

+ invite others to the conversation



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Juan Benet








juan@benet.ai



Initiator










Sharing Menu a panel to invite others to conversations, trails, etc.

It begins with the content initiator (creator) already populated. in the menu, and allows others to be invited. important to select sharing permissions

+ invite others to the conversation		 Read ▼	
	Juan Benet juan@benet.ai		Initiator
	Francis Crick fcrick@mrc.ac.uk		Write ▼
	James Watson jwat@mrc.ac.uk		Read ▼


Sharing Menu a panel to invite others to conversations, trails, etc.


When people are already shared-with, they appear under the initiator (or maybe it is sorted alphabetically or something).
Permissions are blue, so that they are perceived as links.


	<input type="text" value="Rosalind Franklin"/>	<div>+ Add</div>	<div> Read ▼</div>
	Rosalind Franklin rosalind@mrc.ac.uk		<div> Initiator</div>
	Robert Weinberg weinberg@wit.mit.edu		<div> Write ▼</div>
	Robert Sauer		
	James Watson jwat@mrc.ac.uk		<div> Read ▼</div>







Sharing Menu a panel to invite others to conversations, trails, etc.


Add another person or group by entering their name or associated email account. Field auto-completes giving user a contextual menu of options to select from. arrow keys should work. (add appears)


Rosalind Franklin

 Add











 Share ▼

 <div> Juan Benet juan@benet.ai </div>	<div>  Read can view conversation <div>Initiator</div> </div>
 <div> Francis Crick fcrick@mrc.ac.uk </div>	<div>  Write can add content to the conversation <div>Participate ▼</div> </div>
 <div> James Watson jwat@mrc.ac.uk </div>	<div>  Share can invite others to the conversation </div>

 Read ▼












Sharing Menu a panel to invite others to conversations, trails, etc.

Permission selection works similarly, and includes clear, simple explanation of the abilities granted. (permission menu opened automatically after selecting a user, to encourage awareness of permission selection.

	Rosalind Franklin	+ Add	 Share ▼
	Juan Benet juan@benet.ai		 Initiator
	Francis Crick fcrick@mrc.ac.uk		 Write ▼
	James Watson jwat@mrc.ac.uk		 Read ▼

Sharing Menu a panel to invite others to conversations, trails, etc.


Pressing enter (or clicking “Add”) adds the selected user to the list of permissions. This grants permissions and sends them a notification.









	Rosalind Franklin	 Share ▼	 Add
	Juan Benet juan@benet.ai	 Initiator	
	Francis Crick fcrick@mrc.ac.uk	 Write ▼	
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Sharing Menu a panel to invite others to conversations, trails, etc.

Alternative view.

+ invite others to the conversation

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	<div>Francis Crick</div> <div>fcrick@mrc.ac.uk</div>	<div> Write ▼</div>
	<div>James Watson</div> <div>jwat@mrc.ac.uk</div>	<div> Read ▼</div>
	<div>Rosalind Franklin</div> <div>rosalind@mrc.ac.uk</div>	<div> Share ▼</div>

Sharing Menu a panel to invite others to conversations, trails, etc.

Once added, the field clears, but sharing selection is remembered

Conversation Title

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Share an insight

+ invite others to the conversation

👁️

Read

▼



Juan Benet

juan@benet.ai

📁

Initiator

Start new conversation

New Conversation a blank “new conversation” view

Re: A Structure for Deoxyribose Nucleic Acid



This paper is really good because it launched a new field after coming up with the [greatest discovery](#) of all time. This paper is really good because it launched a [new field](#) after coming up with the greatest discovery of all time. This paper is really good. Here is a web link, here is a link [into the paper](#), here is a link to a [@Person](#), here is a link to a [Conversation](#), here is a link to [a figure](#), and i think this paper is [#interesting](#).

