

**BIRLA INSTITUTE OF TECHNOLOGY, MESRA, RANCHI
(END SEMESTER EXAMINATION)**

**CLASS: BTECH
BRANCH: BIOTECHNOLOGY**

**SEMESTER : V
SESSION : MO/2025**

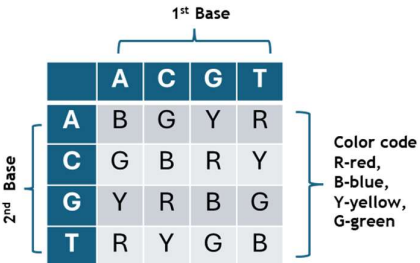
SUBJECT: BE302 FUNCTIONAL GENOMICS AND RDNA TECHNOLOGY

TIME: 3 Hours

FULL MARKS: 50

INSTRUCTIONS:

1. The question paper contains 5 questions each of 10 marks and total 50 marks.
 2. Attempt all questions.
 3. The missing data, if any, may be assumed suitably.
 4. Before attempting the question paper, be sure that you have got the correct question paper.
 5. Tables/Data hand book/Graph paper etc. to be supplied to the candidates in the examination hall.
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	[5]	CO	BL
Q.1(a) Differentiate the mechanisms of genome editing by TALEN and CRISPR/Cas systems. Discuss the advantages of CRISPR/Cas9 over TALEN.	[5]	1	3
Q.1(b) Design an experimental strategy to silence a specific gene in a model plant system, and validation methods you would use to confirm gene silencing.	[5]	1	5
Q.2(a) Describe the significance of Ct value and mention at least two applications where real-time PCR is advantageous over conventional PCR.	[5]	3	4
Q.2(b) Explain the principle and steps involved in replica plating technique.	[5]	2	2
Q.3(a) Explain the principle of SOLiD sequencing technology. Decode the DNA sequence using the color space decoding chart provided. The first dinucleotide is AT and the sequence obtained from fluorescence colors are as follows : <i>Red-green-blue-yellow-yellow-green-red-red-green</i>	[5]	3	5
			
Q.3(b) Describe the principle and workflow of the Sanger chain-termination method of DNA sequencing.	[5]	3	2
Q.4(a) List the characteristic features of an ideal plasmid cloning vector.	[5]	5	1
Q.4(b) Describe the structure and components of a Yeast Artificial Chromosome with a neat labeled diagram.	[5]	4	2
Q.5(a) Explain the principle and workflow of PCR-based diagnostic methods.	[5]	6	3
Q.5(b) Discuss the types and application of gene therapy.	[5]	6	3

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