



PT-003-019403

Seat No. _____

**M. Sc. (Microbiology) (Sem. IV) (CBCS)
(W.E.F. 2010) Examination**

August - 2020

**Micro- 421 : Biomolecular Engineering
(Elective - I) (Old Course)**

Faculty Code : 003

Subject Code : 019403

Time : $2\frac{1}{2}$ Hours]

[Total Marks : 70

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

1 Answer Any Seven : (2 Marks each) 14

- (a) What are the objectives of protein engineering?
- (b) Differentiate tertiary and quaternary structures of the protein.
- (c) What are the basic approaches of the protein engineering?
- (d) Comment on the insoluble fraction of the enzymes in over expression?
- (e) Comment on the inclusion bodies in the recombinant DNA Technology.
- (f) How Real Time PCR differs from the rest of the PCR?
- (g) How recombinant clones are screened?
- (h) What is the significance of transformation in recombinant DNA Technology?
- (i) How molecular tagging is significant in cloning?
- (j) Comment on the fidelity of the PCR.

2 Write comments on : (Any Two) 7×2=14

- (a) Stabilization of various structures of a protein.
- (b) Family shuffling.
- (c) In-vivo strategies to enhance protein solubility.

- 3** Write comments : (7 marks each) **14**
- (a) Assisted Protein folding in the presence of molecular chaperones.
 - (b) Different PCR methods and their significance.
- OR**
- 3** Comment on : (7 marks each) **14**
- (a) Recombinant DNA Technology.
 - (b) Chimeric genes and protein engineering.
- 4** Write comments : (7 marks each) **14**
- (a) Directed evolution and gene shuffling.
 - (b) In-vitro strategies of the protein folding.
- 5** Write comments on : (Any Two) (7 marks each) **14**
- (a) Primer designing
 - (b) Next Generation DNA sequencing
 - (c) Metagenomic Library
 - (d) Error-Prone PCR.
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