

Lab 6: The Central Dogma – From DNA to Protein

BIOL-8

Name: _____ Date: _____

Objectives

By the end of this lab, you will be able to:

- **Model DNA replication** by physically separating and rebuilding complementary strands
- **Perform transcription** by converting a DNA template strand into an mRNA sequence
- **Perform translation** by decoding mRNA codons into an amino acid chain
- **Trace the flow of genetic information** from DNA → RNA → Protein (the Central Dogma)
- **Use a codon table** to identify amino acids from mRNA triplets
- **Explain the significance** of start and stop codons in protein synthesis

Introduction

The **Central Dogma of Molecular Biology** describes the flow of genetic information in living cells:

DNA (Nucleus) → **mRNA** (Nucleus → Cytoplasm) → **PROTEIN** (Ribosome)

In this lab, you will use **paper cutouts** of nucleotides and amino acids to physically walk a short DNA sequence through all three processes: **replication**, **transcription**, and **translation**.

Key Terms:

Term	Definition
Nucleotide	Building block of DNA/RNA (base + sugar + phosphate)
Complementary base pairing	A pairs with T (DNA) or U (RNA); G pairs with C
Template strand	The DNA strand read by RNA polymerase (3'→5')
Coding strand	The DNA strand with the same sequence as mRNA (except T→U)

Term	Definition
Codon	Three-nucleotide sequence on mRNA that codes for an amino acid
Anticodon	Three-nucleotide sequence on tRNA that is complementary to a codon
Start codon	AUG — signals the beginning of translation (codes for methionine)
Stop codon	UAA, UAG, or UGA — signals the end of translation

Materials

- Scissors
 - Tape or glue stick
 - Colored pencils or markers (4 colors recommended)
 - **Appendix A:** Printable DNA nucleotide cutouts (A, T, G, C)
 - **Appendix B:** Printable RNA nucleotide cutouts (A, U, G, C)
 - **Appendix C:** Printable amino acid cards
 - **Appendix D:** Standard genetic code (codon table)
 - Blank paper or lab worksheet for assembling sequences

Safety Considerations

- Use scissors carefully
 - Keep workspace clear and organized

The Sequence We Will Use

Throughout this lab, you will work with the following **18-base DNA sequence** (template strand):

Template strand (read 3' → 5'):

This codes for a small protein: **Start + 4 amino acids + Stop**

The complementary **coding strand** is:

Coding strand (5'→3'):

5'- A T G T T C A A A C G T G G C T A A -3'

Part 1: Preparation – Cutting Out Materials

Learning Goal: Prepare the physical models needed to simulate the Central Dogma processes.

Procedure

1. **Cut out** 18 DNA nucleotide pieces for the **template strand**: T-A-C-A-A-G-T-T-T-G-C-A-C-C-G-A-T-T
 2. **Cut out** 18 DNA nucleotide pieces for the **coding strand**: A-T-G-T-T-C-A-A-A-C-G-T-G-C-T-A-A
 3. **Cut out** RNA nucleotides (from Appendix B) to be used for building the complementary mRNA.
 4. **Cut out** the amino acid cards (from Appendix C). Keep these organized for later use.
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Part 2: DNA Replication

Learning Goal: Model the semi-conservative process of DNA replication using paper nucleotide cutouts.

Background

Before a cell divides, it must copy its DNA. During **DNA replication**:

1. **Helicase** unwinds and separates the two strands (breaks hydrogen bonds)
2. **DNA polymerase** reads each template strand and adds complementary nucleotides
3. **Ligase** seals any gaps in the new strands
4. Result: **Two identical double-stranded DNA molecules**

Procedure

1. **Take your cut-out DNA nucleotides** from Part 1.
2. **Arrange** them as a double-stranded DNA molecule, pairing each base with its complement.
3. **Simulate helicase:** Physically separate the two strands by pulling them apart
4. **Simulate DNA polymerase:** For each separated strand, cut out NEW complementary nucleotides and pair them
5. **Tape** the new nucleotides in place to form two complete double-stranded molecules

Data Collection

DNA Replication Results

#	DNA Molecule	Strand 1 Sequence (top)	Strand 2 Sequence (bottom)	Identical to Original?
1				
2				

Analysis

1. Why is DNA replication called "semi-conservative"?

**2. Each new DNA molecule contains one original strand and one new strand.
Which strand is original and which is new in each copy?**

3. What would happen if DNA polymerase made an error and inserted the wrong nucleotide?

Part 3: Transcription – DNA → mRNA

Learning Goal: Convert the DNA template strand into a complementary mRNA sequence, simulating transcription.