+ invite others to the conversation

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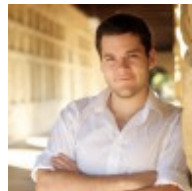
Juan Benet
juan@benet.ai



Initiator

Start new conversation

New Conversation a populated “new conversation” view
(the title should be seeded with where user came from)



Juan Benet



Share an insight

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Comment

New (Blank) Comment at the end of a running conversation



Juan Benet



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Comment

New Comment at the end of a running conversation. mid-edits.



Juan Benet



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alternative view



Juan Benet

10 minutes ago



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Posted Comment in a conversation. if it is current user's, then user can see “edit” and “delete” buttons.



Juan Benet

10 minutes ago




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Update comment

Editing/Updating Posted Comment

Several Comments Together



Juan Benet


10 minutes ago

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Juan Benet


10 minutes ago

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Juan Benet

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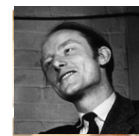
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Comment

Running Conversation with
several comments, and shared
with several participants

Re: A Structure for Deoxyribose Nucleic Acid

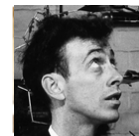


Juan Benet



10 minutes ago

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Juan Benet



10 minutes ago

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Juan Benet



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Running Conversation after clicking the “Add User” button next to participants, Sharing Menu pops up.

Re: A Structure for Deoxyribose Nucleic Acid



+ invite others to the conversation

 Share ▼

	Juan Benet juan@benet.ai		Initiator
	Francis Crick fcrick@mrc.ac.uk		Write ▼
	James Watson jwat@mrc.ac.uk		Read ▼
	Rosalind Franklin rosalind@mrc.ac.uk		Share ▼

Close



Juan Benet


10 minutes ago




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Conversations


Re: A Structure for Deoxyribose Nucleic Acid




updated 20 minutes ago

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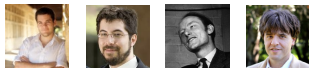
Re: Physical Principles for Scalable Neural Rec...




updated 9 years ago

 234


Re: Millisecond-timescale, genetically targeted ...




updated 9 years ago

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Re: A Structure for Deoxyribose Nucleic Acid




updated 20 minutes ago

 12k


Re: Physical Principles for Scalable Neural Rec...




updated 9 years ago

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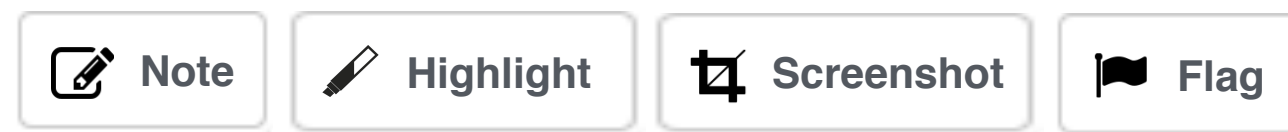
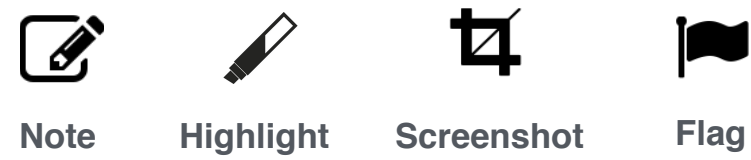
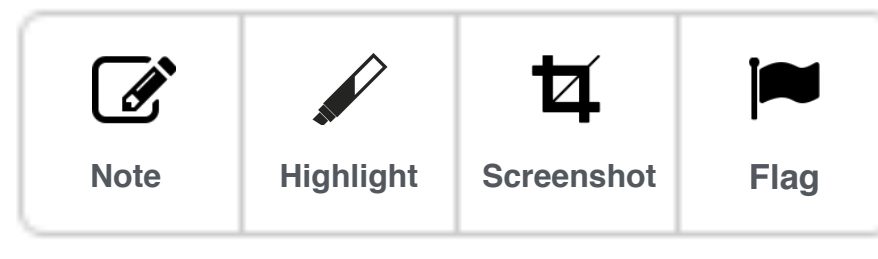
Re: Millisecond-timescale, genetically targeted ...



