

| | | | | | | | | |
|--------|--|--|--|--|--|--|--|--|
| U.S.N. | | | | | | | | |
|--------|--|--|--|--|--|--|--|--|

B.M.S.College of Engineering, Bengaluru-560019

Autonomous Institute Affiliated to VTU

January / February 2025 Semester End Main Examinations

Programme: B.E.

Semester: V

Branch: Biotechnology

Duration: 3 hrs.

Course Code: 23BT5PCIMM /22BT5PCIMM /19BT5DCIMM

Max Marks: 100

Course: Immunotechnology

Instructions: 1. Answer any FIVE full questions, choosing one full question from each unit.
2. Missing data, if any, may be suitably assumed.

| UNIT - I | | | CO | PO | Marks | |
|---|---|----|---|-----------|--------------|----------|
| Important Note: Completing your answers, compulsorily draw diagonal cross lines on the remaining blank pages. Revealing of identification, appeal to evaluator will be treated as malpractice. | 1 | a) | Explain the distinctions between innate immunity and adaptive immunity, emphasizing their unique characteristics and roles in the immune response. | CO1 | PO 1 | 6 |
| | | b) | Compare and contrast T-cells and B-cells in terms of their structure, functions, and roles within the immune system. | CO1 | PO2 | 7 |
| | | c) | Highlight the key differences between innate and adaptive immunity, focusing on their mechanisms and response timelines. | CO1 | PO2 | 7 |
| OR | | | | | | |
| Important Note: Completing your answers, compulsorily draw diagonal cross lines on the remaining blank pages. Revealing of identification, appeal to evaluator will be treated as malpractice. | 2 | a) | Evaluate the effectiveness of passive and active immunity in controlling infectious diseases, and propose strategies to enhance their application in public health. | CO1 | PO2 | 6 |
| | | b) | Propose a model to study the development and activation of primary and secondary lymphoid organs, emphasizing their roles in immune cell maturation and response. | CO1 | PO2 | 7 |
| | | c) | Discuss the relationship between humoral and cellular immunity, and analyze a clinical scenario where both are required to resolve an infection effectively. | CO1 | PO1 | 7 |
| UNIT - II | | | | | | |
| Important Note: Completing your answers, compulsorily draw diagonal cross lines on the remaining blank pages. Revealing of identification, appeal to evaluator will be treated as malpractice. | 3 | a) | Outline the process of antibody gene and protein assembly in humans, focusing on the in vivo mechanisms involved. | CO2 | PO2 | 7 |
| | | b) | How do T-cells contribute to adaptive immunity by recognizing antigens, coordinating with other immune cells, and enhancing immune response effectiveness? | CO2 | PO2 | 6 |

| | | | | | |
|---|----|--|-----|-----|---|
| | c) | Discuss the organization of human antibody genes and how it contributes to the generation of diverse antibody repertoires. | CO2 | PO3 | 7 |
| | | OR | | | |
| 4 | a) | Antigens come in different shapes and types. Justify this statement mentioning various types of antigens. | CO2 | PO3 | 6 |
| | b) | Draw the typical structures of immunoglobulins and discuss their primary functions in the immune defense system. | CO2 | PO3 | 6 |
| | c) | Differentiate between Class I and Class II MHC proteins, detailing their structure and roles in antigen presentation. | CO2 | PO3 | 8 |
| | | UNIT - III | | | |
| 5 | a) | List various autoimmune disorders and describe how they involve the immune system targeting the body's own tissues. | CO3 | PO3 | 8 |
| | b) | How Major Histocompatibility Complex (MHC) molecules contribute to allograft rejection during transplantation. | CO3 | PO3 | 8 |
| | c) | Elaborate on the concept of tumor antigens, discussing their origin, expression patterns, and examples across different cancer types. | CO3 | PO3 | 4 |
| | | OR | | | |
| 6 | a) | What is the primary role of the complement system, and how is it activated to enhance immune responses? | CO3 | PO2 | 8 |
| | b) | Enumerate and describe the different types of hypersensitivity reactions, providing examples for each. | CO3 | PO2 | 8 |
| | c) | Explain the purpose of organ transplantation in medicine and discuss the circumstances that may necessitate such procedures. | CO3 | PO2 | 4 |
| | | UNIT - IV | | | |
| 7 | a) | Design an outline for developing a synthetic vaccine against a novel viral strain, specifying the steps and considerations involved. | CO4 | PO3 | 6 |
| | b) | Critique the role of immuno-toxins in targeted cancer therapy, providing examples of their mechanisms and potential side effects. | CO4 | PO2 | 6 |
| | c) | Discuss the engineering principles behind antibody mimics like adnectins and affibodies, and propose a scenario where they could be more advantageous than traditional antibodies. | CO4 | PO2 | 8 |
| | | OR | | | |
| 8 | a) | Evaluate the therapeutic potential of chimeric and humanized minibodies in treating autoimmune diseases compared to monoclonal antibodies. | CO4 | PO3 | 8 |

| | | | | | | |
|--|----|----|--|-----|-----|----------|
| | | b) | Assess the challenges in producing monoclonal antibodies using hybridoma technology and propose modern alternatives to address these challenges. | CO4 | PO4 | 8 |
| | | c) | Examine the ethical and logistical challenges associated with large-scale production of therapeutic antibodies for global pandemics. | CO4 | PO8 | 4 |
| | | | UNIT - V | | | |
| | 9 | a) | Differentiate between affinity and avidity in antigen-antibody interactions and discuss how these properties influence assay sensitivity. | CO5 | PO4 | 5 |
| | | b) | Design a workflow to detect a rare biomarker in a patient sample using a combination of immunoprecipitation and Western blotting. | CO5 | PO3 | 8 |
| | | c) | Assess the role of cytotoxicity assays in evaluating the efficacy of immuno-therapeutic agents and propose a scenario for their application. | CO5 | PO3 | 7 |
| | | | OR | | | |
| | 10 | a) | Discuss the principle of immuno-electrophoresis and design a protocol to identify a specific antibody in a serum sample. | CO5 | PO3 | 7 |
| | | b) | Compare precipitation reactions and agglutination reactions in terms of their mechanisms and applications in clinical diagnostics. | CO5 | PO2 | 7 |
| | | c) | Suggest an experimental approach to visualize antigen-antibody complexes in tissue sections using immunofluorescence, highlighting the importance of controls. | CO5 | PO3 | 6 |

B.M.S.C.E. -