JAYPEE UNIVERSITY OF INFORMATION TECHNOLOGY, WAKNAGHAT TEST -2 EXAMINATION- 2023

M.Tech-1 Semester (BT)

		WI, I COII-1 A	SCIII	ester (D.	.,		
COURSE CODE (CREDITS): 13M11BT112 (3) MAX							RKS: 25
COURS	SE NA	ME: Advanced Bioinformatics					
COURSE INSTRUCTORS: Dr. Raj Kumar MAX. TIME: 1 Hour 30 N							Minutes
Note: (d	ı)All q	uestions are compulsory.					
(b)Marks are indicated against each question in square brackets.							
(c) The candidate is allowed to make Suitable numeric assumptions wherever required for							
solving problems							
Q1. Wri	ite dov	wn a R-programming code to crea	ite g	iven data	frame	es. Combine the given	———— data
frames using bind() function?							[3]
Data Frame -1					Tajira L	Data Frame -2	r- 3
Gene	ID	Common Name		Gene	_ID	Common No	ате
ATR	231	ATR serine-threonine kinase		BCL2	69	B-cell lymphoma 2	
BAK1	278	BCL2-antagonist 1		BDNF	191	Brain-derived neurot	rophic fact
BAX	119	BCL2-associated X protein		BLM	68	RecQ helicas	e-like
Q2. Describe the principle of chain termination using dideoxynucleotides.							[3]
Q3. Give a brief overview of a typical NGS data analysis pipeline.							[3]
Q4. Describe the following steps involved in performing quality control analysis on NGS data:							
							$[2 \times 4]$
a) Per base sequence quality							[2 ^ +]
b) Per sequence quality scores							
c) Per sequence GC content							
d) Base calling							
194	Massilla Silangan						
Q5. Give a brief account of SRA accession types.							[2]
Q6. What is the Phred quality score scale, and how does it relate to NGS quality scores?							[2]
Q7. Wh	at is tl	ne significance of the Q30 quality	scoi	re in NG	S data	, and why is it often us	sed as a
benchmark for data quality?							[2]

[2]

Q8. Discuss fundamental approaches to genome assembly.