

1058

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 Efflux transporters.
- b) What is pharmacokinetics?
- c) What is high throughput screening?
- d) What are prodrugs?
- e) What is acute toxicity?
- f) Define clinical trials?
- g) What is ANDA?
- h) What is trial drug packaging? (8x2)

UNIT - I

- II.
 - a) Discuss the factors affecting drug excretion.
 - b) Discuss the factors contributing to 2D QSAR. (2x8)
- III.
 - a) Discuss the significance and methods of high throughput screening.
 - b) Discuss the procedure of ligand based drug design. (2x8)

UNIT - II

- IV.
 - a) Discuss the derivation of AUC and elimination half life from C vs T plot.
 - b) Discuss the different theories of coordinate bonding. (2x8)
- V.
 - a) Discuss the significance and methods to study chronic toxicity.
 - b) Write short notes on:-
 - i) Pharmacodynamics
 - ii) Clearance of drug (2x8)

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(2)

UNIT – III

- VI. a) Describe the procedure of drug approval by FDA.
b) Discuss the regulations governing the conduct of clinical trials. (2x8)
- VII. a) Explain the parameters and role of preclinical testing in drug approval.
b) Describe the considerations for blinding of drug products. (2x8)

UNIT – IV

- VIII. a) Discuss the mechanism of action for soft drugs.
b) Explain the applications of microparticles in drug delivery. (2x8)
- IX. a) Discuss the diffusion controlled drug delivery systems.
b) Describe the ligand appended approach to drug delivery. (2x8)

x-x-x