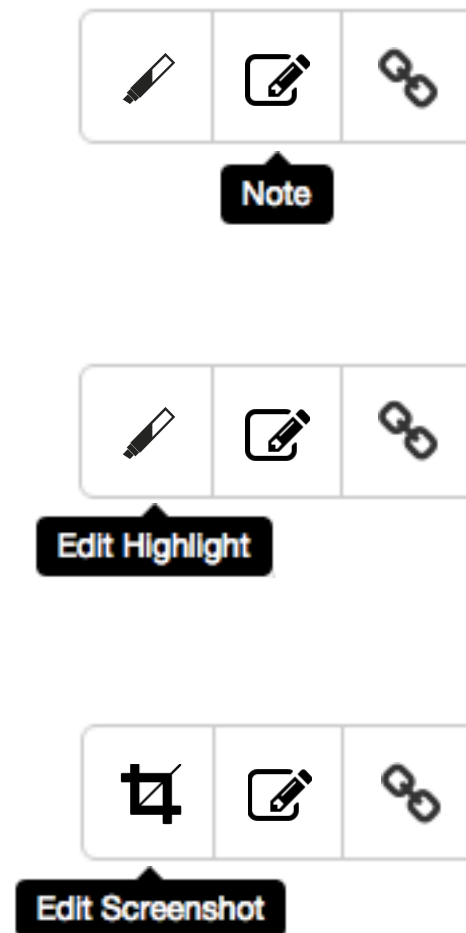
updated 9 years ago

 1

Conversations this view shows a set of conversations.



Home Toolbar this are things to do with the current entity.



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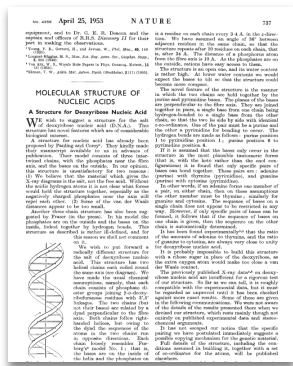
We wish to put forward a radically different structure for the salt of deoxyribose nucleic acid. This structure has two helical chains each coiled round the same axis (see diagram). We have made the usual chemical assumptions, namely, that each chain consists of phosphate diester groups joining β -D-deoxy-ribofuranose residues with 3',5' linkages. The two chains (but not their bases) are related by a dyad perpendicular to the fibre axis. Both chains follow right-handed helices, but owing to the dyad the sequences of the atoms in the two chains run in opposite directions. Each chain loosely resembles Furberg's² model No. 1; that is, the bases are on the inside of the helix and the phosphates on



Annotation Toolbars this is a toolbar for floating annotations.

(Edit, Note, Link) — link copies the link to the annotation for use in conversations (smart linking) or anywhere else (link to the annotation only)

