

PT-003-1194003

Seat No.

M. Sc. (Microbiology) (Sem. IV) (CBCS) (W.E.F. 2016) Examination

August - 2020

Bimolecular Engineering: Paper - 421
(Elective - I)

Faculty Code: 003

Subject Code: 1194003

Time: $2\frac{1}{2}$ Hours] [Total Marks: 70]

Instruction: All questions are compulsory. Provide suitable illustrations where necessary.

1 Answer Any Seven: (2 Marks each)

14

- (a) Why should we change/modify the macromolecules?
- (b) How tertiary structure of a protein differs from that of the quaternary structure ?
- (c) Enlist basic approaches of the protein engineering.
- (d) What are soluble and insoluble fractions of the enzymes under cellular conditions?
- (e) How the inclusion bodies are significant in recombinant DNA Technology ?
- (f) What is the basic difference between Real Time PCR and rest of the PCR?
- (g) Enlist various methods of screening the recombinant clones.
- (h) What is the significance of co-expression of genes in recombinant DNA Technology ?
- (i) What is the significance of molecular tagging?
- (i) State how the fidelity of the PCR can be altered?

2	Write comments on Any Two:		′×2=14	
	(a)	Molecular forces necessary to stabilize various structure	es	
		of a protein.		
	(b)	Family shuffling and Error-Prone PCR in gene shufflin	g.	
	(c)	In-vitro protein folding.		
3	Write comments: (7 marks each)		14	
	(a)	Protein folding and molecular chaperones		
	(b)	Various methods of PCR and their significance		
		OR		
3	Comment on: (7 marks each)		14	
	(a)	Molecular cloning and gene library		
	(b)	Methods and significance of construction of the chimer	ic	
		genes		
4	Write comments: (7 marks each)		14	
	(a)	Directed evolution: The concept and Methods		
	(b)	Various in-vivo strategies to promote solubilization	of	
		the expressed protein		
5	Write comments on Any Two: (7 marks each)		14	
	(a)	Strategies for primer designing		
	(b)	DNA sequencing: New Approaches		
	(c)	Metagenomic Library		
	(d)	Overlapping PCR and its significance		