JAYPEE UNIVERSITY OF INFORMATION TECHNOLOGY, WAKNAGHAT TEST -2 EXAMINATIONS APRIL-2025

M.Tech-II Semester (BT)

COURSE CODE (CREDITS): 14M11BT213 (3)

MAX. MARKS: 25

COURSE NAME: FUNCTIONAL GENOMICS

COURSE INSTRUCTOR: DR JATA SHANKAR

MAX. TIME: 1 Hour and 30 minutes

Note: (a) All questions are compulsory.

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

Q. No.	Question	Marks
Q1 Q2	Explain with an around by the state of the s	
	Explain with an example how the identification and discovery of SNPs	3
	led to the creation of a marker to differentiate between healthy and	
	diseased individuals considering TP53	
Q2 Q3	Describe how the genes of the patients affect the effectiveness of the	3
	medication 'Gefitinib'. Explain the pharmacokinetics and	
	pharmacodynamics factors/ terms.	
Q4.	Explain the principle of Restriction Fragment Length Polymorphism	3
	(RFLP). Describe the steps involved in conducting an RFLP analysis	
	and discuss its significance in distinguishing homozygous,	
	heterozygous and homozygous mutant disease alleles.	
ү	Describe the concepts of sequence alignment and mapping in	3
	genomics. Explain the sequence alignment techniques, including	
	global and local alignment, and discuss their applications.	
	Additionally, elaborate on the importance of sequence mapping in	
	genomics and how it facilitates genome assembly, variant detection,]
05	and functional annotation.	
Q5	What are the main genomics data retrieved from the human genome	4
	Project, which is a scientific landmark? What is a reneating element	
	in the genome, and is there a need for it? Estimate gene density using	
36	the number of genes and genome size?	
Q6	Describe the mechanism and methodology of Serial Analysis of Gene	4
	Expression. How SAGE helps to identify the expression pattern of two	
37	different cell types	
Q7 ·	Define	5
	a. Phred score	
	b. Transcriptome	
	c. Transcript	
	d. Transcription	
	e. Alternate splicing	