Molecular Basis of Inheritance & Bacterial Genetics

PREFACE

We just finished evolution and defined it as a "change in allele/**gene** frequency in a population over time". We also discussed the importance of mutations as they generate new **genes**.

- **In this powerpoint we will explore these genes further.**
- What are genes.
- How were genes discovered?
- How do genes work? What do they do?
- How do genes mutate?

What determines an organism's traits?

DNA → **Genes** → Proteins → Traits

<u>OK, What exactly is a gene?</u>

- (Basic Definition) A unit of inheritance that controls a phenotypic character.
- (Better Definition) A nucleotide sequence along a molecule of DNA that codes for a protein.
- **(Best Definition) save for another time.**

DNA…"The Blueprints of Life"

DNA → **Genes** → Proteins → Traits

DNA is the molecule of inheritance!

- This idea was hotly contested through half of the 20th century.
- Finally in 1953 James Watson and Francis Crick not only confirmed that DNA was in fact the molecule of inheritance but they described its structure and hypothesized a replicating mechanism.
- **The story begins in the early 1900's...**

DNA…"The Story"

After the work of Darwin and Mendel the race was on to clarify the vague meaning of "units of heredity"…What exactly were the units of heredity?

- Two leading suspects were nucleic acids and proteins.
- Most thought proteins were more likely suspect.
  - with 20 subunits making up protein and only 4 subunits making up nucleic acids, protein diversity was enormous

DNA…"The Story"

Early 20th Century Circumstantial Evidence

- Biologists had noted that the amount of DNA in a cell prior to cell division was "X", prior to mitosis the amount was "2X" and after cell division the amount of DNA returned to "X" amount.
- It was a fact that biologists could not explain at the time, with their current understandings.
- Obviously later with new perspectives this makes perfect sense

Makes you wonder… what knowledge today is floating around, going unnoticed, waiting to help us answer one of our many unanswered questions.

DNA…"The Story"

1928 Frederick Griffith

- While trying to develop a vaccine against *Streptoccocus pneumoniae,* he made a rather interesting observation.
- He later explained his observation by a term he coined...
- ***Transformation*** a chemical component moved from one cell to another and changed the phenotype of the recipient cell.
- Today we know that external DNA was taken up and assimilated by the bacteria whose phenotype was altered

Griffith's Experiment is described on the next slide

DNA…"The Story"

1928 Frederick Griffith

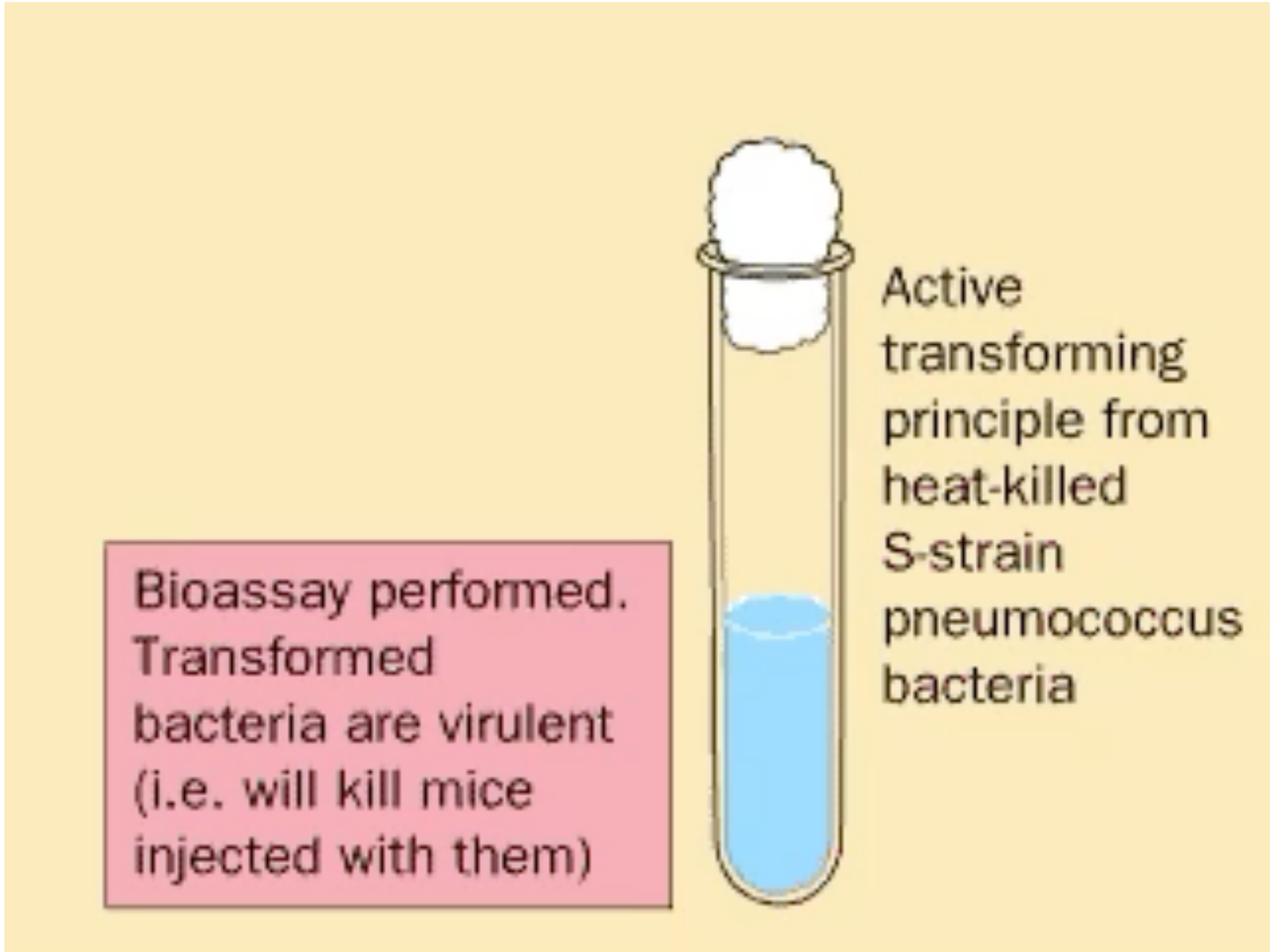
DNA…"The Story"

1944 Avery, McCarty & Macleod

- For 14 years Oswald Avery tried to determine the identity of Griffith's "transforming" agent.
- Avery's work centered around purifying molecules from the heat killed bacteria
- Finally in 1944 he and his colleagues identified the agent... DNA!

DNA…"The Story"

1944 Avery, McCarty & Macleod



DNA…"The Story"

1947-1950 Erwin Chargaff

- A biochemist, Chargaff was analyzing and comparing DNA from different species.
- From his work he two observations emerged which later became known as "Chargaff's Rules"
  - (ironically there was no basis for them at the time)
- **Rule 1: nucleic acid bases vary between species**
  - this was somewhat unexpected
- **Rule 2: within a species the number of A bases are equal to T bases and C bases are equal to G bases**

DNA…"The Story"

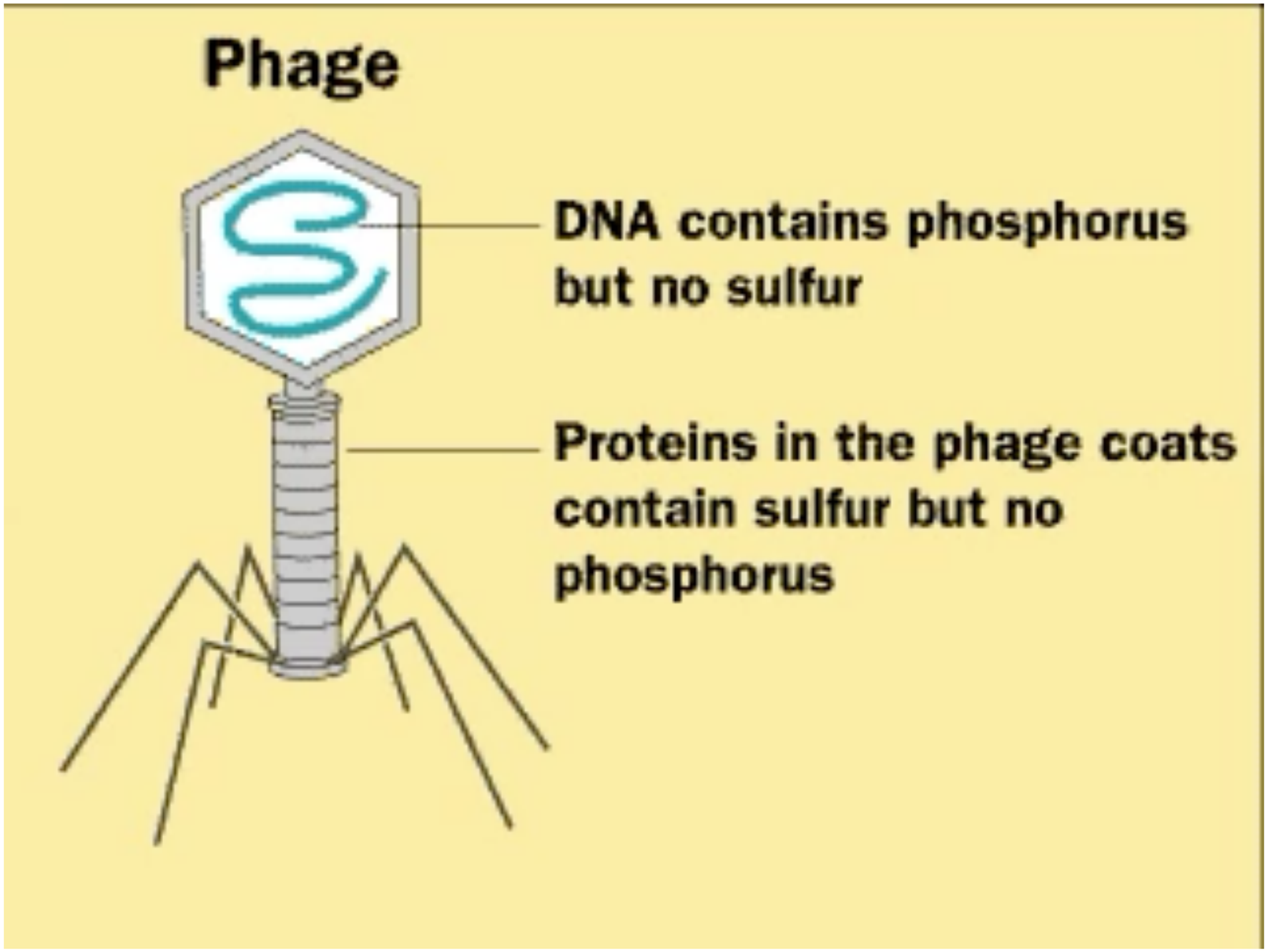
1952 Alfred Hershey & Martha Chase

- Hershey and Chase continued the search for the elusive "unit of heredity".
- Their experiment definitively showed that DNA was in fact the unit of heredity used by viruses
  - many began to contemplate the idea that DNA may be the unit of heredity for all living organisms
  - The tide was changing!

Hershey and Chase's Experiment is described on the next slide

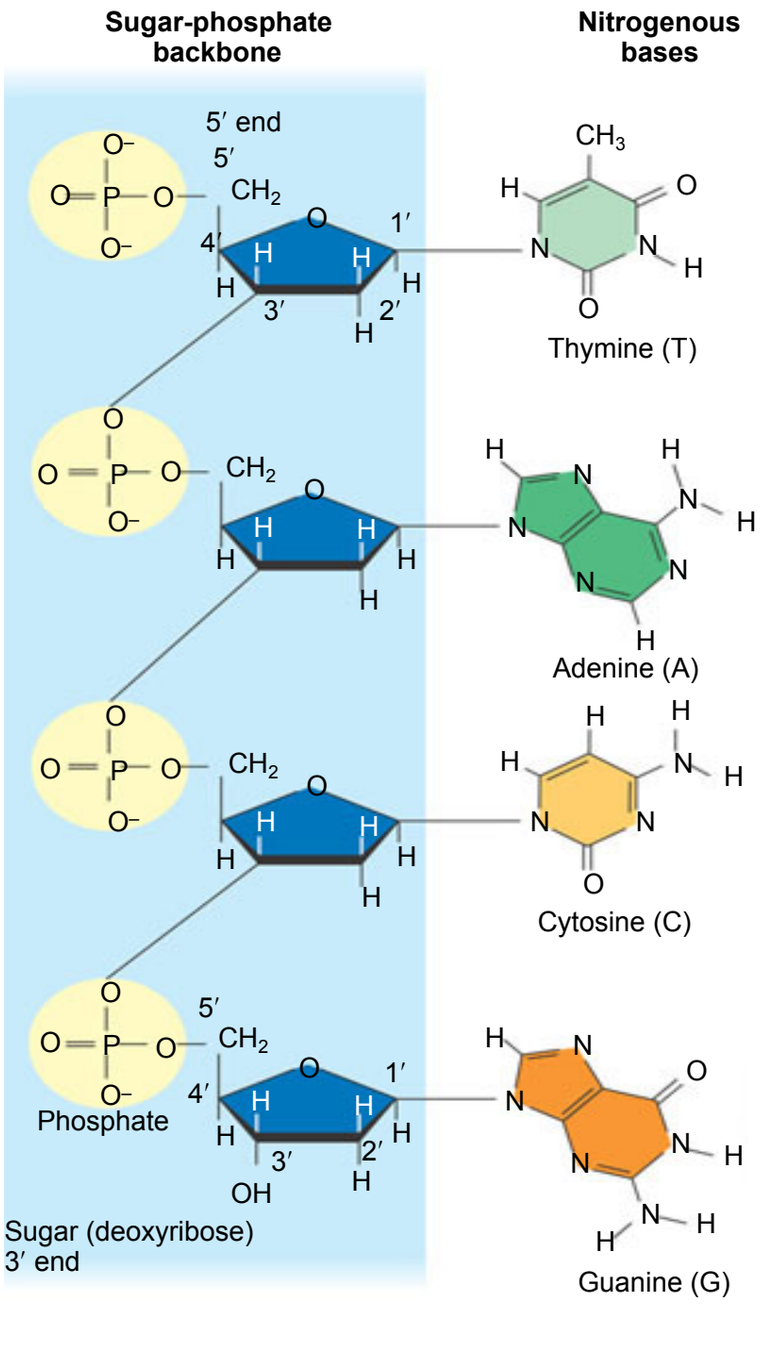
DNA…"The Story"

1952 Alfred Hershey & Martha Chase



DNA…"The Story"

1950's Rosalind Franklin, Linus Pauling, Maurice Wilkins



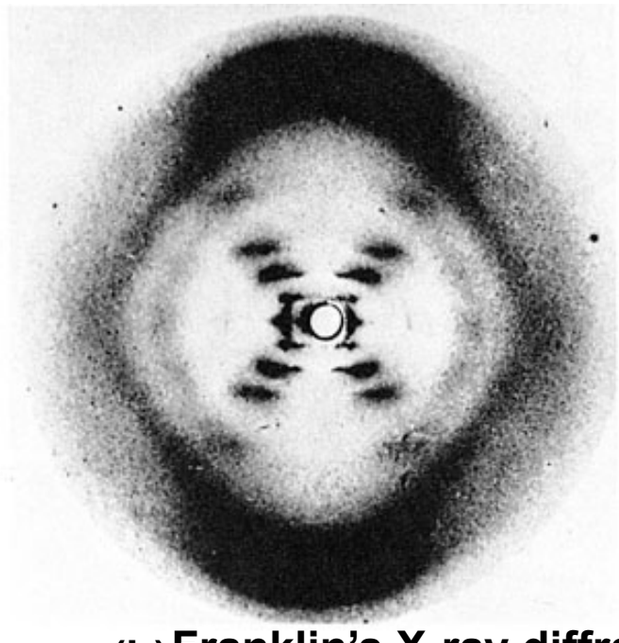
- By now many felt that DNA was the elusive "unit of heredity" and the next step would be to determine its structure.
- Prior to the 1950's chemists already knew that DNA is a polymer of nucleotides, each consisting of three components: a nitrogenous base, a sugar, and a phosphate group.

DNA…"The Story"

1950's Rosalind Franklin, Linus Pauling, Maurice Wilkins

- Rosalind Franklin wrote that the sugar-phosphate groups made up the backbone on DNA.
- Wilkins and Franklin used X-ray crystallography to determine DNA's 3-D shape but could interpret the images

(a) Rosalind Franklin

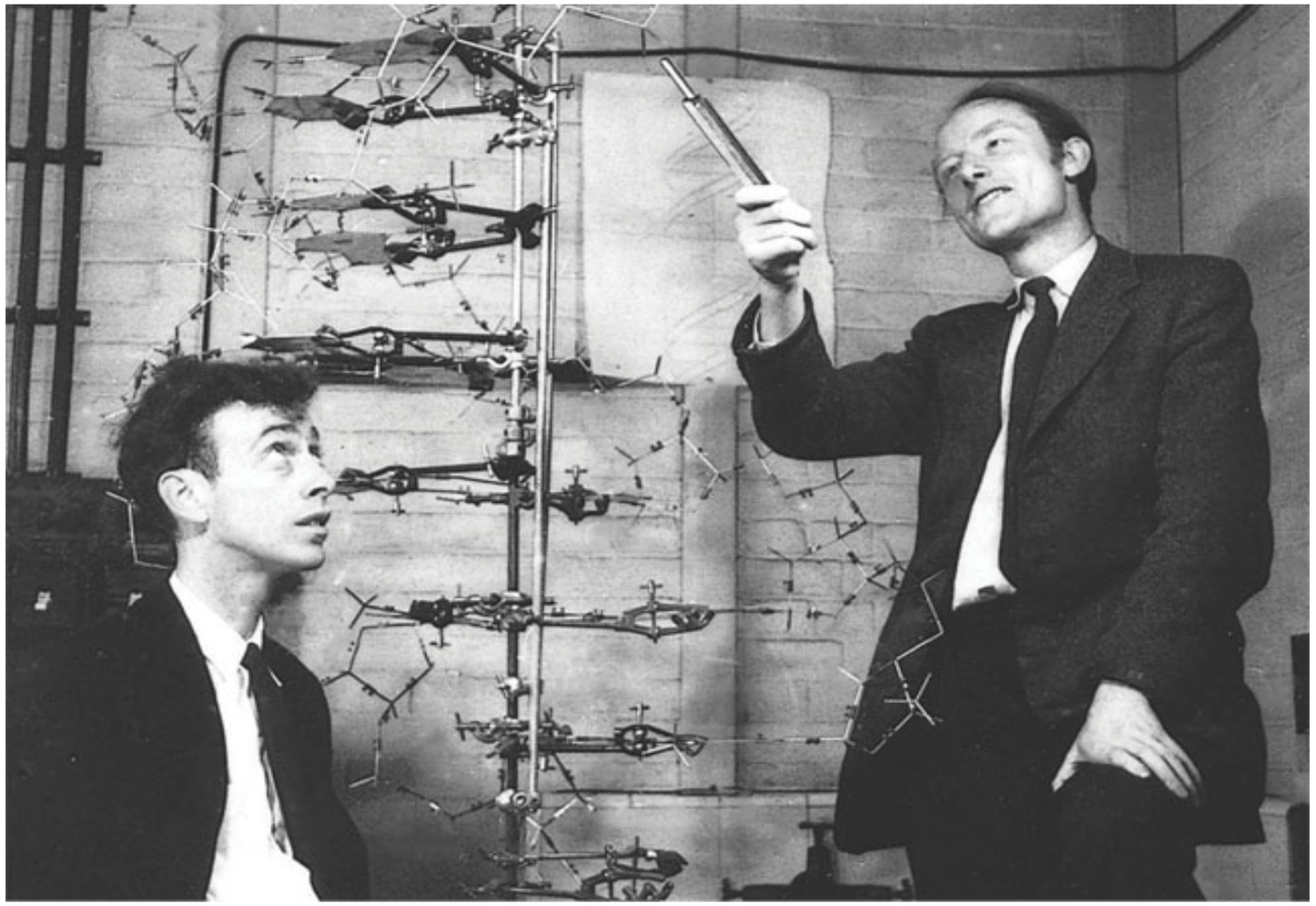


(b) Franklin's X-ray diffraction Photograph of DNA

DNA…"The Conclusion"

