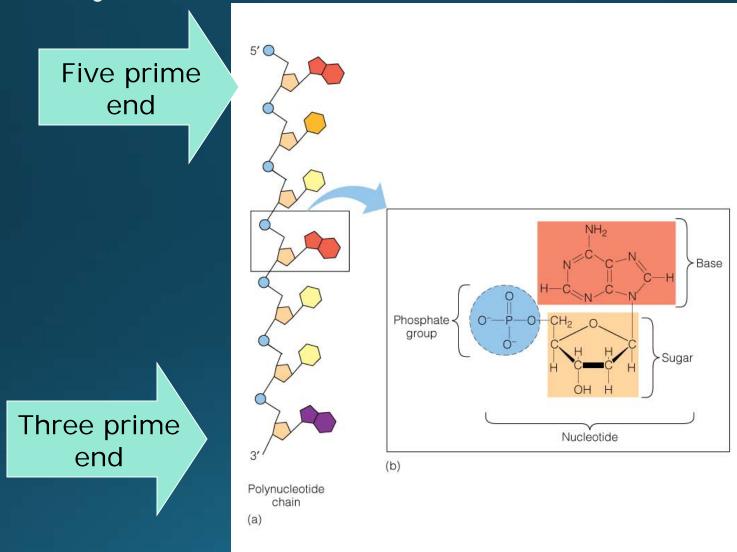
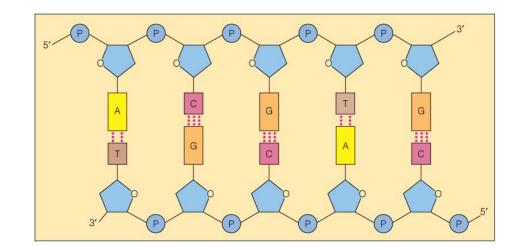
DNA Structure

Polynucleotides Are Directional

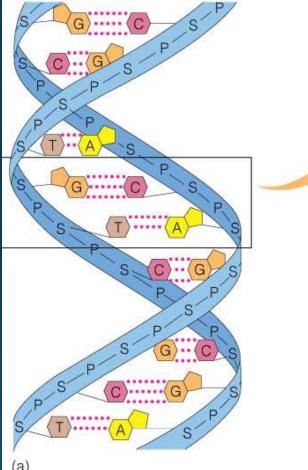


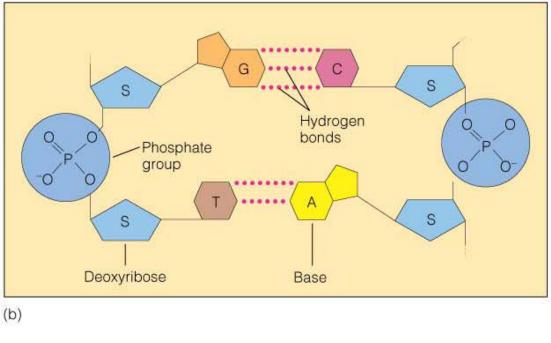
Erwin Chargaff Rule of Base Composition of DNA Amount of A = Amount of T Amount of G = Amount of C

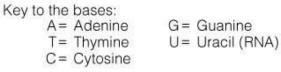
- <u>A forms 2</u>
 <u>hydrogen</u>
 <u>bonds with T</u>
- <u>G forms 3</u>
 <u>hydrogen</u>
 <u>bonds with C</u>



DNA Is a Double-Stranded Helix







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Important Properties of the Model

• <u>Genetic information is stored in the sequence of</u> bases in the DNA

• <u>The model offers a molecular explanation for</u> <u>mutation</u>

• <u>Complementary strands of DNA can be used to</u> <u>explain how DNA copies itself</u>

Molecular Model of DNA

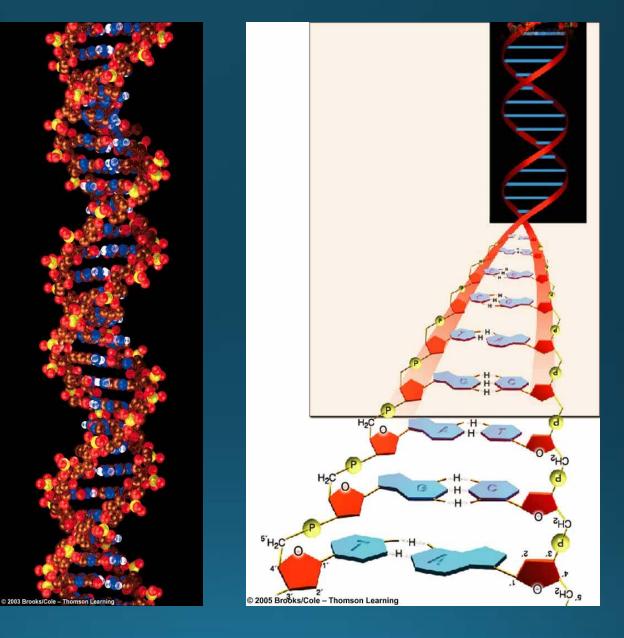
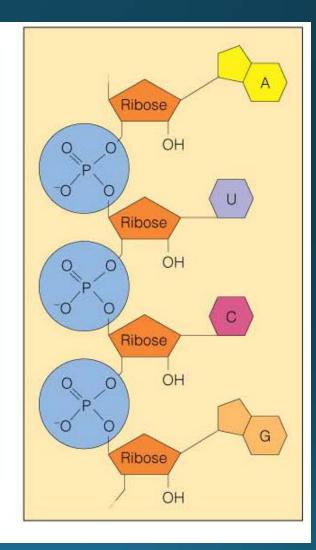


Fig. 8.9

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RNA Is Single Stranded

- <u>Transfers genetic information</u> from the nucleus to the <u>cytoplasm</u>
- <u>Participates in protein</u> <u>synthesis</u>
- It is a component of ribosomes
- <u>Contains ribose sugar and A, U,</u> <u>G, and C</u>



DNA to Chromosomes

DNA stores the genetic information

Mitochondrial chromosome

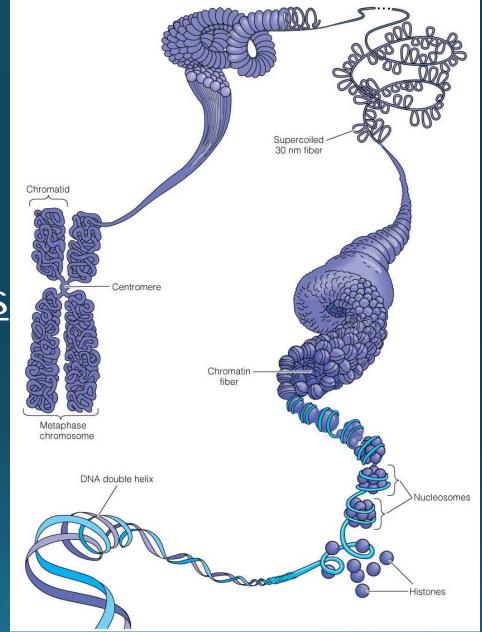
- It is a <u>circular</u> DNA molecule
- It is not compacted
- <u>Similar to prokaryotic chromosomes</u>
- <u>Reflects evolutionary history of mitochondria</u>