Publication



A Structure for Deoxyribose Nucleic Acid



Francis Crick



James Watson



234



10634

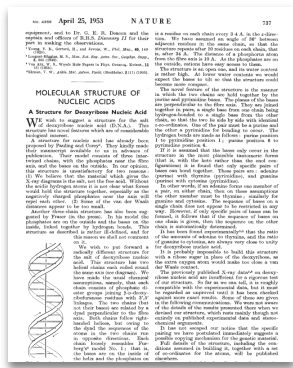


Tools



Publication this view shows publication metadata

Publication



A Structure for Deoxyribonucleic Acid



Francis Crick



James Watson



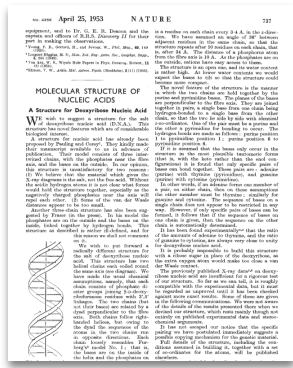
234



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Tools



A Structure for Deoxyribonucleic Acid



Francis Crick



James Watson



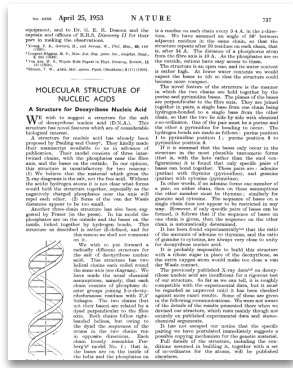
234



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Tools



A Structure for Deoxyribonucleic Acid



Francis Crick



James Watson



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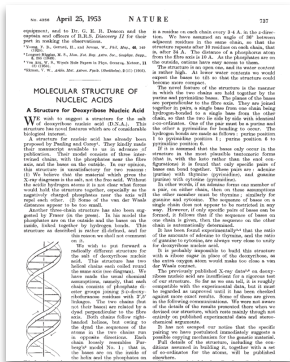


Tools



Publication List

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A Structure for Deoxyribose Nucleic Acid



Francis Crick



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Tools



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10.1038/171737a0



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Publication this view shows publication metadata
Clicking the tool icon opens a contextual menu with many useful
tools that we don't want to clutter the UI with.

equipment, and to Dr. G. E. R. Deacon and the captain and officers of R.R.S. *Discovery II* for their part in making the observations.

¹ Young, F. B., Gerrard, H., and Jevons, W., *Phil. Mag.*, **40**, 149 (1920).

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⁴ Ekman, V. W., *Arkiv. Mat. Astron. Fysik. (Stockholm)*, **2** (11) (1905).

MOLECULAR STRUCTURE OF NUCLEIC ACIDS

A Structure for Deoxyribose Nucleic Acid

WE wish to suggest a structure for the salt of deoxyribose nucleic acid (D.N.A.). This structure has novel features which are of considerable biological interest.

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We wish to put forward a radically different structure for the salt of deoxyribose nucleic acid. This structure has two helical chains each coiled round the same axis (see diagram). We have made the usual chemical assumptions, namely, that each chain consists of phosphate di-ester groups joining β -D-deoxy-ribofuranose residues with 3',5' linkages. The two chains (but not their bases) are related by a dyad perpendicular to the fibre axis. Both chains follow right-handed helices, but owing to the dyad the sequences of the atoms in the two chains run in opposite directions. Each chain loosely resembles Furberg's² model No. 1; that is, the bases are on the inside of the helix and the phosphates on

is a residue on each chain every 3-4 Å. in the z-direction. We have assumed an angle of 36° between adjacent residues in the same chain, so that the structure repeats after 10 residues on each chain, that is, after 34 Å. The distance of a phosphorus atom from the fibre axis is 10 Å. As the phosphates are on the outside, cations have easy access to them.

The structure is an open one, and its water content is rather high. At lower water contents we would expect the bases to tilt so that the structure could become more compact.

The novel feature of the structure is the manner in which the two chains are held together by the purine and pyrimidine bases. The planes of the bases are perpendicular to the fibre axis. They are joined together in pairs, a single base from one chain being hydrogen-bonded to a single base from the other chain, so that the two lie side by side with identical z-co-ordinates. One of the pair must be a purine and the other a pyrimidine for bonding to occur. The hydrogen bonds are made as follows: purine position 1 to pyrimidine position 1; purine position 6 to pyrimidine position 6.

If it is assumed that the bases only occur in the structure in the most plausible tautomeric forms (that is, with the keto rather than the enol configurations) it is found that only specific pairs of bases can bond together. These pairs are: adenine (purine) with thymine (pyrimidine), and guanine (purine) with cytosine (pyrimidine).

In other words, if an adenine forms one member of a pair, on either chain, then on these assumptions the other member must be thymine; similarly for guanine and cytosine. The sequence of bases on a single chain does not appear to be restricted in any way. However, if only specific pairs of bases can be formed, it follows that if the sequence of bases on one chain is given, then the sequence on the other chain is automatically determined.

It has been found experimentally^{3,4} that the ratio of the amounts of adenine to thymine, and the ratio of guanine to cytosine, are always very close to unity for deoxyribose nucleic acid.

It is probably impossible to build this structure with a ribose sugar in place of the deoxyribose, as the extra oxygen atom would make too close a van der Waals contact.

