

Q1. In the space provided next to each definition or description, clearly write the letter of the appropriate term from the list of terms given on the last page. [20 × 0.5 =10]

- i) _____ A short, single-stranded DNA that serves as the necessary starting material for the synthesis of the new DNA strand in PCR
- ii) _____ The synthesis of DNA using DNA as a template
- iii) _____ The building blocks of DNA and RNA
- iv) _____ The synthesis of protein using information encoded in mRNA
- v) _____ The location in a eukaryotic cell where the electron transport chain occurs
- vi) _____ The major component of cell membranes
- vii) _____ The genetic composition of an organism
- viii) _____ An organism without membrane-bound organelles
- ix) _____ A cell with 1n chromosomes
- x) _____ The building blocks of proteins
- xi) _____ A cell with 2n chromosomes
- xii) _____ A major source of energy that has the general formula (CH₂O)_n
- xiii) _____ An enzyme needed for completion of lagging strand synthesis, but not leading strand synthesis
- xiv) _____ The synthesis of RNA using one strand of DNA as a template
- xv) _____ A circular DNA molecule that can be used to move foreign DNA in or out of a cell
- xvi) _____ The membrane that surrounds the cell
- xvii) _____ The DNA from a eukaryote formed by the enzyme reverse transcriptase; this DNA lacks introns.
- xviii) _____ An organism with 2 identical alleles for the same gene
- xix) _____ An organism with genetic material inside a nucleus
- xx) _____ A technique for the rapid production of millions of copies of a particular region of DNA

Q2. You have discovered a new enzyme, enzyme X, which breaks down proteins by cleaving peptide bonds after tyrosine or phenylalanine. Enzyme X is the product of gene X that encodes a protein with the molecular weight of 50 kilodaltons (50 kD). You purify active enzyme X and find it has a molecular weight of 250 kilodaltons (250 kD). Why is active enzyme X larger than the product encoded by gene X? What you can interpret about its structure? [2]

Q3. The following double-stranded DNA contains sequence of an eukaryotic gene:

 a b

5' -ATGGCCTTCACACAGGAAACAGCTATGGCCATGAGCACGCCAGTCTCGGCATTATCCTATTAAAGGGAAGTCTGAGGTGA-3'

3' -TACCGGAAGTGTGTCCTTTGTCGATACCGGTACTCGTGCGGTCAGAGCCGTAATAGGATAATTTCCCTTGACTCCACT-5'

a) Transcription begins at the underlined A/T at base pair 17 (**a**) and proceeds to the right. What are the first 12 nucleotides of the resulting mRNA? Indicate the 5' and 3' ends of the mRNA. [2]

b) The first 7 amino acids of the protein encoded by this gene are:

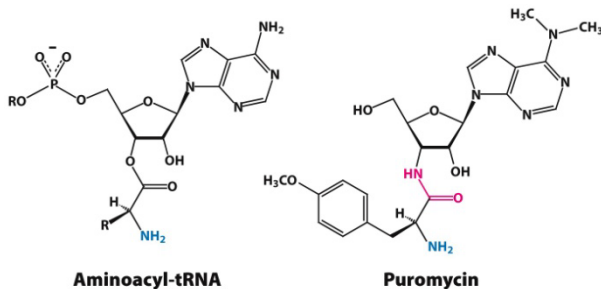
NH₃⁺ -met-ala-met-ser-pro-his-tyr....COO⁻

Draw a box around the intron region in this gene. [3]

c) What would be the preferred secondary structure of the initial amino acids in this sequence? [2]

d) How would the resulting protein change if the underlined G/C base pair at position 22 (**b**) was deleted from the DNA sequence? Briefly explain. [2]

Q4. Puromycin, which is structurally similar to the aminoacyl terminus of an aminoacyl-tRNA (see diagram), inhibits protein synthesis. Based on these information, can you give a hypothesis how puromycin can affect protein synthesis? [1]



