

2072

M.Sc. (Biotechnology) Fourth Semester
MBIO-402: Drug Designing and Drug Delivery

Time allowed: 3 Hours

Max. Marks: 80

NOTE: Attempt five questions in all, including Question No. I which is compulsory and selecting one question from each Unit.

x-x-x

I. Attempt the following:-

- a) Define pharmacophore.
- b) What is virtual screening?
- c) What is therapeutic index?
- d) What is pharmacodynamics?
- e) What is subacute toxicity?
- f) Define selection bias?
- g) What is NDA?
- h) What is trial drug packaging? (8x2)

UNIT - I

- II. a) Discuss the factors affecting drug absorption.
- b) Discuss the factors contributing to 3D QSAR. (2x8)
- III. a) Discuss the process and role of high throughput screening.
- b) Discuss the process of lead identification. (2x8)

UNIT - II

- IV. a) Discuss clearance, volume of distribution and AUC as pharmacokinetic parameters.
- b) Discuss the factors affecting coordination complex stability and its measurement. (2x8)
- V. a) Discuss the significance and methods to study *in vitro* toxicity,
- b) Discuss c versus t plot for intravenously administered drug. (2x8)

UNIT - III

- VI. a) Give an overview of drug approval process by FDA.
 - b) Discuss the regulations governing the conduct of clinical trials. (10,6)
- P.T.O.

(2)

- VII. a) Explain the procedure of preclinical testing in drug approval.
b) Describe the considerations for blinding drug product for clinical trial packaging. (2x8)

UNIT - IV

- VIII. a) Discuss the mechanism of action for prodrugs.
b) Explain the applications of liposomes in drug delivery. (2x8)
- IX. a) Discuss the osmotic controlled drug delivery systems.
b) Describe the approaches towards effective protein and peptide drug delivery. (2x8)

x-x-x