

We crossed virgin *Antp* females to a multiply-marked *D. simulans* stock carrying the recessive visible mutations *forked* (*f*: 1-56), *net* (*nt*: 2-0), *plum* (*pm*: 2-100), *scarlet* (*st*: 3-49), *ebony* (*e*: 3-63). (Third chromosome map positions from Jones and Orr, 1998). F1 *Antp* males were then backcrossed to virgin *f; nt pm; st e* females. Their male and female progeny were scored for *Antp* and each of the five markers. The results showed that *Antp* is not on the X chromosome as both male and female progeny showed the mutant phenotype. We further found that while 17.4% (*n* = 218) of *nt pm* progeny also showed *Antp*, no *st e* progeny (*n* = 168) showed *Antp*. Therefore, like *D. melanogaster*, *Antp* in *D. simulans* is on the third chromosome.

We determined *Antp*'s map position on the *D. simulans* third by backcrossing F1 *Antp* females to *f; nt pm; st e* males. Nearly 2000 progeny were then scored for the presence of *Antp* and the two third chromosome markers, *st* and *e* (Table 1). It should be noted that gene orders in *D. melanogaster* and *D. simulans* are essentially the same with the exception of loci included in a known paracentric inversion on the right arm of the third chromosome (Ashburner, 1989). In fact we found that the order of the three genes in *D. simulans* (*st e Antp*) differs from that in *D. melanogaster* (*st Ant e*) indicating that *Antp*, along with *e*, is included in the 3R inversion of *D. simulans*. Distances between markers are presented in Table 2. These map distances place *Antp* at 3-78.7. After accounting for the inversion

Table 1. Backcross progeny from F1 *Antp* females x *f, nt pm; st e* males.

Genotype	Progeny
<i>st + e</i>	750
<i>st Antp e</i>	1
<i>st Antp +</i>	63
<i>st + +</i>	93
<i>+ + e</i>	149
<i>+ Antp e</i>	8
<i>+ Antp +</i>	355
<i>+ + +</i>	514

Table 2. Map distances (cM).

Intervals	<i>D. simulans</i>	<i>D. melanogaster</i> †
<i>st e</i>	16.2	26.7
<i>e Antp</i>	31.9	23.2
<i>st Antp</i>	48.1*	3.5

**st Antp* distance is the sum of the smaller distances.

† Data are from Lindsley and Zimm (1992)

difference between species, the map distances between the three loci in *D. simulans* are 27% higher than those in *D. melanogaster*. This is consistent with the known average 30% genome-wide increase in recombination rate in *D. simulans* relative to *D. melanogaster* (Ashburner, 1989).

These findings indicate that the *Antp* mutation reported here is likely a lesion at the *D. simulans* homologue of the *Antennapedia* locus in *D. melanogaster*.

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References: Ashburner, M., 1989, *Drosophila: A Laboratory Handbook*. Cold Spring Harbor Laboratory Press; Carrol, S.B., 1996, *Nature* 376: 479-485; Jones, C.D., and H.A. Orr 1998, *Dros. Inf. Serv.* 81: 137-138; Lindsley, D.L., and G.G. Zimm 1992, *The Genome of Drosophila melanogaster*. Academic Press, New York.

Recurrence of *yellow* mutation in *Drosophila subobscura*.

Solé, E., and F. Mestres. Dept. Genètica. Facultat de Biologia. Universitat de Barcelona. 08071 - Barcelona (Spain).

Two males of *yellow* phenotype were detected in a homokaryotypic stock (*O₃₊₄₊₂₂/O₃₊₄₊₂₂*) of *D. subobscura*. These males were crossed with virgin females of the *yellow* stock. All offspring individuals were *yellow*, confirming that the original males presented the *yellow* mutation. This mutation has appeared many times in our laboratory stocks of *D. subobscura* (Mestres, 1996; Solé, 1997). In all cases the *yellow* mutation has arisen in the *cherry curled* strain or in stocks obtained by means of genetic crosses with this strain. This suggests that the *yellow* mutation originates in the *ch cu* strain, and that some transposable genetic element could be involved.

References: Mestres, F., 1996, *Dros. Inf. Serv.* 77: 148; Solé, E., 1997, *Dros. Inf. Serv.* 80: 105.