THE MAP DISTANCE BETWEEN THE e AND wlo LOCI

Murfet, I.C. Botany Department, University of Tasmania, Hobart, Australia

The presence of locus e\_ on chromosome 6 was first indicated by linkage between e and p (2). Subsequently, testcross data involving genes e, p and pl revealed the gene order to be e-p-pl (4) The observed crossover value of 28% between e\_ and p placed the e locus at the upper extremity of chromosome 6 above the existing uppermost marker, wlo. The joint segregation data in Table 1 indicate a distance of 26 units between e and wlo. The map of chromosome 6 as drawn by Blixt (1) is therefore extended by that amount. The data in Table 1 were obtained by crossing line 60 (lf E Sn hr Wlo) to segregates with the genotype lf e Sn hr wlo in the F, of cross 256 (L31 x L58).

Table 1. Joint segregation data for genes  $\underline{e}$  and  $\underline{wlo}$  (coupling phase) obtained from cross 284 F<sub>2</sub>.

Phenotype				Total	Joint seg. $\chi^2_1$	$Cr0\% \pm S.E.$
E Wlo	E wlo	e Wlo	e wlo	2010.28 20 11		
62	10	14	15	101	17.19***	$26.3 \pm 5.3$

Although the position of the e locus is now established, the locus itself has certain disadvantages as a marker. Firstly, the E/e segregation is only apparent with certain combinations of the major flowering genes, e.g. it is obscured by the epistatic action of the genes Lf and Lf (2,5). These higher alleles in the <u>lf</u> series are in fact very common. Secondly, the variable penetrance of Sn(with respect to flowering node) means that with certain polygenic backgrounds genotype lf e Sn hr can flower at a low node and may therefore be confused with lf E- Sn hr (2). Thirdly, a single dose of E is sometimes insufficient to overcome the latening action of Sn and segregates of genotype lf Ee Sn hr may flower at a high node and thus be confused with the penetrant lf e Sn hr plants (5). The last two situations can often be resolved by growing  $\mathsf{F3}$  . Such was the case in Cross 284 where, fortunately, the penetrance of Sn was very close to one but about 25% of Ee heterozygotes escaped from the early class. These escapes were readily identified in F.. Finally, as with the other flowering genes, observation of an E/e segregation is facilitated by the use of controlled environment conditions. However, special conditions are not necessarily obligatory (3).

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