

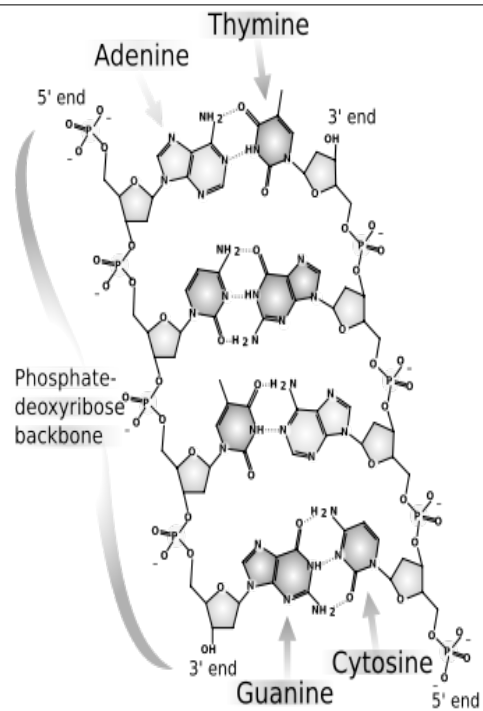
# DNA

## Structure & Replication

### A. Structure

#### 1. General

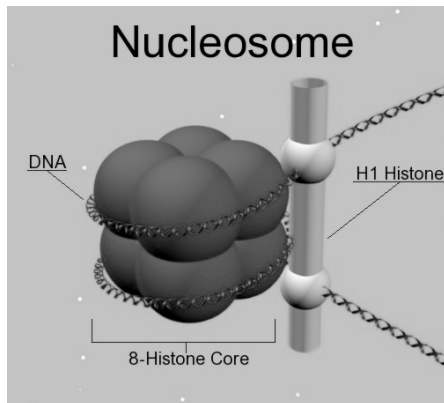
- Double helix:
  - Two anti-parallel strands (one is 5' to 3' carbon, the other 3' to 5' carbon)
- Nucleotides:
  - Nitrogenous base
    - Purines (adenine and guanine) hydrogen bond with pyrimidines (thymine and cytosine)
    - A-T, G-C
  - Phosphate group & sugar
    - Phosphate covalently attached to the C<sub>5</sub> of the deoxyribose
- 3' - 5' linkage: phosphate of next nucleotide attaches to the C<sub>3</sub>



## A. Structure, continued

### 2. Nucleosome structure

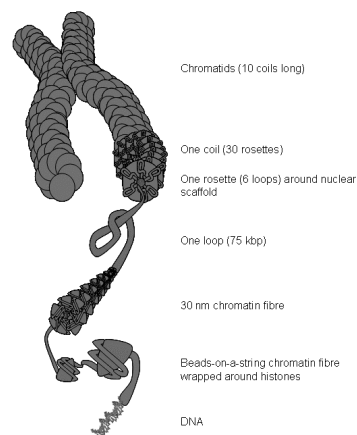
- In eukaryotes, DNA is associated with proteins to form nucleosomes.
- 8 histones wrapped twice with DNA, held together by another histone.



## A. Structure, continued

### 3. Nucleosome function

- Helps to supercoil chromosomes for mitosis and meiosis
- Regulate transcription
  - Mark genes to promote gene expression by transcription & translation
  - Cause silencing of a gene by preventing transcription



## A. Structure, continued

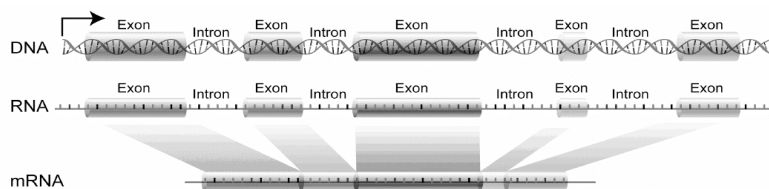
### 4. Repetitive Sequences

- Much of Eukaryotic DNA is repetitive - and is NOT translated.
- Sometimes called satellite DNA (5-300 bases, may repeat as many as 10,000 times)
- Function not known.
- Constitute 5-45% of typical eukaryote DNA
- Single copy, or unique genes 1.5% of human genome. These carry our genetic information.

## A. Structure, continued

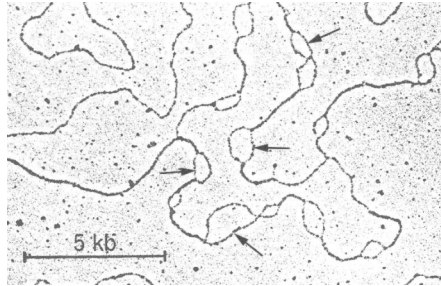
### 5. Introns & Exons

- Introns - sequences of bases that are transcribed, but not translated.
- Exons - sequences of bases that are transcribed AND translated.
- After transcription of the whole gene, the introns are cut out. (called "post-transcriptional modification")
- Prokaryotes do not have introns in their genes.



