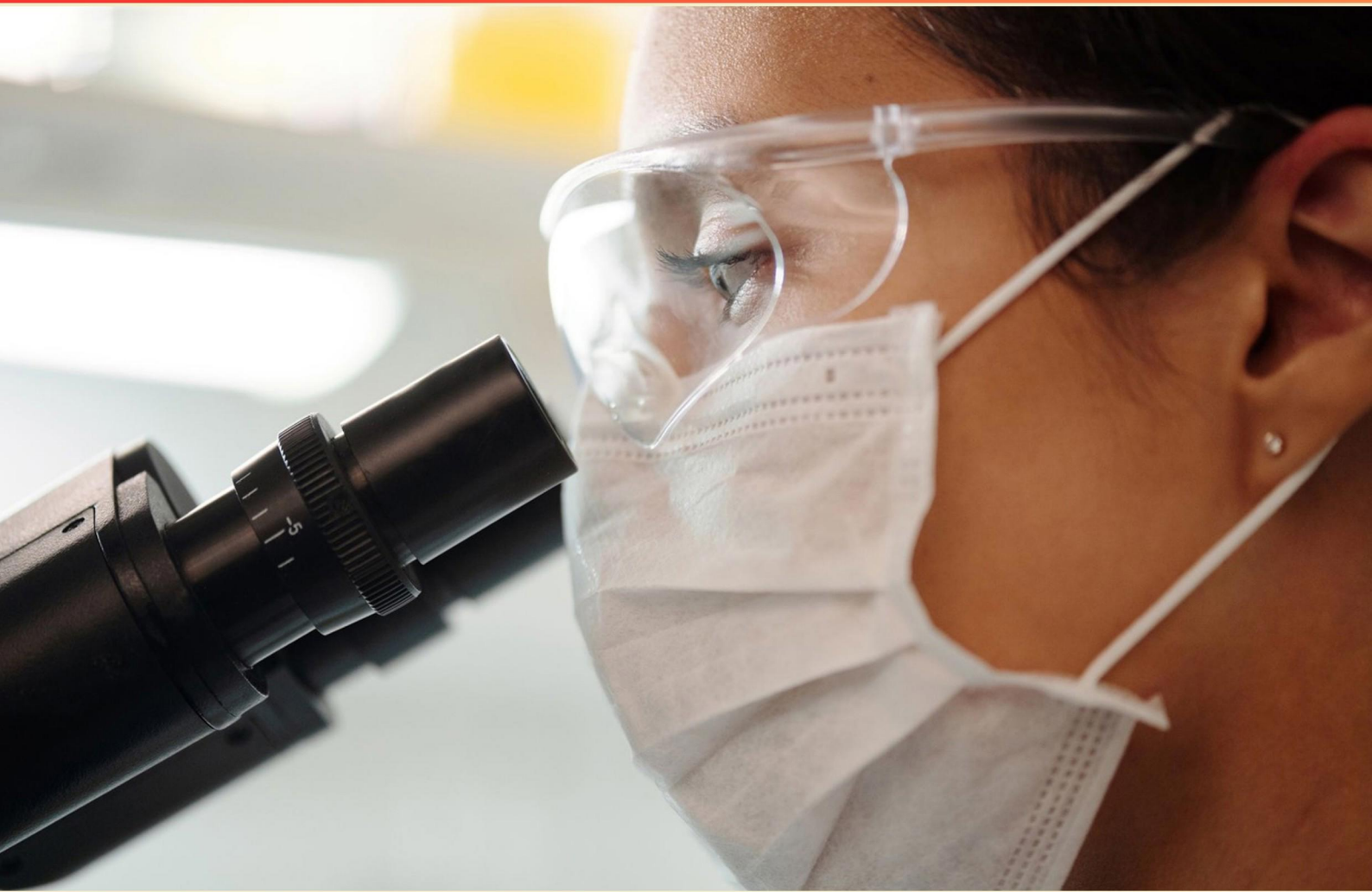


TEXAS A&M UNIVERSITY (TAMU) BIOL111 INTRODUCTORY BIOLOGY I EXAM 3 PRACTICE EXAM (SAMPLE)

