

JAYPEE UNIVERSITY OF INFORMATION TECHNOLOGY, WAKNAGHAT

END SEMESTER EXAMINATION-2015

M.Tech Biotechnology II Semester

COURSE CODE: 14M11BT212

MAX. MARKS: 45

COURSE NAME: Immunotechnology

COURSE CREDITS: 4

MAX. TIME: 3 HRS

Note: All questions are compulsory. Carrying of mobile phone during examinations will be treated as case of unfair means.

Section A

Q1. Explain each of the following in short: [1X 5 = 5]

- a. Herd Immunity
- b. Need for humanized antibodies
- c. Isotype Switching
- d. Negative selection during T cell development
- e. Basis of BrdU proliferation assay

Q2. Differentiate each of the following: [2 X5 = 10]

- a. De novo pathway and Salvage pathway for DNA synthesis
- b. Apoptosis versus Necrosis
- c. Allograft versus Autograft
- d. MHCI and MHCII
- e. B cell Mitogens and T Cell mitogens

Section B

Q1. Answer each of the following. [3 X5 = 15]

- a. Describe the logistic behind "ADAPT".

- b. What are T reg cells? Discuss their relevance in health and disease?
- c. "Diabetes and Immunobiology" Is there a link? If yes, how?
- d. Antibodies are potential drugs in context of different diseases? Do you agree? If yes, give one example to support your answer.
- e. Information derived from Flowcytometry and Immunohistochemistry based studies complement each other to better understand the model system under study. Do you agree? Why or why not?

Section C

Q1. Answer each of the following. [5 X3 = 15]

- a. Vaccine development against different pathogenic group require different sets of optimized parameters. Do you agree? If yes, what are the parameters to be considered?
- b. Why immunological surveillance against developing tumors is an infrequent phenomenon?
- c. What are the different Immune-evasion strategies used by extracellular bacteria?