

Evidence supporting the revision and integration of the pea chromosome 7 linkage map

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The application of biochemical and molecular technology to the development of the pea linkage map has not only increased the overall number of available genetic marker loci, but has also revealed linkage associations that sometimes differ markedly from the previously accepted relationships presented by Blixt (1). Based on their own studies, as well as the findings of a number of other geneticists and cytogeneticists, Weeden and Wolko (8) have suggested substantial rearrangements for chromosomes 2, 5 and 7. The most recent proposal extending from this work is summarized on the cover of volume 23 of *Pisum Genetics*; a portion of this revised map is included below in Fig. 1.

The observation that much of the former chromosome 7 linkage group (most notably, *r* and *tl*) resides on chromosome 5 has already gained additional support (2, 5). The proposal that the (upper) nucleolar organizer region of chromosome 7 is linked with the upper arm segment of former chromosome 2, however, requires further corroboration.

In Table 1 and Fig. 1, evidence is presented of linkage associations that span these previously unrelated groups of genes assigned to chromosome 7. Data obtained for the F₂ progeny from two independent crosses, SLOW x JI1794 and A1078-234 x PI179449, verify linkage between genetic loci formerly assigned to chromosome 7 (*Skdh*, *Cab*, *oh* and *Aat-m*) and genes associated with the nucleolar organizer region of chromosome 7 (*Pgm-c*, *Pgd-p* and *Rrn-2*). (Please note that designations for both *Rrn-2* and *Cab* have been modified to conform with standard nomenclature).

The linkage distances shown in Fig. 1 display internal consistency and they are reasonably close to values generated by other mapping studies, with the obvious exception of the large recombinant fraction ($19.6 \pm 5.0\%$) for *Skdh* and *oh*. Previous investigations (3, 7) have characterized these loci as very tightly linked, more in keeping with the map distances shown here for *Skdh* - *Cab* and *Cab* - *oh*. Notwithstanding this discrepancy, the data clearly corroborate the revision of chromosome 7 and assign both of its segments to the same linkage group.

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Table 1. Joint segregation analyses for chromosome 7 genetic marker loci.

Loci	Number of F ₂ progeny in each genotypic class ^a									n	χ^2	P	Recombinant fraction ^g
	11.11	11.12	11.22	12.11	12.12	12.22	22.11	22.12	22.22				
	SLOW X JI 1794												
<i>Pgm-c, Pgd-p^b</i>	21	4	0	6	43	1	0	1	21	97	133.5	<<0.001	6.4 ± 1.8
<i>Pgd-p, Rrn-2^c</i>	15		10	3		41	0		20	89	34.5	<<0.001	15.6 ± 4.1
<i>Pgm-c, Rrn-2^c</i>	14		11	5		40	0		20	90	26.3	<<0.001	19.0 ± 4.5
<i>Skdh, Rrn-2^c</i>	9		5	10		45	0		22	91	22.0	<<0.001	19.0 ± 4.5
<i>Pgd-p, Skdh^b</i>	9	15	3	5	31	13	2	11	9	98	11.4	<0.05	34.4 ± 4.4
<i>Skdh, Cab^b</i>	10	1	0	1	34	2	0	1	14	63	96.0	<<0.001	4.1 ± 1.8
<i>Cab, Rrn-2^c</i>	5		3	5		29	1		12	55	10.9	<0.01	21.7 ± 6.2
<i>Cab, oh^d</i>	9		0	21		0	1		10	41	36.1	<<0.001	2.4 ± 2.4
<i>Rrn-2, oh^e</i>	16		1				39		14	70	3.2	<0.1	27.1 ± 10.9
<i>Skdh, oh^d</i>	15		1	36		4	8		12	76	22.2	<<0.001	19.6 ± 5.0
	A1078 -234 X PI 179449 ^f												
<i>Pgm-c, Pgd-p^b</i>	11	1	1	3	22	5	0	2	15	60	61.1	<<0.001	11.4 ± 3.1
<i>Pgd-p, Rrn-2^b</i>	10	0	0	4	11	4	0	7	12	48	38.3	<<0.001	16.5 ± 4.2
<i>Pgm-c, Rrn-2^b</i>	7	1	2	5	12	7	2	4	8	48	13.6	<0.01	29.9 ± 5.8
<i>Rrn-2, Cab^b</i>	7	1	0	2	13	4	0	4	13	48	31.4	<<0.001	16.7 ± 4.2
<i>Cab, Aat-m^d</i>	11		0	22		4	7		8	52	11.9	<0.01	22.2 ± 6.4
<i>Rrn-2, Aat-m^d</i>	13		1	14		3	8		6	45	5.5	<0.1	30.9 ± 8.0

^aGenotypic designations: 11 - JI 1794 or PI 179449 homozygotes, 22 - Slow or A1078-234 homozygotes, and 12 - heterozygotes (for *Cab*, 1 and 2 represent haplotypes).

^b1:2:1:2:4:2:1:2:1 expected ratio.

^c1:3:2:6:1:3 expected ratio; in this cross, *Rrn-2* band scored as present or absent.

^d3:1:6:2:3:1 expected ratio; due to dominance, *oh* heterozygote listed as 11; *Aat-m* band scored as present or absent.

^e3:1:9:3 expected ratio; in this cross, *Rrn-2* band scored as present or absent; due to dominance, *oh* heterozygote listed as 11.

^fReciprocal cross.

^gCalculated using the LINKAGE-1 computer program (4).

