Perk ins, D. D. Gene order in the albino region of linkage group [.

The marker arg-6: arginine-6 is usually shown left of a[-2; a] bino-2 in published maps. This location was bared largely on data with the incompletely albino mutant aur: aurescent and involved the assumption that al-2 and arg-6 are contiguous. The data pre-

rented here suggest that this assumption may be incorrect. arg-6 is almost certainly right of al-2, and is probably located between al-2 and our. The results in Table | indicate the order ad-9 act-1 al-2 arg-6 aur hr. (The alternative ad-9 act-1 al-2 org-6 hs aur is possible but less likely.) act-1: actidione-1 would thus be the best flanking marker left of al-2.

Methods: Crosses were made at 25 °C. For the first two crosses, ascospores were isolated to agar slants in 75 mm tubes and heat-shocked the tubes. Glycerol complete medium was used in cross 1. Vogel's minimal + L-arginine (OS mg/ml) + DL-homoserine (0.2 mg/ml) was used in cross 2. Scoring was by transfer to slants of minimal medium plus appropriate supplements. Actidione (cycloheximide) was used for testing at 10 µg/ml, with normal autoclaving. Tests were unambiguous.

Although act-1 (Hsu 1963 J. Gen. Microbiol. 32:341) is not as close to albino as might be desired, it has the advantages of showing excellent viability and scorability and requiring no supplement. hr: homoserine is also a good marker, though caution must be taken to use supplemented minimal medium rather than complete medium, so as to avoid inhibition by unidentified constituents of the latter. arg-6 presents no difficulties of culture or scoring.

For crosses 3 to 5, ascospores were suspended, heatshocked and plated in minimal medium containing 1% sorbose + 0.05% glucose + 0.05% fructose, and incubated 48 hours at 34°C before colonies were isolated to minimal slants.

Results: The preferred order, a = 2 left of arg = 6, is clear when the least frequent single-crossover class is compared with the double crossovers, considering three loci at a time in crosser | and 2, and assuming first one and then the other of the two alternative orders (a) arg-6 right of arg = 6 left of a

Postulated order	- Singles			
	Parentals	Region	Region 2	Doubles
Wa <u>d</u> -9 al-2 <u>org−6</u>	66	2 4	1	0
(b) ad-9 arg-6 al-2	6 6	2 4	0	1
(a)_act-lal-2_ org-ó	359	35	6	1
(b) act-l grg-6 al-2	359	35	1	6

The order al-2 arg-6 hs was indicated by cross 2 on the basis of 2 singles between al-2 and hs versus 0 doubler. This order was supported by cross 3, where the coupling phase is reversed. Ascospores were plated in minimal sorbose medium. Of 67 prototrophic progeny, 62 were albino. The remaining five isolates (from atypical slow colonies) developed orange pigment, but all five produced albino progeny when crossed by al. They thus originated as pseudowild types or mixtures, and were excluded from the tabulation. Only prototrophs from fast-growing colonies were isolated in the succeeding crosses.

in 1969 (Genetics 44, second cross on p. 1194) arg-6 was placed left of <u>qur</u> on the basis of <u>q small</u> number of isolates picked visually as prototrophic germinating <u>ascospores</u>. The identical parental strains have been preserved, and were crossed to obtain the results listed under cross 4. Seventy fort-growing prototrophic colonies were isolated to slants, and only one among them was rejected as <u>qn</u> apparent aneuploid that darkened the medium and grew atypically. The 69 bong fide recombinants were <u>aur</u>, consistent either with the order arg-6 <u>qur</u> hs or <u>arg-6</u> hs <u>qur</u>, but not with the order <u>aur</u> arg-6 hr.

Cross 4 results indicate that <u>qur</u> is much closer to <u>hs</u> than to arg-6. This was borne out in cross 5, where <u>qur</u> separated from hs in only one prototrphic recombinant among **Q** total of 6) between erg-6 and hs. The single <u>qur</u>-hs recombinant favors locating-our left rather than right of hr.

There results suggest that <u>al-2</u> and <u>our are</u> separate genes located on opposite cider of <u>arg-6</u>, consistent with what is known both as to recombination and complementation between <u>al-2</u> and <u>our</u>. Obviously further critical data ore needed. Confirmation of the order <u>al-2</u> arg-6 <u>our</u> would support the proposal of <u>Barratt e+ al.</u> (1954) that the <u>our</u> locus be designated <u>al-1</u>: albino-1.

The assistance of R. J. Lloyd and R. E. Padilla is appreciated. (Support: PHS Grants AI 01462 and K6-GM-4899.) = = = Department of Biological Sciences, Stanford University, Stanford, California 94305.

Table 1

Marker

isolation

numbers

29997

34508

51504

91 Y154M37 + + al-2 em-6 36 1 0 ad-9 ace-l aly (917)29 KH52 15.4 13.2 1.1 15300 ALS4 29997 be 148 310 KH52 (78%)ace-1 al-2 arg-6 + 15 2 131 15300 7.7 1.9 0.6 29997 51504 0 62 15300 3 62 prototrophs 29997 (all al) 51504 69 0 69 29997 aur prototrophs 34508 (all aur) 51504 4 1959 results: 5 3 prototrophs (5 **aur**) aur he 60 61 15300

Table 1. Data from random segregants, establishing the order shown. Crosses 1 and 2 involved total isolation, crosses 3-5, selective plating.

Singles

1

Region

Parental

combinations

Zygote Genotype

a nd

Recombination %

Cross

No.

5

Recombination

Singles

Region

Doubles

Regions

1 and 2

Total and %

germination

prototrophs

(60 wild type,

1 aur)

Singles

Region

2

(The top number in each pair represents the class that has the + allele of the leftmost marker.)

act-l = resistant, act-l+ = sensitive.