



# Lecture 7:

## DNA: Replication Experiments

Course 281

# Lessons for life

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"Don't lose the moon  
while counting the stars."

# AIMS

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- Understand why DNA needs to be replicated.
- Understand the model of DNA replication and the experiment that led to the discovery.
- Understand Meselson and Stahl experiment.

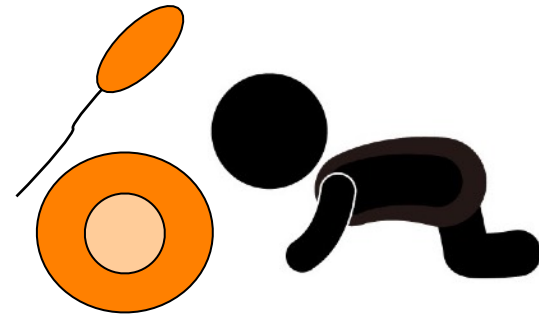
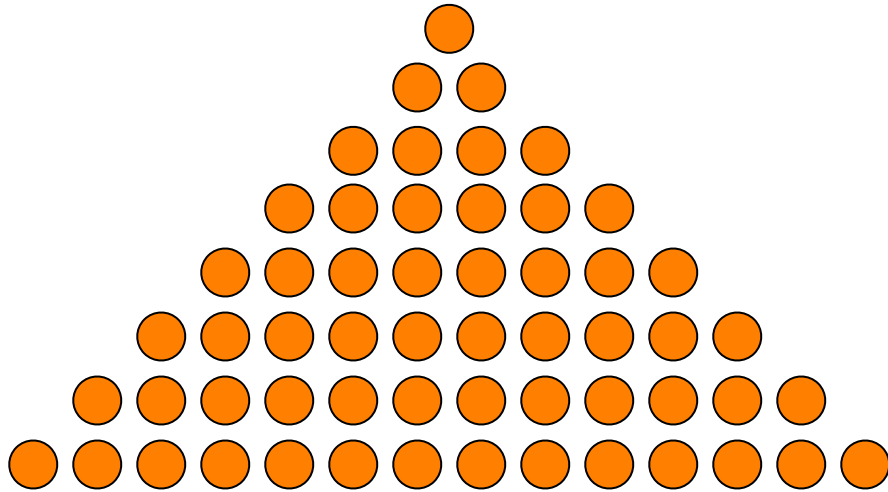
# Why DNA replication?

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- DNA is the genetic material.
  - Pass genetic material from one generation of cells to the next (growth/prokaryotes).
  - Pass genetic material from one generation of organisms to the next (multicellular).

# Why DNA replication?

**DNA replication is transmission and growth!**



# DNA structure and replication



- The discovery of DNA structure hinted to a mechanism of DNA replication.
- Watson and Crick mentioned this in their publication.
- **What parts of the DNA structure give hints?**
  - Double strands.
  - Complementary base-pairing.

# DNA structure and replication

No. 4356 April 25, 1953

NATURE

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## 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 Corey<sup>1</sup>. 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 diester groups joining  $\beta$ -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 Furlberg's<sup>2</sup> model No. 1; that is, the bases are on the inside of the helix and the phosphates on the outside. The configuration of the sugar and the atoms near it is close to Furlberg's 'standard configuration', the sugar being roughly perpendicular to the attached base. There



This figure is purely diagrammatic. The two ribbons symbolize the two phosphate-sugar chains, and the horizontal rods the pairs of bases holding the chains together. The vertical line marks the fibre axis.

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 need to be stated 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<sup>3,4</sup> 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<sup>5,6</sup> 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 unpublished experimental data and stereo-

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.

We are much indebted to Dr. Jerry Donohue for constant advice and criticism, especially on inter-atomic distances. We have also been stimulated by a knowledge of the general nature of the unpublished experimental results and ideas of Dr. M. H. F. Wilkins, Dr. R. E. Franklin and their co-workers at

King's College, London. One of us (J. D. W.) has been aided by a fellowship from the National Foundation for Infantile Paralysis.

J. D. WATSON  
F. H. C. CRICK

Medical Research Council Unit for the Study of the Molecular Structure of Biological Systems, Cavendish Laboratory, Cambridge, April 2.