The previously published X-ray data^{5,6} on deoxyribose nucleic acid are insufficient for a rigorous test of our structure. So far as we can tell, it is roughly compatible with the experimental data, but it must be regarded as unproved until it has been checked against more exact results. Some of these are given in the following communications. We were not aware of the details of the results presented there when we devised our structure, which rests mainly though not entirely on published experimental data and stereochemical arguments.

It has not escaped our notice that the specific pairing we have postulated immediately suggests a possible copying mechanism for the genetic material.

Full details of the structure, including the conditions assumed in building it, together with a set of co-ordinates for the atoms, will be published elsewhere.



Note



Highlight



Screenshot



Flag

Publication



A Structure for Deoxyribose Nucleic Acid



Francis Crick



James Watson



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Tools

Conversations

Re: A Structure for Deoxyribose Nucleic Acid



updated 20 minutes ago

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Re: Physical Principles for Scalable Neural Rec...



updated 9 years ago

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Re: Millisecond-timescale, genetically targeted ...



updated 9 years ago

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Re: A Structure for Deoxyribose Nucleic Acid



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Francis Crick



James Watson



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Tools



Conversations

Re: A Structure for Deoxyribose Nucleic Acid



updated 20 minutes ago

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Re: Physical Principles for Scalable Neural Rec...



updated 9 years ago

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Re: Millisecond-timescale, genetically targeted ...



updated 9 years ago

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Re: A Structure for Deoxyribose Nucleic Acid



updated 20 minutes ago

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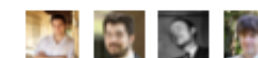
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Francis Crick



James Watson



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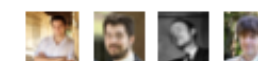
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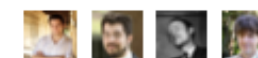
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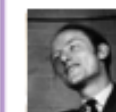
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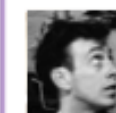
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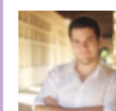
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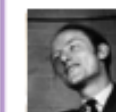
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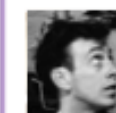
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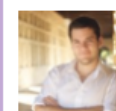
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A Structure for Deoxyribose Nucleic Acid



Francis Crick



James Watson



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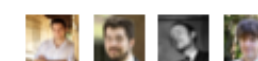
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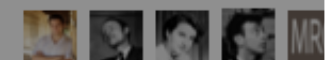
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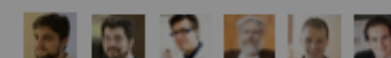
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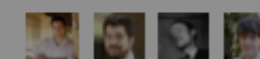
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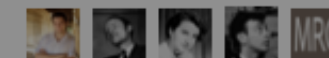
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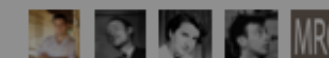
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is a residue on each chain every 3-4 Å. in the z-direction. We have assumed an angle of 36° between adjacent residues in the same chain, so that the

Figure from: A Structure for Deoxyribose Nucleic Acid

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See figure [Fig 1](beagle.io/AT2gyy5okh/Figure-from-A-Structure-for-Deoxyribose-Nucleic-Acid)

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equipment, and to Dr. G. E. R. Deacon and the captain and officers of R.R.S. *Discovery II* for their part in making the observations.

¹ Young, F. B., Gerrard, H., and Jevons, W., *Phil. Mag.*, **40**, 149 (1920).

² Longuet-Higgins, M. S., *Mon. Not. Roy. Astro. Soc., Geophys. Supp.*, **5**, 285 (1949).

³ Von Arx, W. S., *Woods Hole Papers in Phys. Oceanog. Meteor.*, **11** (3) (1950).

⁴ Ekman, V. W., *Arkiv. Mat. Astron. Fysik. (Stockholm)*, **2** (11) (1905).