1953 James Watson & Francis Crick

- Watson & Crick put all the puzzle pieces together in a 1 page paper that described the structure of DNA.
  - They won Nobel Prize



DNA…"The Conclusion"

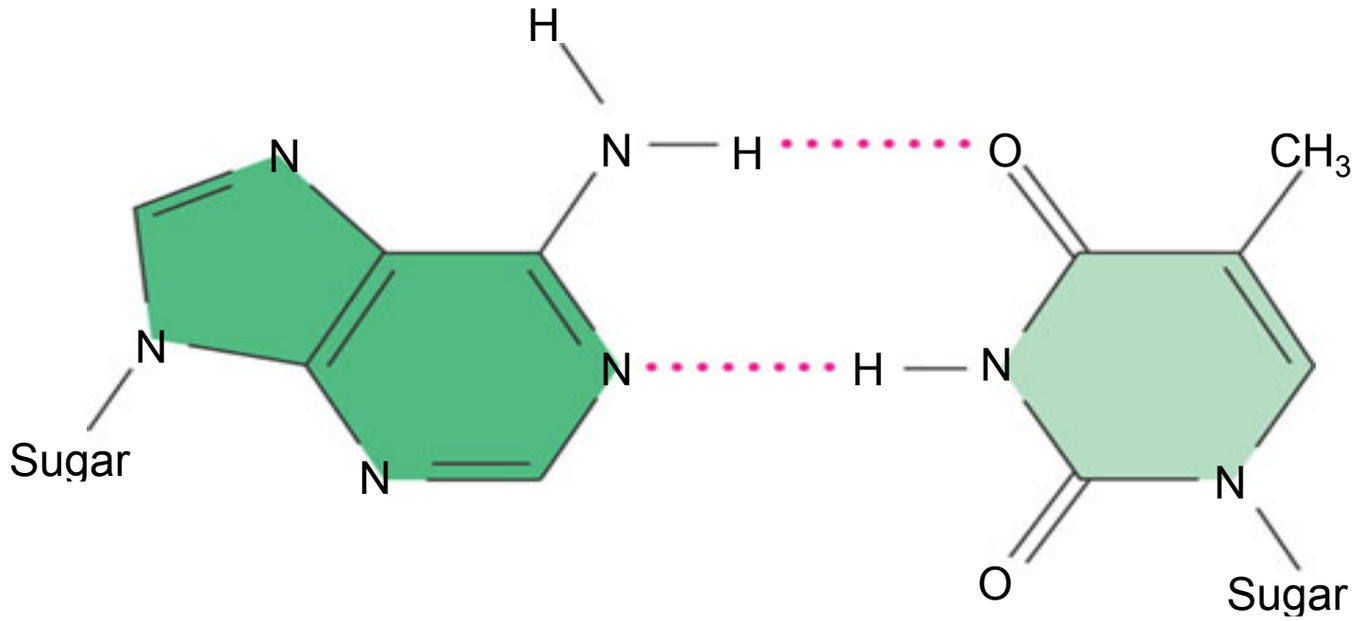
The Puzzle Pieces

- Watson recognized the x-ray image from Wilkin's lab as a helix.
- They used what chemists already knew about DNA
- They used Chargaff's observations
- They read Franklin's paper suggesting a sugar-phosphate backbone
    - From these pieces Watson & Crick deduced the following structure of DNA as seen on the next few slides

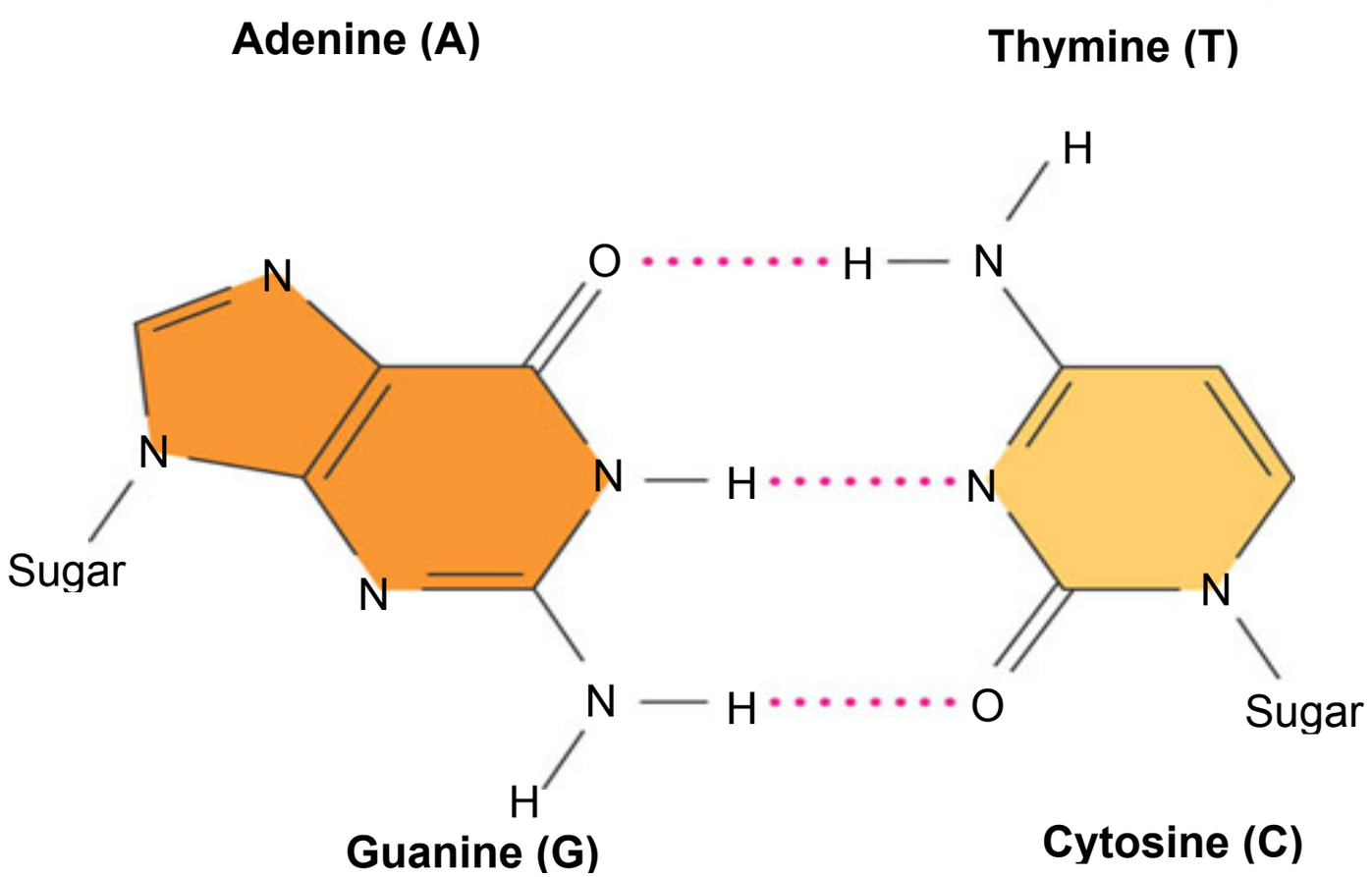
DNA

They knew...

base pairing rules



bonds between the bases



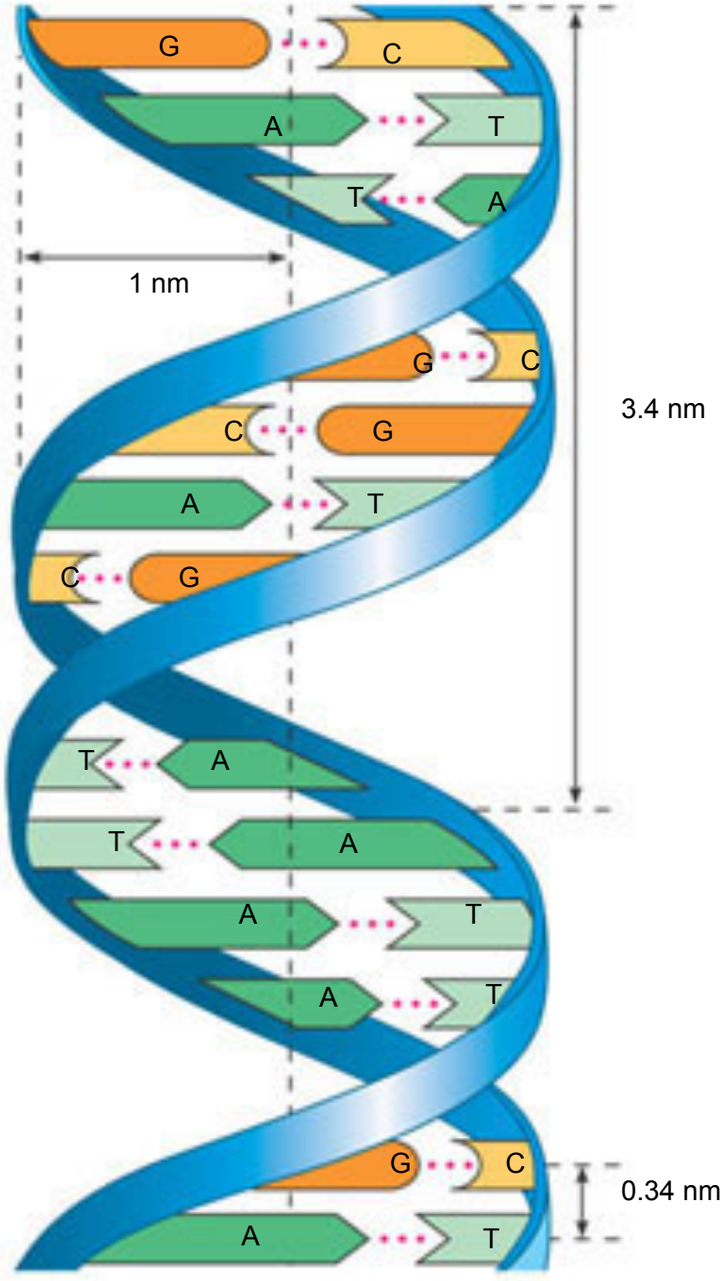
DNA

They knew...

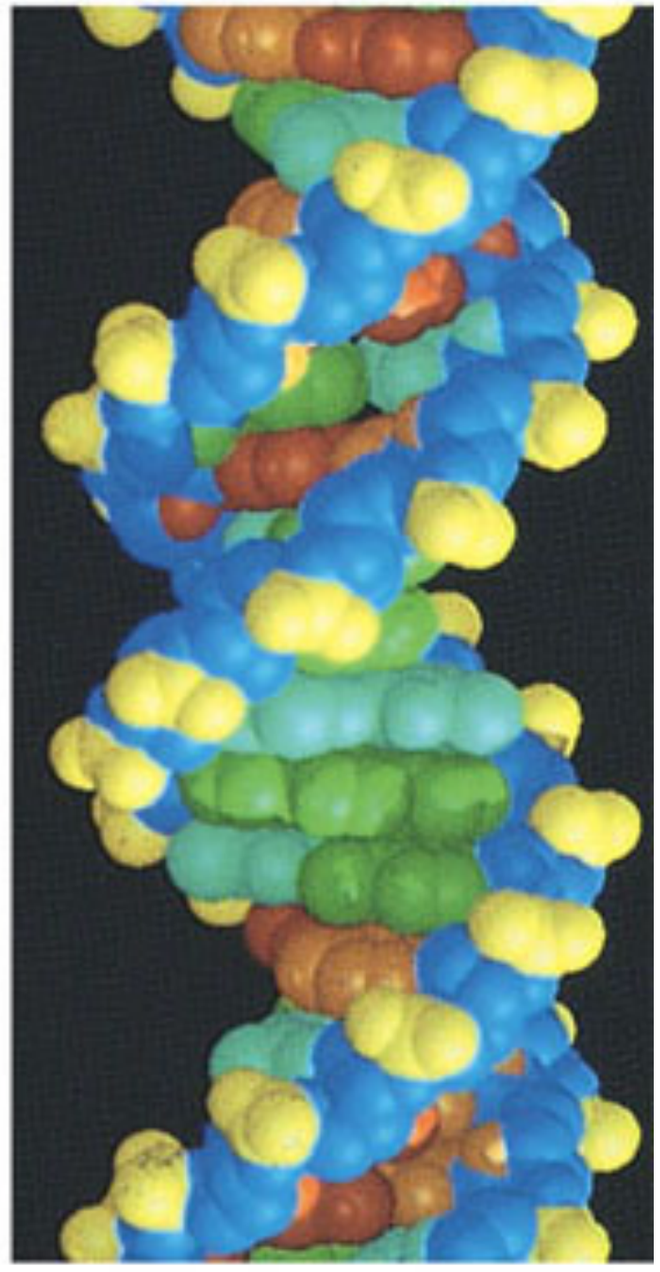
its width

its length

its shape



(a) Key features of DNA structure



(c) Space-filling model

Structure of DNA

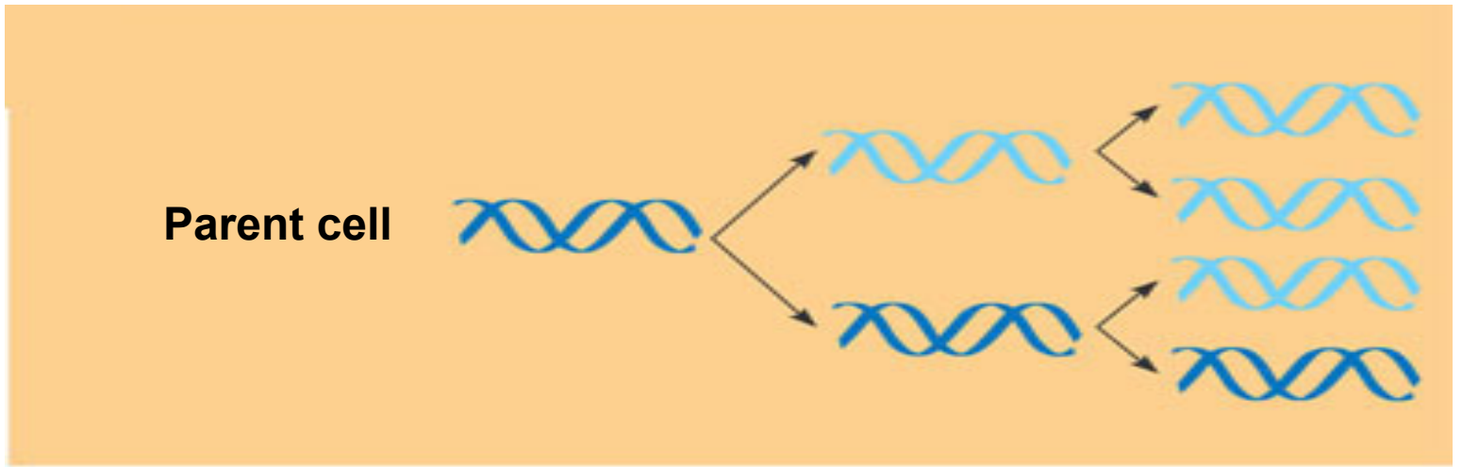
DNA Replication

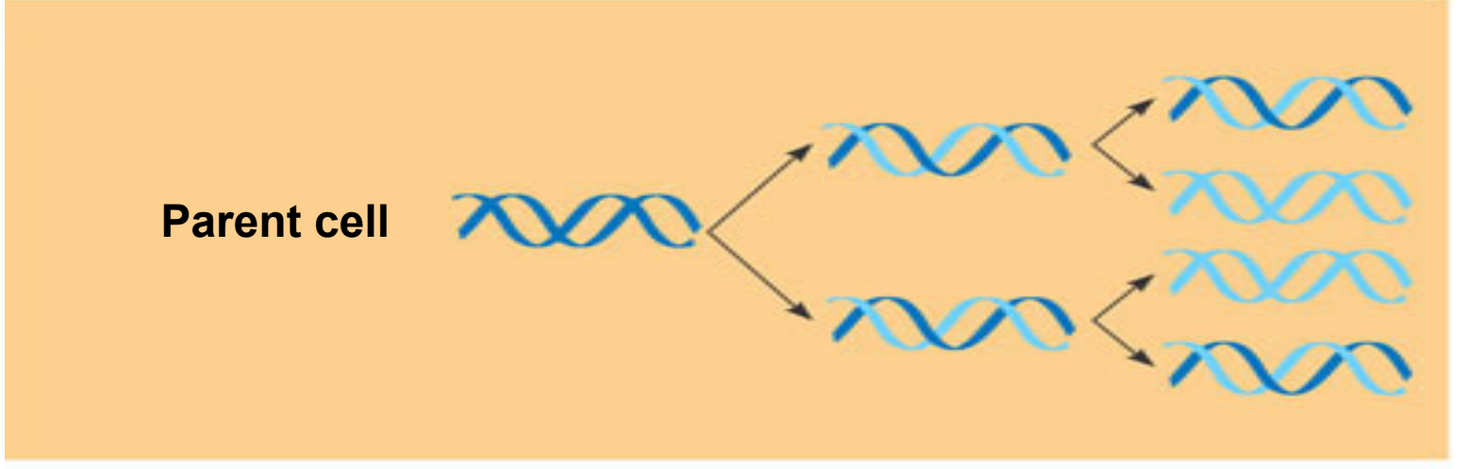
3 possible copying mechanisms

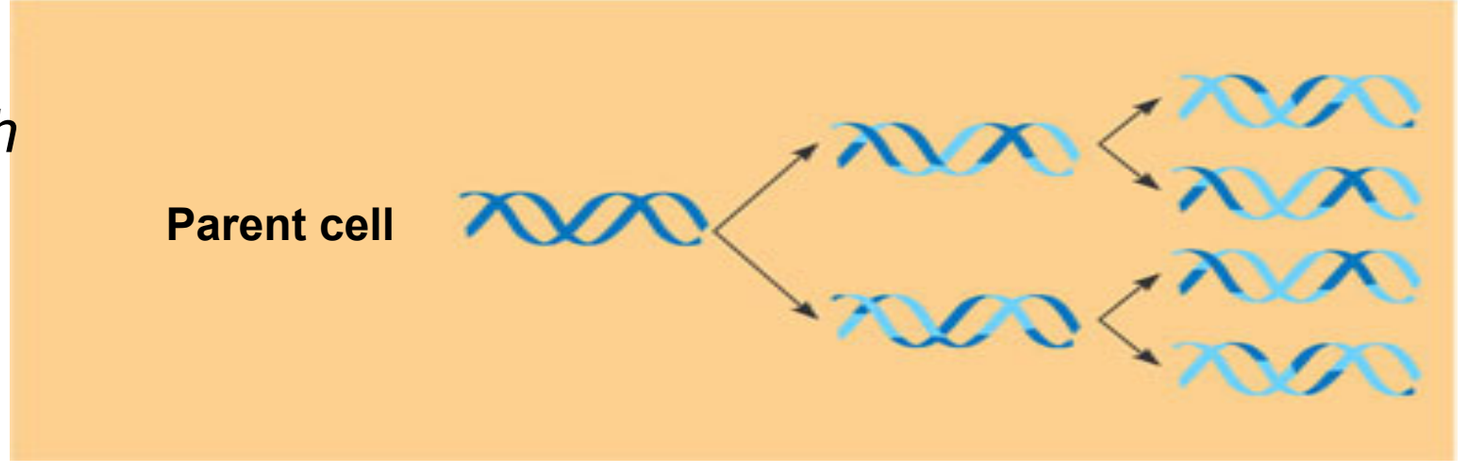
Conservative model. The two parental strands reassociate after acting as templates for new strands, thus restoring the parental double helix.

Semiconservative model. The two strands of the parental molecule separate, and each functions as a template for synthesis of a new, complementary strand.

Dispersive model. Each strand of *both* daughter molecules contains a mixture ofold and newly synthesized DNA.