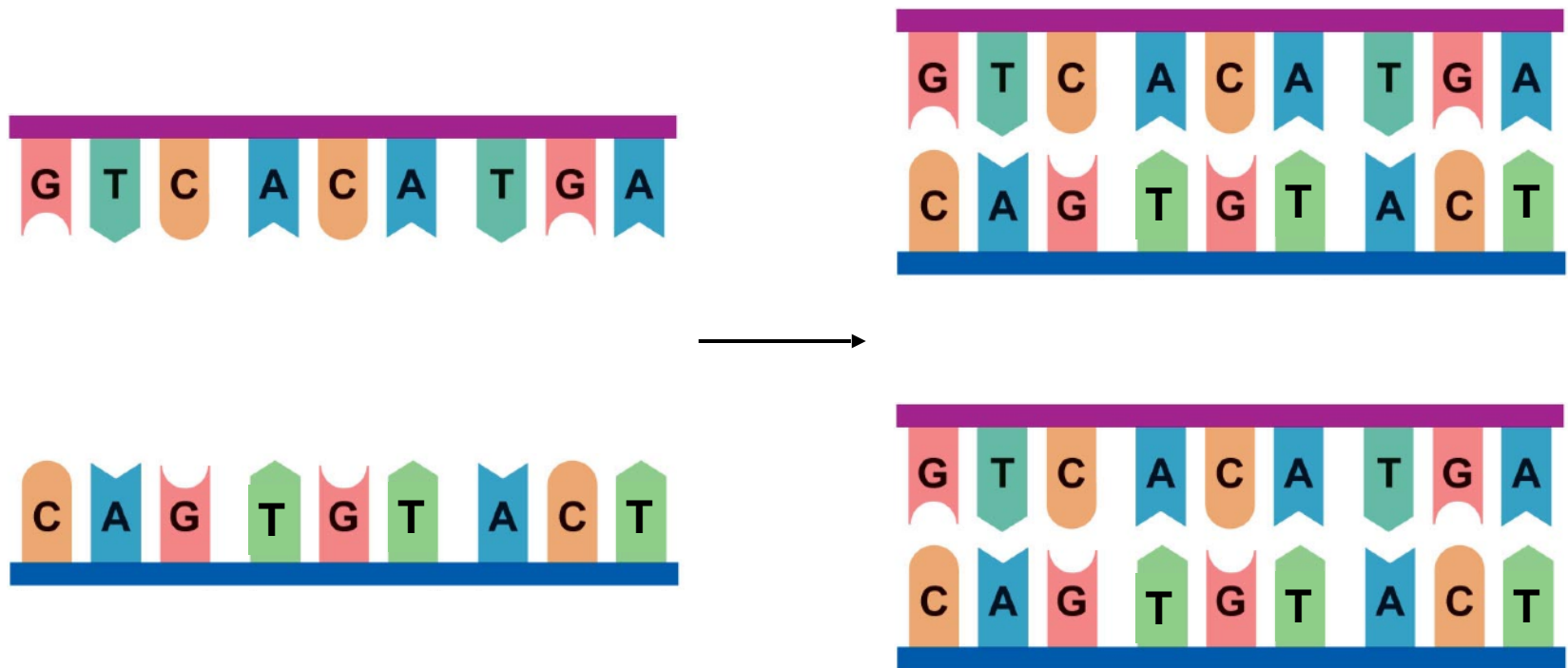
<sup>1</sup> Pauling, L., and Corey, R. B., *Nature*, 171, 346 (1953); *Proc. U.S. Nat. Acad. Sci.*, 39, 84 (1953).  
<sup>2</sup> Furlberg, S., *Acta Chem. Scand.*, 6, 434 (1952).  
<sup>3</sup> Chargaff, E., for references see Zamenhof, S., Braverman, G., and Chargaff, E., *Biochim. et Biophys. Acta*, 8, 402 (1952).  
<sup>4</sup> Wyatt, G. R., *J. Gen. Physiol.*, 36, 201 (1952).  
<sup>5</sup> Astbury, W. T., *Symp. Soc. Exp. Biol.*, 1, Nucleic Acid, 66 (Camb. Univ. Press, 1947).  
<sup>6</sup> Wilkins, M. H. F., and Randall, J. T., *Biochim. et Biophys. Acta*, 10, 132 (1953).

