

Code No: R05322305

R05**Set No. 2**

III B.Tech II Semester Examinations, December 2010

IMMUNOLOGY

Bio-Technology

Time: 3 hours

Max Marks: 80

Answer any FIVE Questions
All Questions carry equal marks

1. Explain the chemical nature of haptens? Discuss how they differ from adjuvants? [16]
2. Write short answers for the following:
 - (a) Epitope
 - (b) Affinity of antibodies
 - (c) Polyclonal antibody
 - (d) Fc region of antibody. [4×4]
3. Discuss how endogenous antigens processed & presented through MHC. [16]
4. Explain the various Ag-Ab reactions that are used in clinical diagnosis. [16]
5. What is lymphocyte trafficking? Explain the steps involved in the process of lymphocyte trafficking. [16]
6. (a) Discuss the importance of Haematopoietic microenvironment in B Cell development.
(b) Discuss the negative selection process of B cell development. [8+8]
7. Describe the different immunosuppressive protocols used during transplantation programmes. [16]
8. What are primary and secondary organs of the immune system? Explain them? [16]

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R05

Set No. 4

III B.Tech II Semester Examinations, December 2010

IMMUNOLOGY

Bio-Technology

Time: 3 hours

Max Marks: 80

Answer any FIVE Questions
All Questions carry equal marks

1. Describe the different immunosuppressive protocols used during transplantation programmes. [16]
2. Discuss how endogenous antigens processed & presented through MHC. [16]
3. Explain the chemical nature of haptens? Discuss how they differ from adjuvants? [16]
4. Explain the various Ag-Ab reactions that are used in clinical diagnosis. [16]
5. Write short answers for the following:
 - (a) Epitope
 - (b) Affinity of antibodies
 - (c) Polyclonal antibody
 - (d) Fc region of antibody. [4×4]
6. (a) Discuss the importance of Haematopoietic microenvironment in B Cell development. [8+8]
(b) Discuss the negative selection process of B cell development.
7. What is lymphocyte trafficking? Explain the steps involved in the process of lymphocyte trafficking. [16]
8. What are primary and secondary organs of the immune system? Explain them? [16]

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R05

Set No. 1

III B.Tech II Semester Examinations, December 2010

IMMUNOLOGY

Bio-Technology

Time: 3 hours

Max Marks: 80

Answer any FIVE Questions
All Questions carry equal marks

1. (a) Discuss the importance of Haematopoietic microenvironment in B Cell development.
(b) Discuss the negative selection process of B cell development. [8+8]
2. Explain the various Ag-Ab reactions that are used in clinical diagnosis. [16]
3. Explain the chemical nature of haptens? Discuss how they differ from adjuvants? [16]
4. Discuss how endogenous antigens processed & presented through MHC. [16]
5. Write short answers for the following:
(a) Epitope
(b) Affinity of antibodies
(c) Polyclonal antibody
(d) Fc region of antibody. [4×4]
6. What are primary and secondary organs of the immune system? Explain them? [16]
7. Describe the different immunosuppressive protocols used during transplantation programmes. [16]
8. What is lymphocyte trafficking? Explain the steps involved in the process of lymphocyte trafficking. [16]

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R05

Set No. 3

III B.Tech II Semester Examinations, December 2010

IMMUNOLOGY

Bio-Technology

Time: 3 hours

Max Marks: 80

Answer any FIVE Questions
All Questions carry equal marks

1. Write short answers for the following:

- (a) Epitope
- (b) Affinity of antibodies
- (c) Polyclonal antibody
- (d) Fc region of antibody.

[4×4]

2. Explain the chemical nature of haptens? Discuss how they differ from adjuvants? [16]

3. (a) Discuss the importance of Haematopoietic microenvironment in B Cell development.

(b) Discuss the negative selection process of B cell development. [8+8]

4. Discuss how endogenous antigens processed & presented through MHC. [16]

5. Explain the various Ag-Ab reactions that are used in clinical diagnosis. [16]

6. What is lymphocyte trafficking? Explain the steps involved in the process of lymphocyte trafficking. [16]

7. What are primary and secondary organs of the immune system? Explain them? [16]

8. Describe the different immunosuppressive protocols used during transplantation programmes. [16]
