

# Lab 10: Module 07–10 Review — Genetics, Cell Division, Tissues & Inheritance

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BIOL-8

Name: \_\_\_\_\_ Date: \_\_\_\_\_

Name:  Date:

Section:  Lab Partner(s):

## Purpose

This lab is a structured, paper-based review to prepare you for **Exam 02**, which covers:

Module Topic	Key Concepts
07	Genetics & DNA DNA structure, replication, transcription, translation, mutations
08	Cell Division Mitosis, meiosis, the cell cycle, comparison of both processes
09	Tissues Four tissue types, structure-function relationships, locations
10	Inheritance Mendelian laws, Punnett squares, pedigrees, sex-linked traits

**How to use this lab:** Work through each section systematically. The activities progress from recall → application → analysis → synthesis. Complete all fill-in tables and reflection questions honestly — this is practice for the exam.

## Part 1: Module 07 — Genetics & DNA

### 1A. Core Vocabulary Check

Fill in the blank with the correct term:

## Module 07 Vocabulary

#	Definition	Term
1		
2		
3		
4		
5		
6		
7		
8		
9		
10		

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### 1B. DNA Structure Diagram

In the space below (or on scratch paper), **sketch a short segment of DNA** (at least 4 base pairs) and label:

- Sugar-phosphate backbone
- Hydrogen bonds
- At least two correctly paired bases (A-T and G-C)

**After sketching, answer: Why is the DNA double helix described as "antiparallel"?**

**Why does accurate base pairing matter for DNA replication?**

### 1C. Central Dogma Flow

Complete the flow chart by filling in the blanks:

DNA  $\xrightarrow{\hspace{2cm}}$   $\xrightarrow{\hspace{2cm}}$   
(Transcription) (Translation)

Location:  $\xrightarrow{\hspace{2cm}}$  Location:  $\xrightarrow{\hspace{2cm}}$

**A mutation changes a codon from CAU to CAA. Both code for different amino acids. What type of mutation is this (silent, missense, or nonsense)?**

**Why can a single point mutation sometimes have no effect on the final protein?**

## Part 2: Module 08 — Cell Division

### 2A. The Cell Cycle

Label the phases of the cell cycle in the diagram below by completing the table:

## Cell Cycle Phases

#	Phase	What Happens	Is the Cell Dividing?
1			
2			
3			
4			
5			
6			

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### 2B. Mitosis Phase Sequencing

**Cut and arrange** (or simply number 1–6 in order) the following events:

## Order the Events of Mitosis

#	Event	Order (1–6)
1		
2		
3		
4		
5		
6		
7		

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## 2C. Mitosis vs. Meiosis Comparison

### Comparing Mitosis and Meiosis

#	Feature	Mitosis	Meiosis
1			
2			
3			
4			
5			
6			
7			
8			
9			

**Why must meiosis produce haploid cells? (Think about what happens at fertilization.)**

**Name two ways meiosis generates genetic variation:**

1.

2.

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## 2D. Chromosome Counting Practice

Use the following starting cell: **diploid organism,  $2n = 6$**  (3 homologous pairs).

### Chromosome Counts Through Division

#	Stage	Number of Chromosomes per Cell	Number of Cells	Total Chromatids
1				
2				
3				
4				
5				

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## Part 3: Module 09 — Tissues

### 3A. The Four Tissue Types

#### Four Primary Tissue Types

#	Tissue Type	Primary Function	Key Characteristic	Example Location in Body
1				
2				
3				
4				
5				

### 3B. Epithelial Tissue Classification

Epithelial tissue is classified by two features: **shape** (squamous/cuboidal/columnar) and **layers** (simple/stratified/pseudostratified).



Epithelial Tissue Types

#	Epithelial Type	Shape	Layers	Location / Function
1				
2				
3				
4				
5				
6				

3C. Connective Tissue Variety

Match each connective tissue type to its defining feature and location:

## Connective Tissue Types

#	Connective Tissue	Defining Feature	Location
1			
2			
3			
4			
5			
6			

### 3D. Tissue Identification Scenarios

For each description, identify the most likely tissue type and subtype:

**1. You observe cells that are flat and scale-like, arranged in a single layer lining the inside of a blood vessel. What tissue is this?**

**2. You observe long, cylindrical, multinucleated cells with visible striations (stripes). This tissue is:**

**3. You observe cells scattered in a gel-like matrix with abundant collagen fibers running in all directions. This is most likely:**

**4. A tissue sample shows branching, involuntary cells with intercalated discs connecting them. This is:**

**5. A tissue with cells that have long processes (axons and dendrites) embedded in a matrix of support cells (glia). This is:**

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### **3E. Structure-Function Synthesis**

**Why does the intestinal lining use simple columnar epithelium rather than stratified squamous epithelium? (Hint: think about absorption vs. protection.)**

**The trachea (windpipe) is lined with pseudostratified ciliated columnar epithelium. What function do the cilia serve, and why is this important for lung health?**

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## **Part 4: Module 10 — Inheritance**

### **4A. Mendel's Laws**

Fill in the blanks:

## Mendel's Laws

#	Law	Statement	What It Predicts
1			
2			
3			

### 4B. Monohybrid Cross Practice

**Scenario:** Freckles (F) are dominant over no freckles (f). Two heterozygous parents (Ff × Ff) have children.

#### 1. Complete the Punnett square:

	F	f
F		
f		

What fraction of offspring will have freckles?