## Each strand can serve as the template to be copied

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

# Do you remember DNA structure?

If we separate the strands, we can you predict the complementary sequence!



**Each strand can serve as the template to be copied**



# Replication

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- We will go over DNA replication in prokaryotes and eukaryotes covering:
  - Process.
  - Enzymes involved.

**BUT**

**Before that when need to know what is the model of DNA replication**

# How is DNA replicated?

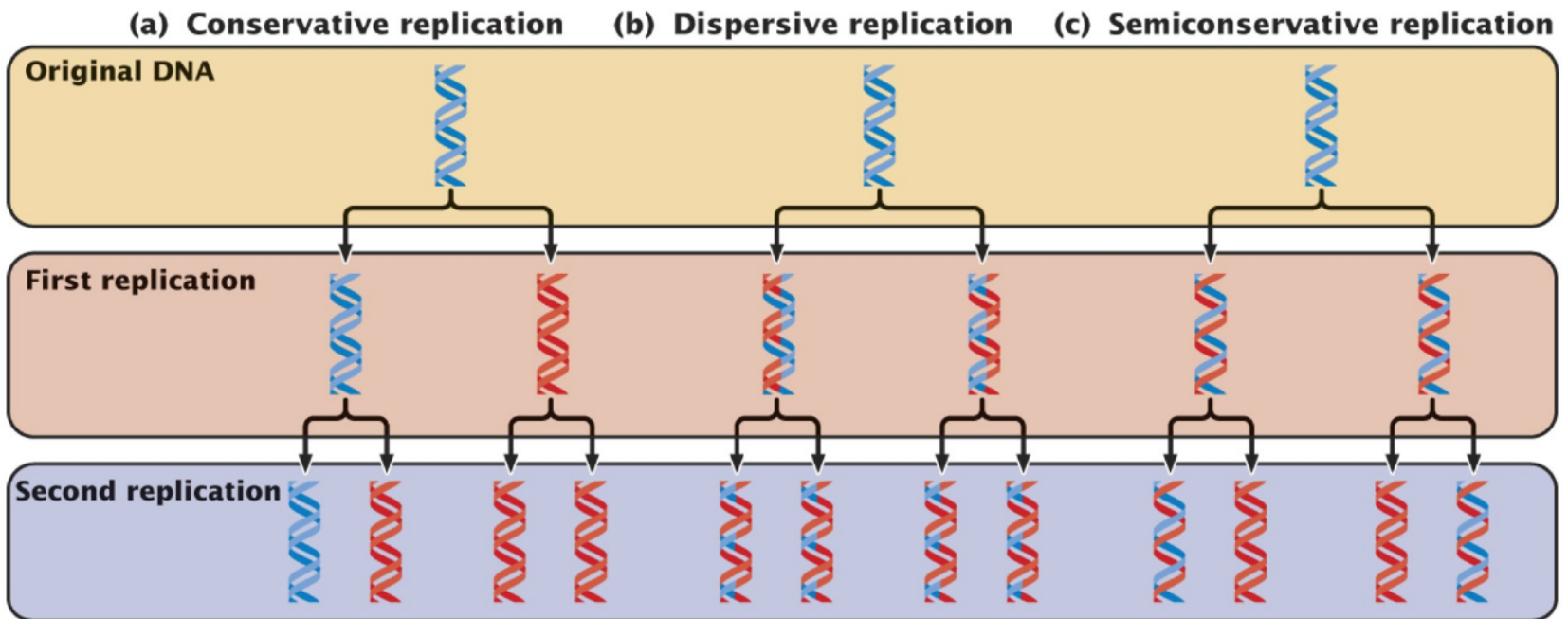


## Three hypotheses:

1. Conservative replication: replicated DNA results in one old double strand DNA and one new double strands of DNA.
2. Dispersive replication: replicated DNA results in each double strands with segments of old and new DNA.
3. Semi-conservative replication: each double strand has one old strand and one new strand.