<u>Nuclear chromosomes</u>

• Have a <u>complex structure</u>

<u>Nuclear</u> Chromosomes

 DNA is packed into chromosomes by several levels of coiling and <u>compaction</u>



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Nuclear Chromosomes

<u>Chromatin</u>

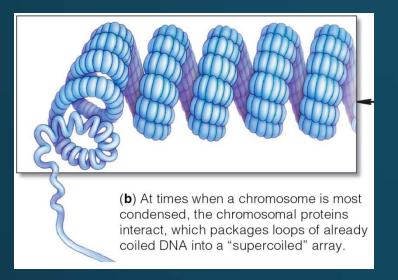
- makes up chromosomes
- it is a <u>complex of DNA and protein</u>

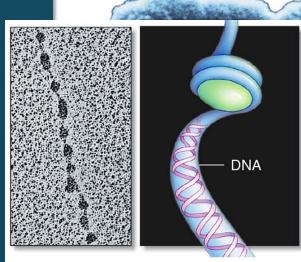
• <u>Histones</u>

- **DNA-binding proteins**
- They <u>assist in compacting and folding</u> DNA into a chromosome
- <u>Shorten</u> the DNA <u>length by a factor of 6 or 7</u>

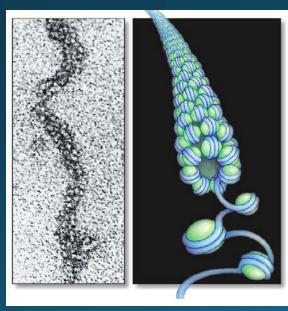
<u>Nucleosomes</u>

• <u>bead-like structures composed of histones wrapped</u> <u>with DNA</u>

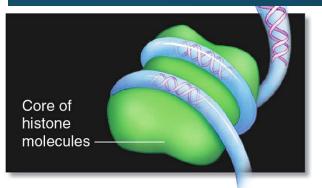




(d) Immerse a chromosome in saltwater and it loosens up to a beads-on-a-string organization. The "string" is one DNA molecule. Each "bead" is a nuclesome.



(c) At a deeper level of structural organization, the chromosomal proteins and DNA are organized as a cylindrical fiber.



(e) A nucleosome consists of part of a DNA molecule looped twice around a core of histones.

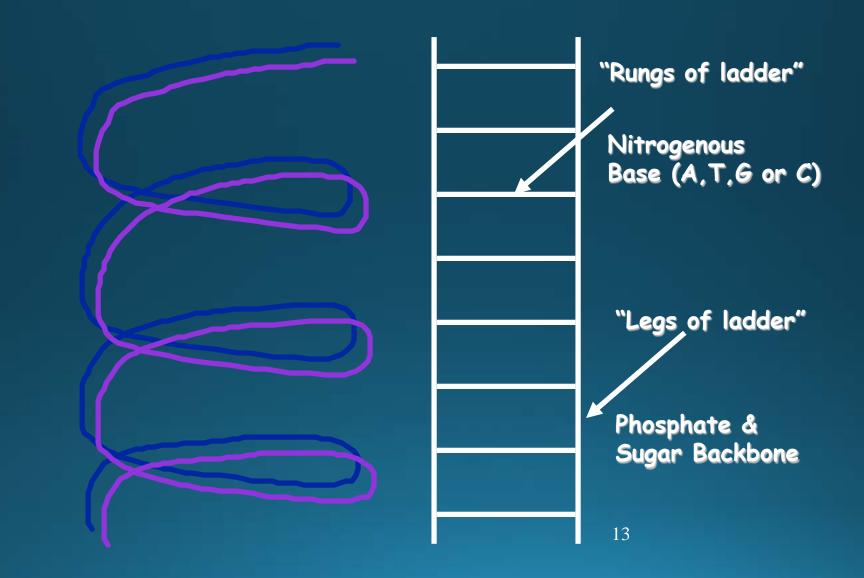
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<u>Two strands coiled called a double helix</u>
<u>Sides made of a pentose sugar Deoxyribose</u>
<u>bonded to phosphate (PO₄) groups by</u>
<u>phosphodiester bonds</u>

•<u>Center</u> made of <u>nitrogen bases</u> bonded together by weak <u>hydrogen bonds</u>

DNA Double Helix



Helix

Most DNA has a <u>right-hand twist</u> with <u>10</u> <u>base pairs in a complete turn</u>

• <u>Left twisted</u> DNA is called <u>Z-DNA</u> or southpaw DNA

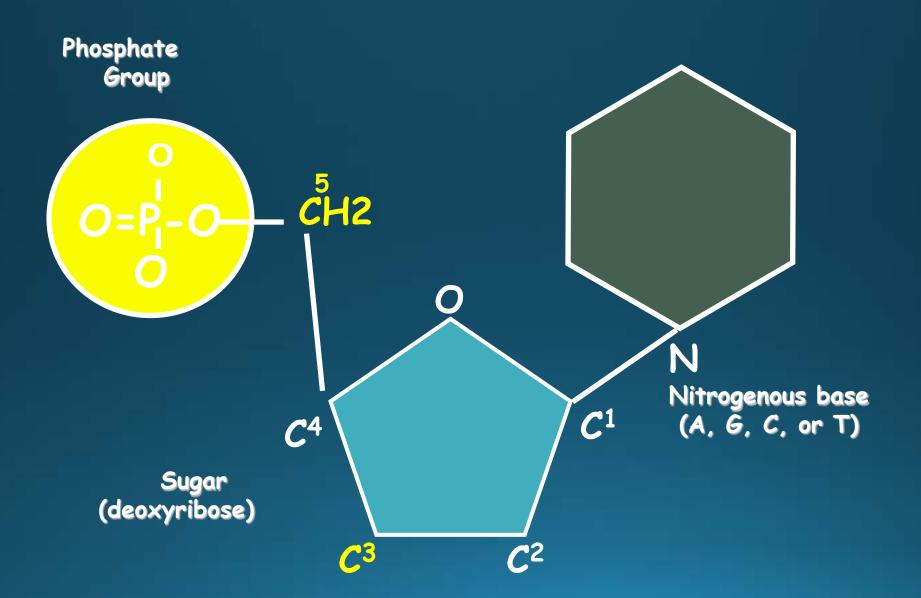
• <u>Hot spots occur where right and left twisted</u> <u>DNA meet producing mutations</u>

