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chromosomes.

Crossing over in the male between sterile  
Y chromosomes (derived from  $y^+$  Y, Bros-  
seau, '60 Genetics 45:257) and an  $X \cdot Y^L$   
chromosome was examined in the process of  
generating  $X \cdot Y^{KL} \cdot y^+$  chromosomes for

another purpose.  $ycv\bar{f}car \cdot Y^L / y^+ Y^{KL} \cdot Y^S$  ♂♂ were mated to RA  $y\bar{f} / Y^{BS}$  ♀♀. The expected  
cross overs, unmarked fertile Y's and  $X \cdot Y^{KL} \cdot y^+$  were recovered as  $y\bar{f}$  ♀♀ (RA  $y\bar{f} / 0$  is lethal)  
and  $cv\bar{f}car$  B ♂♂ respectively. The data are presented in Table 1.

Table 1. Exchange between marked free Y's and an attached  $X \cdot Y^L$  chromosome.

RA  $y\bar{f} / Y^{BS}$  ♀♀ x  $ycv\bar{f}car \cdot Y^L / y^+ Y^{KL} \cdot Y^S$  ♂♂

Stock	Loci Affected	Total No. Flies	Crossovers*		Frequency of Crossing over x 10 <sup>-4</sup>
			$y\bar{f}$	$cv\bar{f}carB$	
L 3	k1 5	37,865	6	4	2.6
L 19	k1 3,4,5	11,023	2	2	3.6
L 24	k1 3,4,5	29,963	20	7	9.0
L 28+	k1 5	21,515+	34	12	10.7*
L 36	k1 4,5	34,865	9	5	4.0
L 38	k1 3,4	14,471	14	7	14.4
L 41	k1 3	36,882	17	6	6.2
4-15	k1 1,2	38,751	20	23	11.1
4-65	k1 3,4,5	29,383	11	12	7.8
4-93	k1 3	14,928	10	7	11.4

\*Corrected for nondisjunctional progeny.

+The  $ycv\bar{f}car \cdot Y^L$  was found to contain a  $y^+$  suppressed lethal therefore only female  
and recombinant male survivors were recorded.

\*Calculated by doubling the number of females as an estimate of the total number of  
flies.

All of the free Y's used, except 4-15, show a variegation for male fertility (V-type  
position effects, Brosseau personal communication). It is therefore probable that these  
chromosomes contain intrachromosomal rearrangements which may account for the very different  
frequencies of crossing over as well as the differential recovery of the reciprocal recombi-  
nant classes of some chromosomes. Nonrandom recovery may reflect the occurrence of nonran-  
dom disjunction of recombinant chromosomes.

It is likely that at least some of the crossing over occurs during premeiotic mitoses  
(Brosseau, '58 DIS 32:115). Since more than one male parent was used in each bottle the  
present data do not provide conclusive evidence for clustering.

Exchange between the free Y and the attached  $X \cdot Y^L$  occurs with frequencies similar to  
the frequencies of exchanges between X heterochromatin and a Y (Lindsley '55 Genetics 40:24).  
One might expect that with more extensive regions of homology between the attached  $X \cdot Y^L$   
(probably containing a Y centromere and part of  $Y^S$  in addition to  $Y^L$ ) and a free Y exchanges  
could occur in more than one place. This might be manifested by a higher frequency of  
exchange, however the frequencies are not substantially enhanced. The existence of multiple  
places of exchange is suggested by the additional finding that though the derived attached  
 $X \cdot Y^L$  position effect chromosomes as a group often show different fertility levels from that  
of the free Y position effect chromosome, in some cases there exist two distinct classes of  
fertility levels among the recombinant chromosomes, perhaps reflecting qualitative differ-  
ences in their basal heterochromatin.

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