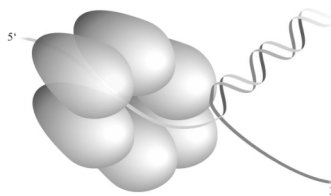
## B. Replication

1. Replication is initiated at many points on the eukaryotic chromosome.
  - Called replication “bubbles”



## B. Replication, continued

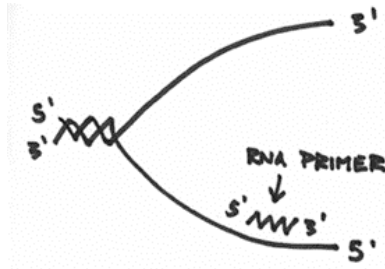
2. Helicases (enzymes)
  - Unwind the double helix
  - Breaks the hydrogen bonds between the nitrogenous bases
  - Splits it into two separate template strands
  - One strand is 3' to 5' while the other is 5' to 3'.



## B. Replication, continued

### 3. RNA Primer

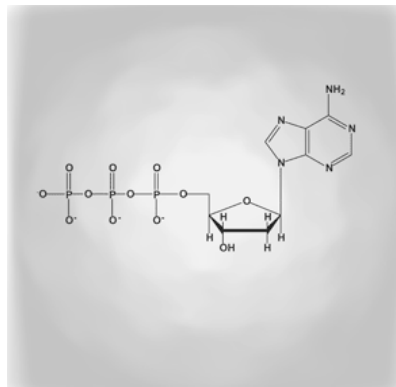
- A few RNA nucleotides bind to the old DNA strand
- Hydrogen bonds between bases
- The enzyme RNA Primase will bind these nucleotides together (covalent bonds).
- This primer must be there to start the replication.



## B. Replication, continued

### 4. The cell has many free nucleotides

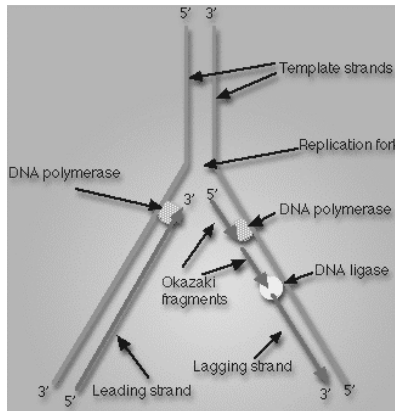
- “deoxyribonucleoside triphosphates” (dATP, dCTP, dGTP, dTTP)
- each has 3 phosphate groups
- 2 phosphate groups are removed during replication to release energy (as in ATP)
- The rest binds to the DNA as a nucleotide building block.



## B. Replication, continued

### 5. DNA Polymerase III

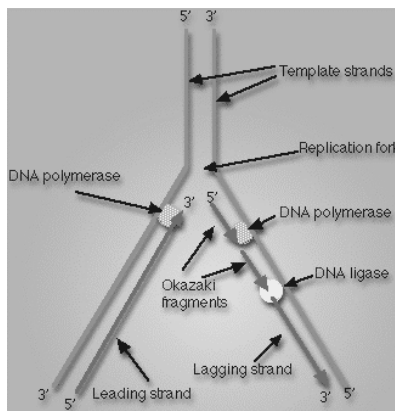
- The free nucleotides bond to template in 5' to 3' direction
- New strand is complementary to template
- Enzyme makes covalent bonds between nucleotides
- DNA Polymerase III only works in one direction
  - Leading strand made in the same direction as the replication fork
  - The lagging strand takes a little more work.



## B. Replication, continued

### 6. Lagging strand built

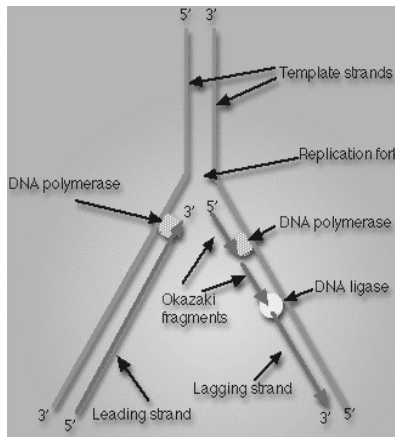
- RNA primase adds RNA Primer
- Deoxyribonucleoside triphosphates hydrogen bond with the complementary bases in the template
- DNA Polymerase adds nucleotides in the 5' to 3' direction to form short pieces called Okazaki fragments.



## B. Replication, continued

### 7. Finishing up

- DNA Polymerase I removes RNA Primer and replaces it with DNA
- DNA Ligase forms bonds between Okazaki fragments to create ONE strand.



## A SUMMARY OF DNA REPLICATION