DNA

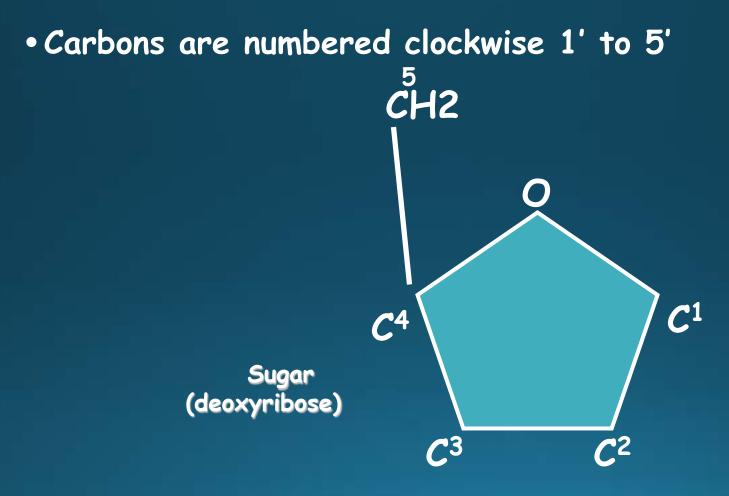
- Stands for Deoxyribonucleic acid
 Made up of subunits called nucleotides
 <u>Nucleotide made of:</u>

 <u>Phosphate group</u>
 - 2. <u>5-carbon sugar</u>
 - 3. Nitrogenous base

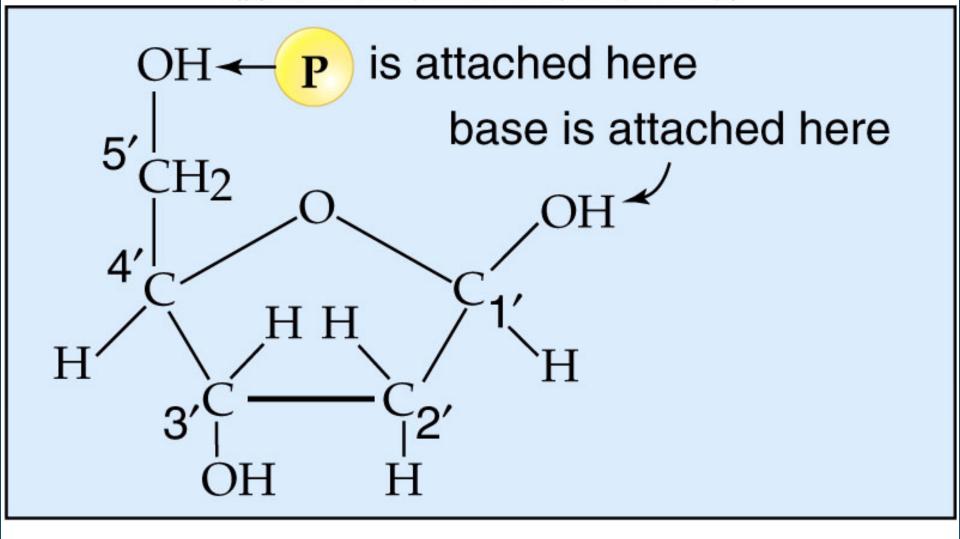
DNA Nucleotide



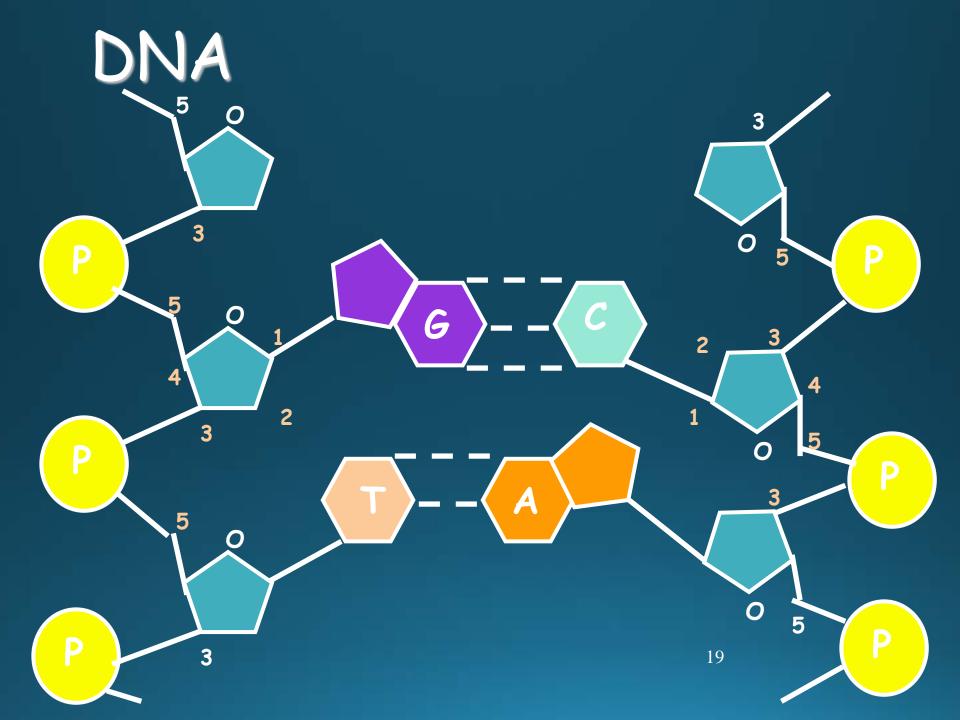
Pentose Sugar



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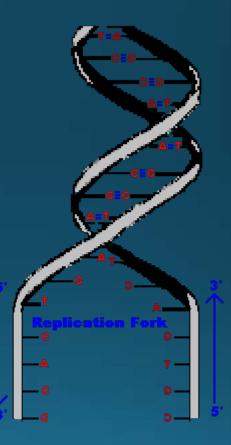
a.



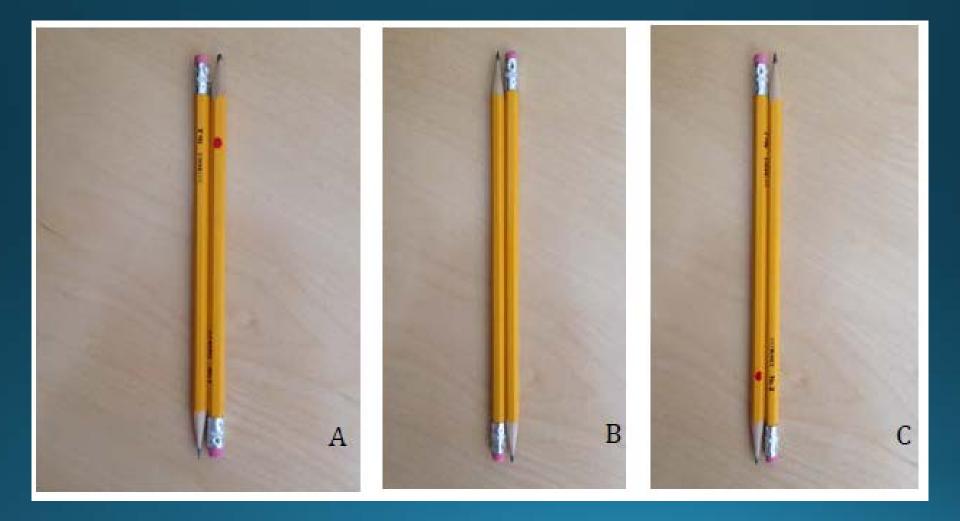
Antiparallel Strands

 One strand of DNA goes from 5' to 3' (sugars)

• The <u>other strand is opposite in</u> <u>direction going 3' to 5' (sugars)</u>



<u>DNA strands have **DYAD symmetry**</u>-pretend two pencils are two strands of DNA. When flipped upside down and backwards, it looks the same as the original.