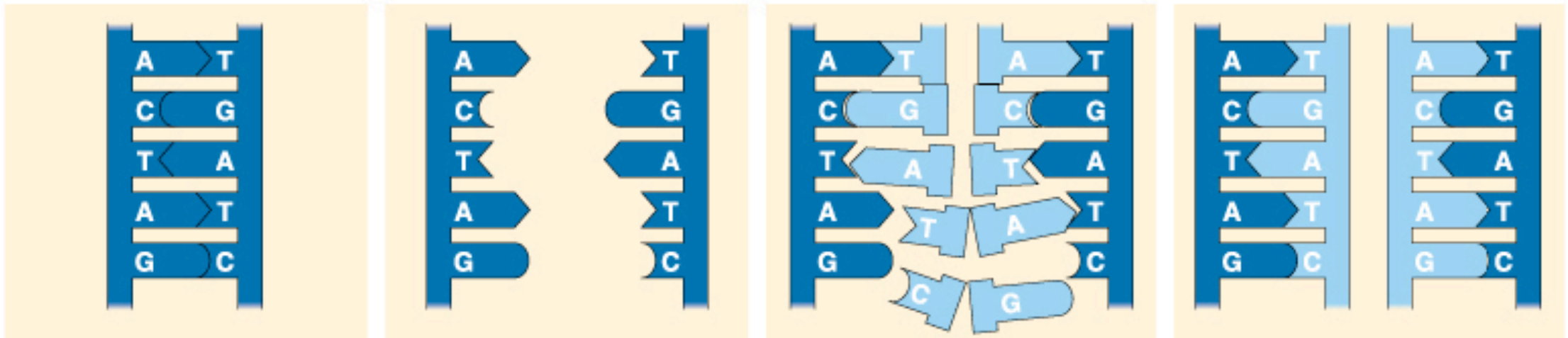






DNA Replication

The basic principle behind DNA replication is that each of the two complimentary strands serves as a template for the replication of new strands



(a) Before replication, the parent molecule has two complementary strands of DNA. Each base is paired by hydrogen bonding with its specific partner, A with T and G with C.

(b) The first step in replication is separation of the two DNA strands.

(c) Each "old" strand now serves as a template that determines the order of nucleotides along "new" complementary strands. Nucleotides plug into specific sites along the template surface according to the base-pairing rules.

(d) The nucleotides are connected to form the sugar-phosphate backbones of the new strands. Each DNA molecule now consists of one "old" strand and one "new" strand. We have two DNA molecules identical to the one molecule with which we started.

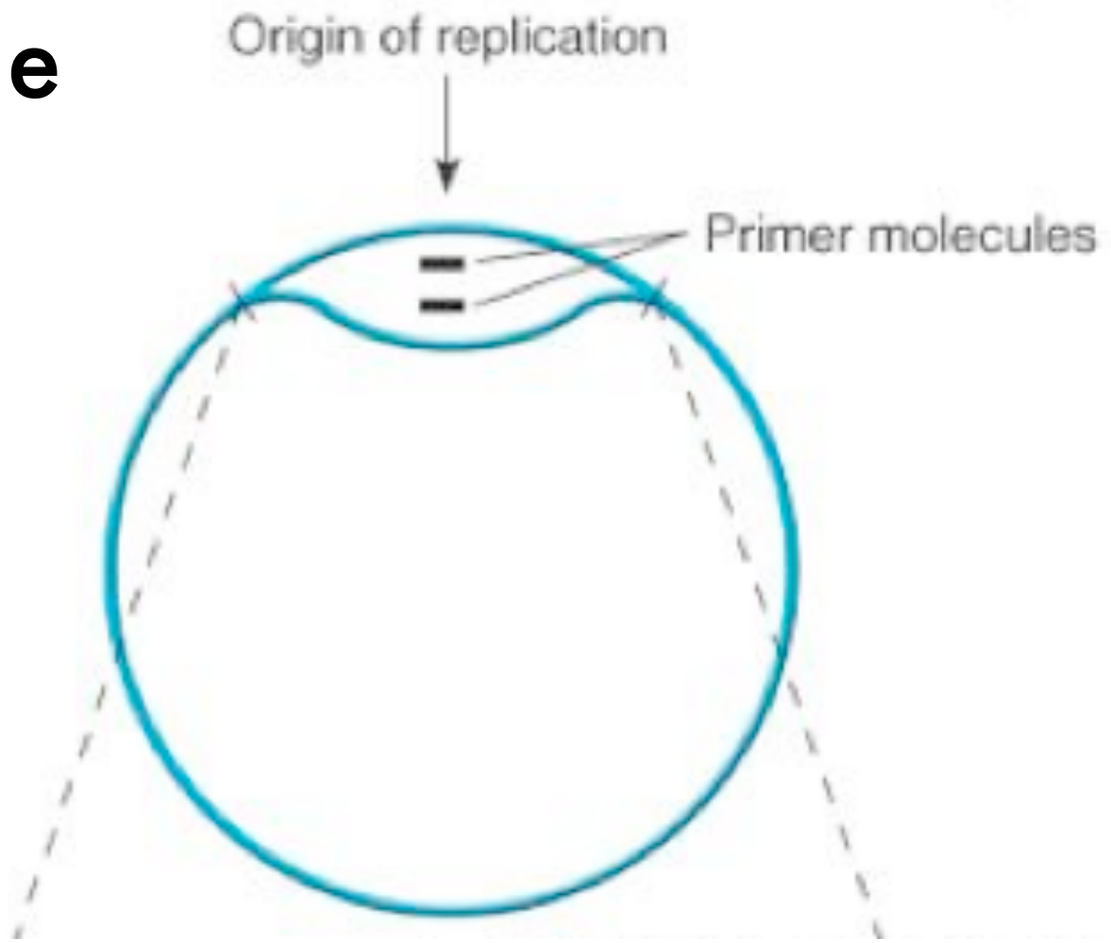
DNA Replication

- Many proteins work together in DNA replication and repair
- The parent molecule unwinds, and two new daughter strands are built based on base-pairing rules
- Copying DNA Is done with remarkable in its speed and accuracy
- The replication of DNA begins at special sites called **origins of replication**, where the two strands are separated

DNA Replication (prokaryotes)

Bacteria have only one origin of replication

1 Replication origin. Short RNA primers are positioned to start replication.

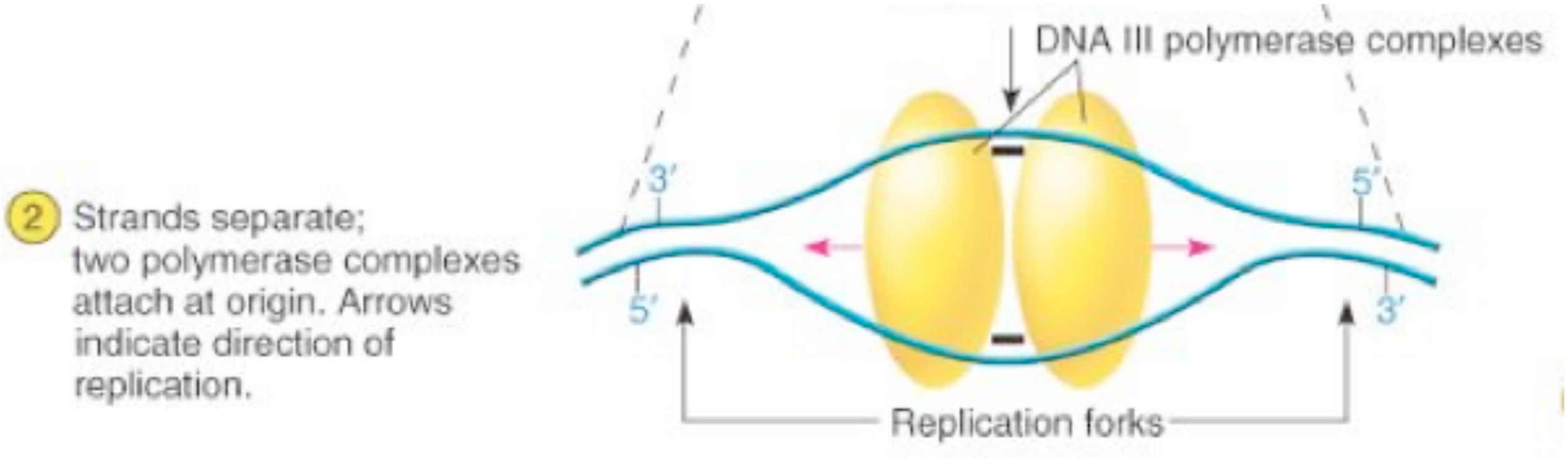


DNA Gyrase- *relaxes the supercoils and unwinds DNA*

Helicase- *seperates DNA strands (requires ATP)*

Single Strand Binding Proteins- holds the two strands apart

DNA Replication (prokaryotes)



Primase- *special RNA polymerase that lays down 15-50 nucleotides, that serve as a starting point for replication*

DNA Polymerase III- *large protein/enzyme complex that synthesizes new DNA strands from the template strands by adding one nucleotide at a time according to base pair rules*

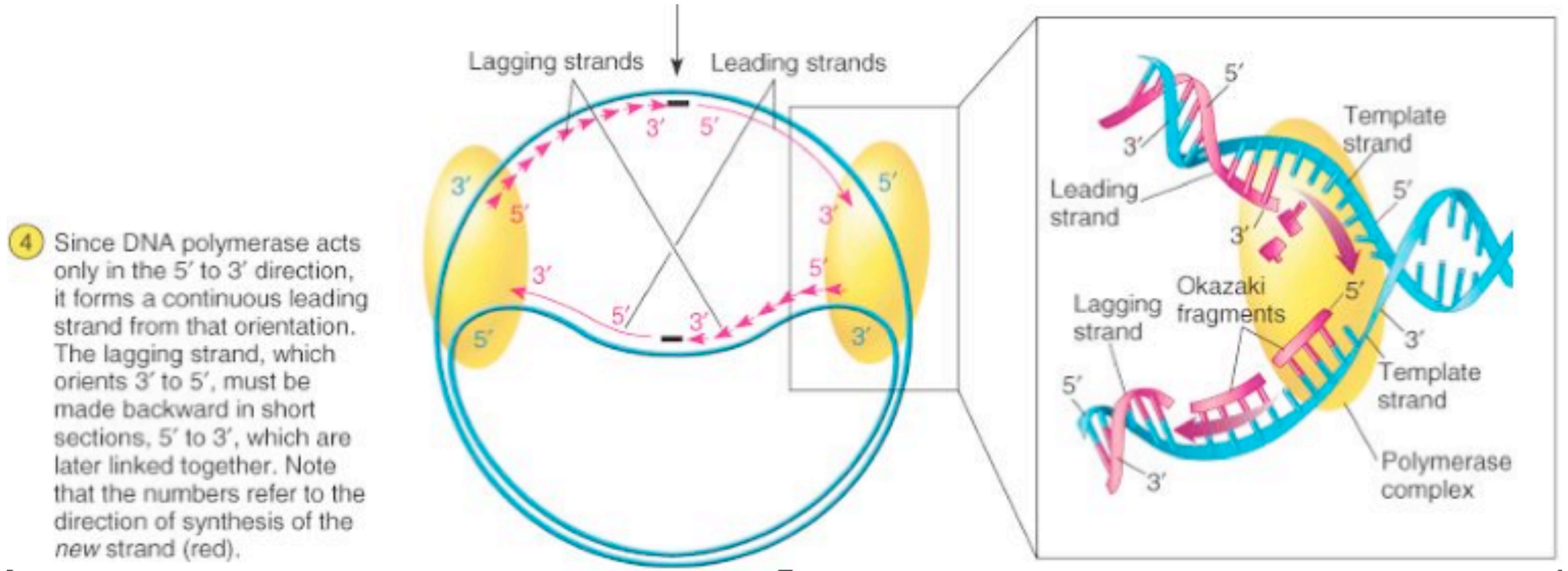
DNA Replication (prokaryotes)

3 At primer sequence, each polymerase complex synthesizes two strands at the replication forks.

Primase- *special RNA polymerase that lays down 15-50 nucleotides, that serve as a starting point for replication*

DNA Polymerase III- *large protein/enzyme complex that synthesizes new DNA strands from the template strands by adding one nucleotide at a time according to base pair rules*

DNA Replication (prokaryotes)



4 Since DNA polymerase acts only in the 5′ to 3′ direction, it forms a continuous leading strand from that orientation. The lagging strand, which orients 3′ to 5′, must be made backward in short sections, 5′ to 3′, which are later linked together. Note that the numbers refer to the direction of synthesis of the *new* strand (red).

Ligase- *forms covalent bonds between Okazaki fragments in the lagging strand*

Proteins Involved in DNA Replication

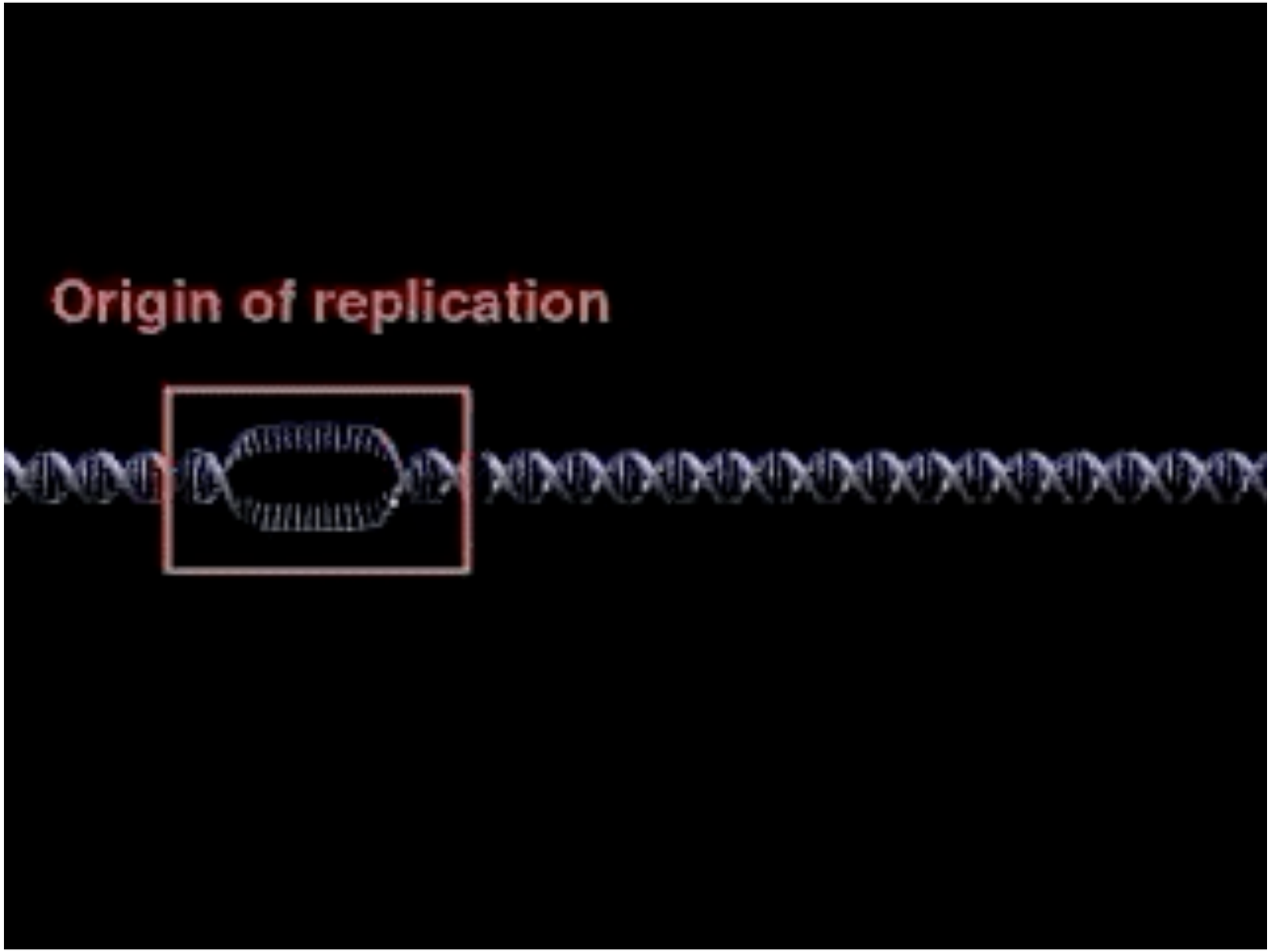
Bacterial DNA replication proteins and their functions

Protein	Function for Leading and Lagging Strands
Helicase	Unwinds parental double helix at replication forks
Single-strand binding protein	Binds to and stabilizes single-stranded DNA until it can be used as a template
Topoisomerase	Corrects "overwinding" ahead of replication forks by breaking, swiveling, and rejoining DNA strands

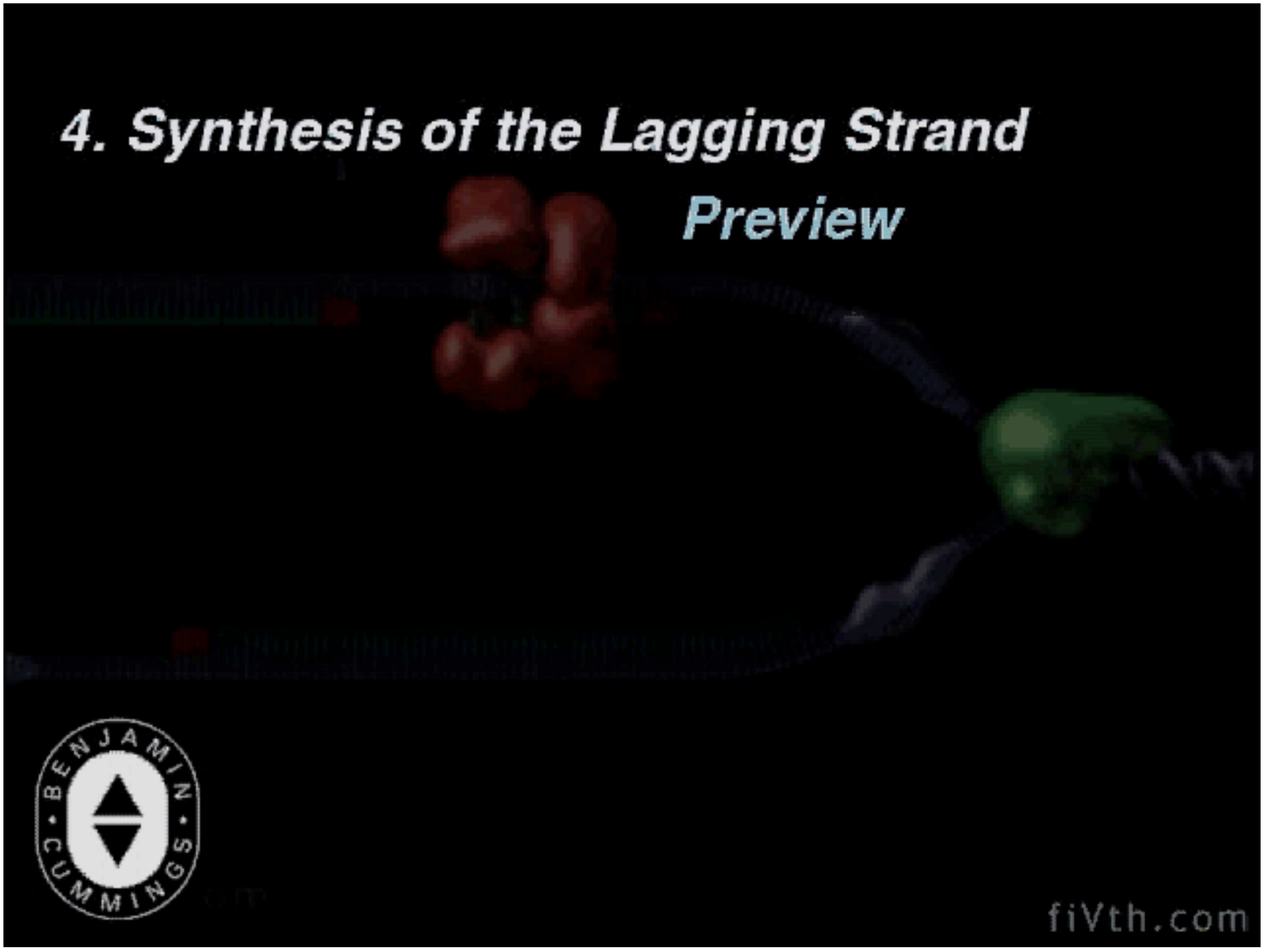
Gyrase

	Function for Leading Strand	Function for Lagging Strand
Primase	Synthesizes a single RNA primer at the 5′ end of the leading strand	Synthesizes an RNA primer at the 5′ end of each Okazaki fragment
DNA pol III	Continuously synthesizes the leading strand, adding on to the primer	Elongates each Okazaki fragment, adding on to its primer
DNA pol I	Removes primer from the 5′ end of leading strand and replaces it with DNA, adding on to the adjacent 3′ end	Removes the primer from the 5′ end of each fragment and replaces it with DNA, adding on to the 3′ end of the adjacent fragment
DNA Ligase	Joins the 3′ end of the DNA that replaces the primer to the rest of the leading strand	Joins the Okazaki fragments

