16 Cotober 17, 1940

My dear Taylor,

I have just completed the job of work on scoring of which I told you earlier, and enclose you comies of a few tables, (s) scores for 56 combinations based now on both A and B readings, between which there is no sign of a significent difference. This may be important as indicating remarkable physical similarity between the physical systems determining the a sorbtion of & and /3 on red cells. For the moment its chief roint is that I can use the whole body of data to give a rather accurate set of scores. The second table (b) gives nearly equivalent titrations, the effects differing by .1 of a two fold dilution, i.e., by about 7% antibody concentration, the standard error of a single titration, using I give a third table (c) which these scores, being just I use to calculate the variance between these three successive readings used in each titration, regarding these as independent estimates - which they may or may not be, for the analysis shows that there is less variation between these even than the small residual found by comparing different cells with different sera, as would be expected even with independent readings or if there were any specific residual affinity between cells and sera, or any

variation in technical procedure capable of affecting the whole dilution series similarly. I do not imagine that there is here anything more to follow up, but the analysis so far is of some interest.

I have further rescored the 60 titrations which you gave in your paper on <u>Weak A reaction</u>. Here the residuals may be obtained from the three expressions

$$A_1 - A_2 + A_1B - A_2B$$
 or, briefly, $A_1 - A_2$
 $A_1 + A_2 - A_1B - A_2B$, or, briefly $A_1 - A_2B$

and, finally,/interaction

 $A_2 + A_1B - A_1 - A_2B$

On the enclosed mess sheet I give these in three columns and derive from each the total mean and sum of squares of deviations from the mean, i.e. the contribution to the residual variance for which these 14 degrees of freedom are responsible; finally the variance of a single titration inferred from each set. These are all rather large, i.e. .20, .15, and .26, as compared with .11 from the very large material obtained two years ago. Why they are large may be worth discussing; I mean they may be readings by different people instead of all by Miss Prior as was the case with the material on which the e.s. Thus are may be worthed that the contrast A - A is not more variable than the others, as I think it should be if the 15 different sera differed at all in the proportion of and diwhich they contain. It suggests, in fact, that possibly these two substances occur naturally very nearly in a fixed

proportion, so far, that is to say, as really rather refined tests can show. On the view that these 15 sera differed appreciably in the ratio in which these two substances occur, we should have expected something like a variance .30 for A - A2, and perhaps .11 from the others. I believe we never used an A2 in the titrations two years ago, but if we did there would be some further data for checking up this point, which I think you will agree has some theoretical importance. Perhaps also you have now more data of the kind which you published, which could be included in the test.

Even if & and & occur in fixed proportion, the impression you have that a preliminary titration will give an idea that a particular serum is specially suitable for absorption to produce any be well founded relative to the individual cells to be used for absorption. It may, in fact, depend on such residual affinities as might also explain the greater agreement of readings in the same titration as compared with readings when several different sems and cells are compared.

I think this aspect of the subject is ripe for discussion between us, with a view to publishing what has been learned so far.

If you check my scorings, as I hope you will, or have them checked, you will notice that in three cases only two readings wer are available, and in those I have inferred what the one behind would be from what happens in other cases. In fact, before the readings # () - I find in seven cases a # and in one case a V, and give a compromise score accordingly. Before a sequence + () - I have seven cases all #, and before the reading () w - I have 19 cases, of which 8 are #, 10 + and one (). These frequencies are taken from the 60 titrations concerned, so I imagine they afford a rational basis for compiling the score in these three doubtful cases.

Yours sincerely,

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c.	Sums of	Squares	Jo Gomb	vidual r inations tetralion	, + fre , w	for each		= 2.91018 1.96374 1.61448 1.13949 .57027
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(48) 0.508	(23) 0.643	# + (12) 1.383	# (†) (47) 0.694	(98) 0.260	# (95) 0.267	(13) 0.804	# () w (78) 0.195	
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	d/f Sum of Squares		Reduced to		Mean square			
	2760 525.759 1025			175.253 112.825		.063497 Within titrations .11007 Residual between titrations.		