

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*.

The $su(w^a)$ locus is known to be in the
 $su(s)^2$ - dor interval (M. M. Green,
unpublished). As an initial step in two
independent studies of genetic fine
structure in this region of the X chromo-
some, it was felt that ambiguities and
inconsistencies in the published reports

of gene order and cytological localization ought to be cleared up. Utilizing several pre-
viously 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 chromo-
some 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 recom-
binant 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 communi-
cation), *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)							$1(1)EN2$
			$su(s)^2$	<i>tw</i>	$su(w^a)$	<i>sta</i>	<i>dor</i>	<i>hfw</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> - $su(w^a)$	1E2-4	$su(w^a)$ in 1D4-1E3
$Dp(1;f)101$	<i>y</i> - $su(w^a)$	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).