# How is DNA replicated?

- **Three hypotheses:**
  - Conservative replication
  - Dispersive replication
  - Semi-conservative replication



# Meselson and Stahl experiment

- Matt Meselson and Frank Stahl designed **the most beautiful experiment in biology.**
- The experiment tests the DNA replication models.
- They used bacteria grown in a media of a heavy isotope of nitrogen .



# Meselson and Stahl experiment

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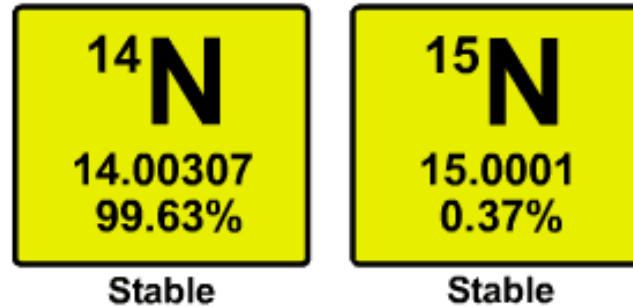
**Do you know heavy isotopes?**

# Meselson and Stahl experiment



- In nature, some elements have different isotopes which they differ in the number of neutrons.
- The difference in the number of neutrons makes the isotopes' atoms heavier.

# Meselson and Stahl experiment



- The abundant nitrogen is  $\text{N}^{14}$  where there are 7 protons and 7 neutrons and it weights 14g/mole.
- $\text{N}^{15}$  is a stable isotope of nitrogen and has 7 protons and 8 neutrons and weights 15g/mole.

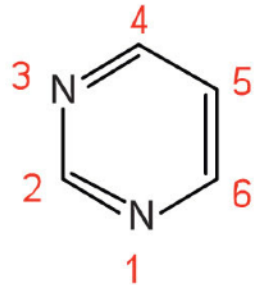
# Meselson and Stahl experiment

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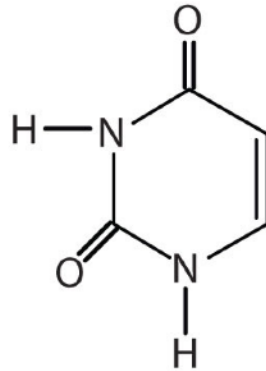
**Why heavy nitrogen isotope N<sup>15</sup>?**



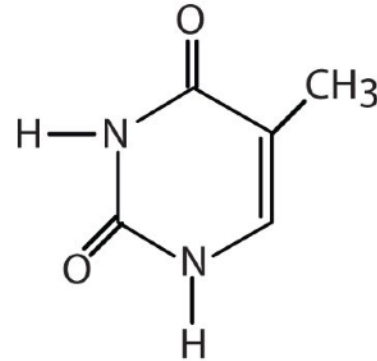
# Meselson and Stahl experiment



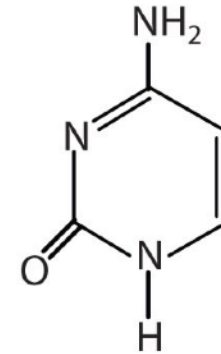
Pyrimidine



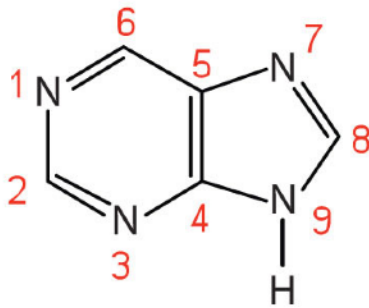
Uracil (U)  
RNA only



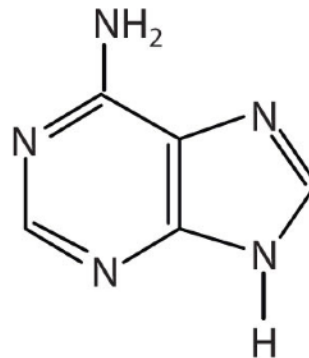
Thymine (T)  
DNA only



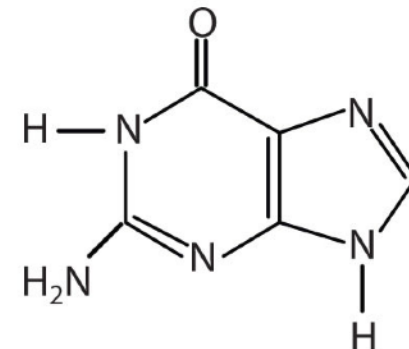
Cytosine (C)  
both DNA and RNA



Purine



Adenine (A)



Guanine (G)

Remember DNA bases contain **Nitrogen**

# Meselson and Stahl experiment

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## The experiment:

1) Grow bacteria in a media containing the heavy  $N^{15}$  for many generations.