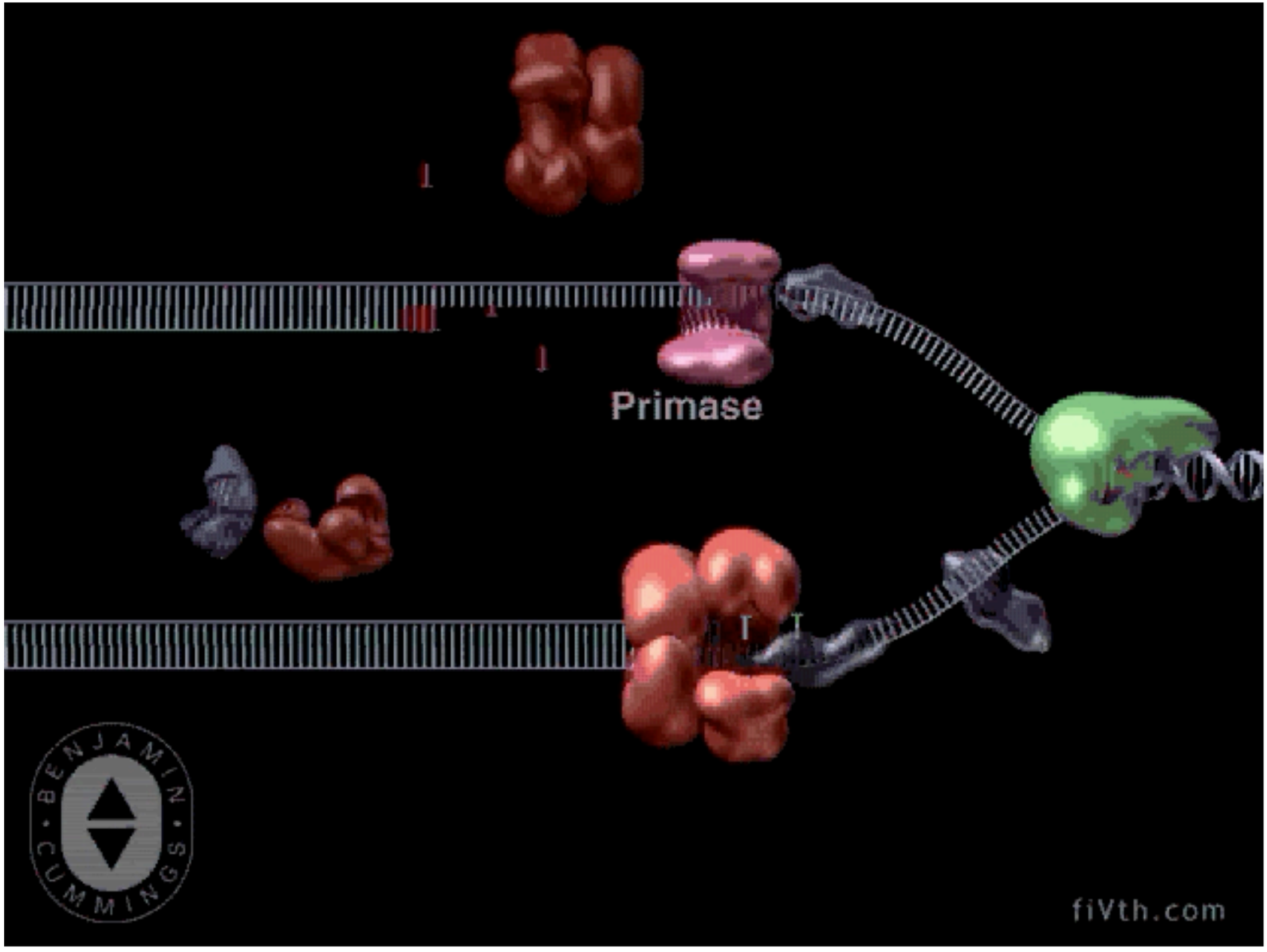
DNA Replication: Leading Strand



DNA Replication: Lagging Strand



DNA Replication: Video Review



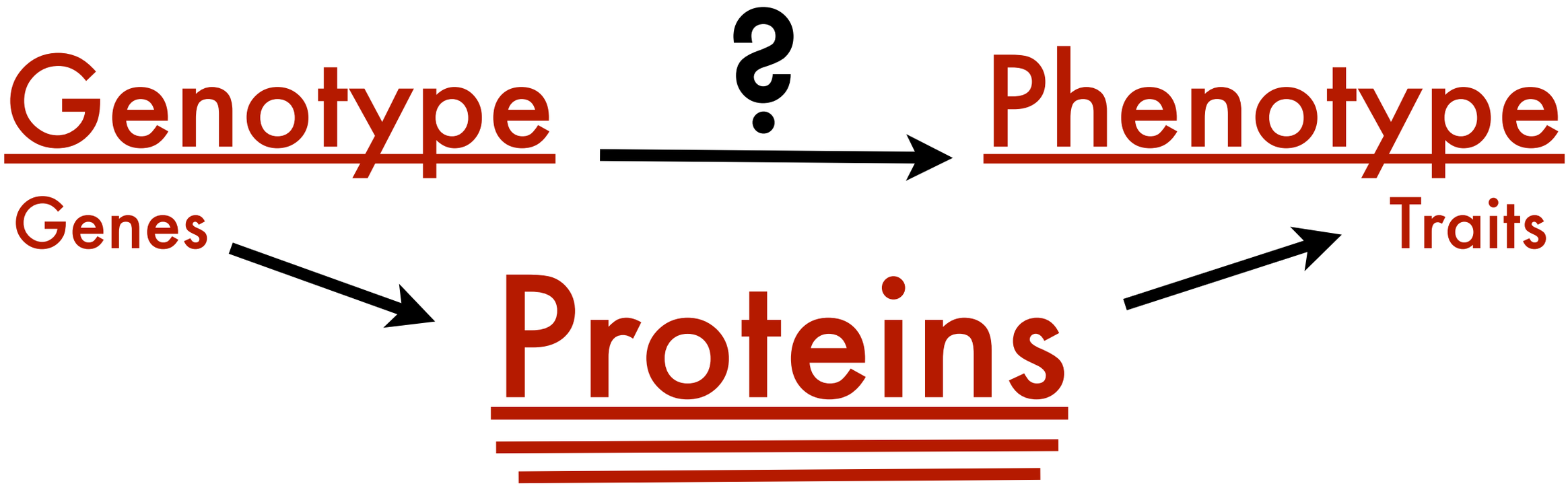
Revisit this idea…

DNA → **Genes** → Proteins → Traits

<u>OK, What exactly is a gene?</u>

- (Basic Definition) A unit of inheritance that controls a phenotypic character.
- (Better Definition) A nucleotide sequence along a molecule of DNA that codes for a protein.
- **(Best Definition) A region of DNA that can be expressed to produce a final functional product that is either a polypeptide or an RNA molecule.**

How do genes produce traits?



Proteins are the link between genotypes and phenotypes
Proteins are the link between genotypes and phenotypes
Proteins are the link between genotypes and phenotypes
Proteins are the link between genotypes and phenotypes

How do genes produce traits?

Global Flow of Information

DNA → RNA → Protein

- The flow of genetic information involves two processes.
  - Transcription
  - Translation
- Together these two processes represent gene expression.

Gene Expression…"The Story"

Early 1900's Archibald Garrod

- **In 1902, British physician Archibald Garrod hypothesized that "genes" control enzymes which in return control phenotypes.**

Early 1940's George Beadle & Edward Tatum

- **Beadle & Tatums's experimental results supported their *one gene - one enzyme hypothesis* (which states that one dictates the production of a specific enzyme).**

Early 1940's Adrian Srb & Norman Horowitz

- **Srb and Horowitz's experimental results provided additional support for the *one gene - one enzyme hypothesis*.**

Gene Expression…"The Conclusion"

Decades Later

- As researched continued the *one gene - one enzyme hypothesis* was modified as our understanding grew and technologies evolved.
  - First we realize not all proteins are enzymes thus it became
    - *one gene - one protein hypothesis*
  - Later we learn that many proteins are constructed from multiple polypeptides thus it becomes
    - *one gene - one polypeptide hypothesis*
  - Today we know that genes also code for RNA molecules…thus

A region of DNA that can be expressed to produce a final functional product that is either a polypeptide or an RNA molecule.

RNA

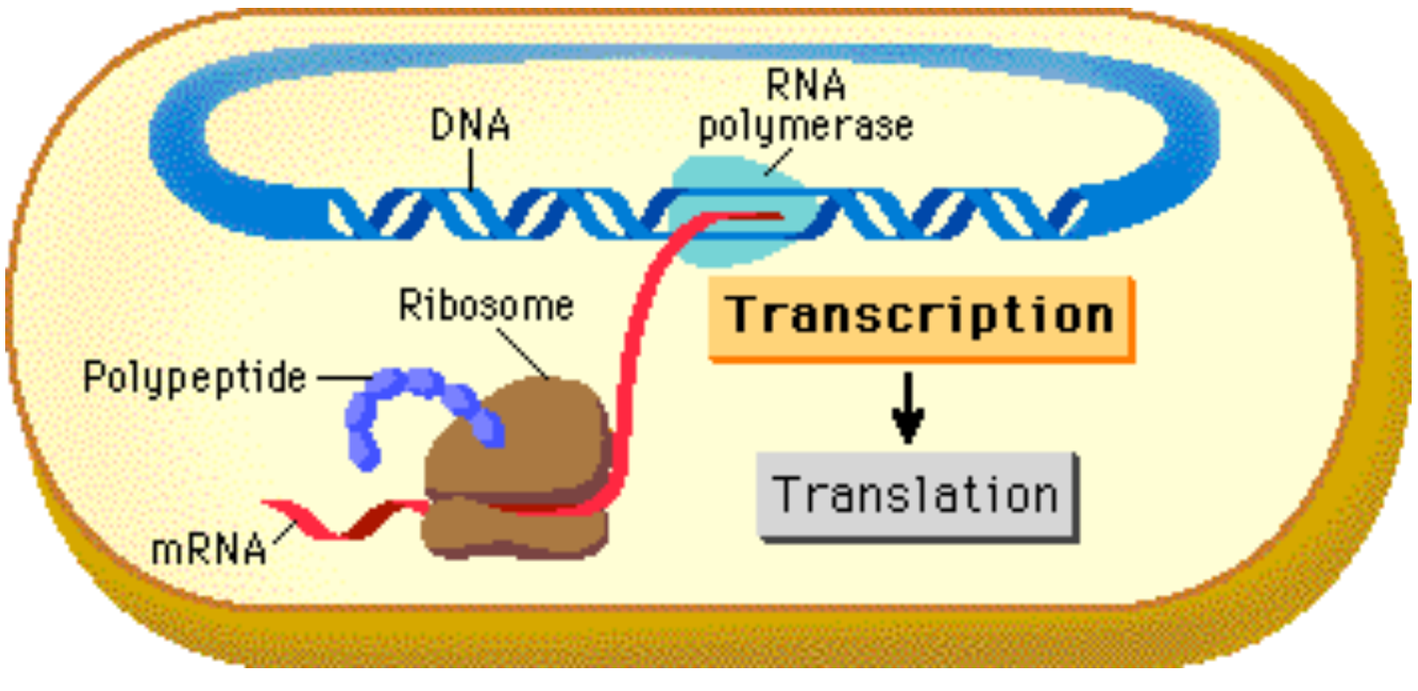
Protein Synthesis (The Basics)

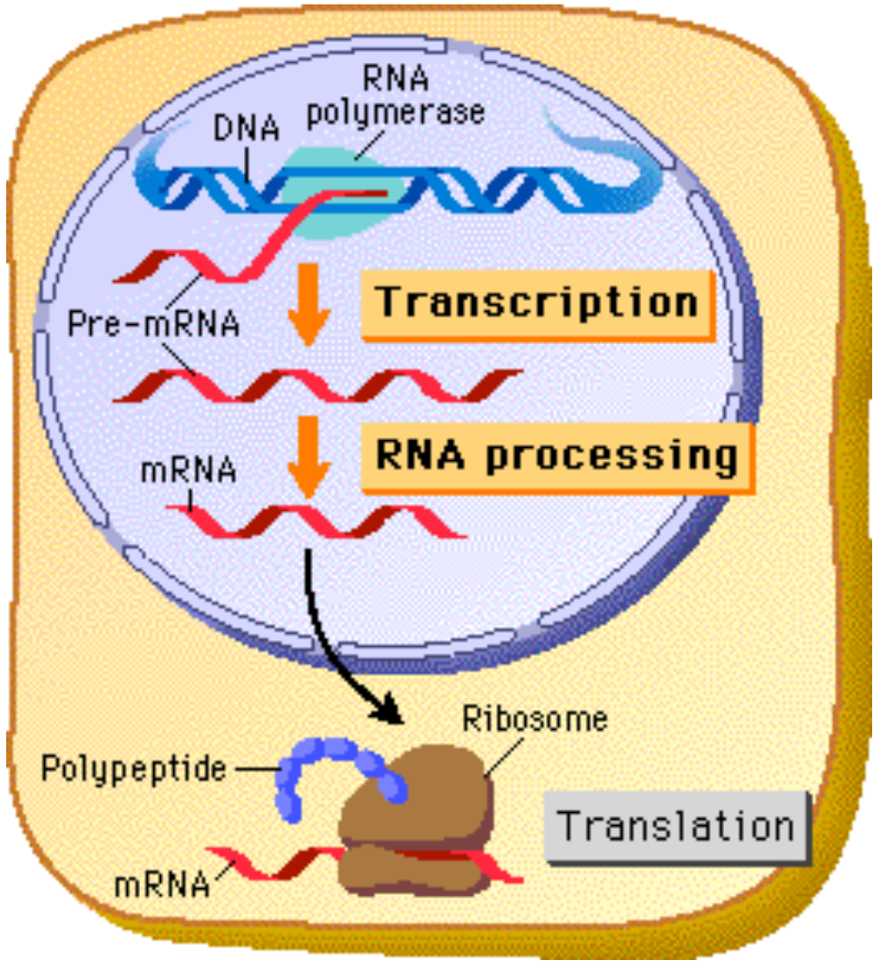
- The flow of genetic information involves two processes.
  - ***Transcription*, the synthesis of RNA using info stored in the DNA**
    - DNA serves as a template for mRNA
    - Their forms differ but their language is the same
  - ***Translation*, is the building of a polypeptide using the info stored in mRNA**
    - The language differs between nucleic acids and proteins
    - The cell must translate a nucleotide sequence into an amino acid sequence of the polypeptide

The Central Dogma

DNA → RNA → Protein

- **Transcription & Translation occurs in every organism.**
  - ***The mechanics are the same or very similar in all cells***
  - ***However, one very important difference exists between prokaryotes and eukaryotes***





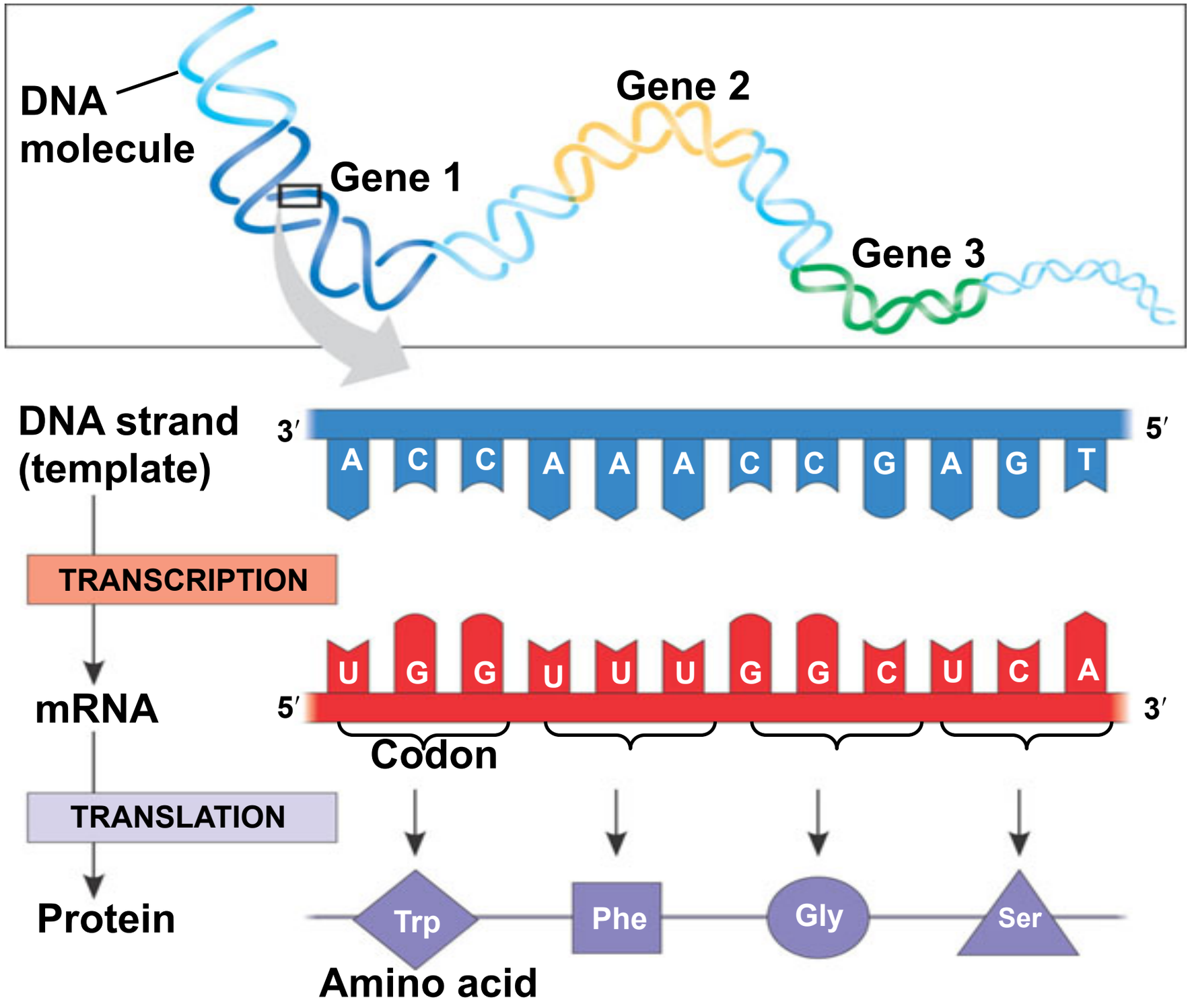
The Genetic Code

- **Once science agreed that DNA was the elusive "unit of inheritance"the next was to "crack the code".**
- **Both nucleic acids and proteins are long polymers made of molecular subunits BUT nucleic acids are built with only 4 nucleotides (subunits) and proteins are built using 20 amino acids (subunits).**
- **How does a language with 4 characters translate into a language with 20 characters?**

The Genetic Code

- 1 nucleotide could not code for 1 amino acid it would not be enough. $4^1 = 4 < 20$
- 2 nucleotides could not code for 2 amino acids it would not be enough. $4^2 = 16 < 20$
- 3 nucleotides could not code for 3 amino acids it would be more than enough. $4^3 = 64 > 20$
  - We know that the language of life (nucleic acids) is written in a ***triplet code***.
  - ***DNA uses three non-overlapping nucleotides to code for three non-overlapping nucleotides (codons) of mRNA which in turn codes for a single amino acid.***

The Genetic Code



How many nucleotides would it take to build a protein with 250 amino acids?