What fraction will be homozygous recessive (ff)?

What is the genotypic ratio (AA : Aa : aa)?

What is the phenotypic ratio (dominant : recessive)?

#### 4C. Dihybrid Cross

**Scenario:** Smooth skin (S) is dominant over rough (s). Brown eyes (B) is dominant over blue (b). Two parents heterozygous for both traits cross: **SsBb × SsBb**.

List all possible gametes each parent can produce:

Using the **9:3:3:1** ratio, out of 16 offspring, how many would you expect to be:

- Smooth skin AND brown eyes:  / 16
- Smooth skin AND blue eyes:  / 16
- Rough skin AND brown eyes:  / 16
- Rough skin AND blue eyes:  / 16

If the genes for skin texture and eye color are on the same chromosome (linked), would independent assortment still apply? Explain:

#### 4D. Pedigree Analysis

The following family has a history of cystic fibrosis (CF). CF is **autosomal recessive** (let C = normal allele, c = CF allele).



Generation III: ☐ ☐

*Legend: □ = unaffected male, ○ = unaffected female, ● = affected female. Lines between individuals indicate mating; vertical lines indicate offspring.*

**The shaded individual in Generation II has CF. What is her genotype?**

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**What must the genotypes of her parents (Generation I) be?**

\_\_\_\_\_

**Her unaffected brother in Generation II marries an unaffected woman with no family history. What is the probability that their child (Generation III) will have CF? Show your reasoning.**

**Could CF appear in Generation III even if neither parent is known to be a carrier?**  
**Explain:**

#### 4E. Sex-Linked Inheritance

**Key Rule:** Genes on the X chromosome follow X-linked inheritance. Males (XY) only have one copy of X-linked genes, so they express whatever allele is present.

**Scenario:** Color blindness is **X-linked recessive** ( $X^B$  = normal vision,  $X^b$  = color blind).

**Cross:** Carrier female ( $X^B X^b$ )  $\times$  Normal male ( $X^B Y$ )

**Complete the Punnett square:**

$X^B$	$Y$

X <sup>B</sup>		
X <sup>b</sup>		

**What is the probability of a color-blind son?**

**What is the probability of a color-blind daughter?**

**Why are males much more commonly affected by X-linked recessive conditions than females?**

**In hemophilia (also X-linked recessive), could a daughter be affected? Under what circumstances?**

## Part 5: Integration — Connecting the Modules

### 5A. Cross-Module Concept Map

On your scratch paper, **draw a concept map** (or complete the partial map below) that shows how the four modules are connected. Key relationships to include:

- DNA (Module 07) → controls cell function → required for cell division (Module 08)
- Cell division (Module 08) → meiosis → produces gametes → enables inheritance (Module 10)
- Inheritance (Module 10) → determines which genes are expressed → affects tissue development (Module 09)

- Mutations in DNA (Module 07) → can disrupt the cell cycle (Module 08) → can alter tissue type or function (Module 09)

**In your own words, explain how a single mutation in a DNA repair gene (Module 07) could ultimately lead to uncontrolled cell division (Module 08). What is the medical term for this condition?**

**Sickle cell disease is caused by a single point mutation (Module 07). It is inherited recessively (Module 10). The abnormal hemoglobin causes red blood cells to sickle and clog capillaries. This affects which tissue types (Module 09)?**

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## **5B. Exam Readiness Self-Assessment**

Rate your confidence on each topic (1 = need more study, 5 = very confident):



## Exam Readiness Check

#	Topic	Confidence (1–5)	One Question I Still Have
1			
2			
3			
4			
5			
6			
7			
8			
9			
10			

### 5C. Final Synthesis Questions

**1. What is the relationship between meiosis and Mendel's Law of Independent Assortment? (Which stage of meiosis directly explains this law?)**

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**2. A cell has  $2n = 46$  chromosomes. After meiosis I is complete, how many chromosomes are in each cell? After meiosis II? Are these chromosomes duplicated (as sister chromatids) or single-stranded at those points?**

**3. Two parents are both carriers (heterozygous) for an autosomal recessive condition. They have 4 children. Predict how many children you would expect to be affected. Does this mean exactly that number will be affected? Why or why not?**

**4. Explain why stratified squamous epithelium is well-adapted for the skin (outer epidermis) but would be a poor choice for lining the lungs' alveoli (air sacs).**

**5. BONUS — Critical Thinking:** A woman is a carrier for an X-linked recessive condition. Her sister is also a carrier. They each have one son. For each son, what is the probability of being affected? If both sons are affected, is that evidence of a genetic link between the brothers, or just chance? Explain.

## Quick Reference: Key Formulas & Rules

Concept	Rule / Formula
Independent assortment combinations	$2^n$ (where $n$ = number of chromosome pairs)
Punnett square offspring types	Monohybrid: 4 boxes; Dihybrid: 16 boxes
Diploid ( $2n$ ) after mitosis	Still $2n$
After meiosis I	$n$ chromosomes (duplicated)
After meiosis II	$n$ chromosomes (single-stranded)
X-linked recessive in males	Only one X needed for expression

**Concept****Rule / Formula**

X-linked recessive in females

Two recessive alleles needed

Carrier (heterozygous) phenotype

Dominant phenotype (does not show recessive trait)

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**Connection to Exam 02:** This lab directly maps to all content on Exam 02 (Modules 07–10). If you can answer every question on this worksheet without looking at your notes, you are well prepared.

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