**Result:** all bacteria has  $N^{15}$  in their DNA.

# Meselson and Stahl experiment



## The experiment:

2) Transfer some  $N^{15}$  bacteria to grow in  $N^{14}$  media and allow to grow for several generations.

**Result:** DNA of newly divided bacteria will have  $N^{14}$  instead of  $N^{15}$ .

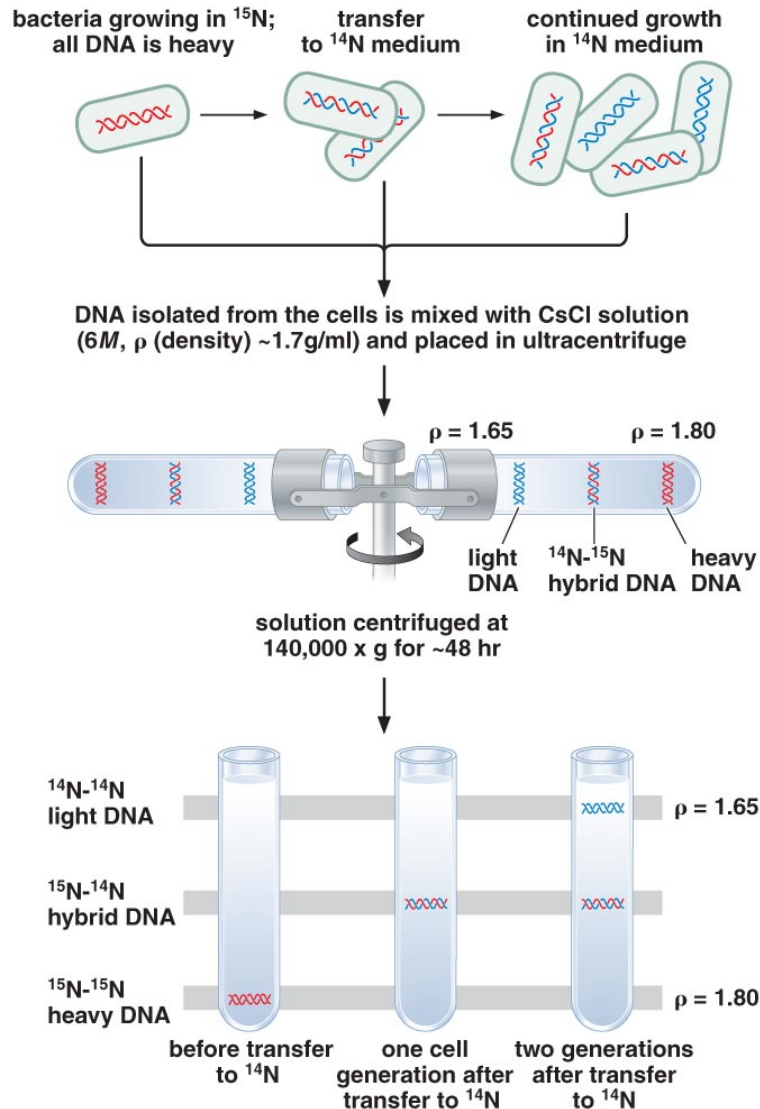
# Meselson and Stahl experiment

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## The experiment:

3) Take samples from the growing bacteria at different time and study the density of the DNA of the cells.

# Meselson and Stahl experiment



Simply the idea is to mark the old DNA and see what happens to the newly synthesized DNA.