750 (at least)

The Genetic Code

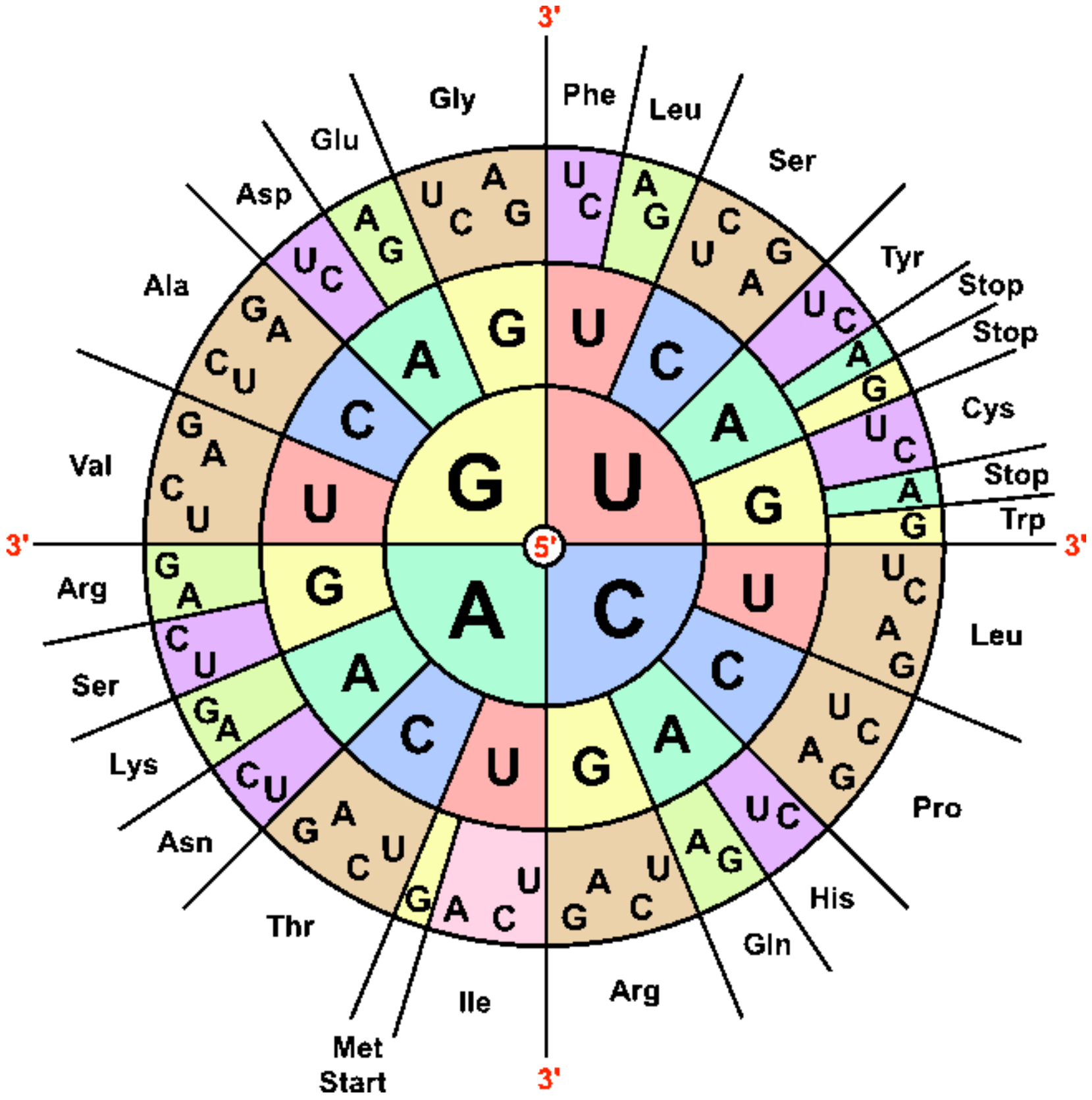
1961 Marshall Nirenberg

- Determined that UUU coded for the amino acid phenyalanine.
  - By the mid 1960's all 64 codons were deciphered.

Second Position

First Position	U		C		A		G		Third Position
U	UUU	Phe / F	UCU	Ser / S	UAU	Tyr / Y	UGU	Cys / C	U
	UUC		UCC		UAC		UGC		C
	UUA	Leu / L	UCA		*UAA*	*STOP*	*UGA*	*STOP*	A
	UUG		UCG		*UAG*	*STOP*	UGG	Trp / W	G
C	CUU	Leu / L	CCU	Pro / P	CAU	His / H	CGU	Arg / R	U
	CUC		CCC		CAC		CGC		C
	CUA		CCA		CAA	Gln / Q	CGA		A
	CUG		CCG		CAG		CGG		G
A	AUU	Ile / I	ACU	Thr / T	AAU	Asn / N	AGU	Ser / S	U
	AUC		ACC		AAC		AGC		C
	AUA		ACA		AAA	Lys / K	AGA	Arg / R	A
	AUG	Met / M	ACG		AAG		AGG		G
G	GUU	Val / V	GCU	Ala / A	GAU	Asp / D	GGU	Gly / G	U
	GUC		GCC		GAC		GGC		C
	GUA		GCA		GAA	Glu / E	GGA		A
	GUG		GCG		GAG		GGG		G

Another Amino Acid Look Up Table



The Genetic Code

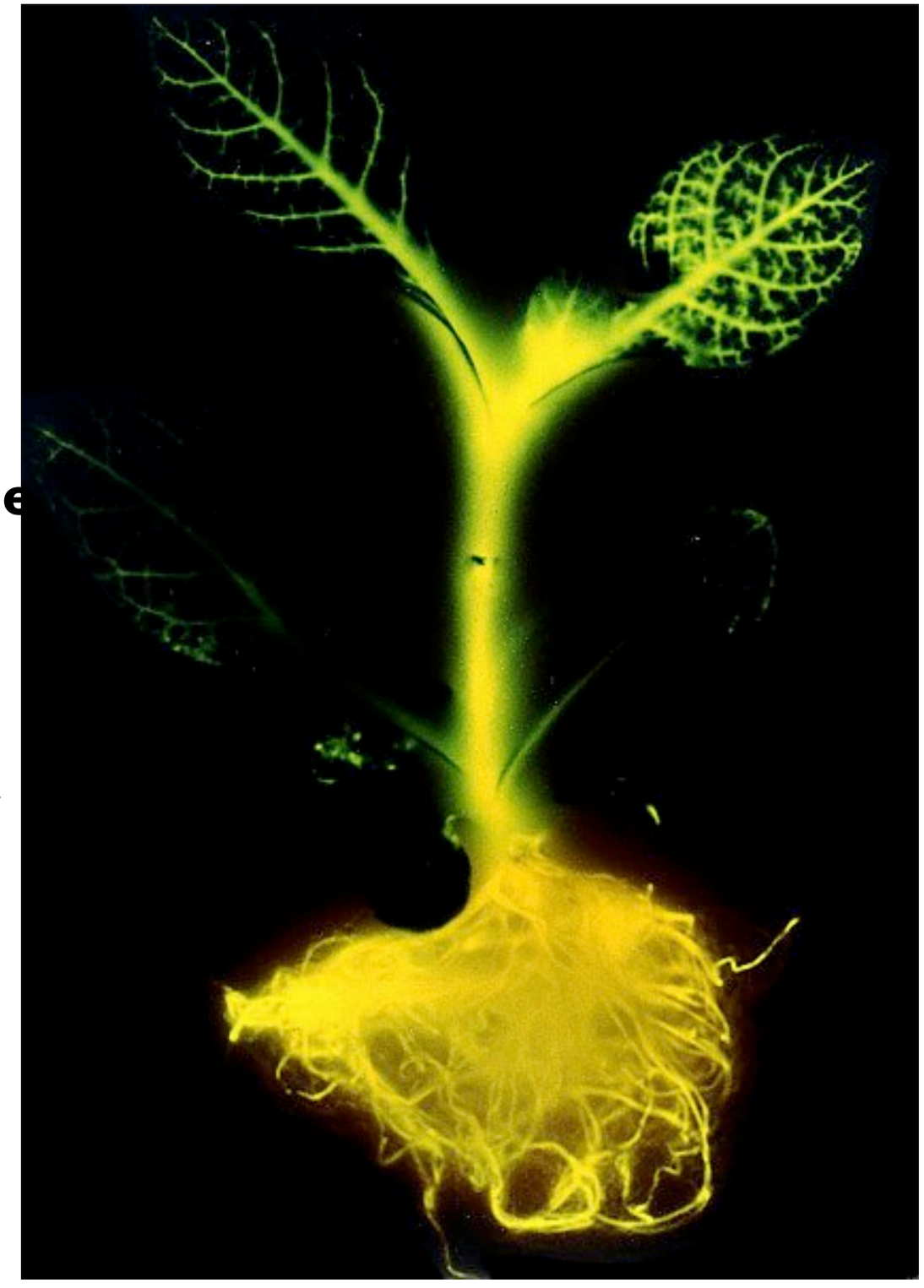
- The genetic code has some noteworthy characteristics.
  - **Redundancy**
    - AGU = serine, AGC = serine, multiple codons exist for the same amino acid
  - **No Ambiguity**
    - AGU = serine, any codon always codes for the same amino acid, it never changes
- **Universal* (nearly)**
  - This code is identical from bacteria to blue whales!

*A shared genetic code supports the idea common ancestry among all living organisms

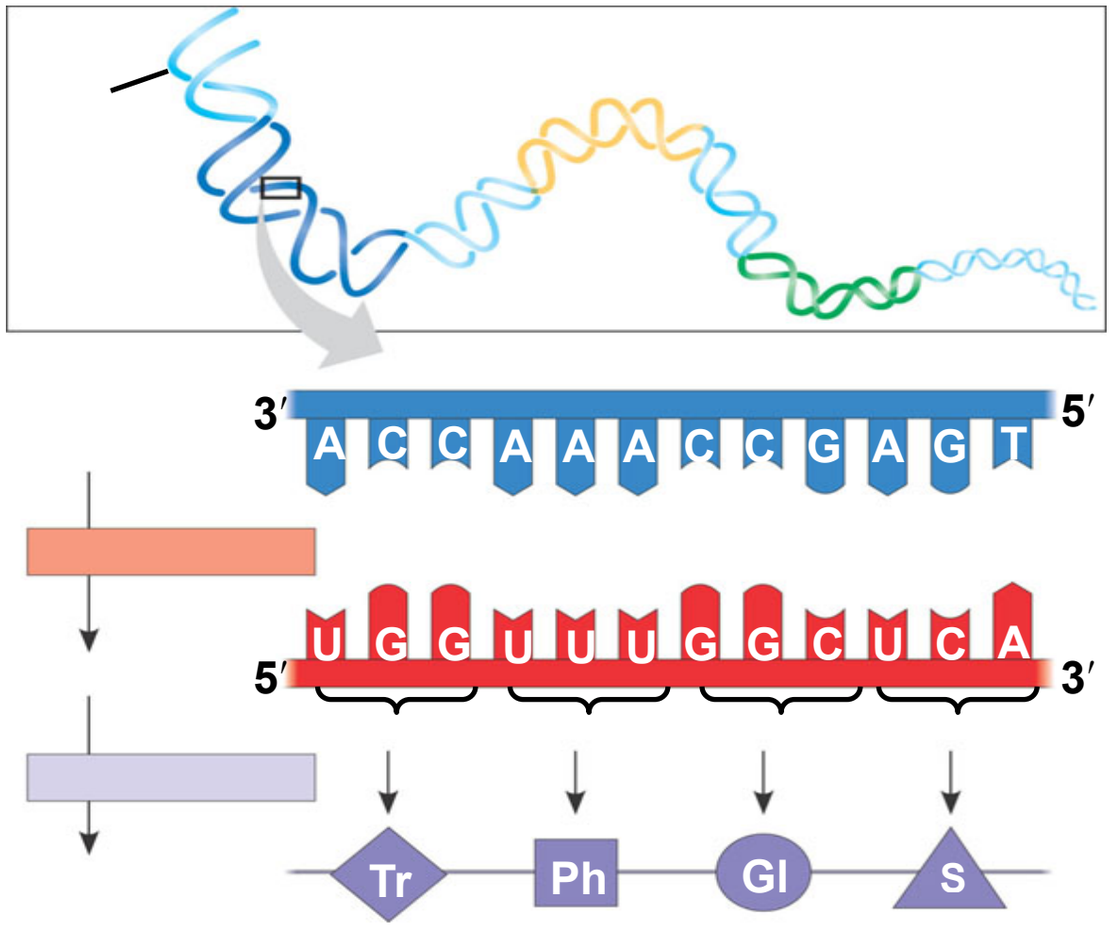
Universal Genetic Code

Luciferase Gene

Tobacco



A Short Side Trip…

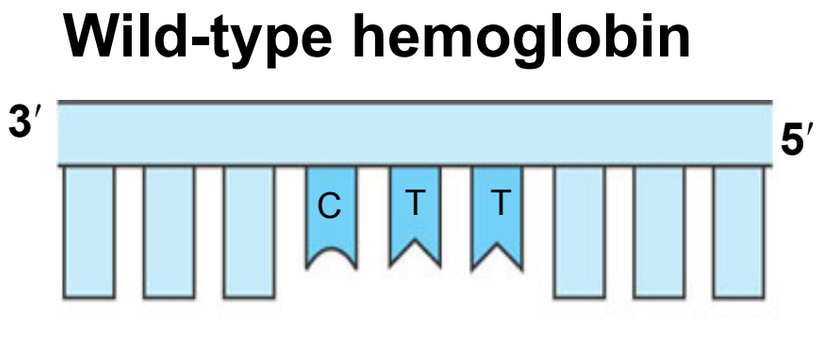


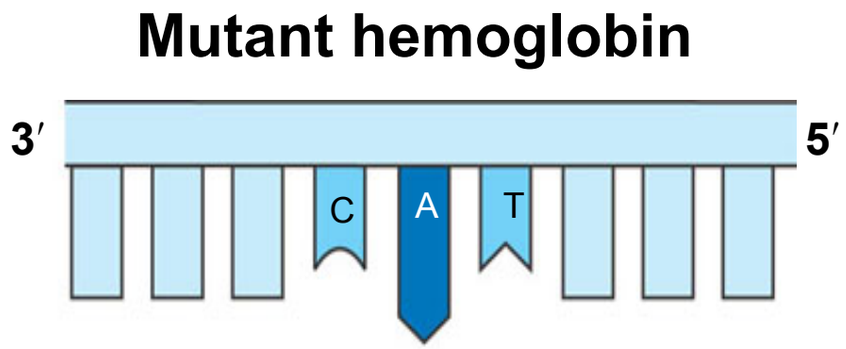
Quick Reminder…

What happens when one or more of these nucleotides changes?

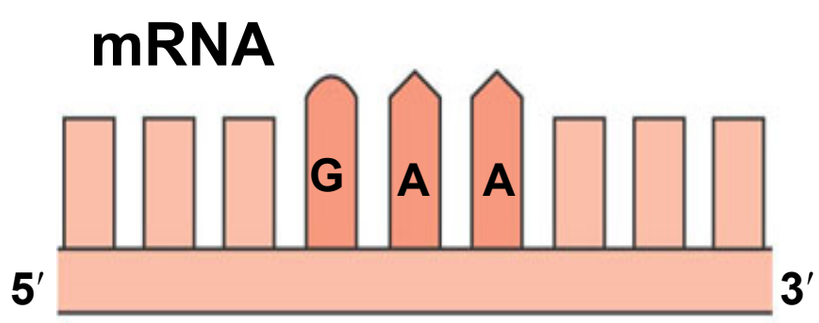
Point Mutations

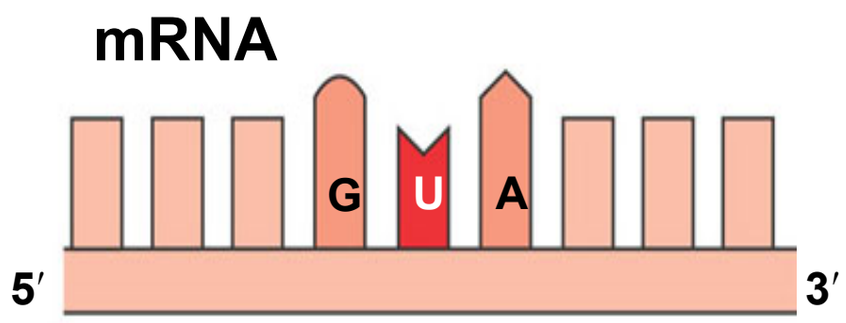
Changes in one base pair of a gene.



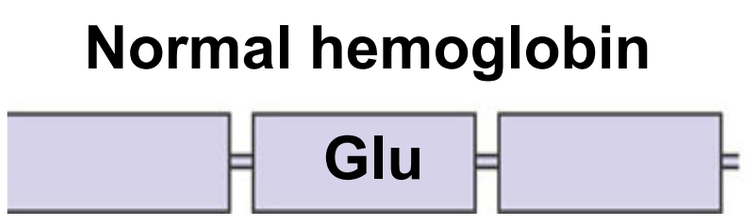


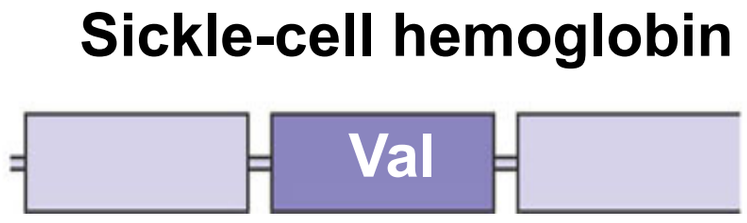
In the DNA, the mutant template strand has an A where the wild-type template has a T.





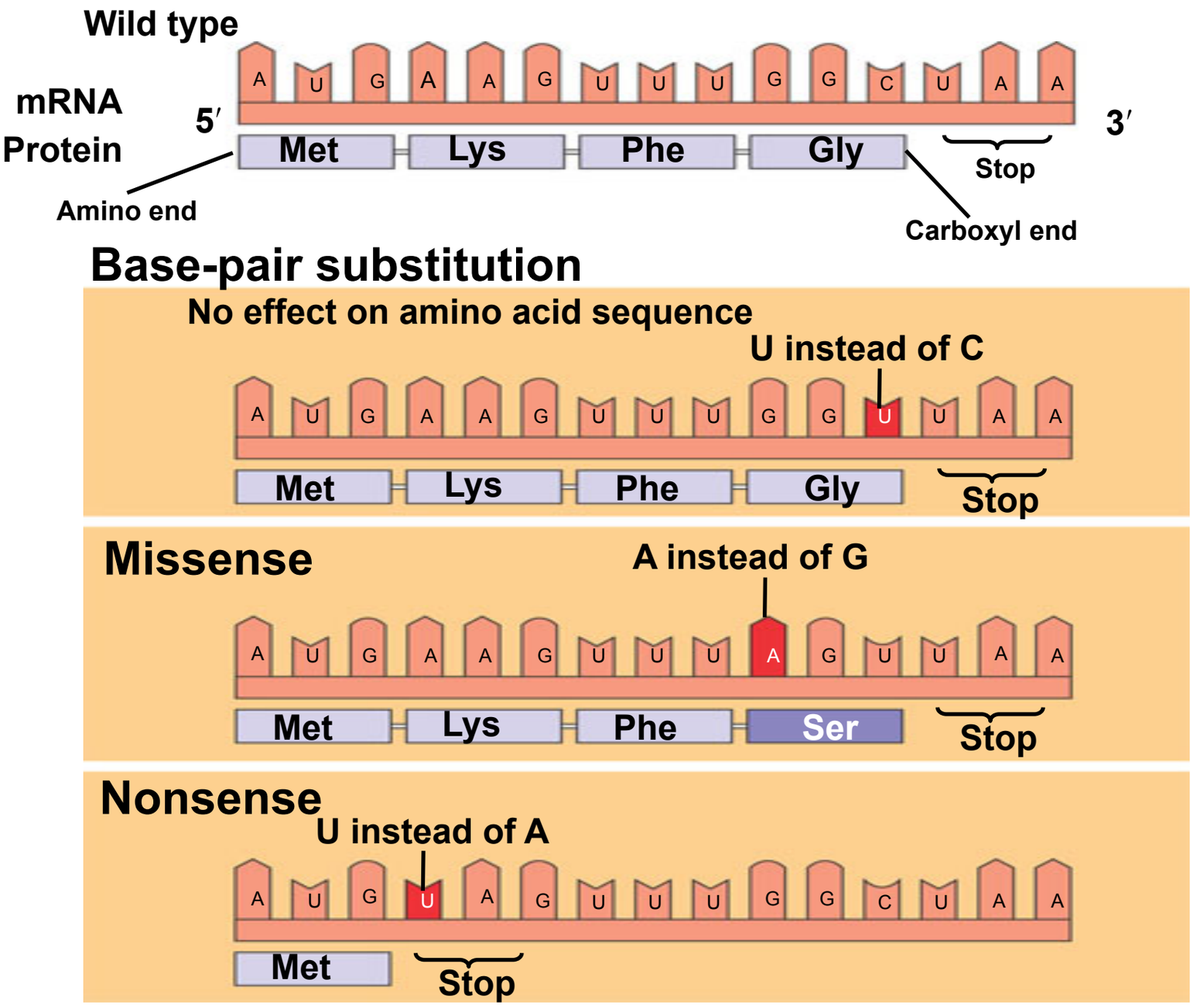
The mutant mRNA has a U instead of an A in one codon.





