JAYPEE UNIVERSITY OF INFORMATION TECHNOLOGY, WAKNAGHAT TEST-3 EXAMINATIONS MAY-2025

M.Tech-II Semester (BT)

COURSE CODE (CREDITS): 14M11BT213 (3)

MAX. MARKS: 35

COURSE NAME: FUNCTIONAL GENOMICS

COURSE INSTRUCTOR: DR JATA SHANKAR

MAX. TIME://2 Hours

Note: (a) All questions are compulsory.

(b) The candidate is allowed to make Suitable numeric assumptions wherever required for solving problems

Q.	Question	Marks
No.		
Q1	Explain the underlying principle and procedural steps involved in Serial	5
	Analysis of Gene Expression (SAGE). How does this technique facilitate the	
	comparison of gene expression profiles between two distinct cell types?	
Q2	Explain how genetic variations in patients influence the therapeutic efficacy	5
	and safety of the anticoagulant drug Warfarin, Discuss the key	
	pharmacokinetic and pharmacodynamic factors involved in its metabolism	
Transport 200	and action.	
Q3	What is RNA interference (RNAi) technology? Provide an overview of gene	5
	silencing mechanisms, including the role of microRNAs (miRNAs). Why is	
	RNAi considered a valuable tool in functional genomics research? Illustrate	
	its relevance with an example of miRNA function in a model organism.	/
Q4	Describe DNA microarray and its workflow. Explain the importance of	5
	normalisation in correcting systematic variations and biases. Differentiate	
	between pre-normalized and post-normalised data, highlighting how	
	normalisation helps in accurately identifying true biological differences in	
	gene expression.	
Q5	Discuss the concept of gene density and compare the genomic organisation	5
	of E. Coli with that to Saccharomyces cerevisiae (yeast) and the human	
	genome. How do differences in gene density reflect the complexity and	
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Q6,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Mustrate, with a suitable example, how the identification and	5
"IIIIIIII	characterisation of single-nucleotide polymorphisms (SNPs) contributed to	
411	the development of genetic markers for distinguishing between healthy and	
	diseased individuals, with specific reference to the TP53 oncogene.	
Q7	What is quantitative real-time PCR? Differentiate between qualitative and	5
	quantitative assessment of biomarkers? How does it help to identify the	
	pathogen or an unhealthy condition, describe keeping viral infection as an	
	example in cancer patients?	
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