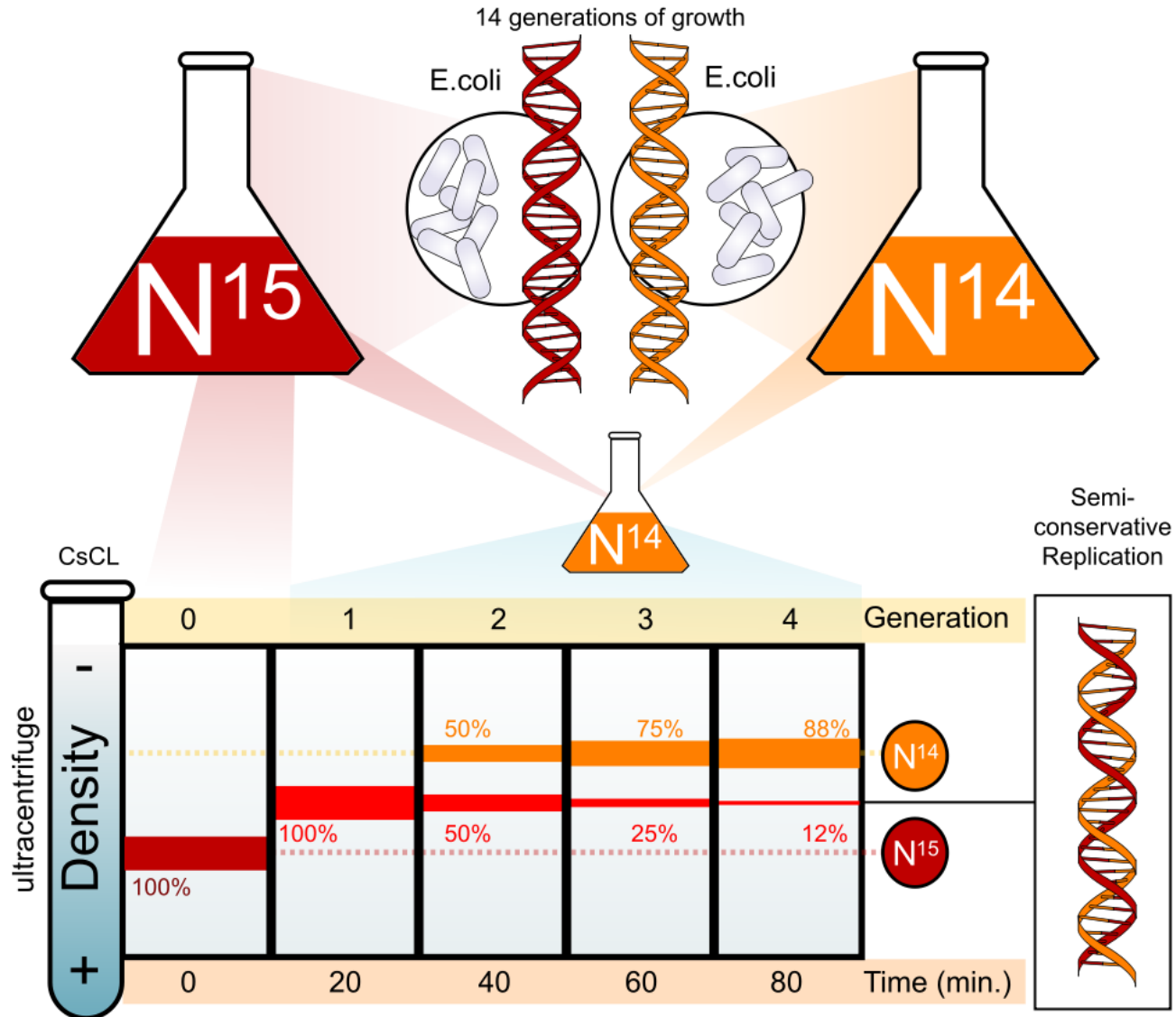
the location of DNA molecules within the centrifuge cell can be determined by ultraviolet optics