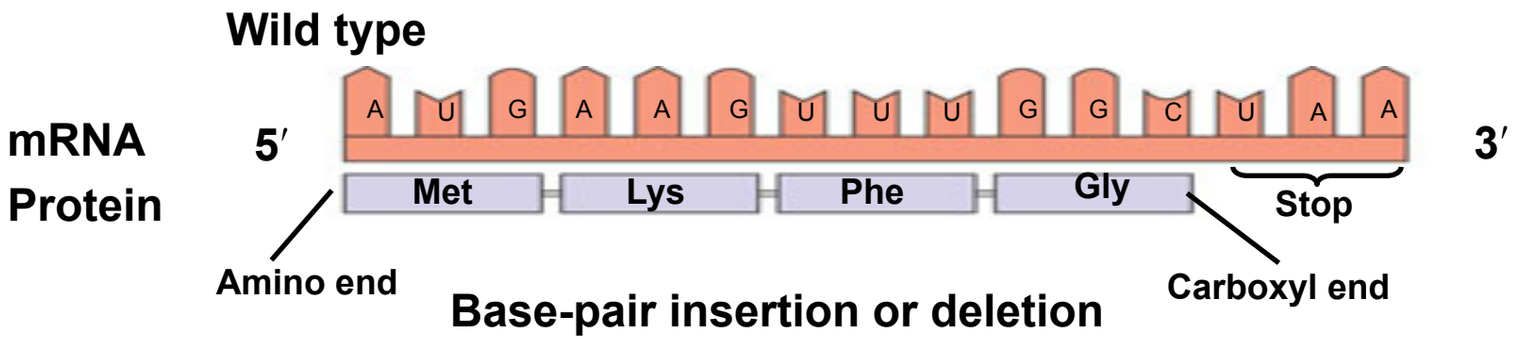
The mutant (sickle-cell) hemoglobin has a valine (Val) instead of a glutamic acid (Glu).

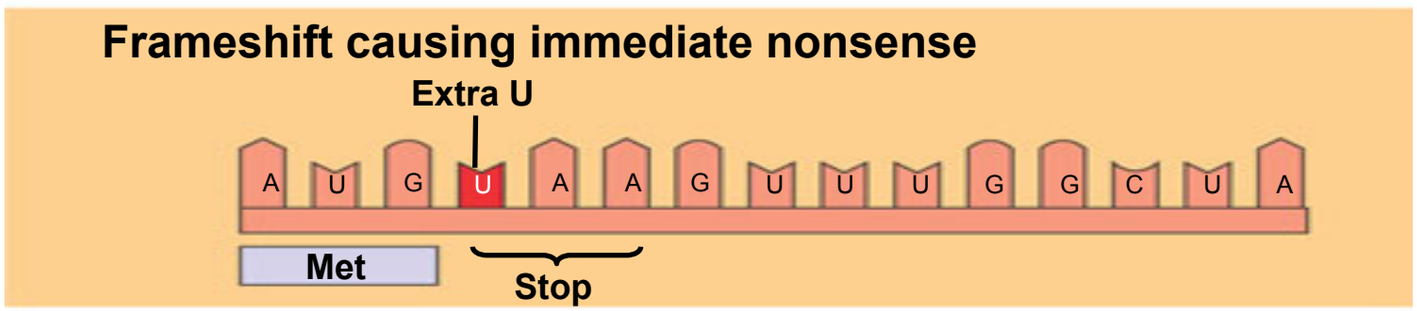
Point Mutations- Base Pair Substitutions

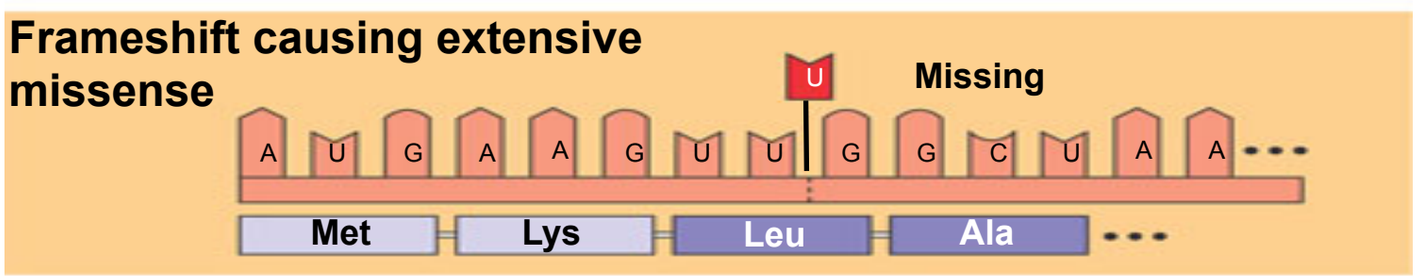


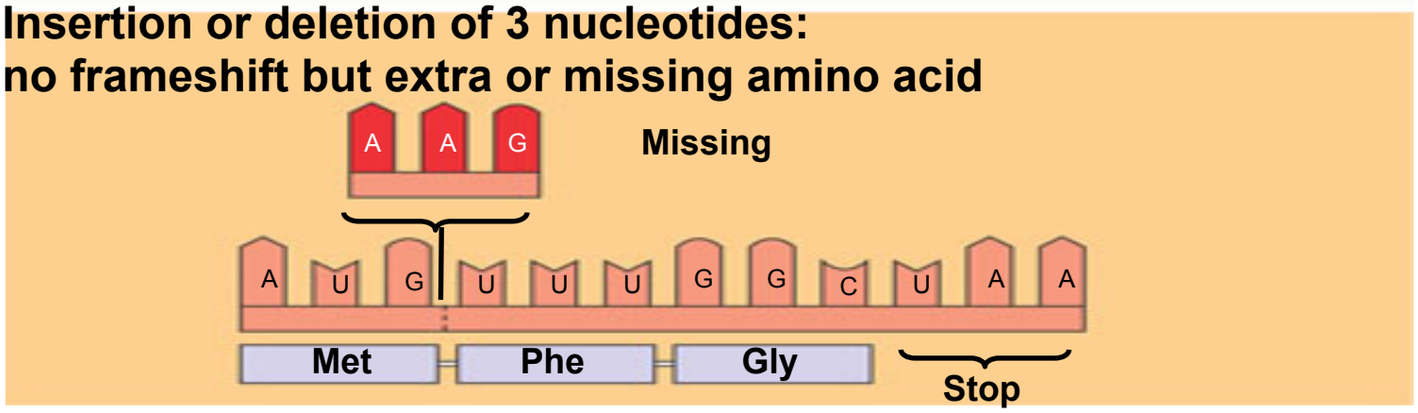
Replacement of one nucleotide and its partner with another pair of nucleotides, can result in missense or nonsense mutations

Point Mutations-
Base Insertions & Deletions



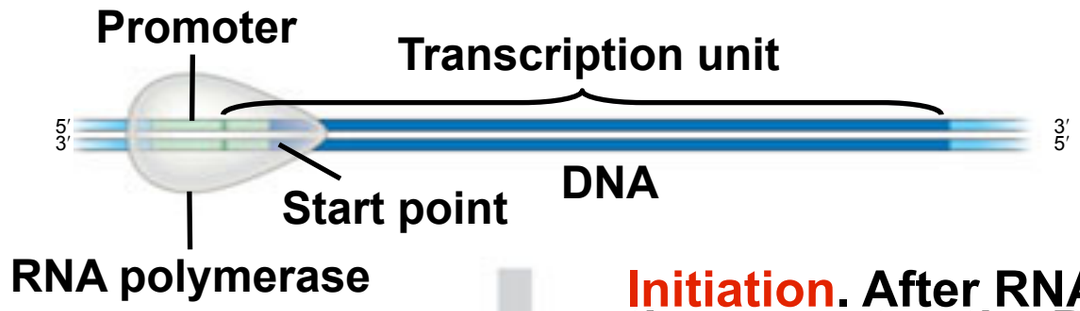




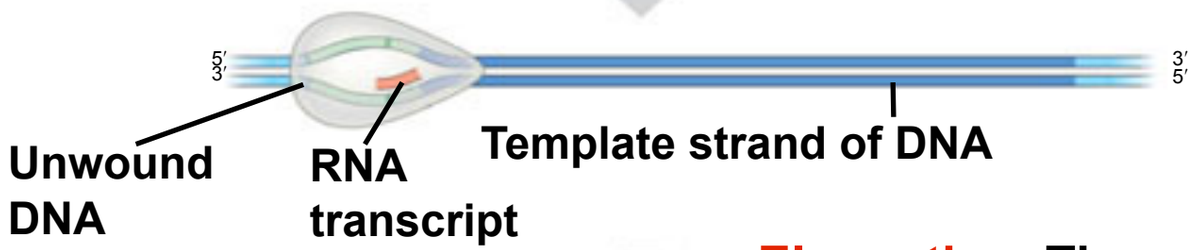


Additions and losses of nucleotide pairs in a gene, can cause frameshifts

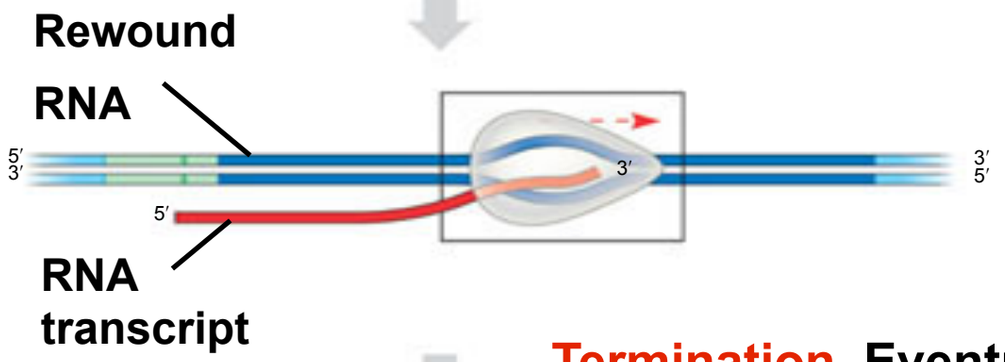
Prokaryotic Transcription



Initiation. After RNA polymerase binds to the promoter, the DNA strands unwind, and the polymerase initiates RNA synthesis at the start point on the template strand.



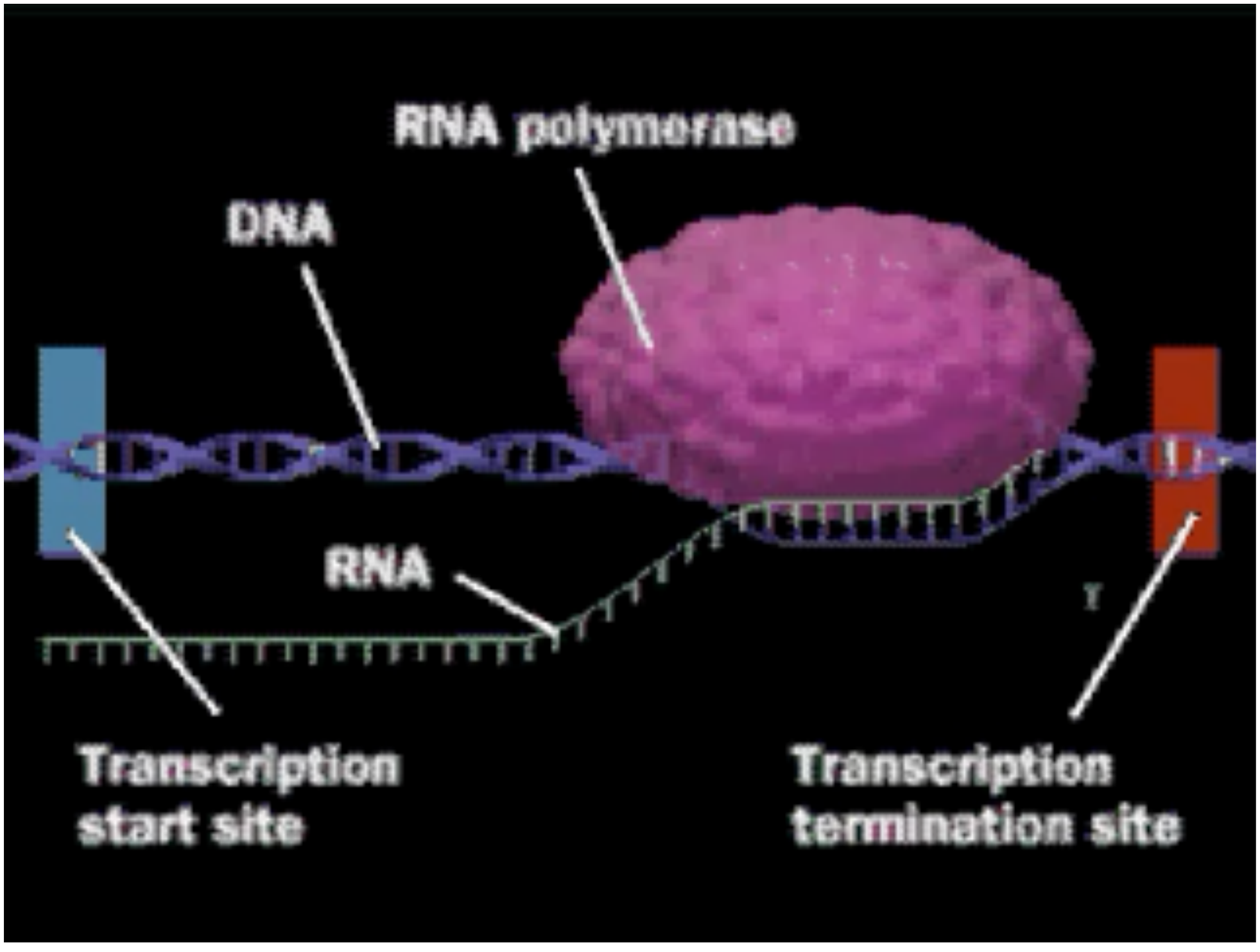
Elongation. The polymerase moves downstream, unwinding the DNA and elongating the RNA transcript 5′ → 3′. In the wake of transcription, the DNA strands re-form a double helix



Termination. Eventually, the RNA transcript is released, and the polymerase detaches from the DNA.

As soon as one RNA polymerase moves off of promoter another one can bind and so forth like a convoy of trucks