Nitrogenous Bases

•<u>Double ring **PURINES**</u> <u>Adenine (A)</u> <u>Guanine (G)</u>

•<u>Single ring PYRIMIDINES</u> <u>Thymine (T)</u> <u>Cytosine (C)</u> T or C

A or G

Base-Pairings •Purines only pair with Pyrimidines

Three hydrogen bonds
required to bond Guanine
& Cytosine

3 H-bonds

•Two hydrogen bonds are required to bond Adenine & Thymine



•If there is 30% Adenine, how much Cytosine is present?

Answer:

• There would be 20% Cytosine • Adenine (30%) = Thymine (30%)• Guanine (20%) = Cytosine (20%) • Therefore, 60% A-T and 40% C-G Your turn: On a sheet of paper 1. How many As if there are 23% Gs? 2.How many Cs if there are 44% Ts? **3.How many Ts if there are 31% As?** 4. How many Gs if there are 16% Ts?

DNA Replication

Replication Facts

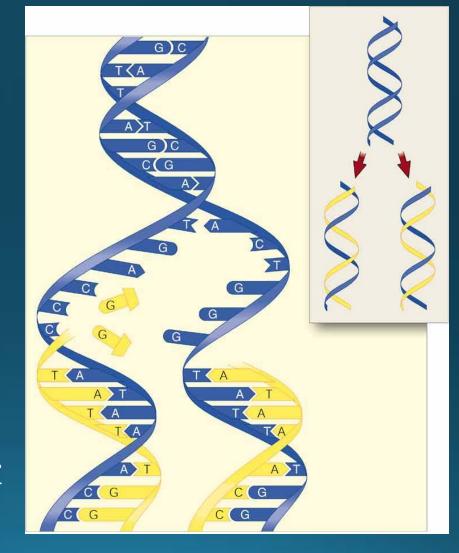
•<u>DNA has to be copied before a cell</u> <u>divides</u>

•<u>New cells will need identical DNA</u> <u>strands</u>

Semi-Conservative Replication of DNA

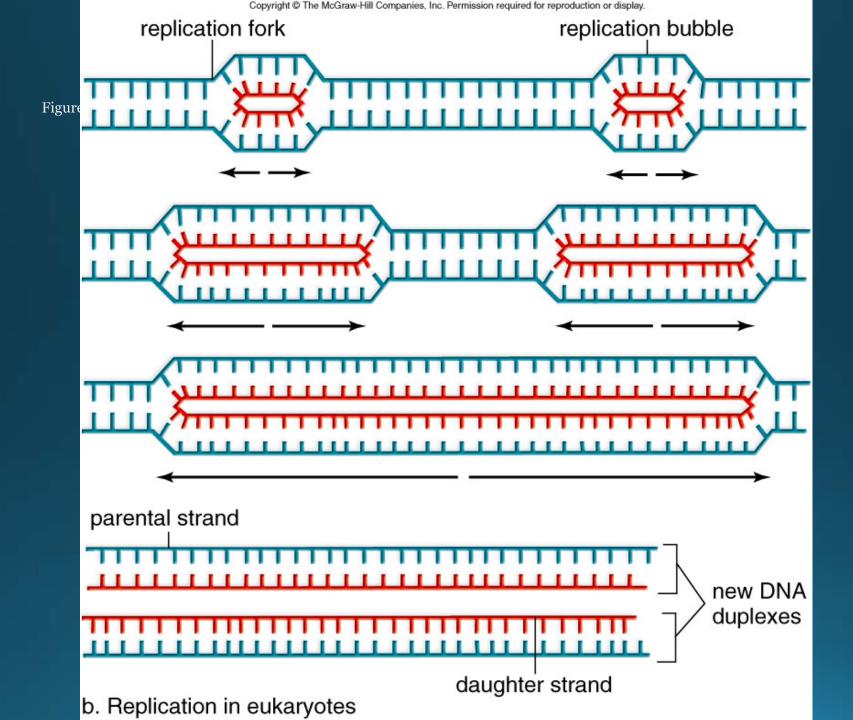
DNA Polymerase
 catalyzes the synthesis of
 DNA using a template of
 DNA and nucleotides

<u>One original strand is</u>
 <u>conserved in each daughter</u>
 <u>molecule</u>



Replication: Prokaryotic vs. Eukaryotic

- <u>Eukaryotic Replication</u>
 - DNA replication <u>begins at numerous points along</u> <u>linear chromosome</u>
 - DNA <u>Unwinds and unzips into two strands</u>
 - <u>Old strand</u> of DNA <u>serves as template for new strand</u>
 - <u>Complementary base-pairing forms new strand</u> on each old strand
 - <u>Replication bubbles spread bi-directionally until they</u>
 <u>meet</u>

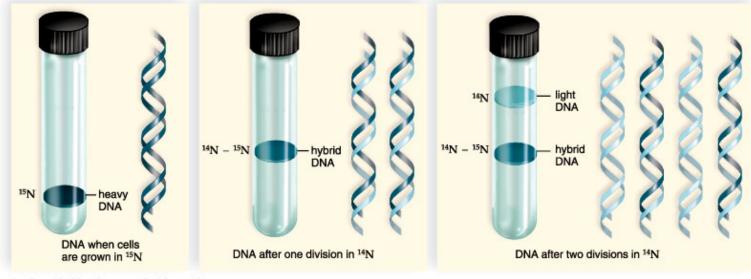


Meselson and Stahl's DNA replication experiment

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a. Possible results when DNA is centrifuged in CsCl

