be recognized. Nevertheless, it was realized that \mathbf{w}^{ch} is located to the left of \mathbf{w} , which in turn signifies that \mathbf{w}^{i} is located to the left of \mathbf{w} .

In the present crossing over tests with w^i and other mutants of a subsite, the frequencies of recombinational derivatives were comparatively low in general when compared to that of the mutants of two different subsites. This may be explained most plausibly due to close sharing of location of w^i with the employed mutants, rather than that w^i inactivates recombination in this genetic interval. This supplements our earlier findings (Hazra et al. 1978) that w^i is a point mutation rather than a duplication as hypothesized by Bowman (1965).

References: Bowman, J.T. Jr. 1965, Genetics 52:1069; Hazra, S.K., J. Banerjee and S.K. Sen 1978, Heredity 40:299; Judd, B.H. 1959, Genetics 44:34; Lewis, E.B. 1959, Genetics 44:522; Lindsley, D.L. and L. Sandler 1963, in: Methodology in Basic Genetics, p. 390.

Hedrick, P.W. and E. Murray. University of Kansas, Lawrence, Kansas. Competition between D. melanogaster and D. simulans from natural populations.

Lawrence, Kansas, melanogaster was the most common of the two species in both 1977 and 1978 (Table 1). At all sampling times, the proportion of melanogaster was around 90% and the overall proportion is exactly 90.0%.

Table 1. The number of melanogaster and simulans males trapped in Lawrence, Kansas.

Date	me1	sim	% me1
7/77	292	25	92.1
9/77	218	35	86.2
7/78	50	1	98.0
9/78	86	11	88.7
Total	646	72	90.0

D. melanogaster and D. simulans are sympatric over much of their distributions and they appear to have similar ecological niches. Generally, melanogaster is the more common species where they coexist although there are some exceptions. In samples captured in a single location in

The proportions of two species observed in nature may be a reflection of a number of factors, such as predation, interspecific competition, sampling techniques, habitat selection, etc. Therefore, an interspecific competition experiment was set up to examine whether the results of interspecific competition in the laboratory were consistent with the field data. As a result only males could be scored, since females of the two species are very difficult to distinguish. At least 50 males were scored for each replicate every generation except in a few generations where there were slightly less than 50 males in a rep-

licate. The two lines of melanogaster, mel 1 and mel 2, were randomly selected isofemale lines caught at the Lawrence location and had been in culture for approximately six months. One simulans line, sim 1, was initiated from approximately 10 females caught in a Kansas City, Kansas park about 35 miles from Lawrence and had been in culture for approximately 30 months. The other simulans line, sim 2, was an isofemale line that was caught at the Lawrence location and had been in culture for approximately 18 months.

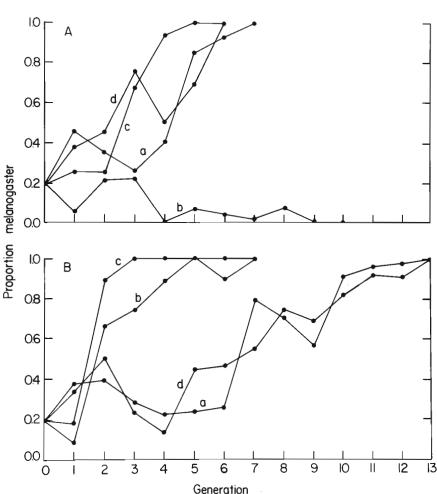
Since melanogaster was in higher proportion in local natural populations and generally outcompetes simulans in laboratory tests, the experiments were initiated with 6 pairs of melanogaster and 24 pairs of simulans, giving an initial frequency of 20% melanogaster. Four replicates of each of the four combinations of the melanogaster and simulans lines were set up. Generations were discrete and kept at 14-day intervals with the adults allowed to lay eggs for four days. Other details are as in Hedrick (1973).

The results of competition for sim 1-mel 1 and sim 1-mel 2 are given in Figures 1A and 1B, respectively. In only one replicate did simulans outcompete melanogaster and become fixed, replicate (b) of the sim 1-mel 1 competition. In all other replicates, melanogaster eventually became 100% of the culture. There is variation between replicates, however, with replicates (a) and (d) of the mel 1-sim 2 competition containing a few simulans even after 12 generations. When sim 2 was competing with mel 1 and mel 2, simulans was elminated within five generations in five of the replicates. In the other three replicates, no melanogaster were ever scored. It appears that for some unexplained reason, the initial six females in these replicates did not produce any progeny.

