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BRAINWARE UNIVERSITY

Term End Examination 2024-2025 Programme - M.Sc.(MB)-2024 Course Name - Biology for Macromolecules Course Code - MMB10105 (Semester I)

Full Marks: 60 Time: 2:30 Hours

[The figure in the margin indicates full marks. Candidates are required to give their answers in their own words as far as practicable.]

Group-A

1 x 15=15

	(Multi	iple Choice Type Question)
1.	Choose the correct alternative from	the following :
(i)	Which of the following is not an example of post-translational modification?	
(ii)	a) Myristoylationc) UbiquitinationChoose the factor, that contributes	b) Polyadenylation d) Alkylation to a decrease in gene density.
(iii)	a) Intergenic sequencec) Exon sequenceIdentify from the following that is n	b) Cell sized) Operonoot a crucial requirement during DNA Cloning.
(iv)	a) Protein expressionc) Molecular cutterThe process of RNA inactivation by	b) Vector d) DNA insert siRNAs is termed as
(v)	a) RNA silencingc) Short RNA inactivationIf a DNA sample is composed of 305	b) RNA interferenced) RNA disfunction% adenine, what is the percentage of guanine in this

(vi) A student is conducting an experiment on DNA replication and finds that replication is proceeding normally, but the DNA strands are highly coiled ahead of the replication fork. Which enzyme is most likely defective?

a) DNA polymerase

b) Helicase

b) 0.3 d) 0.5

c) Topoisomerase a 👙 🌁 d) primase

(vii) If DNA polymerase III in prokaryotes encounters a mismatched base during replication, what mechanism allows it to correct the error?

a) Priming

sample? a) 0.2

c) 0.4

b) Exonuclease activity to the last page and figure and the last page and the last p

c) Telomere extension

d) Okazaki fragment repair

(3)

(viii) A researcher finds that a cell line has an increased mutation rate and a tendency to Barasat, Kokata -700125 develop cancer. Defects in which DNA repair pathway could lead to this phenotype? b) BER a) Mismatch repair d) None of these c) NER (ix) A mutation causes a cell to lose its ability to repair DNA breaks during mitosis. Which repair mechanism is most likely affected? b) BER a) NHEJ d) Mismatch repair c) NER (x) Analyze how transcription factors can act as both activators and repressors depending on their binding sites. Which of the following is an accurate explanation? a) Transcription factors activate transcription b) The outcome depends on the cofactors they recruit at different binding sites. when bound to enhancers but repress when bound to silencers d) Transcription factors do not influence c) Transcription factors are always either transcription directly but only through activators or repressors, depending on their mediator complexes. (xi) How does the presence of small interfering RNAs (siRNAs) affect transcriptional regulation? b) siRNAs lead to degradation of target mRNA, a) siRNAs promote transcription by stabilizing reducing gene expression mRNA d) siRNAs modify histones to increase c) siRNAs enhance the translation of mRNA transcription into protein (xii) In eukaryotic cells, the binding of a transcriptional activator to a promoter region increases transcription. How would the deletion of the corresponding co-activator affect gene expression? b) Gene expression would decrease, as the coa) The transcriptional activator would activator is necessary for transcriptional compensate by directly recruiting RNA activation d) The gene would become constitutively c) There would be no effect, as co-activators expressed due to loss of co-activator are non-essential for transcription repression (xiii) Analyze the potential outcome if DNA methylation in a promoter region is blocked. What would be the likely effect on gene transcription? b) Transcription would increase due to the a) Transcription would be repressed. lack of repressive methylation marks. d) The gene would undergo alternative c) RNA polymerase binding would be splicing. impaired. (xiv) In post-transcriptional gene silencing (PTGS), how does the stability of the mRNA impact the level of gene expression? b) Decreased stability of mRNA through a) Increased stability of mRNA leads to gene degradation leads to gene silencing. silencing. d) The stability of mRNA promotes histone c) mRNA stability does not affect gene acetylation expression (xv) Which post-translational modification is crucial for bacterial chemotaxis? a) Glycosylation of flagella proteins b) Acetylation of pili proteins d) Ubiquitination of surface proteins c) Methylation of chemotaxis receptors Group-B (Short Answer Type Questions) 3 x 5=15

2. Evaluate the role of phosphorylation as a post-translational modification in molecular signaling pathways.

4. How does the cell detect DNA damage? 5. Describe general properties of Genetic code. 6. Analyze the function of Dicer in gene regulation. OR Analyze how changes in RNA transport pathways can impact the regulation of gene expression and its broader aspect on cellular functions. Group-C (Long Answer Type Questions) 5 x 6=3 7. Compare the process of transcription between prokaryotes and eukaryotes. Explain the process of RNA splicing and its significance in gene expression regulation. 9. Briefly describe the experimental proof of Semiconservative model of DNA replication. 10. Analyse the roles of different inhibitors in Bacterial DNA replication. 11. Analyze the errors occurred due to the malfunctioning of aminoacyl-tRNA synthetases during bacterial translation. 12. Compare between the termination of transcription between prokaryotes and eukaryotes. OR Outline the process of spliceosome mediated intron removal of mRNA. (5)	3)
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