

Course : B. Pharmacy

Sem: VI

Subject Name : Pharmaceutical Biotechnology

Subject Code : BP605T

Max Marks : 75

Duration : 3 Hr.

Instructions:

1. All questions are compulsory
2. Draw diagrams / figures wherever necessary
3. Figures to right indicate full marks

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Q. 1. Objective Type Questions (Answer all the questions)

(10 x 2) = 20

- i) Draw neat labelled diagram of structure of immunoglobulin.
- ii) What are mutants? Give their examples. <https://pharmacyindia.co.in/>
- iii) Differentiate between genetic organization of eukaryotes and prokaryotes.
- iv) Define biotechnology. Give its scope in pharmaceutical sciences.
- v) State the uses of microbes in industry.
- vi) What are the functions of DNA ligase and restriction endonucleases?
- vii) Write the importance of aeration and stirring in fermentation.
- viii) Define: Transformation and transduction.
- ix) Differentiate between cellular and humoral immunity.
- x) What is cold chain storage? State the storage conditions of vaccines.

Q. 2. Long Answers (Answer 2 out of 3)

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(2 x 10) = 20

- i) What are biosensors? Explain the working and applications of biosensors in Pharmaceutical sciences.
- ii) Illustrate the principle of rDNA technology with neat labelled diagram. Give the detailed account on human insulin production by rDNA technology.
- iii) Explain the production and purification of monoclonal antibodies by hybridoma technology. Give their applications in pharmaceutical industry.

Q. 3. Short Answers (Answer 7 out of 9)

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(7 x 5) = 35

- i) Explain in detail ELISA technique with neat labelled diagram.
- ii) Illustrate the design of large scale production fermenter and explain its various controls.
- iii) Explain the Collection and Storage of whole human blood. Extend the note on plasma substitutes.
- iv) Give comparative explanation of hypersensitivity reactions.
- v) Explain the structure and functions of Major Histocompatibility Complex (MHC).
- vi) Write a note on Polymerase Chain Reaction (PCR). <https://pharmacyindia.co.in/>
- vii) Explain the principle and methods of protein engineering.
- viii) Explain the methods of enzyme immobilization.
- ix) Write a note on cloning vectors in rDNA technology.

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