dyson and Wright 2005; Cordier et al. 2006). The lack of intrinsic structure has a number of advantages: increased specificity, because binding of a protein to a target is dependent on an energy-induced transition change, and tight control over protein accumulation, because unstructured proteins are quickly degraded (Dyson and Wright 2005). Structural analyses of a number of different transcriptional activation domains (TAD) have revealed that many TADs are unstructured until bound to a target protein, whereupon they fold into amphipathic α-helices (Zor et al. 2002; Dyson and Wright 2005). A role of the SCR N-terminus in transcriptional activation has been proposed before. Deletion of a 17-amino-acid region, encompassing the octapeptide and LASCY motifs, decreased ectopic transcriptional activation of the SCR targets, forkhead and CrebA, in Drosophila embryos (Tour et al. 2005). Interestingly, a search for potential TADs found that the SCR14 N-terminus encodes a 9aa TAD that is commonly found in a number of yeast and animal transcription factors (Piskacek et al. 2007). This predicted 9aa TAD motif in SCR14 is MSLYQFVNS, which is the sequence of the octapeptide motif. Although a 9aa TAD motif was not found in the SCR+ N-terminus, it is possible that this region may encode a TAD with different biochemical properties.

**SCR N-terminus as an interaction interface**

The SCR14 mutation maps to a universally conserved octapeptide region that has been maintained since the last common bilateran ancestor, whereas the adjacent LASCY motif is a more recently derived sequence and is present in only protostome SCR homologs. The LASCY motif may have been conserved to stabilize protein–protein interactions required for the octapeptide to function as a TAD. The fortuitous identification SCR14 has allowed us to capture a potential function of the SCR N-terminus in mediating protein–protein interactions. Evidence suggesting that the octapeptide and LASCY motifs function as a TAD implies that this region must at least make contacts with components of the transcriptional machinery. In an extension of this idea, the ability of SCR14 to inhibit endogenous SCR+ activity also presents a novel mechanism of SCR regulation by leucine zipper encoding proteins. This suggests that the region of the SCR N-terminus encoding the octapeptide and LASCY motifs may serve an evolutionarily conserved function as a general interaction interface. One class of proteins known to regulate HOX function and also encode leucine zippers are the polycomb group (Pc-G) and trithorax group (Trx-G) classes of chromatin-associated proteins (Stankunas et al. 1998; Crosby et al. 1999; Kal et al. 2000). It remains to be determined whether proteins of either group modulate SCR activity by making intermolecular interactions with the SCR N-terminus.

**The antimorphic–hypomorphic Scr14 allele**

In this study, we reclassify Scr14 as an antimorphic–hypomorphic allele. From the time of initial sequencing of the Scr14 (Sivanantharajah and Percival-Smith 2009) to the time of this publication, there appears to be a significant change in phenotypic behavior of this allele. We initially reported wild-type behavior of Scr14 when hemizygous; however, we now observe that SCR14 has reduced protein activity demonstrating that it is hypomorphic. When we first observed wild-type phenotypes in the proboscis, prothoracic leg, and salivary gland, we sequenced the Scr14 allele (Figure 1F) to ensure that the genotype was correct and found that it was. Our conclusion is that the phenotypic change observed in this study relative to the study by Sivanantharajah and Percival-Smith (2009) is not the result of changes to the Scr14 locus. A possible explanation may be the loss of a phenotypic modifier of Scr activity, which we were not successful at recovering from the stock in 2013, from the original Scr14 stock used for phenotypic analysis in 2009. Despite the potential loss of this hypothetical modifier, the Scr14 allele used here in all our analyses retains its key antimorphic activity.

Scr14 is also a hypomorphic allele, meaning that it has reduced wild-type function. This property allows determination of the tissues in which the octapeptide is normally required and determination of whether this motif is uniformly or differentially required. If each region of SCR were uniformly required in all tissues, then the same allelic series would be expected for each tissue. In a previous characterization of highly conserved motifs of SCR, it was found that the DYTMQ motif, YPWM motif, and CTD were differentially required for three tissues assayed (Sivanantharajah and Percival-Smith 2009). To determine how the octapeptide was required for Scr function, the position of Scr14 was determined in an allelic series with previously characterized hypomorphic alleles of Scr in the study by Sivanantharajah and Percival-Smith (2009). The rank order, from weakest to strongest Scr phenotype, in the salivary glands was Scr14 > Scr15 = Scr12 = Scr8 = Scr5 = Scr7 = Scr8 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7; and in the prothoracic segment, the order was Scr14 > Scr15 = Scr12 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7 = Scr8 = Scr5 = Scr7. The rank order of alleles varied for each of the three tissues examined, indicating a differential requirement of each protein region in each tissue. Particularly important (indicated in bold) was the placement within the order of the allelic series with changes in the octapeptide (Scr14), DYTMQ motif (Scr15), YPWM motif (Scr12), and CTD (Scr8), demonstrating that these four conserved protein regions are differentially pleiotropic.

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