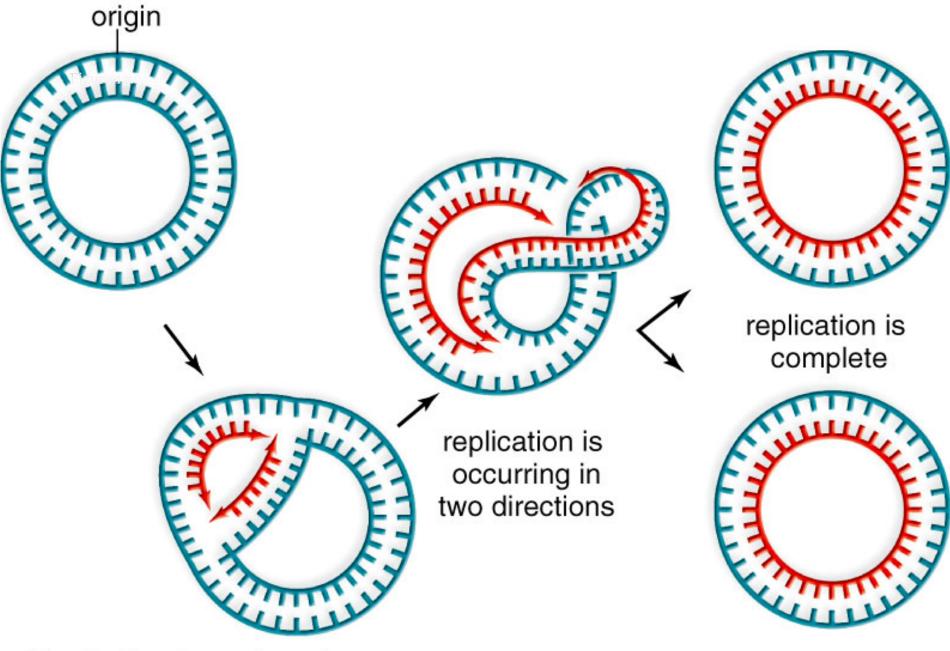


b. Steps in Meselson and Stahl experiment

Replication: Prokaryotic vs. Eukaryotic

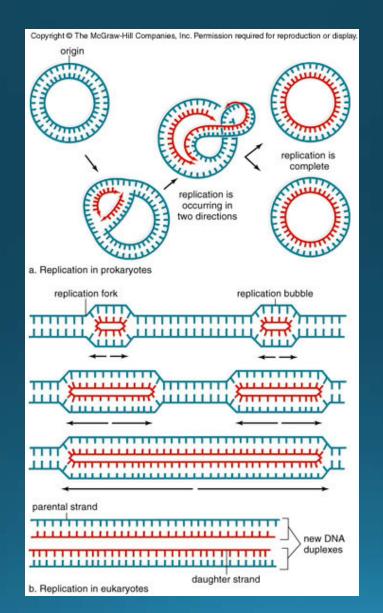
- <u>Prokaryotic Replication</u>
 - <u>Bacteria have a single circular loop</u>
 - <u>Replication moves</u> around the circular DNA molecule <u>in both directions</u>
 - Produces two identical circles
 - <u>Cell divides between circles, as fast as every 20</u> <u>minutes</u>





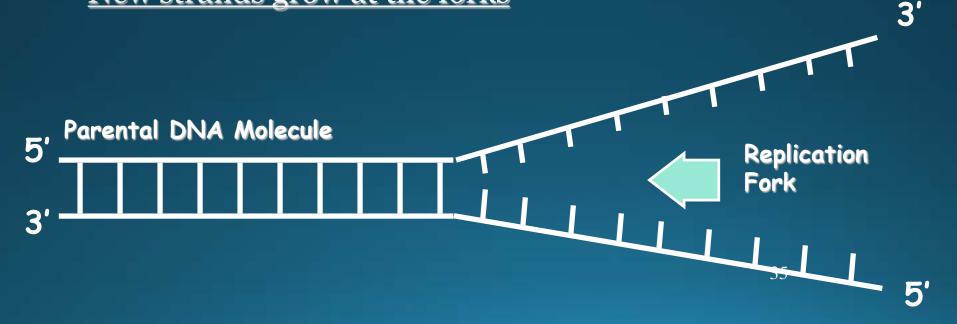
a. Replication in prokaryotes

Replication: Prokaryotic vs. Eukaryotic



DNA Replication

- <u>Begins at Origins of Replication</u>
- <u>Two strands open forming Replication Forks</u> (Yshaped region)
- <u>New strands grow at the forks</u>



DNA Replication

- As the 2 DNA strands open at the origin, <u>Replication</u>
 <u>Bubbles form</u>
- <u>Eukaryotic chromosomes have MANY bubbles</u>
- Prokaryotes (bacteria) have a single bubble



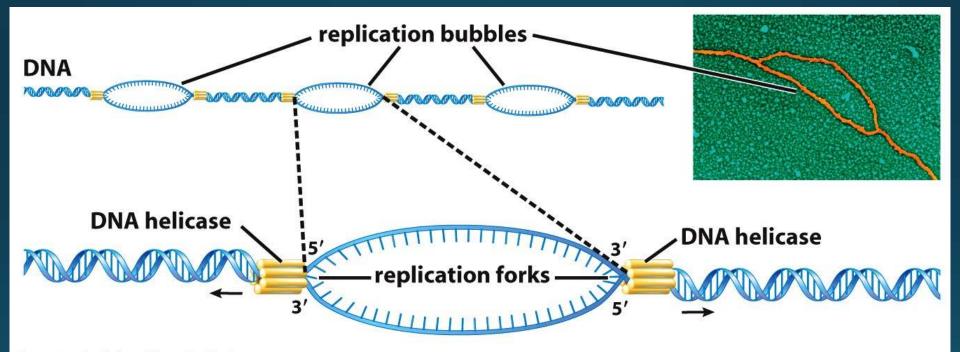
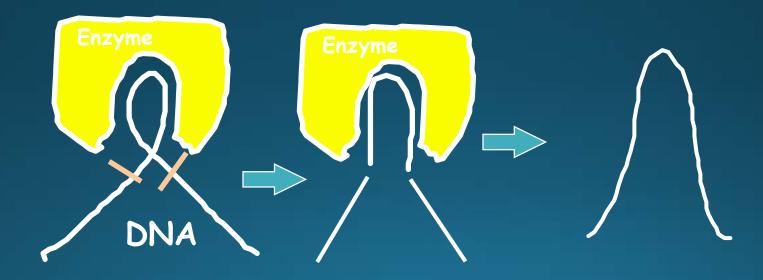


Figure E9-7ab Biology: Life on Earth, 8/e © 2008 Pearson Prentice Hall, Inc.