STUDY GUIDE



SAMPLE STUDY GUIDE WITH LIMITED QUESTIONS

VISIT US ONLINE FOR
THE FULL VERSION STUDY GUIDE

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Table of Contents

Copyright	1
Table of Contents	2
Introduction	3
How to Use This Guide	4
Questions	5
Answers	8
Explanations	10
Next Steps	15

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Introduction

Preparing for a certification exam can feel overwhelming, but with the right tools, it becomes an opportunity to build confidence, sharpen your skills, and move one step closer to your goals. At Examzify, we believe that effective exam preparation isn't just about memorization, it's about understanding the material, identifying knowledge gaps, and building the test-taking strategies that lead to success.

This guide was designed to help you do exactly that.

Whether you're preparing for a licensing exam, professional certification, or entry-level qualification, this book offers structured practice to reinforce key concepts. You'll find a wide range of multiple-choice questions, each followed by clear explanations to help you understand not just the right answer, but why it's correct.

The content in this guide is based on real-world exam objectives and aligned with the types of questions and topics commonly found on official tests. It's ideal for learners who want to:

- Practice answering questions under realistic conditions,
- Improve accuracy and speed,
- Review explanations to strengthen weak areas, and
- Approach the exam with greater confidence.

We recommend using this book not as a stand-alone study tool, but alongside other resources like flashcards, textbooks, or hands-on training. For best results, we recommend working through each question, reflecting on the explanation provided, and revisiting the topics that challenge you most.

Remember: successful test preparation isn't about getting every question right the first time, it's about learning from your mistakes and improving over time. Stay focused, trust the process, and know that every page you turn brings you closer to success.

Let's begin.

How to Use This Guide

This guide is designed to help you study more effectively and approach your exam with confidence. Whether you're reviewing for the first time or doing a final refresh, here's how to get the most out of your Examzify study guide:

1. Start with a Diagnostic Review

Skim through the questions to get a sense of what you know and what you need to focus on. Your goal is to identify knowledge gaps early.

2. Study in Short, Focused Sessions

Break your study time into manageable blocks (e.g. 30 – 45 minutes). Review a handful of questions, reflect on the explanations.

3. Learn from the Explanations

After answering a question, always read the explanation, even if you got it right. It reinforces key points, corrects misunderstandings, and teaches subtle distinctions between similar answers.

4. Track Your Progress

Use bookmarks or notes (if reading digitally) to mark difficult questions. Revisit these regularly and track improvements over time.

5. Simulate the Real Exam

Once you're comfortable, try taking a full set of questions without pausing. Set a timer and simulate test-day conditions to build confidence and time management skills.

6. Repeat and Review

Don't just study once, repetition builds retention. Re-attempt questions after a few days and revisit explanations to reinforce learning. Pair this guide with other Examzify tools like flashcards, and digital practice tests to strengthen your preparation across formats.

There's no single right way to study, but consistent, thoughtful effort always wins. Use this guide flexibly, adapt the tips above to fit your pace and learning style. You've got this!

Questions

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- 1. What is the largest structural organization of DNA?**
 - A. Linear DNA strand
 - B. 30 nm fiber
 - C. 700 nm fiber
 - D. Chromatin
- 2. Which of the following statements best defines DNA torsion during replication?**
 - A. A twisting force that can hinder the process
 - B. A necessary condition for the separation of strands
 - C. A method for stabilizing the DNA structure
 - D. A process of DNA repair
- 3. What is a telomere?**
 - A. A nucleotide sequence at the end of chromatids
 - B. A type of protein that stabilizes DNA
 - C. A region where RNA binds to DNA
 - D. A segment of coding DNA
- 4. What is the distinguishing feature of haploid cells?**
 - A. They have two sets of chromosomes
 - B. They have one set of chromosomes
 - C. They undergo mitosis
 - D. They are always gametes
- 5. How many replication origins can eukaryotes have per chromosome compared to prokaryotes?**
 - A. One
 - B. Thousands
 - C. Multiple
 - D. No origins

