YALE UNIVERSITY

OSBORN BOTANICAL LABORATORY NEW HAVEN, CONNECTICUT

KK March 3, 1947.

Dear Dr. Mather:

This should perhaps have been written as a postscript to my letter of the 8th ult., since certain remarks therein will have to be revised.

The results of the cross B-P-X T-L-B₁-Lac-V^r do not support the map order B₁; BM:Lac:V:P:TL which seemed to be suggested by the comparisons of the crosses BM X TLB₁V^r and BM X TPV^r. The data are:

B-P-T+L+B₁+Lac+V^S X B+P+T-L-B₁-Lac-V^T. (The P here does not recombine with P of TP).

-R +R -S +S Protetrophs: 25 0 27 4

 $B_1 - + B_1 + 62 2 41 1$

(these are types from B_1 plates. B_1 ca lox B_1 +

B- + B+ 74 10 69 12.

(from B plates but B not more than about 1/2 as frequent as B+)

must the k b, +?

There are two respects in which these data do not fit the map order assumed above:

1. -S is much too frequent

2. B- is much too infrequent:

Why wash in B- w afrequent in thems

However, these data are not fatal to the hypothesis of linear linkage, since the recombination value for V in the cross BM X TP may depend on the map order BM--P-V-T, on the assumption that the recombinations between V and T...P depend on two crossovers. The numerical data are not inconsistent with the theory (even in the absence of interference) particularly if T is to some extent to the right of L.

In this cross then, the recom crossovers required are:

Pretetrophs: A.DE (A.D is -S; A.E is -R; D=E) or A.B.C.DE A.B.C.D is +S; A.B.C.E is +R/

B- F.DE W.F.A, C.DE ABC. PE

What down ? - R ABR Cerns on the your. + R C.E.

dia.

It would not appear to be too worthwhile to attempt to make too precise estimations of absolute distances from such data as these until we learn more about the possibilities of intereferences, which could certainly throw one wide of the mark.

Another approach to the problem of linearity has suggested itself, and I am working on it now.

It should be possible to find 'cross-over suppressor' stocks after intense irradiation. These would be presumable based on the occurrence of inversions. It should be a relatively simple matter to detect inversions covering region A of the map, since crosses involving such stocks could yield B1-but no protrophs. (using, eg BM X TLB1.) If such can be found, one can also determine whether, if attall, of the regions B,C,D,or E are involved; a group of such stocks could provide evidence as to the linear plotting of regions of suppressed interchange and thereby indicate the linearity of the gene arrangement. First, the inversions will have to be detected.

What news on the new addition of your 'Statistical Analysis?'

Sincerely,

oshua Lederberg.