•Enzyme **DNA Helicase** unwinds and separates the 2 DNA strands by breaking the weak hydrogen bonds

•<u>Single-Strand Binding Proteins (SSBs)</u> attach and <u>keep the</u> 2 DNA <u>strands</u> <u>separated and untwisted</u>

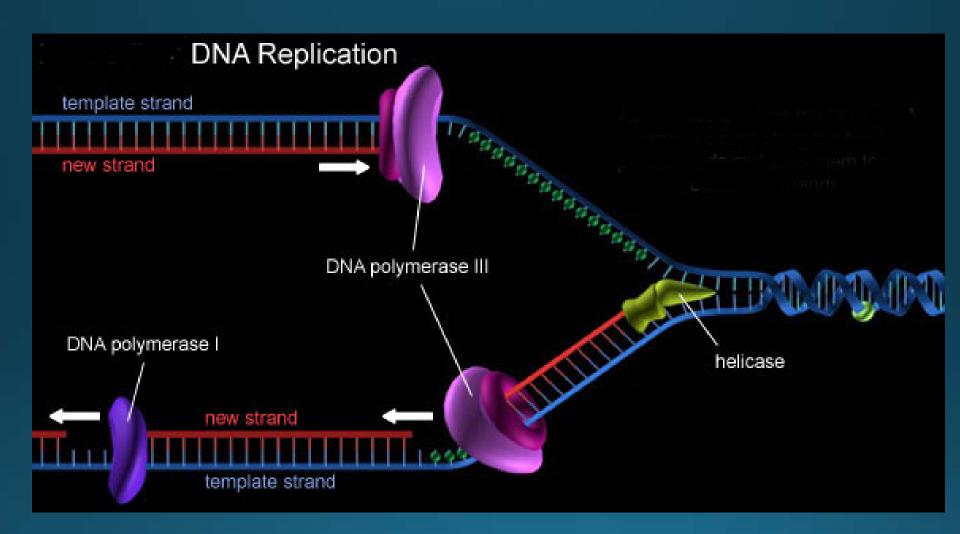
• <u>Enzyme **Topoisomerase**</u> attaches to the 2 forks of the bubble to <u>**relieve stress**</u> on the DNA molecule as it separates



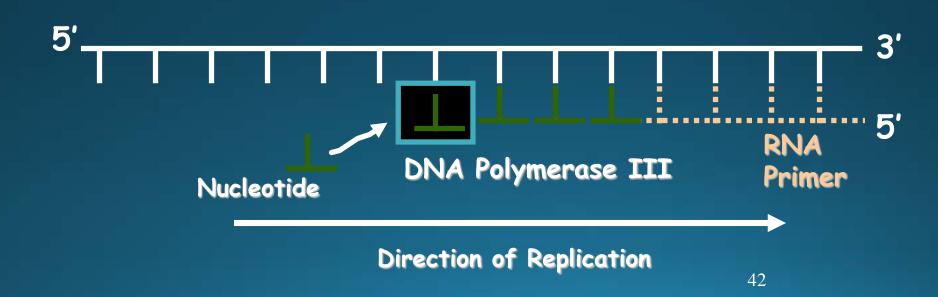
• **Primase** is an enzyme that <u>synthesizes a</u> <u>complimentary RNA Primer</u>

• **<u>RNA primers</u>** bind to serve as a starting point for the addition of new nucleotides

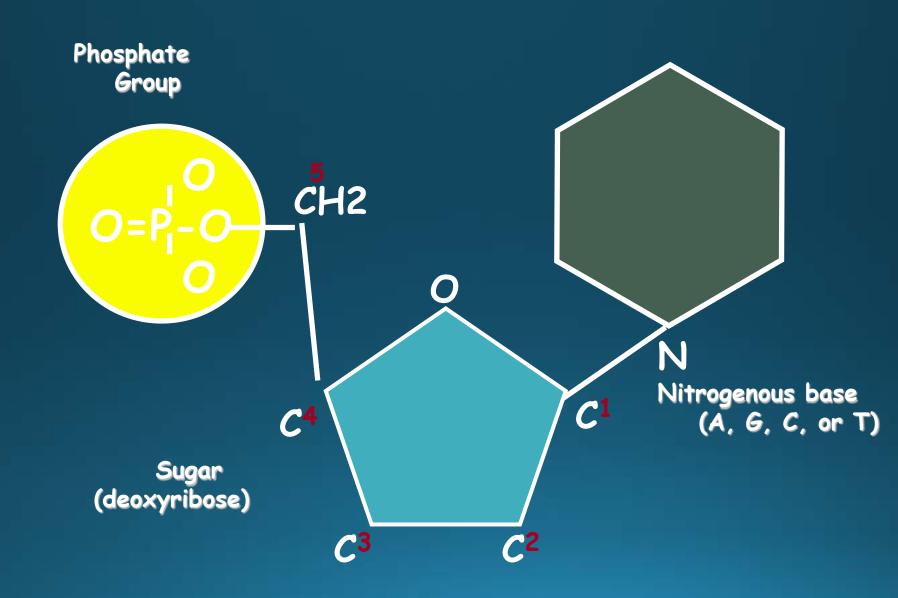
• **DNA polymerase III** can then <u>add the new</u> <u>nucleotides</u>



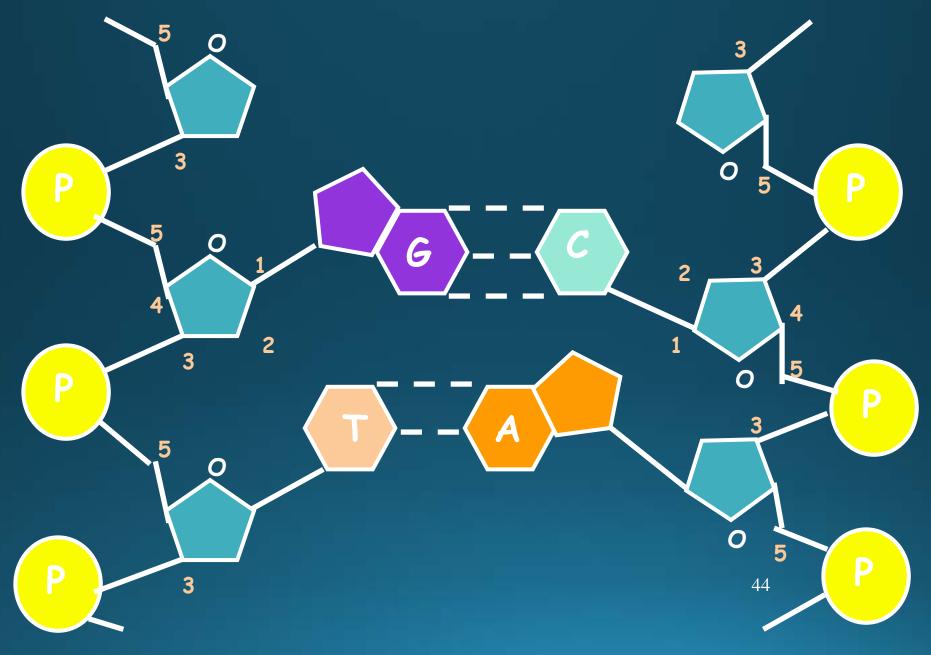
- DNA <u>polymerase</u> III <u>can only add nucleotides to the</u> <u>3' end of the DNA</u>
- This causes the <u>NEW strand to be built in a 5' to 3'</u>
 <u>direction</u>



Remember HOW the Carbons Are Numbered!