Q5. The following is the sequence of two DNA helices. Which DNA helix (*helix 1* or *helix 2*) needs a higher temperature to denature (uncoiling)? **Explain** why you selected this option. [1]

Helix 1: 5' ATGCGGGAGA3' **Helix 2:** 5' ATGTTTTAGA3'
 3' TACGCCCTCT5' 3' TACAAAATCT5'

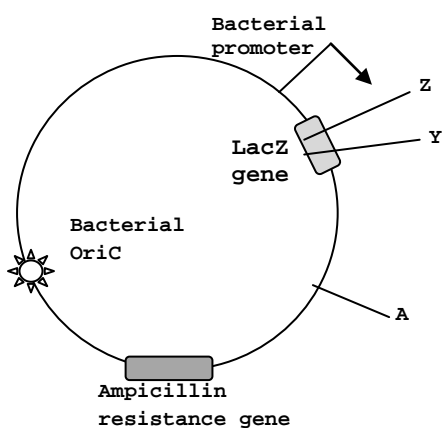
Q6. You have identified a mutant cell line that shows a **single mutation** in the **X** gene. You find that although the **X** gene has the same protein coding sequence in the wild-type and mutant cells, it is **NOT transcribed in mutant cells**. Based on this information, select the sequence from the choices below that might have a mutation and explain why you selected this sequence. [2]

- i) Promoter ii) Introns iii) Exons iv) 3' Untranslated region v) 5'Cap vi) Ribosome binding site

Q7. The **X** cDNA has the recognition site for restriction enzymes **R** and **A**. You want to clone **X** cDNA into the following plasmid that has recognition sites for restriction enzyme **Z**, **Y** and **A** as shown. *Please note: A slash (/) represents the cutting site for each restriction enzyme.*

Z	Y	R	A
5' GT/TATT AC3'	5' AG AATT/CT3'	5' AT AATT/GC3'	5' TG CCTT/CC3'
3' CA ATAA/TG5'	3' TC/TTAA GA5'	3' TA/TTAA CG5'	3' AC/GGAA GG5'

Which enzyme (*R/A*) would you use to cut the cDNA to insert into the plasmid? Which enzyme will you use to cut the plasmid for a successful and detectable transfection? Explain. [2.5+2.5]



You may detach this page during the exam.

List of terms for Question 1.

- a) allele
- b) amino acids
- c) autosomal gene
- d) carbohydrate
- e) cDNA
- f) competitive inhibitor
- g) diploid
- h) endoplasmic reticulum
- i) eukaryote
- j) G protein
- k) genotype
- l) haploid
- m) heterozygote
- n) homozygote
- o) mitochondria
- p) non-competitive inhibitor
- q) nucleotides
- r) DNA ligase
- s) phenotype
- t) phospholipids
- u) plasma membrane
- v) plasmid
- w) polymerase chain reaction
- x) primer
- y) prokaryote
- z) DNA polymerase
- aa) replication
- bb) repressor protein
- cc) sex-linked gene
- dd) transcription
- ee) translation

Genetic Code - Table

		Second Letter					
		U	C	A	G		
1st letter	U	UUU Phe UUC UUA Leu UUG	UCU UCC Ser UCA UCG	UAU Tyr UAC UAA Stop UAG Stop	UGU Cys UGC UGA Stop UGG Trp	U C A G	3rd letter
	C	CUU Leu CUC CUA CUG	CCU CCC Pro CCA CCG	CAU His CAC CAA Gln CAG	CGU CGC Arg CGA CGG	U C A G	
	A	AUU Ile AUC AUA AUG Met	ACU ACC Thr ACA ACG	AAU Asn AAC AAA Lys AAG	AGU Ser AGC AGA Arg AGG	U C A G	
	G	GUU Val GUC GUA GUG	GCU GCC Ala GCA GCG	GAU Asp GAC GAA Glu GAG	GGU GGC Gly GGA GGG	U C A G	

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6