Background

During **transcription**, the enzyme **RNA polymerase** reads the DNA template strand and builds a complementary **mRNA** molecule. Key differences from DNA replication:

- RNA uses **uracil (U)** instead of **thymine (T)**
- RNA is **single-stranded**
- Only **one strand** of DNA (the template strand) is read

Base Pairing Rules used for Transcription:

DNA Template Base mRNA Base

A	U
T	A
G	C
C	G

Procedure

1. Take your template strand from earlier:

3' – T A C A A G T T T G C A C C G A T T – 5'

1. Use your **RNA nucleotide cutouts** from Part 1.
2. Apply the base pairing rules: A→U, T→A, G→C, C→G
3. Arrange the **mRNA nucleotides** in a single row next to the DNA template, reading 3'→5' on the template to produce the mRNA 5'→3'

4. **Tape** the mRNA strand together in order

Build Your mRNA

Transcription: DNA → mRNA

#	Position	DNA Template Base	→	mRNA Base
1				
2				
3				
4				
5				
6				

Write your complete mRNA sequence here:

5'— —3'

Which base in RNA replaces thymine (T) from DNA?

Analysis

1. Transcription occurs in which part of the cell?

2. Why does the cell make an mRNA copy rather than using the DNA directly for protein synthesis?

For more information about the study, please contact Dr. John Smith at (555) 123-4567 or via email at john.smith@researchinstitute.org.

3. Compare the mRNA sequence you built to the coding strand of DNA. What do you notice?

For more information about the study, please contact the study team at 1-800-258-4263 or visit www.cancer.gov.

Part 4: Translation – mRNA → Protein

Learning Goal: Decode the mRNA message into a chain of amino acids, simulating translation at the ribosome.

Background

During **translation**, the ribosome reads the mRNA **three nucleotides at a time** (codons). Each codon specifies an amino acid. **tRNA** molecules carry amino acids to the ribosome and match their **anticodon** to the mRNA **codon**.

mRNA:	5' – A U G	U U C	A A A	C G U	G G C	U A A	–3'
	↓	↓	↓	↓	↓	↓	
Amino acids:	Met	Phe	Lys	Arg	Gly	STOP	
	(start)						(release)

Use the **Codon Table** (Appendix) to decode your sequence.

Procedure

1. **Take your mRNA strand** from Part 3.
 2. **Divide it into codons** — group the mRNA into sets of three nucleotides.
 3. **For each codon**, look up the amino acid in the Codon Table (Appendix).
 4. **Select the corresponding amino acid cards** from your prepared set.
 5. **Assemble the polypeptide chain** — line up the amino acid cards in order, left to right.

6. Stop when you reach the stop codon — do not add an amino acid for it.

Decode Your mRNA

Translation: mRNA → Amino Acids

#	Codon #	mRNA Codon	Amino Acid	tRNA Anticodon
1				
2				
3				
4				
5				
6				

Write your final amino acid sequence (polypeptide) here:

How many amino acids are in your polypeptide (not counting the stop)?

Analysis

1. What is the start codon, and what amino acid does it always code for?

2. What happens at the ribosome when a stop codon is reached?

3. The codon table shows that multiple codons can code for the same amino acid (e.g., UUU and UUC both code for Phe). Why might this "redundancy" be beneficial?

4. What is the role of tRNA in translation?

Part 5: Mutation Simulation

Learning Goal: Observe how a single base change in DNA can alter the protein produced.