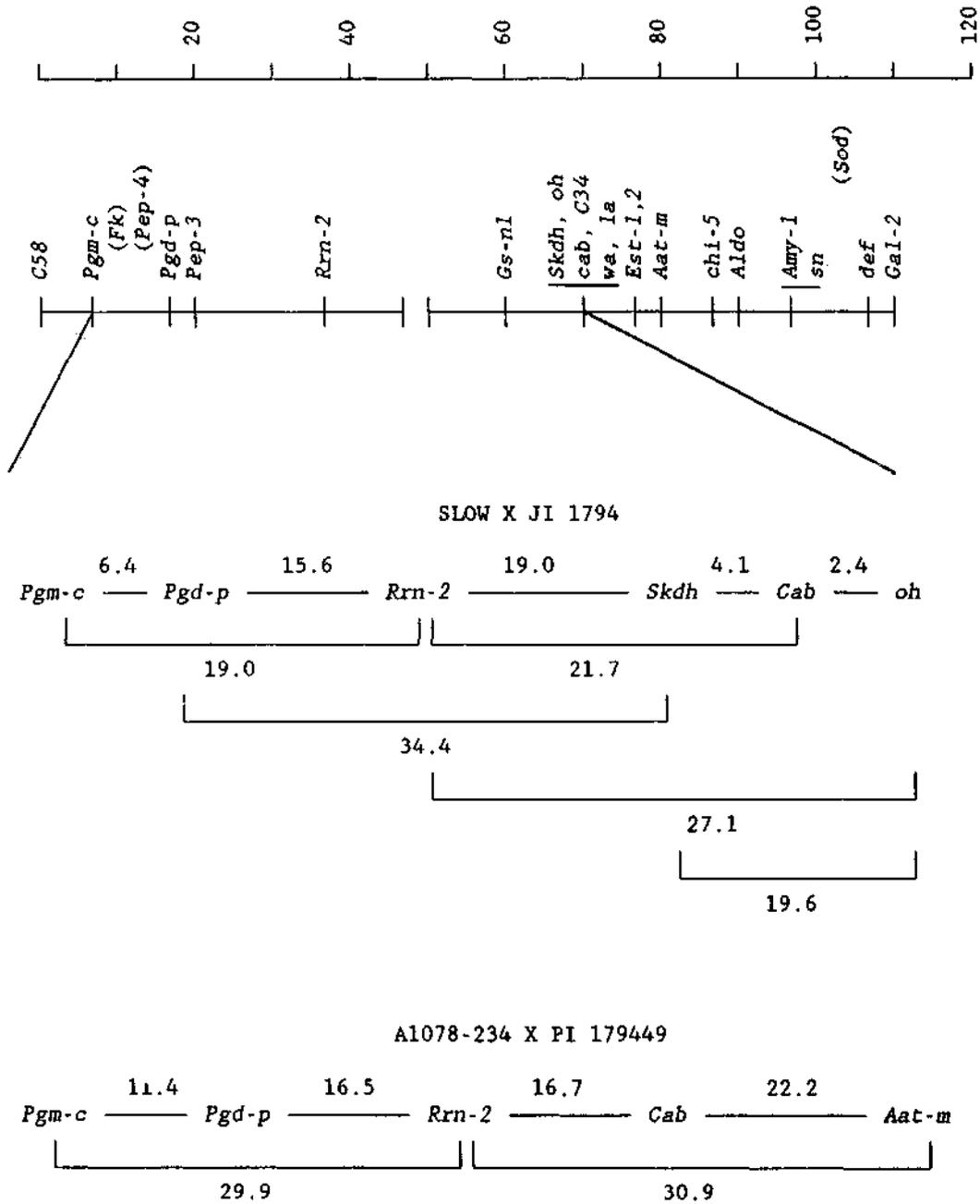


Fig. 1. Linkage group relationships for chromosome 7. The upper diagram represents the putative arrangement of linkage groups recently proposed by Weeden et al. (6), the space between the upper and lower chromosome arms indicating an unconfirmed association. The scale is in centimorgans (cM). The middle and lower diagrams depict mapping relationships generated from data in Table 1 that support the proposed association. The middle linkage map shows the intrachromosomal assignments among a group of genetic loci segregating in F₂ progeny from the cross SLOW x JI1794, while the lower linkage map shows similar results from the cross A1078-234 x PI179449. Other genetic loci shown in the upper diagram do not segregate in the crosses used and, thus, are not included in these analyses.

1. Blixt, S. 1974. *In Handbook of Genetics*, Ed. R.C. King, Plenum Press, pp. 181-221.
2. Murfet, I.C. 1990. *Pisum Newsl.* 22:38-40.
3. Polans, N.O., Weeden, N.F. and Thompson, W.F. 1985. *Proc. Natl. Acad. Sci. USA* 82:5083-5087.
4. Suiter, K. A., Wendel, J.F. and Case, J.S. 1983. *J. Heredity* 74:203-204.
5. Swiecicki, W.K. 1990. *isum Newsl.* 22:62-63.
6. Weeden, N.F., Ambrose, M. and Swiecicki, W.K. 1991. *Pisum Genet.* 23:cover.
7. Weeden, N.F. and Marx, G.A. 1984, *J. Heredity* 75:365-370.
8. Weeden, N.F. and Wolko, B. 1990. *In Genetic Maps*, Ed. S. O'Brien, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York, pp. 6106-6112.