# Meselson and Stahl experiment

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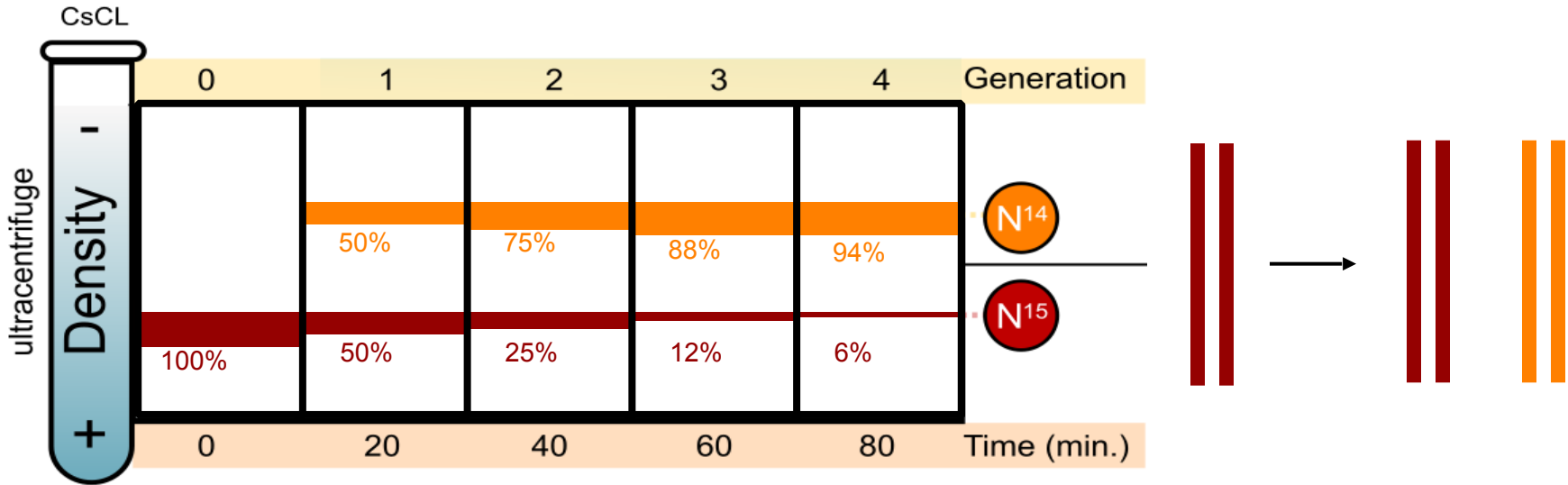
**Let's look in details at the experiment and the expected results of each hypothesis**

