

Alonso, C. University of Nijmegen, The Netherlands. The effect of gibberellic acid on the development of *D. hydei*.

The effect of injection into larvae of *D. hydei* of various concentrations of gibberellic acid (GA<sub>3</sub>) on the chromosomal puffing pattern, the moment of puparium formation and the number of flies emerging from pupae was investigated.

It was observed that GA<sub>3</sub> can induce a new puff, 4-72B, in 15% of the larvae injected at 140 hours following oviposition. This puff can be observed 9 - 10 hours after injection. If GA<sub>3</sub> is injected at 153 h. a new puff develops within 3 - 4 hours at region 2-21B. These puffs have never been observed in normal development and are absent in animals injected with the solvent (Ringer's). The frequency with which puff 2-21B occurs depends on the concentration GA<sub>3</sub> applied, but never reaches a 100%. Apart from the newly induced puffs, GA<sub>3</sub> appears to affect the occurrence and size of puffs which are characteristic for the period prior to puparium formation. Injection of 1.5 µg/larva delays the appearance of the ecdysone-specific puff 4-78B and inhibits its full development. The same applies for puff 4-77BC. Mixtures of ecdysone and GA<sub>3</sub> injected into intermolt larvae gave a similar effect, a delayed development of the puffs 4-78B, and 4-77BC as compared with larvae injected with ecdysone only (2.10<sup>-1</sup> µg/larva).

Also puparium formation was delayed if GA<sub>3</sub> was injected at 153 h. or later (pup. formation occurs normally around 160 h.). The delay was increased with increasing concentration of GA<sub>3</sub>.

The percentage of flies emerging from pupae developed from GA<sub>3</sub> injected larvae decreased considerably with increasing GA<sub>3</sub> concentration. Injection of 6 µg/larva resulted in 54% lethality.

Schalet, A.\* University of Connecticut, Storrs, Connecticut. Two modified crossover-selector systems of general application to fine structure analysis.

Scheme 1 has been used in a fine structure analysis of the vermilion locus, but is applicable to other X-linked and autosomal loci. For the *v* locus a cross of the following type was used:

$$\begin{array}{l} \text{♀♀ (A) } \underline{11} + v^x \underline{13} + x \quad \text{♂♂ (C) } + \underline{12} v \underline{13} + ;bw^D \\ \text{(B) } + \underline{12} vY + \underline{14} \quad \text{(D) } - + v + - Y \end{array}$$

From this cross the only ♂♂ that survive are 1/2 of the crossovers between the outside lethals, 11 & 14. The only ♀♀ that survive, aside from non-disjunctionally produced individuals, are 1/2 of the crossovers between the inside lethals, 12 & 13. If v<sup>x</sup> is located to the left of v<sup>y</sup>, then v<sup>+</sup> recombinant ♂♂ will survive. If v<sup>x</sup> is located to the right of v<sup>y</sup>, then v<sup>+</sup> recombinant ♀♀ will survive.

Chromosome (B) was introduced into the cross thru parental ♂♂ carrying a Y chromosome that covered the region from 12 thru 14 (Schalet DIS 44: 123). This chromosome, as well as chromosome (D), was derived from the v<sup>+</sup>Y y<sup>+</sup> chromosomes synthesized by Chovnick, DIS 43: 170. The presence of the y<sup>+</sup> region on the Y, chromosome (D), with its Hw effect, and the partial suppression of the bw<sup>D</sup> phenotype in ♀♀ carrying a Y chromosome, permitted the detection of XXY ♀♀. Such ♀♀ appeared at an estimated rate of 1 for every 1,500 regularly produced zygotes. Linkage relationships determined from other crosses were as follows: ras--11, 0.1; 11--12, 0.4; 12--v, 0.2; v--13, 0.7; 13--14, 2.0. Note that the value of 0.7 for the interval between ras and v is closer to the value of 0.59 reported by Lefevre, DIS 45: 40, than the standard value of 0.2.

In the table below the total number of zygotes sampled has been calculated on the basis that each regularly produced ♀ represented approximately 1/444 of the number of eggs laid, (2/1,000 & 7/1,000)/4. Although only 1/2 of the eggs laid represent sampled chromosomes, this scheme provides the advantage that for any two potentially separable alleles, whatever their left-right orientation, only a single cross is required. Consequently, each allele need be inserted in or induced on only one of the two types of lethal bearing chromosomes.

Test	v <sup>+</sup> /Total	Order	Test	v <sup>+</sup> /Total	Order
2/1	0/307,000		1/E1	1♂/330,000	1-E1
1/k	0/890,000		36f/E1	1♀/460,000	E1-36f
36f/65c	0/250,000		2/E1	3♂♂/167,000	2-E1
1/36f	(Green)	1-36f	36f/2	1♀/195,000	2-36f
48a/36f	(Barish & Fox)	48a-36f	36f/k	2♀♀/350,000	k-36f