MOLECULAR STRUCTURE OF NUCLEIC ACIDS

A Structure for Deoxyribose Nucleic Acid

WE wish to suggest a structure for the salt of deoxyribose nucleic acid (D.N.A.). This structure has novel features which are of considerable biological interest.

A structure for nucleic acid has already been proposed by Pauling and Corey¹. They kindly made their manuscript available to us in advance of publication. Their model consists of three intertwined chains, with the phosphates near the fibre axis, and the bases on the outside. In our opinion, this structure is unsatisfactory for two reasons: (1) We believe that the material which gives the X-ray diagrams is the salt, not the free acid. Without the acidic hydrogen atoms it is not clear what forces would hold the structure together, especially as the negatively charged phosphates near the axis will repel each other. (2) Some of the van der Waals distances appear to be too small.

Another three-chain structure has also been suggested by Fraser (in the press). In his model the phosphates are on the outside and the bases on the inside, linked together by hydrogen bonds. This structure as described is rather ill-defined, and for this reason we shall not comment on it.

We wish to put forward a radically different structure for the salt of deoxyribose nucleic acid. This structure has two helical chains each coiled round the same axis (see diagram). We have made the usual chemical assumptions, namely, that each chain consists of phosphate di-ester groups joining β -D-deoxy-ribofuranose residues with 3',5' linkages. The two chains (but not their bases) are related by a dyad perpendicular to the fibre axis. Both chains follow right-handed helices, but owing to the dyad the sequences of the atoms in the two chains run in opposite directions. Each chain loosely resembles Furbert's² model No. 1; that is, the bases are on the inside of the helix and the phosphates on

is a residue on each chain every 3.4 Å. in the z-direction. We have assumed an angle of 36° between adjacent residues in the same chain, so that the structure repeats after 10 residues on each chain, that is, after 34 Å. The distance of a phosphorus atom from the fibre axis is 10 Å. As the phosphates are on the outside, cations have easy access to them.

The structure is an open one, and its water content is rather high. At lower water contents we would expect the bases to tilt so that the structure could become more compact.

The novel feature of the structure is the manner in which the two chains are held together by the purine and pyrimidine bases. The planes of the bases are perpendicular to the fibre axis. They are joined together in pairs, a single base from one chain being hydrogen-bonded to a single base from the other chain, so that the two lie side by side with identical z-co-ordinates. One of the pair must be a purine and the other a pyrimidine for bonding to occur. The hydrogen bonds are made as follows: purine position 1 to pyrimidine position 1; purine position 6 to pyrimidine position 6.

If it is assumed that the bases only occur in the structure in the most plausible tautomeric forms (that is, with the keto rather than the enol configurations) it is found that only specific pairs of bases can bond together. These pairs are: adenine (purine) with thymine (pyrimidine), and guanine (purine) with cytosine (pyrimidine).

In other words, if an adenine forms one member of a pair, on either chain, then on these assumptions the other member must be thymine; similarly for guanine and cytosine. The sequence of bases on a single chain does not appear to be restricted in any way. However, if only specific pairs of bases can be formed, it follows that if the sequence of bases on one chain is given, then the sequence on the other chain is automatically determined.

It has been found experimentally^{3,4} that the ratio of the amounts of adenine to thymine, and the ratio of guanine to cytosine, are always very close to unity for deoxyribose nucleic acid.

It is probably impossible to build this structure with a ribose sugar in place of the deoxyribose, as the extra oxygen atom would make too close a van der Waals contact.

The previously published X-ray data^{5,6} on deoxyribose nucleic acid are insufficient for a rigorous test of our structure. So far as we can tell, it is roughly compatible with the experimental data, but it must be regarded as unproved until it has been checked against more exact results. Some of these are given in the following communications. We were not aware of the details of the results presented there when we devised our structure, which rests mainly though not entirely on published experimental data and stereochemical arguments.

It has not escaped our notice that the specific pairing we have postulated immediately suggests a possible copying mechanism for the genetic material.

Full details of the structure, including the conditions assumed in building it, together with a set of co-ordinates for the atoms, will be published elsewhere.



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