Transcription

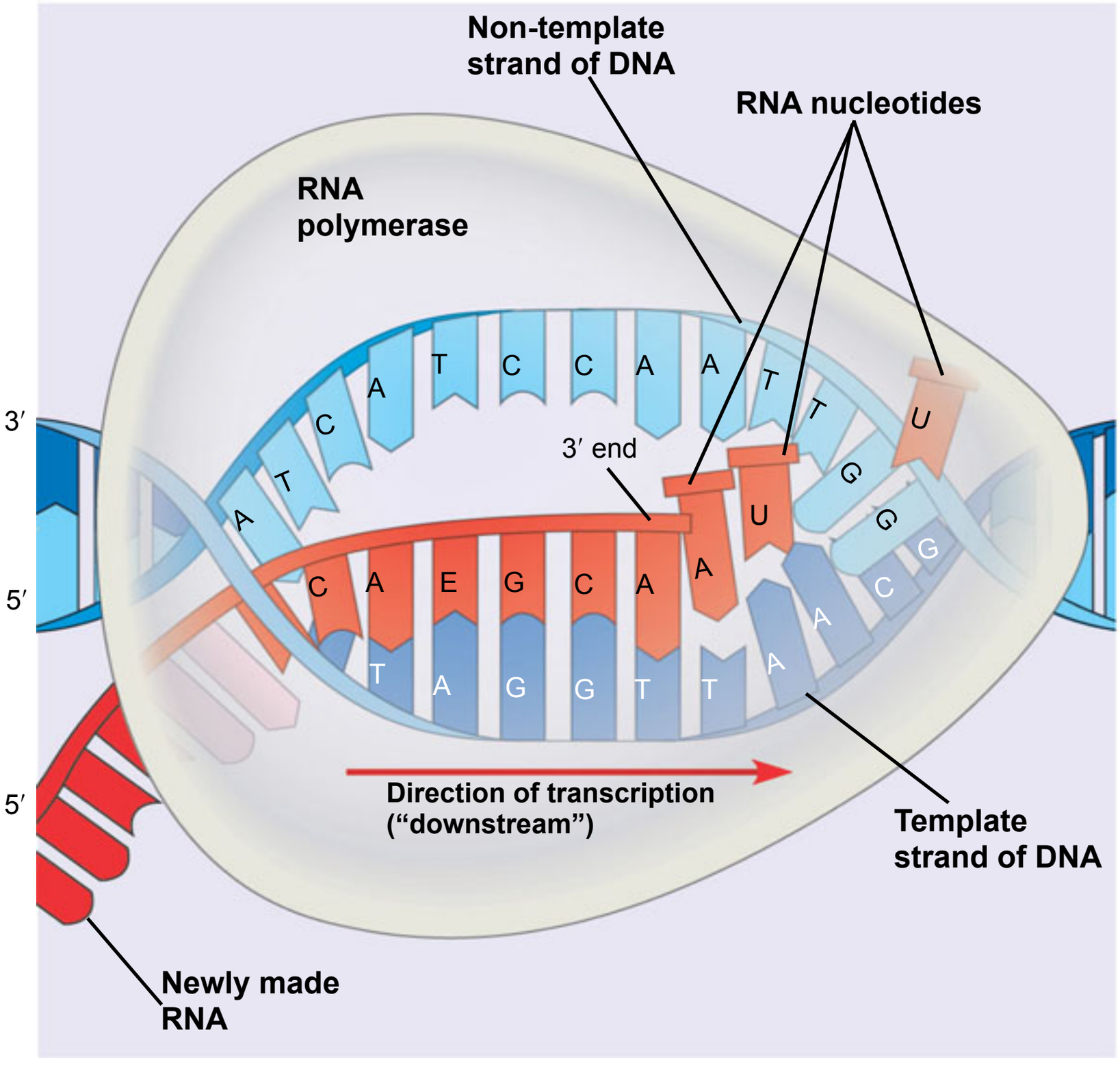


Prokaryotic Transcription

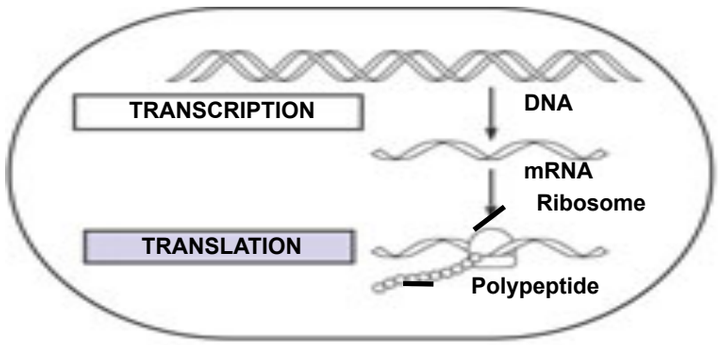
Elongation

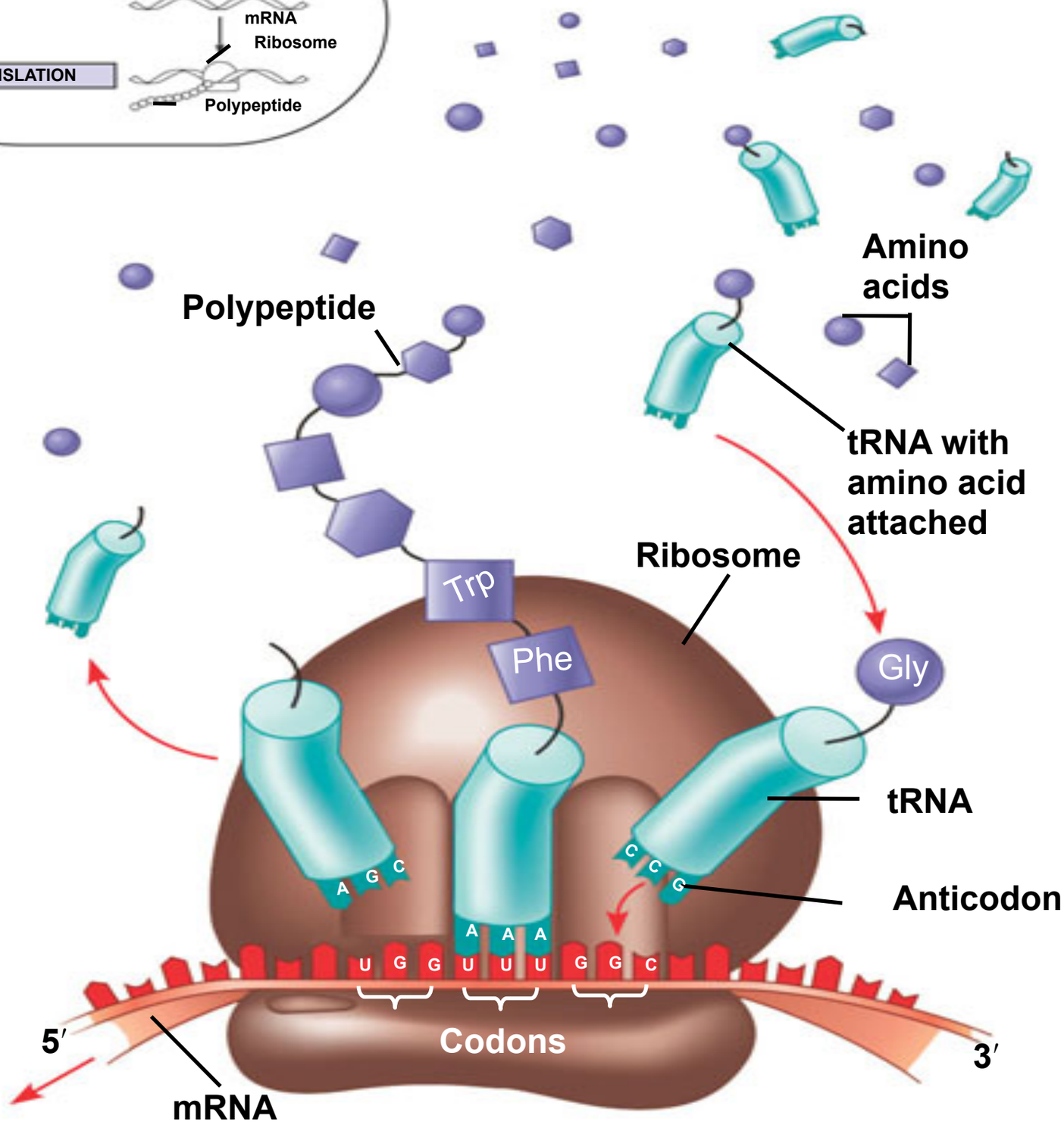
RNA polymerase
-uncoils DNA
-splits DNA
-holds DNA open
-adds RNA nucleotides

proceeds at a rate of approx. 40 nucleotides per second



Prokaryotic Translation





the cytosol is stocked with free floating amino acids

tRNA's are also floating freely

every specific amino acid is carried by a tRNA carrying specific anticodon

Prokaryotic Translation

- Translation involves 3 steps, also named...
  - **Initiation**
  - **Elongation**
  - **Termination**

- Translation involves a number of different “characters”...
  - **tRNA**
  - **ribosomes (small & large subunits)**
  - **mRNA**
  - **amino acids**

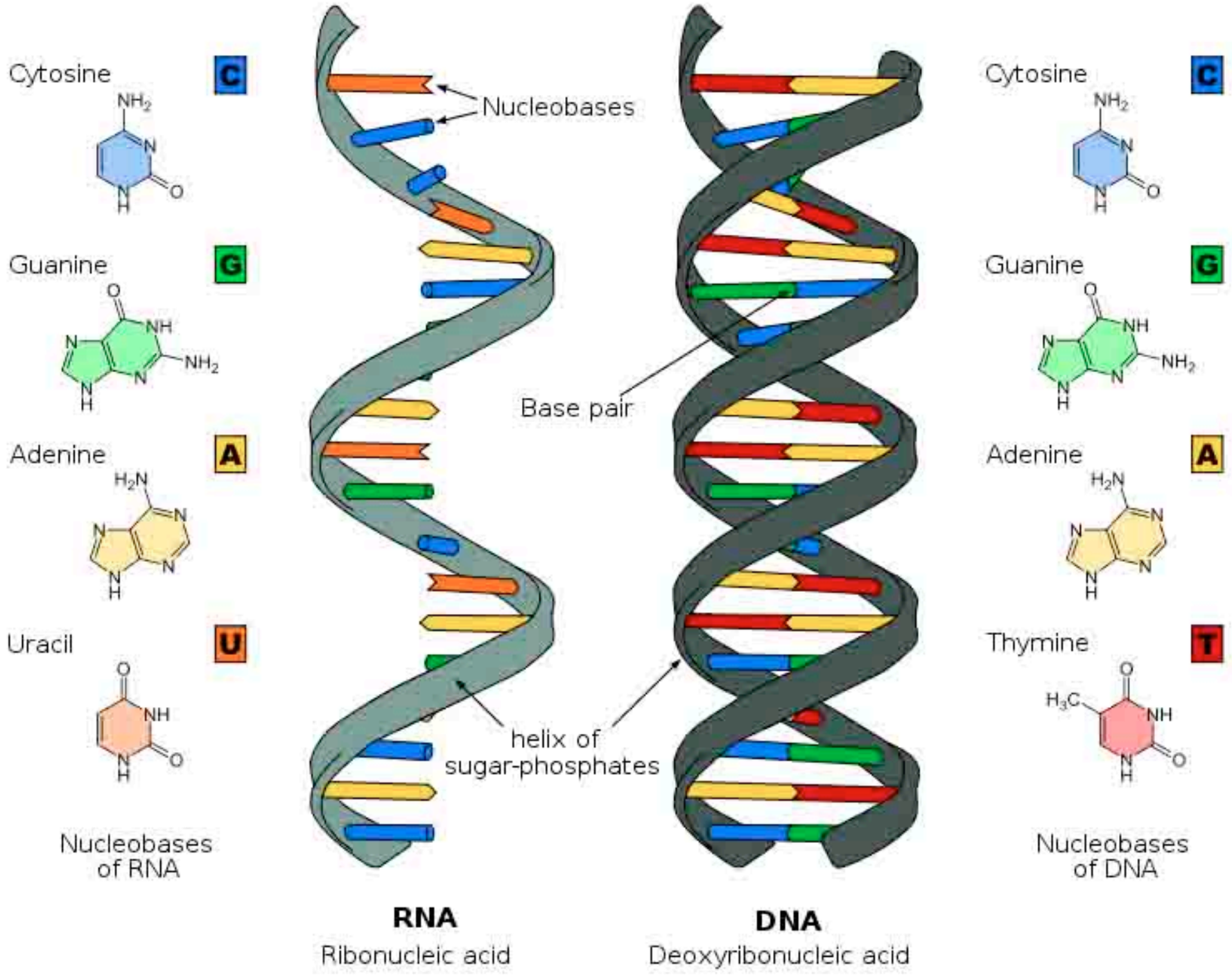
mRNA

"Our Cast of Characters"

single stranded

Uracil

ribose sugar in the backbone

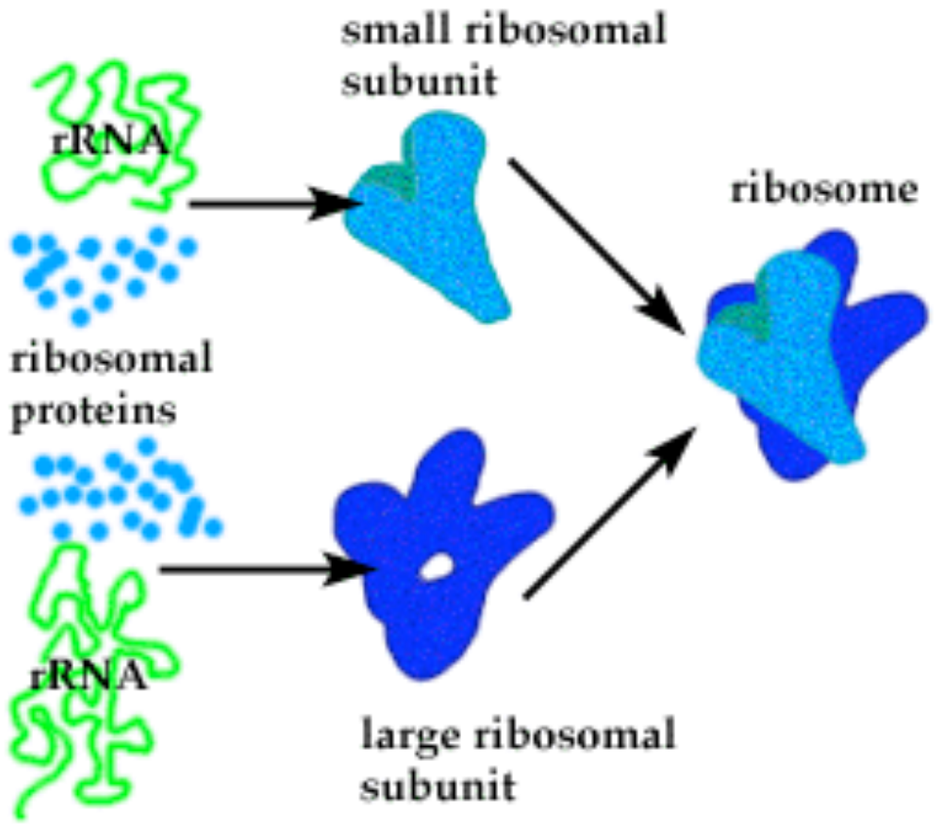


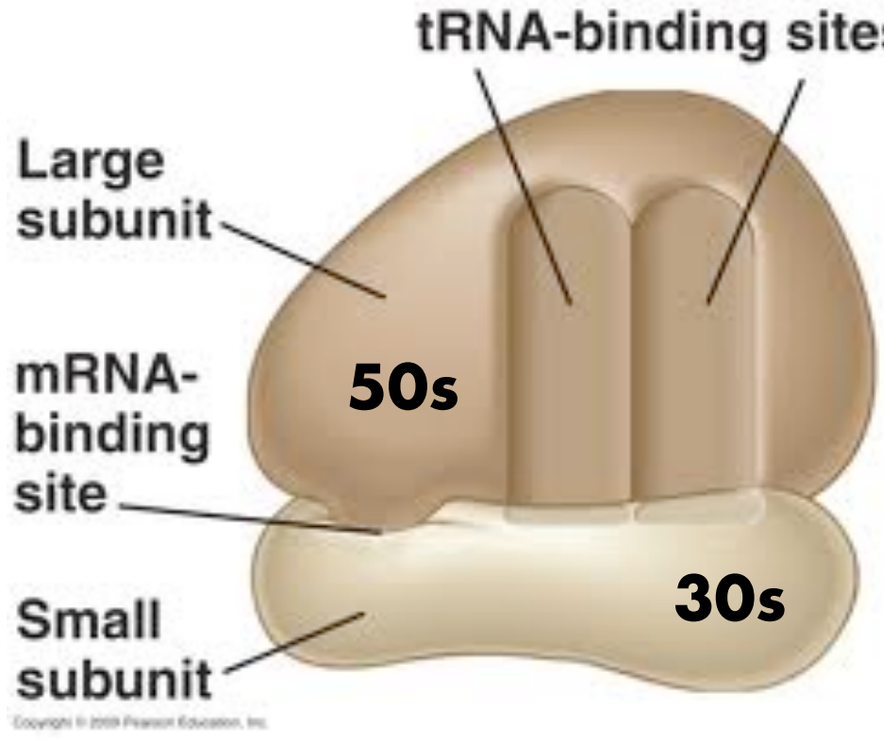
Ribosomes

"Our Cast of Characters"

Translation :

the ribosome



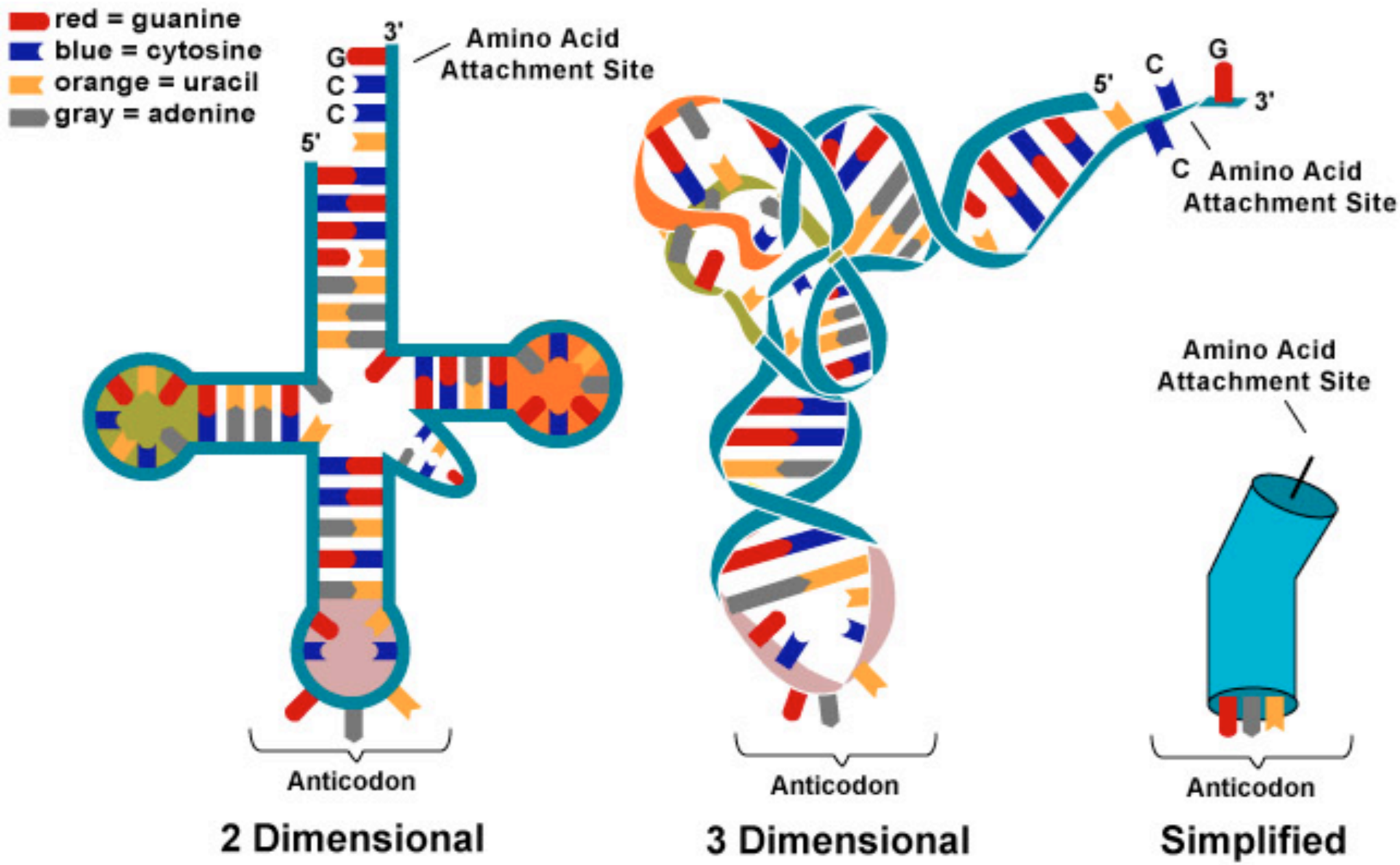


70s

rRNA is the most abundant type of cellular RNA

tRNA

"Our Cast of Characters"