One can measure the relative competitive ability of these two species for the different replicates by finding the "best" numerical fit of the change in proportions over time. Since melanogaster was the winner in all but one replicate, the relative competitive ability of

Table 2. Relative competitive ability estimated from the best fit to the observed change in species composition for each replicate.

	sim	n 1	sim 2			
Replicate	mel 1	me <u>l</u> 2	mel 1	mel 2		
a	0.70	0.80	0.25	0.70		
Ъ	1.30	0.45	_	-		
С	0.65	0.30	0.45	0.65		
d	0.70	0.80	0.25	_		
Mean	0.84	0.59	0.32	0.68		



Figs. 1A and 1B

were established from single females at the same time. On the other hand. sim 1 was different from sim 2 in that it was esta-

melanogaster was assumed to be unity and the simulans competitive ability (x) was varied at 0.05 increments until the smallest sum of squared deviations was found. More specifically, competition was assumed to be analogous to selection in a haploid where p_i and q_i are the proportions of melanogaster and simulans in generation i, respectively. Then the proportion of melano-

$$p_{i+1} = p_i/(p_i + xq_i)$$

gaster in generation i+l is

The difference between these proportions and those estimated in the replicates was squared and summed for different values of

> x. The value of x which gave the minimum sum of squared deviations is given in Table 2. The relative competitive ability for the melanogaster lines can be calculated as 1/x.

An interesting finding from this analysis is that the estimated "relative competitive ability" for the two melanogaster lines are quite similar while those for the simulans lines are quite different. The mean relative competitive abilities for mel 1 and mel 2 were 1.64 and 1.61, respectively, and for sim 1 and sim 2. they are 0.72 and 0.46, respectively. The similarity of the melanogaster lines might have been expected since these lines blished earlier and with a larger initial number of females. Either of these factors may have contributed to the greater competitive ability of sim 1. The earlier establishment date would have allowed

more time for adaptation and the larger initial sample could have given more genetic variation for factors important to competitive ability.

References: Hedrick, P.W. 1973, Evolution 26:513-522.

Gr	O Domestic			Wil	d			S			
Grand Total	Total	Mahaba- leswar	Lonavala	Poona	Total	Mahaba- leswar	Lonavala	Poona	Collection locality	Species group:	Sub-genus:
1	,	740	620	550	١	740	500	550	Altitude (in meters)	up:	s.
903	903	74	684	145	,		1	ı	D. ananassae		
19		1	ı	ı	19	19	ı	1	D. anomelani		
207	4	1	4	ı	203	74	105	24	D. bipectinata		
2	,	ı	i	ŀ	2	1	ı	2	D. coonorensis		
21			1	1	21	13	6	2	D. eugracilis		
956	,			ı	956	484	224	248	D. jambulina	melanogaster	Sophophora
00	٠.	1	ı		∞	00	ı	•	D. kikkawai	noga	noph
690	6		6	ı	684	277	179	228	D. malerkotliana D. melanogaster D. mysorensis		ora
207	207	20	35	152	,			ı			
21	,	1		1	21	12	ı	9			
7		1	ı	1	7	G	2	1	D. pseudoananassae		
ω	١.	•	1		ω	2	1	1	D. montium		
91	,	ı	ı		91	22	36	$\frac{3}{3}$	D. rajasekari		
ω			1		ω	2	1	•	D. suzukii		
273	,	1	,	ı	273	186	41	46	D _o nasuta	immi	Drosc
53		1	ı	ı	53	42	4	7	D. neonasuta	igr	sop
00	,		1	i	∞	00	ı	1	D. immigrans	ophila grans	
4	١,			ı	4	2	2	•	D. nigra	ים ו	- 1
1	١.	1	+	1	<u> </u>	1	1	•	D. krishnamurthyi	phila	Scapto-
26	,	ı	ı	ı	26	26	1		D. varietas		-0.1
2	2	1	ı	2	ı	1	•	ı	D. busckii Dorsilopha		pha
13		•	,	ı	13	4	ı	9	Phorticella striata		
3518	1122	94	729	299	2396	1186	602	608	Total		
22	ر ن	2	4	ω	19	17	1'2	10	Total number of Drosophila species		

Distribution of different Drosophila species in Lonavala, Poona and Mahabaleswar. Table 1.