- 6. Which scientists performed experiments showing that DNA is the genetic material of the T2 phage?**
- A. Watson and Crick
 - B. Wilkins and Franklin
 - C. Hershey and Chase
 - D. Avery, MacLeod, McCarty
- 7. How do prokaryotes handle gene expression differently compared to eukaryotes?**
- A. Prokaryotes require processing of mRNA
 - B. Prokaryotes can start translation before transcription finishes
 - C. Eukaryotes do not utilize RNA polymerase
 - D. Prokaryotes use introns extensively
- 8. How many bases correspond to one amino acid in the genetic code?**
- A. 4
 - B. 3
 - C. 21
 - D. 64
- 9. What is the primary function of the cell membrane?**
- A. Regulates the movement of substances in and out of the cell
 - B. Provides structural support to the cell
 - C. Stores genetic information
 - D. Produces energy for cell functions
- 10. What structural feature is characteristic of the 30-nm fiber in DNA?**
- A. It forms looped domains that are attached to proteins
 - B. It exists solely in a relaxed state
 - C. It consists of multiple parallel strands
 - D. It is primarily found in the nucleus

Answers

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1. C
2. A
3. A
4. B
5. B
6. C
7. B
8. B
9. A
10. A

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Explanations

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1. What is the largest structural organization of DNA?

- A. Linear DNA strand
- B. 30 nm fiber
- C. 700 nm fiber**
- D. Chromatin

The largest structural organization of DNA is best represented by chromatin. Chromatin is a complex of DNA and proteins (histones) that organizes DNA within the nucleus of eukaryotic cells. It exists in two forms: euchromatin, which is loosely packed and accessible for transcription, and heterochromatin, which is more tightly packed and usually inactive. Chromatin undergoes further levels of organization; when it condenses during cell division, it forms chromosomes. The structural hierarchy from the linear DNA strand to the 30 nm fiber represents progressive levels of DNA packaging, but it is the chromatin structure that encompasses these forms and represents the largest and most complex organization of DNA in the functional context of the cell. This makes chromatin the most comprehensive answer regarding the structural organization of DNA.

2. Which of the following statements best defines DNA torsion during replication?

- A. A twisting force that can hinder the process**
- B. A necessary condition for the separation of strands
- C. A method for stabilizing the DNA structure
- D. A process of DNA repair

DNA torsion during replication refers to the twisting tension that builds up in the DNA molecule as the double helix unwinds to allow the replication process to occur. This torque results from the inherent helical structure of DNA and can impede the movement of the replication fork if not properly managed. As the two strands of DNA are separated, they create a positive supercoiling ahead of the fork, which can hinder the progression of helicases and polymerases necessary for replication. While it is true that some torsion is necessary for the separation of strands, excessive torsion can lead to problems, making the management of DNA torsion critically important during replication. Additionally, the torsion is not utilized as a method of stabilizing the DNA structure, nor is it a process directly involved in DNA repair. Instead, it is primarily recognized as a challenge that needs to be addressed for successful replication, highlighting why the defining statement focuses on the hindrance caused by this twisting force.

3. What is a telomere?

- A. A nucleotide sequence at the end of chromatids**
- B. A type of protein that stabilizes DNA
- C. A region where RNA binds to DNA
- D. A segment of coding DNA

A telomere is best defined as a nucleotide sequence at the end of chromatids. Telomeres consist of repetitive sequences of DNA that protect the ends of chromosomes from deterioration or from fusion with neighboring chromosomes. This protective function is crucial because, during DNA replication, the enzymes that duplicate DNA cannot replicate the very ends of the linear chromosomes, leading to progressive shortening of chromosomes with each cell division. Telomeres act as a buffer zone to ensure that important coding regions of DNA are not lost. The other choices do not accurately represent what telomeres are. For instance, while proteins do play a role in DNA stability, they are not specifically what a telomere is. Additionally, the binding of RNA to DNA is related more to processes like transcription, not the protective chromosome ends. Finally, coding DNA refers to segments that code for proteins, which contrasts with the non-coding, repetitive nature of telomeres that function mainly in protection rather than in coding for genes.