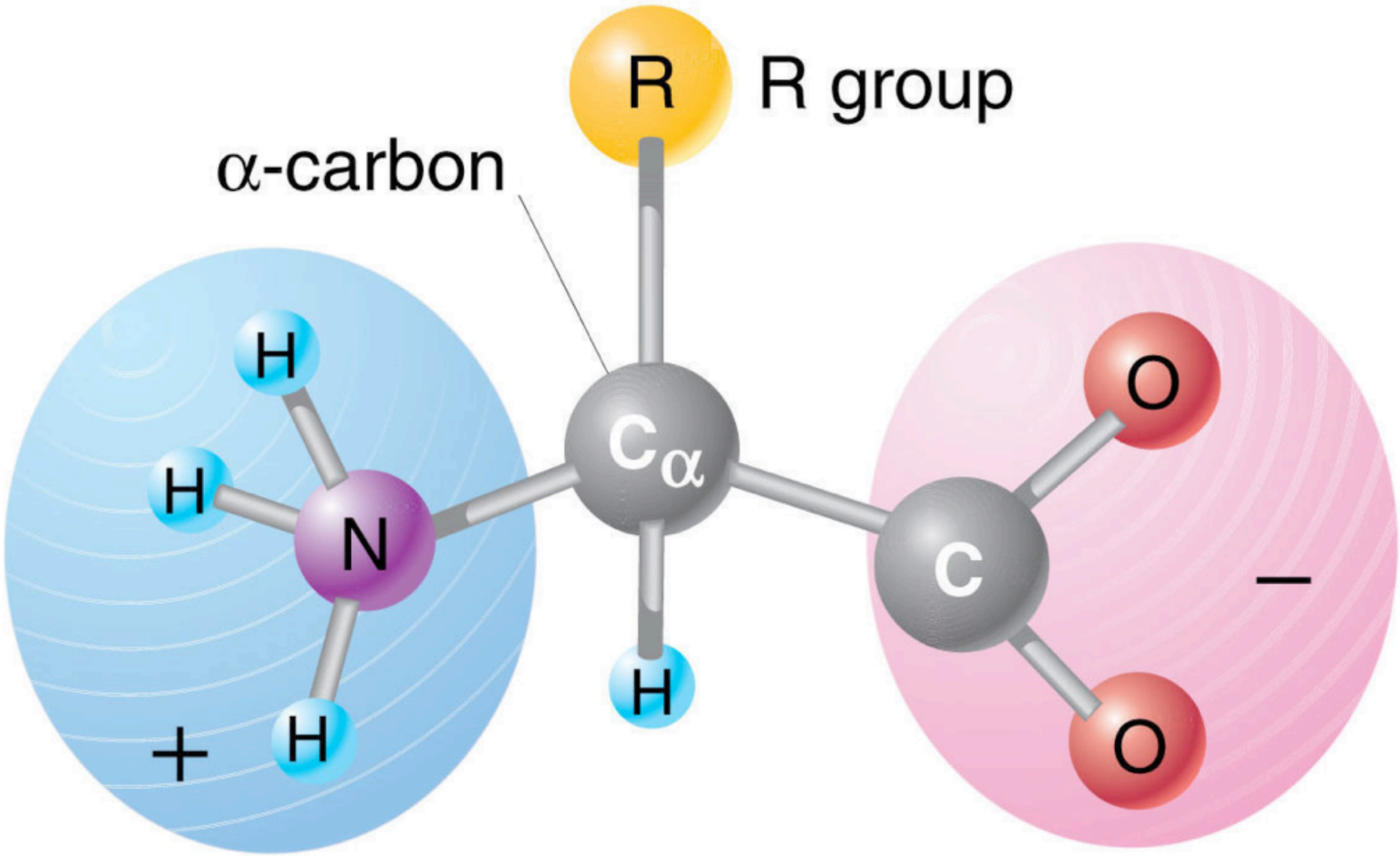


~80 nucleotides

"L" shaped

Amino Acids

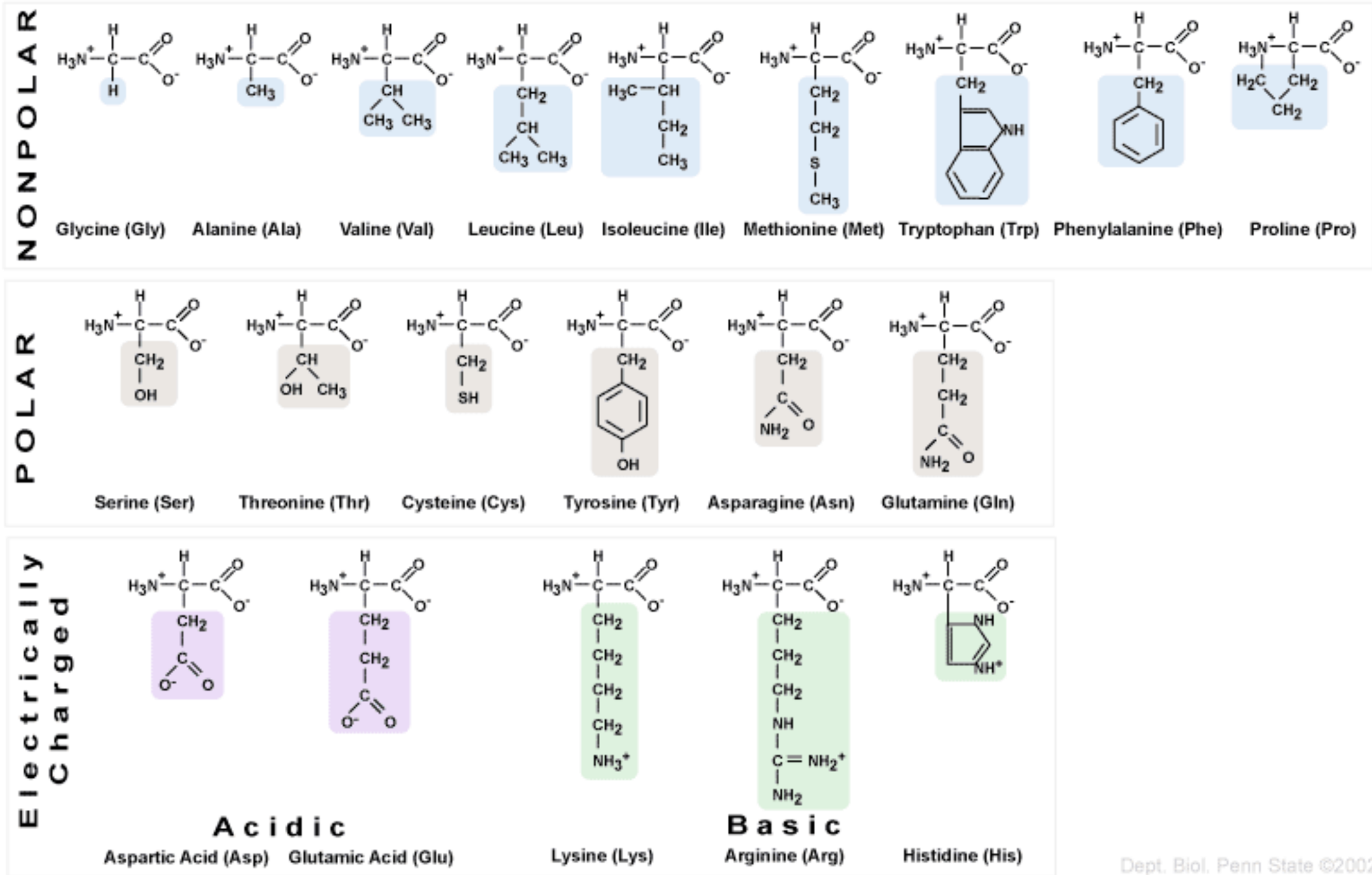
"Our Cast of Characters"



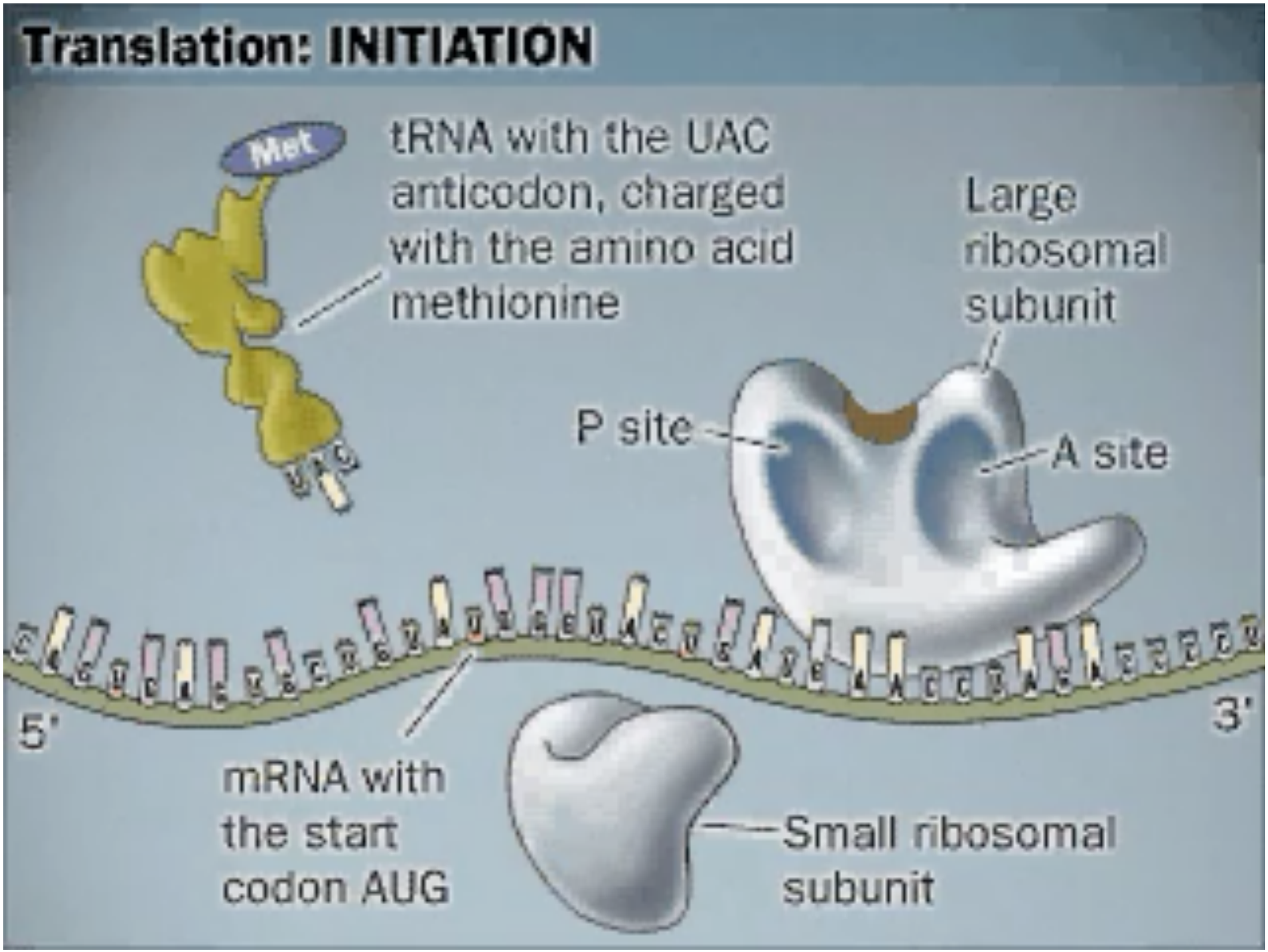
General Amino Acid Structure in Solution

Amino Acids

"Our Cast of Characters"

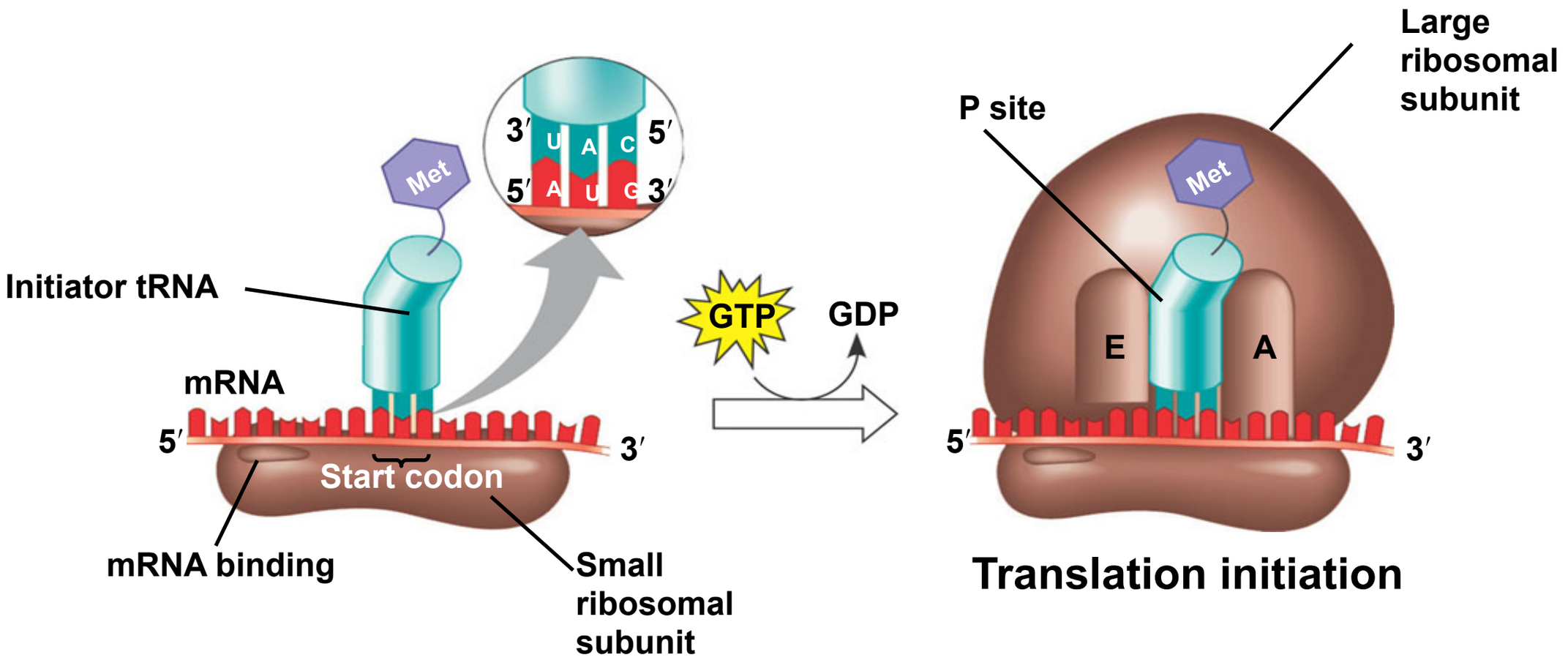


Translation



Translation- Initiation

"The Processes"



Translation initiation

1.

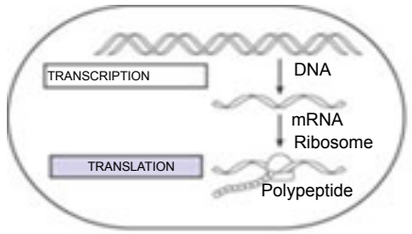
A small ribosomal subunit binds to a molecule of mRNA. In a prokaryotic cell, the mRNA binding site on this subunit recognizes a specific nucleotide sequence on the mRNA just upstream of the start codon. An initiator tRNA, with the anticodon UAC, base-pairs with the start codon, AUG. This tRNA carries the amino acid methionine (Met).

2.

The arrival of a large ribosomal subunit completes the initiation complex. Proteins called initiation factors (not shown) are required to bring all the translation components together. GTP provides the energy for the assembly. The initiator tRNA is in the P site; the A site is available to the tRNA bearing the next amino acid.

Translation- Elongation

"The Processes"



1.

Codon recognition. The anticodon of an incoming aminoacyl tRNA base-pairs with the complementary mRNA codon in the A site. Hydrolysis of GTP increases the accuracy and efficiency of this step.

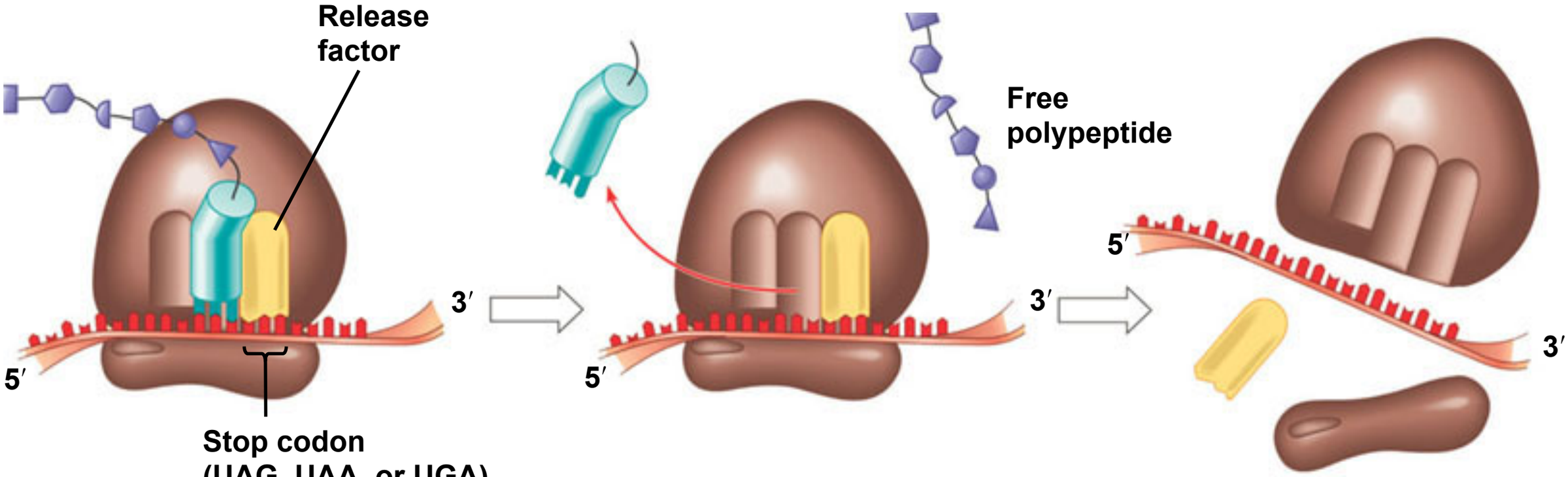
Translocation. The ribosome translocates the tRNA in the A site to the P site. The empty tRNA in the P site is moved to the E site, where it is released. The mRNA moves along with its bound tRNAs, bringing the next codon to be translated into the A site.

Peptide bond formation. An rRNA molecule of the large subunit catalyzes the formation of a peptide bond between the new amino acid in the A site and the carboxyl end of the growing polypeptide in the P site. This step attaches the polypeptide to the tRNA in the A site.

Translation- Termination

"The Processes"

Keep in mind this polypeptide now must fold into a 3-D molecule before it becomes functional



When a ribosome reaches a stop codon on mRNA, the A site of the ribosome accepts a protein called a release factor instead of tRNA.

1.

The release factor hydrolyzes the bond between the tRNA in the P site and the last amino acid of the polypeptide chain. The polypeptide is thus freed from the ribosome.

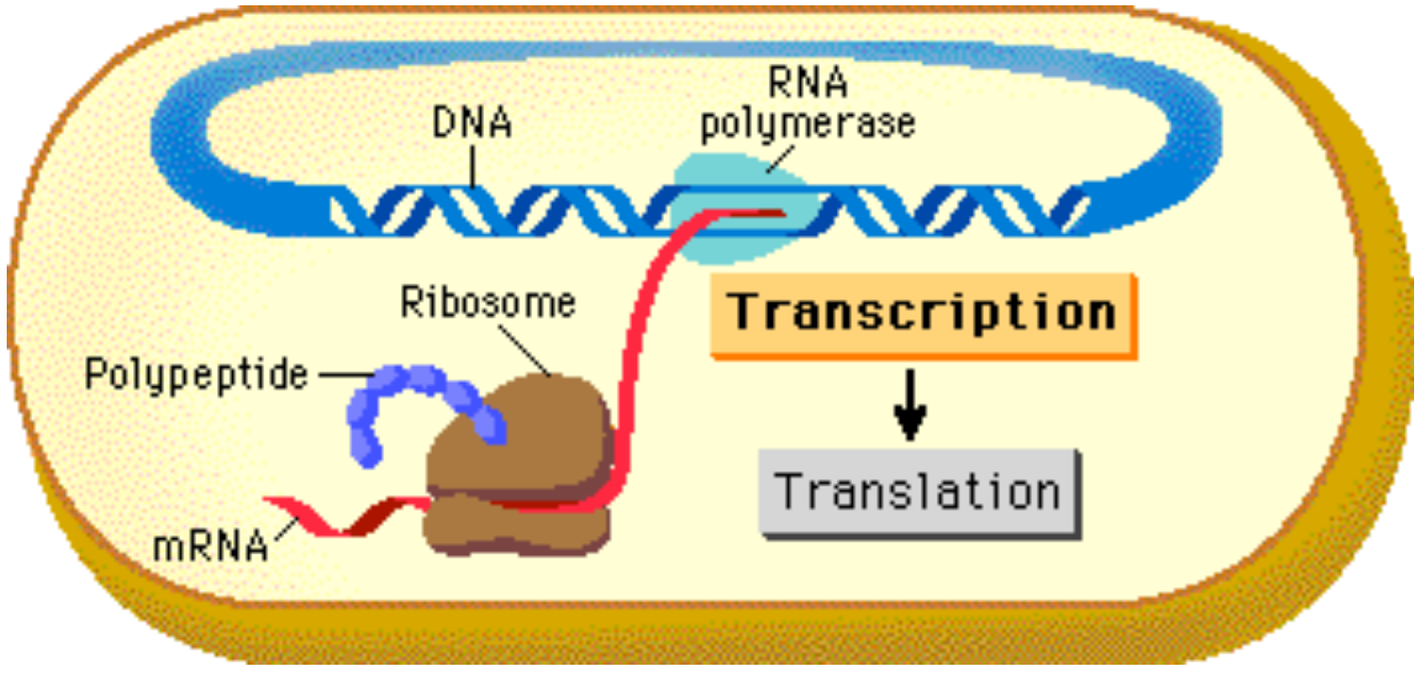
2.

The two ribosomal subunits and the other components of the assembly dissociate. This also requires energy- 2GTP molecules.

3.

A Final Reminder

- **Transcription & Translation occurs in every organism.**
  - ***The mechanics are the same or very similar in all cells***
  - ***However, one very important difference exists between prokaryotes and eukaryotes***



Prokaryotic Transcription and Translation are not separated by time and space

Regulating Gene Expression

- A cell's genome consists of all its genes.
- NOT ALL genes need to be expressed at ALL times.
- ONLY certain genes are expressed at certain times.
- Bacteria turn on and turn off genes in response to the environmental conditions.
- As environmental conditions change so to does gene expression.

Regulating Gene Expression

- A bacteria that can turn its genes on and off in response to environmental changes **will save both resources and energy over time**.
- Natural selection has favored these bacteria over those which have less control.
- *Consider E-coli that live in a human colon, if human meal includes a particular nutrient then they need not produce it (save energy) BUT if human meal does not include a particular nutrient then they need produce it.*
- This fundamentally requires that the E-coli turn on/off certain genes depending on the presence/absence of a particular nutrient.