

JAYPEE UNIVERSITY OF INFORMATION TECHNOLOGY, WAKNAGHAT

TEST -3 EXAMINATIONS- 2026

M.Sc.-II Semester (Biotechnology)

COURSE CODE (CREDITS): 20MS1WBT233 (02)

MAX MARKS: 35

COURSE NAME: Protein Engineering

COURSE INSTRUCTOR: Dr. Saurabh Bansal

MAX. TIME: 2 Hours

*Note: (a) All questions are compulsory.**(b) Use of calculator is allowed.*

Q. No.	Question	Marks
Q1	Explain the principle of directed evolution. Why is it often considered superior to rational design in generating novel protein functions?	2
Q2 a)	What do you understand by screening and selection? Explain each with suitable example.	3
Q2 b)	Explain the importance of HTS in directed evolution.	2
Q3 a)	Explain the principle of phage display. How does it establish a link between genotype and phenotype?	3
Q3 b)	Compare yeast surface display with bacterial display systems. What advantages does yeast offer in terms of protein folding and post-translational modifications?	4
Q3 c)	How is the ternary complex (mRNA-ribosome-polypeptide) stabilized in ribosome display method?	2
Q4 a)	Why are amino acid sequence comparisons generally preferred over nucleotide sequence comparisons?	2
Q4 b)	Name two algorithms used for pairwise sequence alignment.	1
Q4 c)	What is CASP and why is it important?	2
Q5	A recombinant protein expressed in <i>E. coli</i> forms insoluble inclusion bodies. Propose an experimental workflow to obtain soluble and active protein.	2
Q6	Why is structure-based protein engineering considered superior to sequence-only approaches? Discuss the limitations of experimental	4

	structure determination methods.	
Q7	A protein sequence shares 45% identity with a known protein structure. Which structure prediction method would you choose and why?	2
Q8	Discuss the principles of rational design, directed evolution, and semi-rational design. Highlight their advantages, limitations, and typical applications.	6

JUIT TEST-3 EXAMINATIONS- MAY-2026