

CALIFORNIA INSTITUTE OF TECHNOLOGY  
PASADENA

DIVISION OF BIOLOGY  
KERCKHOFF LABORATORIES OF BIOLOGY

December 11, 1954

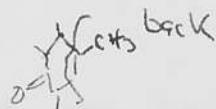
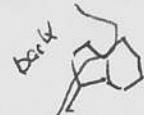
Dear Francis

Your letter and MS came yesterday. The Plant Virus MS seems quite sensible and to the point. I agree that it should be written largely to stimulate Plant Virus Research. As to where it should be published: Why not Advances in Virus Research - A dreadful affair but nevertheless it will be read by the people we want to convert. While at Woods Hole, Hauffer had asked me to write a review on Virus proteins and I turned him down. However I suspect the offer still stands. It's unlikely that I can work on the MS in the immediate future as I will soon be flying East - I leave in a week and beforehand I shall prepare some seminars to give at Harvard, etc. However I shall try to work on it while travelling from place to place.

About protein synthesis. After initial enthusiasm about DUL we are back to being solid RATTIE CLUB members. Idea was the following. Take DNA - break H bonds and

~~shift~~ ~~one chain down so that it forms~~

strings back of the base near back



R  
R<sub>1</sub>  
Mg<sub>2+</sub> bases between the complementary chains. This



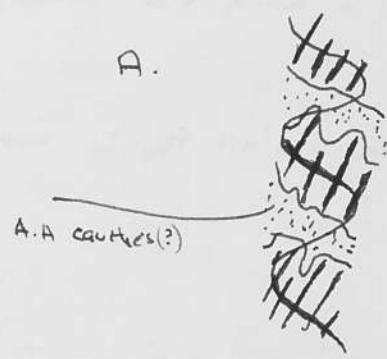
of another base {not necessarily complement}

We thus have a series of holes of 16 base types [more if neighbors count]. However in its disfavor are (1)  $\text{CH}_3$  can be replaced by  $\text{Br}$  - without serious effects - this "shells bad". In antigenic work this usually stops cross reactions. (2) A.A. backbone does not fit in easily. 3) since RNA also makes proteins [is this our downfall?] it seems miraculous that it can provide a similar code without using 5 methyl group - ~~this~~ both RNA and DNA make proteins they must do it by a similar trick.

So back to RNA. At first site this looks just as discouraging as before. However I suspect the answer is staring us in the face. Important fact about RNA picture is

$10 \text{ \AA}$  "metachromatic reflection". Must mean something periodic at this distance. Two possible

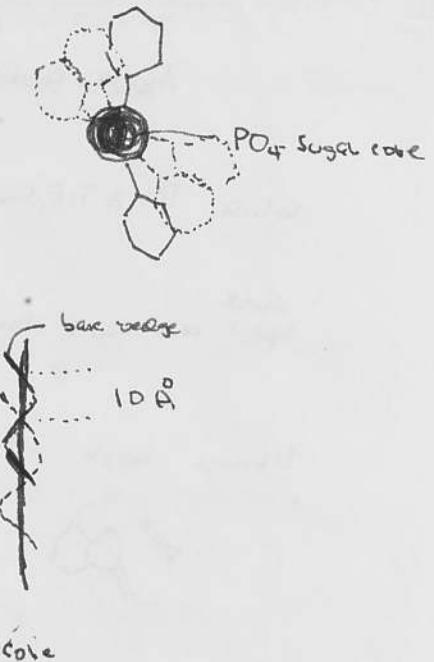
structures. [I neglect multi stranded structures. Origin too difficult if non complementarity exists.]



A.  
one stranded helix  
with pitch of  $\approx 11 \text{ \AA}$   
and containing about  
8-10 residues / turn. In  
this model, bases must be  
seriously tilted. Bases  
inside -  $\text{PO}_4$  out



B.  
one stranded helix  
with pitch  $\approx 4 \text{ \AA}$  or with  
about  $2\frac{1}{2}$  residues / turn  
In this model, bases form  
two wedges



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The main point is that when we tilt the bases ~~about~~ almost  $11^{\circ}$  to fibre axis  
we produce very suitable A.A. cavities. In fact we can say that we were to  
create a A.A. making best we should arrange for a pitch of not greater than  $11.12^{\circ}$   
and probably not less than  $8^{\circ}$  { if shell will be very thick to fit A.A. backbone  
in. So I do not think the X-ray picture is against us. However it will be a nasty job  
to prove existence of 20 specific cavities.

I was in Berkeley over the Thanksgiving Holiday. Pleasant but without scientific  
stimulation. Only complaints about Stanley. Schachman's break story does not convince me.

If they exist, they must be less than 1 in 500.

Did John Platt write you about transfer twists. I will see him while I am in Chicago  
He seems to have misread our articles. I don't think his process really helps the situation

Also, were you invited to Brookhaven? I received an invitation which implied that  
they hoped originally to ask you but that now that you were back in England, you were

no longer available. I could not make out whether they didn't have funds to ask you from  
England or whether you turned them down. I'm not anxious to speak on DNA at the

Mutation process, especially hesitant to write even an abstract on it. So I shall suggest  
they invite Seymour Benzer to talk about his phage crossing over data. [ I believe  
Sidney showed you the Benzer MS at Woods Hole.

I am leaving for Chicago on the 20th. Then to Cambridge Mass for 10 days and  
finally a few days in Washington before flying back. No need to indicate my desire to  
return East.

If I learn anything of interest while in East, I'll write you

With Holiday Greetings to Odile

Tom