

Rayle, R. E. and D. I. Hoar.

University of California, Davis,
California. Gene order and cytological
localization of several X-linked mutants
of *Drosophila melanogaster*.

of gene order and cytological localization ought to be cleared up. Utilizing several previously described duplications and a set of seventeen $Dp(1;Y)$ chromosomes synthesized in this laboratory, it was possible to order the mutants *sta*, $1(1)EN2$, and *tw* with respect to each other and to the other mutants mentioned above (Table 1).

Cytological determination of the breakpoints of the duplications utilized (and several additional duplications and deficiencies) has provided rather precise salivary gland chromosome map locations for *tw*, *su(w^a)*, *sta* and *dor* (Table 2). The mutant *hfw* (Rayle, Genetics 56: 583) could not be ordered with respect to *dor* on the basis of the data of Table 1. Additional tests with *hfw*, the *dor* alleles and *dor*-variegating duplications differing in the direction from which the variegation is exerted (to be described in detail elsewhere) suggested that *hfw* is to the right of *dor*. The outside markers present in the single wild type recombinant so far recovered from *dor/hfw* females are consistent with this interpretation.

Several other observations made during this study should be mentioned, since they support the conclusions reached here. $Df(1)sta$ was found to be deficient for the *su(w^a)* locus. $Dp(1;3)sta$ carries *su(w^a)⁺*. The left breakpoint of $Dp(1;f)1337$ was found to be as described by Krivshenko, as opposed to the description by Gersh. $Dp(1;f)101$ was found not to cover any of the *dor* alleles. Its left breakpoint was shown to be at 1F4-2A2 rather than at 2A2-B1. $T(1;Y)2E$ (Clancy, Genetics 50: 241) was found to differ from the original $Dp(1;Y)2E$ (Masterson, DIS 43) by a loss of the wild type alleles of *su(s)²* (first noted by Clancy, personal communication), *tw*, *su(w^a)* and *sta*. Wild type alleles of *sc*, *dor* and *hfw* are still present in $T(1;Y)2E$.

Table 1. Results of tests for the presence (+) or absence (-) of wild type alleles (y^+ and y^2 both scored as y^{1+}) in duplications used to order mutants in the *su(s)²* - $1(1)EN2$ interval.

Duplication tested	y^1	<i>sc</i>	Loci tested (arranged in correct order)						<i>hfw</i>	$1(1)EN2$
			<i>su(s)²</i>	<i>tw</i>	<i>su(w^a)</i>	<i>sta</i>	<i>dor</i>			
$Dp(1;3)w^{Vco}$	-	-	-	-	-	-	-	-	-	+
Dp Type A	+	+	+	+	+	+	+	+	+	-
Dp Type B	+	+	+	+	+	+	-	-	-	-
Dp Type C	+	+	+	+	+	-	-	-	-	-
$Dp(1;f)3$	+	+	+	+	-	-	-	-	-	-
Dp Type D	+	+	+	-	-	-	-	-	-	-

Dp Type A = seven $Dp(1;Y)$ chromosomes and $Dp(1;f)1337$.

Dp Type B = two $Dp(1;Y)$ chromosomes.

Dp Type C = six $Dp(1;Y)$ chromosomes, $Dp(1;f)18$ and $Dp(1;f)101$.

Dp Type D = two $Dp(1;Y)$ chromosomes.

Note: $Df(1)sta;Dp(1;3)sta = sta$. Tests for coverage of $Df(1)sta$ lethality in absence of $Dp(1;3)sta$ gave results identical to those for coverage of the visible phenotype.

Table 2. Localization of *tw*, *su(w^a)*, *sta*, *dor* and *hfw* on the salivary gland X-chromosome map.

Longest and shortest of each duplication type	Genetic coverage	Left breakpoint	Map location of mutant
$Dp(1;f)3$	<i>y</i> - <i>tw</i>	1D3-1E1	<i>tw</i> in 1C5-1D4*
$Dp(1;Y)60e17.4-3$	<i>y</i> - <i>su(w^a)</i>	1E2-4	<i>su(w^a)</i> in 1D4-1E3
$Dp(1;f)101$	<i>y</i> - <i>su(w^a)</i>	1F4-2A2	
$Dp(1;Y)59k9-1$	<i>y</i> - <i>sta</i>	2A2-B1	<i>sta</i> in 2A1-4
$Dp(1;Y)68h20$	<i>y</i> - <i>sta</i>	2B3-5	
$Dp(1;f)1337$	<i>y</i> - <i>dor</i> , <i>hfw</i>	2B8-9	<i>dor</i> , <i>hfw</i> in 2B4-8

*Based on cytology of $Dp(1;f)3$ and on the published location of *tw* in 1C5-2C10 (Lindsley and Grell, 1968).