4. What is the distinguishing feature of haploid cells?

- A. They have two sets of chromosomes
- B. They have one set of chromosomes**
- C. They undergo mitosis
- D. They are always gametes

Haploid cells are characterized by having one set of chromosomes. This means they possess a single copy of each chromosome, which is distinct from diploid cells that contain two sets, one inherited from each parent. In organisms that undergo sexual reproduction, haploid cells are critical as they represent the gametes (sperm and eggs), which fuse during fertilization to restore the diploid chromosome number in the zygote. While it is accurate that many haploid cells are gametes, not all haploid cells are necessarily gametes. For instance, certain plants and fungi have a life cycle stage where they exist as haploid organisms but are not involved in reproduction at that moment. Therefore, simply identifying haploid cells as being always gametes does not encompass the full range of haploid organisms. The primary defining trait of haploid cells remains the presence of a single set of chromosomes.

5. How many replication origins can eukaryotes have per chromosome compared to prokaryotes?

- A. One
- B. Thousands**
- C. Multiple
- D. No origins

Eukaryotic chromosomes typically have thousands of replication origins, which is significantly more than prokaryotic chromosomes that generally have a single origin of replication. This difference is crucial for understanding how DNA replication efficiently occurs within the complex structures of eukaryotic cells. Eukaryotic cells contain linear chromosomes that are larger and have more sequence complexity, necessitating multiple sites along each chromosome where DNA replication can be initiated. These multiple origins allow replication to proceed simultaneously at various locations, ensuring that the entire genome is replicated within a timely manner, especially considering the larger amount of DNA that needs to be copied. In contrast, prokaryotes, which usually possess circular chromosomes, typically originate DNA replication from a single point due to their simpler genome structure. This single origin is sufficient for the relatively smaller size of the prokaryotic genome, enabling efficient replication within a shorter timeframe. Thus, the correct choice reflects the capacity of eukaryotic organisms to have a far greater number of replication origins compared to their prokaryotic counterparts, highlighting a key difference in their cellular organization and replication strategies.

6. Which scientists performed experiments showing that DNA is the genetic material of the T2 phage?

- A. Watson and Crick
- B. Wilkins and Franklin
- C. Hershey and Chase**
- D. Avery, MacLeod, McCarty

Hershey and Chase conducted landmark experiments in 1952 using the T2 bacteriophage to demonstrate that DNA is the genetic material. They used two separate isotopes to label DNA and proteins in the phage: sulfur, which labels the protein coat, and phosphorus, which labels the DNA. After allowing the T2 phage to infect E. coli, they used a blender to separate the phage coats from the bacterial cells and then measured the presence of the isotopes in the bacteria. The results showed that only the phosphorus (DNA) entered the bacterial cells and directed the formation of new phages, while the sulfur (protein) remained outside. This provided clear evidence that DNA, not protein, carries genetic information, establishing DNA as the hereditary material in this viral model, which was a significant finding in molecular biology.

7. How do prokaryotes handle gene expression differently compared to eukaryotes?

- A. Prokaryotes require processing of mRNA
- B. Prokaryotes can start translation before transcription finishes**
- C. Eukaryotes do not utilize RNA polymerase
- D. Prokaryotes use introns extensively

Prokaryotes handle gene expression differently from eukaryotes in several distinct ways, one of which is that they can begin the process of translation while transcription is still underway. This simultaneous process is possible due to the lack of a nucleus in prokaryotic cells. In prokaryotes, as soon as the RNA polymerase starts synthesizing mRNA from DNA, ribosomes can attach to the growing mRNA strand and start translating it into protein. This is a fundamental feature of prokaryotic biology that allows for rapid protein synthesis, which is essential for their survival and adaptation to varying environmental conditions. In contrast, eukaryotic cells have a defined nucleus where transcription occurs, and mRNA must undergo extensive processing (such as capping, polyadenylation, and splicing) before it can be transported out of the nucleus for translation in the cytoplasm. Therefore, the delayed timeline between transcription and translation in eukaryotic cells is markedly different from the simultaneous action found in prokaryotes. This distinction is crucial for understanding cellular biology, as it highlights the more streamlined and efficient nature of prokaryotic gene expression.