Procedure

1. Start with the original DNA template strand: 3'-TAC AAG TTT GCA CCG ATT-5'
2. Introduce a mutation: Change the 10th base (G) to an A, so position 10–12 becomes ACA instead of GCA
3. Transcribe the mutated template strand into mRNA
4. Translate the mutated mRNA into amino acids

Mutated Sequence

Mutation Analysis

#	Codon Position	Original DNA Template	Mutated DNA Template	Original mRNA	Mutated mRNA	Original AA	Mutated AA
1							
2							
3							
4							
5							
6							

Analysis

1. Did the mutation change the amino acid at position 4? If so, what changed?

2. What type of mutation is this (substitution, insertion, or deletion)?

3. Would a single base change always change the amino acid? Why or why not?

4. How might a changed amino acid affect the function of the protein?

Part 6: Summary – Tracing the Central Dogma

Complete the Flowchart

Fill in each step of the central dogma for your original sequence:

DNA Template Strand: 3'— —5'

↓ Transcription (by RNA polymerase, in the nucleus)

mRNA: 5'— —3'

↓ Translation (by ribosomes, in the cytoplasm)

Amino Acid Sequence:

Number of amino acids in final protein:

Conclusions

1. In your own words, explain the central dogma of molecular biology and why it is important:

2. Why does the cell transcribe DNA into mRNA rather than using DNA directly at the ribosome?

3. If a gene has 300 nucleotides (excluding start and stop), how many amino acids would the resulting protein contain? Show your reasoning:

4. A classmate says "DNA makes protein." Is this statement accurate? How would you correct or clarify it?

5. How does what you learned in this lab connect to the concept of genetic mutations and inherited diseases?

Quick Reference

Central Dogma Summary

Process	Location	Enzyme	Input	Output
DNA Replication	Nucleus	DNA polymerase	1 DNA molecule	2 identical DNA molecules
Transcription	Nucleus	RNA polymerase	DNA template strand	mRNA
Translation	Ribosome (cytoplasm)	—	mRNA + tRNA + amino acids	Polypeptide (protein)

Base Pairing Rules

DNA Replication Transcription

$$A \leftrightarrow T \quad A \text{ (DNA)} \rightarrow U \text{ (mRNA)}$$

$$T \leftrightarrow A \quad T \text{ (DNA)} \rightarrow A \text{ (mRNA)}$$

DNA Replication Transcription

$G \leftrightarrow C$ G (DNA) \rightarrow C (mRNA)
 $C \leftrightarrow G$ C (DNA) \rightarrow G (mRNA)

Connection to Module 07: This lab directly applies the Central Dogma concepts from Module 07 (Genetics). Understanding how DNA encodes proteins through transcription and translation is foundational for genetics, inheritance (Module 09), and understanding how mutations cause disease. Every protein in your body — from hemoglobin to digestive enzymes — is built through this process.

Appendix: Codon Table (Tear-off Sheet)

Standard Genetic Code

		Second Position						
First Position		U	C	A	G			
(5' end)								
U	UUU	Phe	UCU	Ser	UAU	Tyr	UGU	Cys
	UUC	Phe	UCC	Ser	UAC	Tyr	UGC	Cys
	UUA	Leu	UCA	Ser	UAA	STOP	UGA	STOP
	UUG	Leu	UCG	Ser	UAG	STOP	UGG	Trp
C	CUU	Leu	CCU	Pro	CAU	His	CGU	Arg
	CUC	Leu	CCC	Pro	CAC	His	CGC	Arg
	CUA	Leu	CCA	Pro	CAA	Gln	CGA	Arg
	CUG	Leu	CCG	Pro	CAG	Gln	CGG	Arg
A	AUU	Ile	ACU	Thr	AAU	Asn	AGU	Ser
	AUC	Ile	ACC	Thr	AAC	Asn	AGC	Ser
	AUA	Ile	ACA	Thr	AAA	Lys	AGA	Arg
	AUG	Met/START	ACG	Thr	AAG	Lys	AGG	Arg
G	GUU	Val	GCU	Ala	GAU	Asp	GGU	Gly
	GUC	Val	GCC	Ala	GAC	Asp	GGC	Gly
	GUA	Val	GCA	Ala	GAA	Glu	GGA	Gly
	GUG	Val	GCG	Ala	GAG	Glu	GGG	Gly