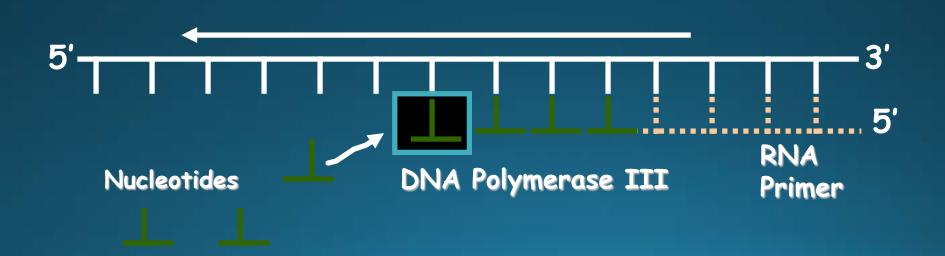


Remember the Strands are Antiparallel



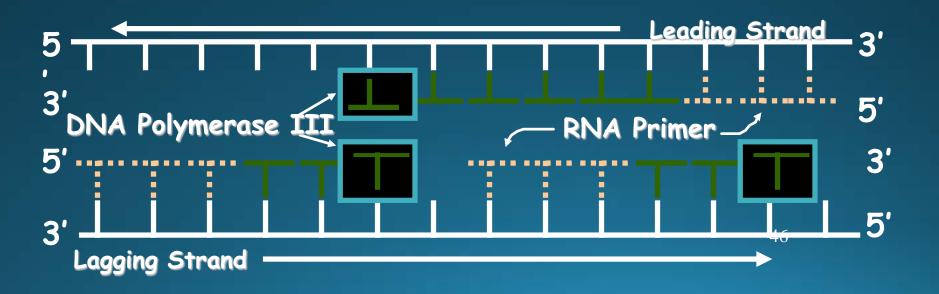
Synthesis of the New DNA Strands

• The <u>Leading Strand</u> is <u>synthesized as a single</u> <u>continuous strand from</u> the point of <u>origin toward</u> the opening <u>replication fork</u>



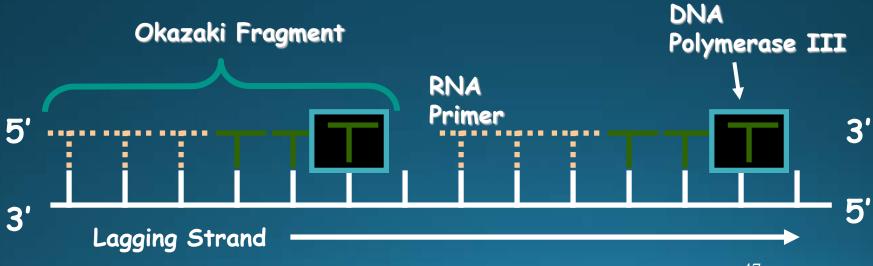
Synthesis of the New DNA Strands

- The <u>Lagging Strand</u> is <u>synthesized discontinuously against</u> overall <u>direction of replication</u>
- This <u>strand is made in MANY short segments</u> It is replicated <u>from the replication fork toward the origin</u>



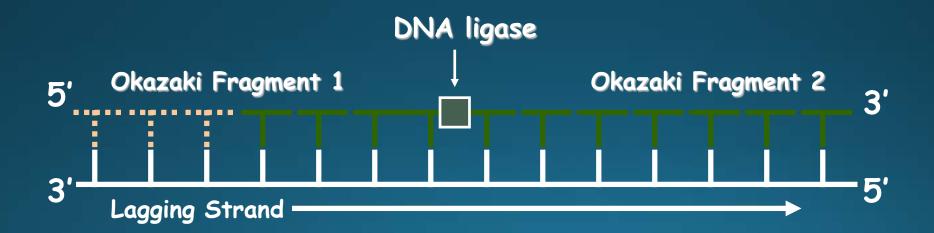
Lagging Strand Segments

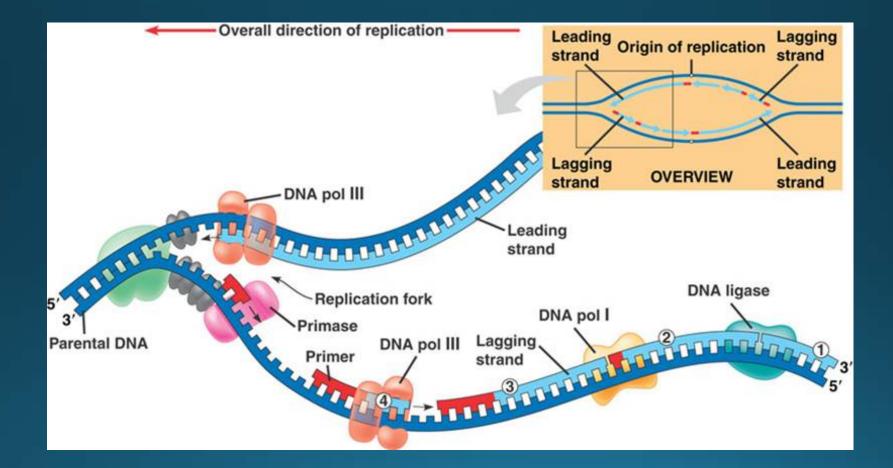
- Okazaki Fragments series of short segments on the lagging strand
- Leaves GAPS between fragments



Joining of Okazaki Fragments

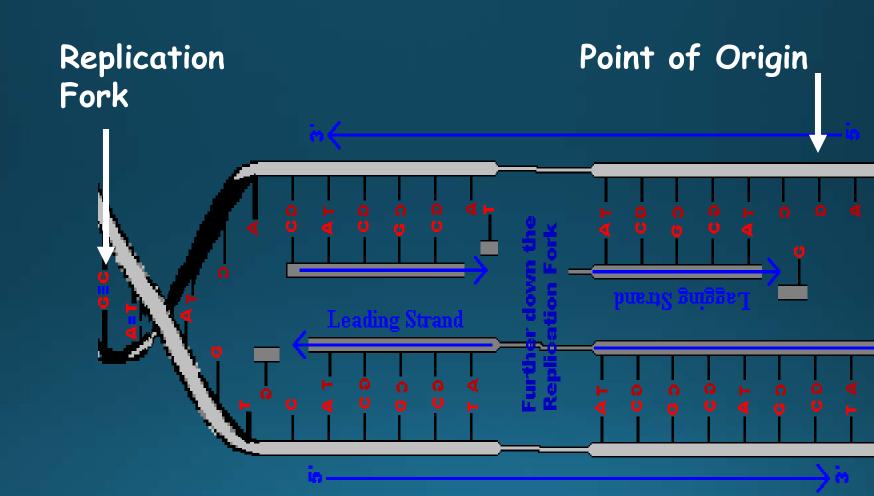
- **DNA Polymerase I** <u>fills in the DNA</u> nucleotides where RNA primers have been removed.
- The enzyme <u>Ligase</u> joins the Okazaki fragments together to make one strand