# Meselson and Stahl experiment



# What would the results be if another model?

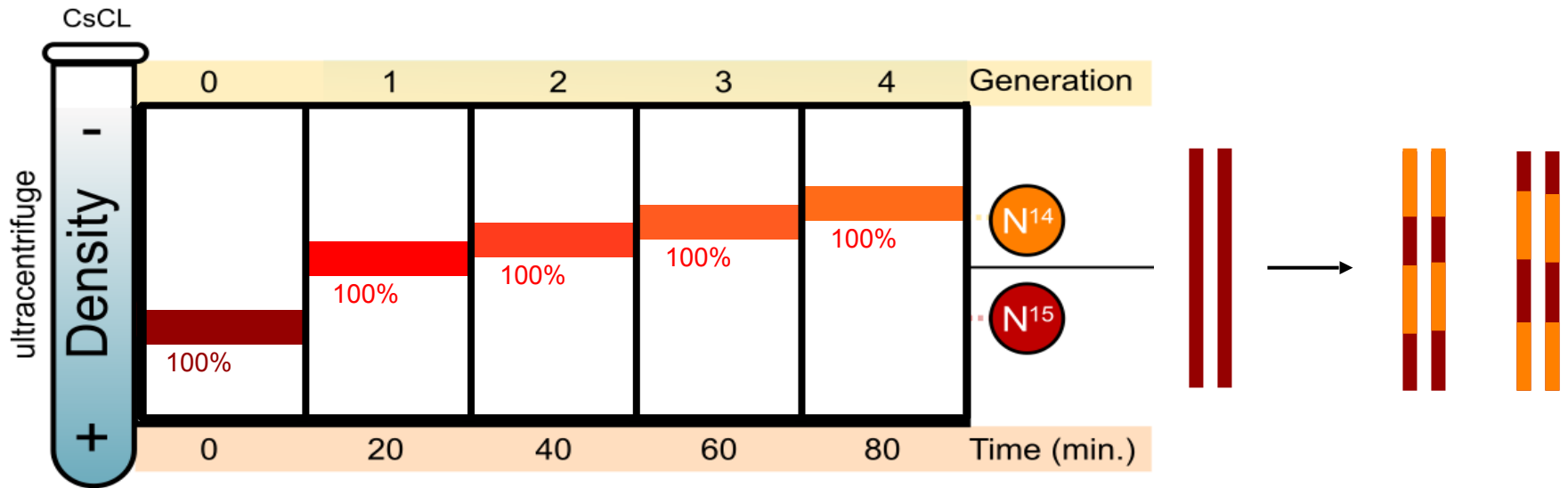
## Conservative replication





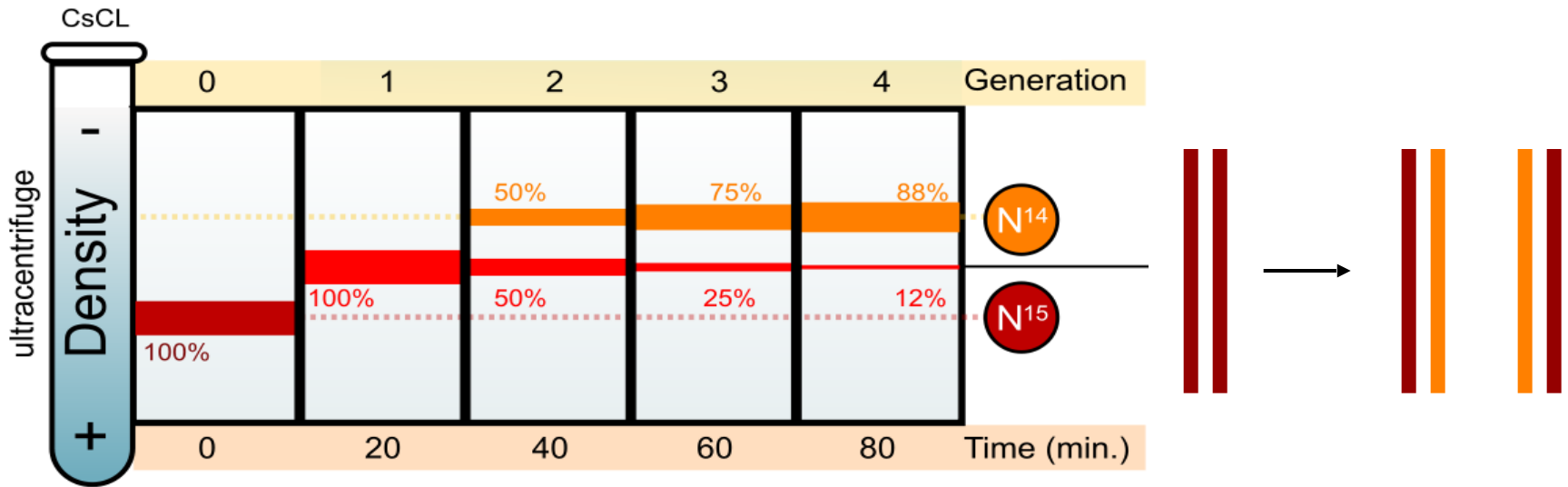
# What would the results be if another model?

## Dispersive replication



# What would the results be if another model?

## Semi-conservative replication



# Quiz

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**Meselson and Stahl's performed the most beautiful experiment in biology and proved that DNA replicates**

- a) Semi-conservatively
- b) Conservatively
- c) Dispersively
- d) (a) and (b) depending on the organism
- e) (a) and (c) depending on the organism

# To study

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N<sup>15</sup>

Conservative model

Dispersive model

Meselson and Stahl experiment

N<sup>14</sup>

Heavy isotope

Semi-conservative model

The most beautiful experiment in biology

DNA replication

# Expectations

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- Know the importance of DNA replication.
- Know the proposed models of DNA replication.
- Know Meselson and Stahl experiment and the proof that DNA replicates in a semi-conservative fashion.

# For a smile