8. How many bases correspond to one amino acid in the genetic code?

- A. 4
- B. 3**
- C. 21
- D. 64

In the genetic code, a group of three nucleotide bases, also known as a codon, corresponds to a single amino acid. This means that one amino acid is specified by a sequence of three consecutive bases in the DNA or RNA sequence. This is a fundamental aspect of how genes are translated into proteins, as the sequence of codons in messenger RNA (mRNA) is read by ribosomes during protein synthesis to assemble the correct sequence of amino acids, ultimately forming proteins. While there are four types of nucleotide bases (adenine, thymine/uracil, cytosine, and guanine), it is the combination of three bases in sequences that results in the coding for 20 standard amino acids. Thus, knowing that three bases are necessary to code for one amino acid is central to understanding the process of translation and protein synthesis.

9. What is the primary function of the cell membrane?

A. Regulates the movement of substances in and out of the cell

B. Provides structural support to the cell

C. Stores genetic information

D. Produces energy for cell functions

The primary function of the cell membrane is to regulate the movement of substances in and out of the cell. This selective permeability is essential for maintaining homeostasis within the cell, allowing it to control its internal environment. The cell membrane is composed of a phospholipid bilayer with embedded proteins that facilitate the transport of molecules. These proteins can function as channels, carriers, or receptors, enabling the cell to respond to changes in its surroundings and communicate with other cells. While the cell membrane does contribute to structural support, that is not its primary function; structural support is mainly provided by the cytoskeleton and cell wall (in plant cells). The storage of genetic information is managed by DNA within the nucleus, and energy production primarily occurs in organelles like mitochondria, not the cell membrane. Understanding these distinctions clarifies the essential role of the cell membrane in maintaining the overall function and health of the cell.

10. What structural feature is characteristic of the 30-nm fiber in DNA?

A. It forms looped domains that are attached to proteins

B. It exists solely in a relaxed state

C. It consists of multiple parallel strands

D. It is primarily found in the nucleus

The correct answer highlights the structural feature of the 30-nm fiber, which is an important level of chromatin organization in eukaryotic cells. This fiber is characterized by its formation of looped domains, which are attached to proteins, particularly to a scaffold of non-histone proteins. The 30-nm fiber represents a more compacted structure of chromatin, which facilitates the efficient packing of DNA within the nucleus, while also playing a crucial role in gene regulation and accessibility. The presence of looped domains allows for further organization and compaction of DNA, which is essential for maintaining chromatin structure during processes such as DNA replication and transcription. This feature is critical for the regulation of gene expression, as it can lead to specific genes being brought into proximity with regulatory elements, thereby influencing their transcriptional activity. The other choices do not accurately represent the characteristic of the 30-nm fiber. The 30-nm fiber does not exist solely in a relaxed state, as it can undergo further compaction and changes in response to varying cellular conditions. Additionally, it does not consist of multiple parallel strands; rather, it is a single, continuous double helix that is folded and organized into a more compact structure. While the 30

Next Steps

Congratulations on reaching the final section of this guide. You've taken a meaningful step toward passing your certification exam and advancing your career.

As you continue preparing, remember that consistent practice, review, and self-reflection are key to success. Make time to revisit difficult topics, simulate exam conditions, and track your progress along the way.

If you need help, have suggestions, or want to share feedback, we'd love to hear from you. Reach out to our team at hello@examzify.com.

Or visit your dedicated course page for more study tools and resources:

<https://tamu-biol111exam3.examzify.com>

We wish you the very best on your exam journey. You've got this!

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