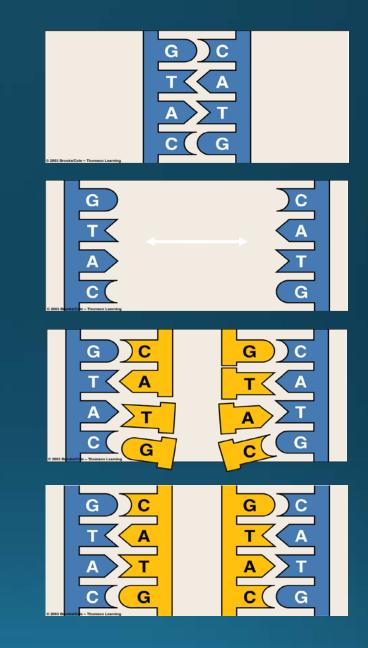
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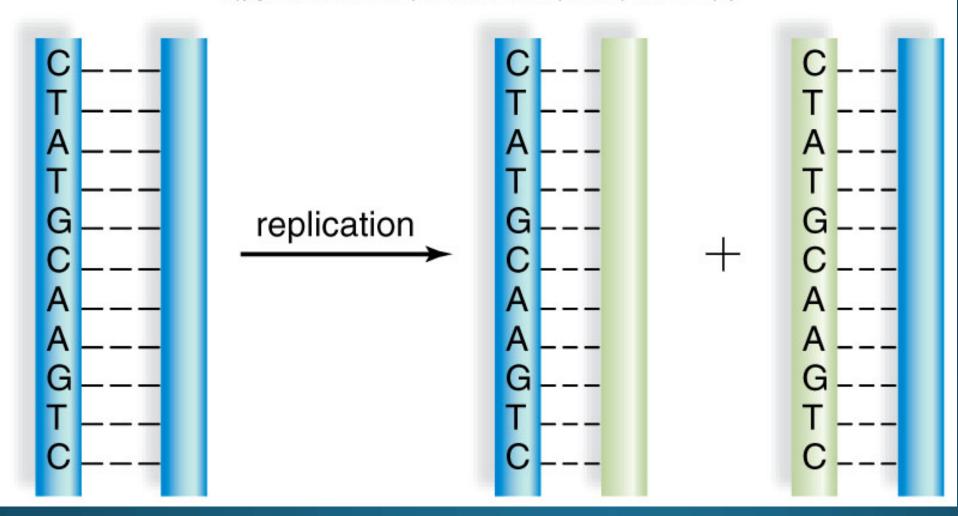
Replication of Strands



50

- Strands are unwound and separate
- New bases pair
 G with C
 T with A
- New DNA composed of one "old" strand and one "new" strand





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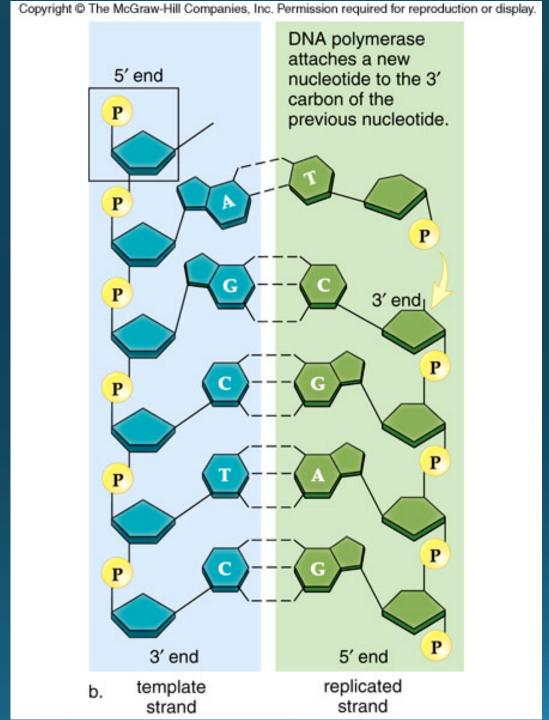
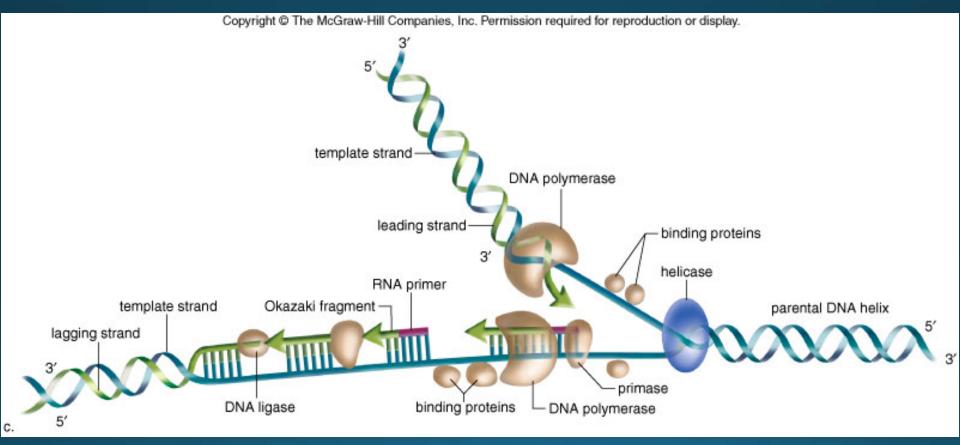


Figure 13Ab

Figure 13Ac



Replication Errors

- <u>Genetic variations are the raw material for evolutionary</u> <u>change</u>
- <u>Mutation</u>:
 - <u>A permanent (but unplanned) change in base-pair sequence</u>
 - Some due to errors in DNA replication
 - Others are due to DNA damage

• <u>DNA repair enzymes</u> are usually available <u>to reverse most errors</u>

Proofreading New DNA

•<u>DNA polymerase initially makes about 1 in</u> <u>10,000</u> base pairing <u>errors</u>

Enzymes proofread and correct these mistakes

•The new error rate for <u>DNA that has been</u> proofread is 1 in 1 billion base pairing errors

DNA Damage & Repair

- <u>Chemicals & ultraviolet radiation</u> damage the DNA in our body cells
- <u>Cells must continuously repair DAMAGED DNA</u>
- <u>Excision repair</u> occurs when any of <u>over 50</u> repair <u>enzymes remove damaged parts of DNA</u>
- DNA polymerase and DNA ligase replace and bond the new nucleotides together

Semiconservative Model of Replication

- Idea presented by Watson & Crick
- The two strands of the parental molecule separate, and each acts as a template for a new complementary strand
- <u>New DNA consists of 1 PARENTAL (original) and 1 NEW</u> <u>strand of DNA</u>





•What would be the complementary DNA strand for the following DNA sequence?

DNA 5'-CGTATG-3'

Answer:

DNA 5'-GCGTATG-3' DNA 3'-CGCATAC-5'

Now you do these AND calculate the

percentage of A/Ts and G/Cs:

1.5'-ATCGGATTTATA-3'

2.5'-ATCGTCAGGCTT-3'

3.5'-CGGCACCTCCGCAGG-3'

4.5'-ATTTACGATTCATTG-3'

5.5'-GGGCGATACGTACATT-3'