

Fourth Year Pharm. D : SUMMER - 2019
SUBJECT: BIOPHARMACEUTICS AND PHARMACOKINETICS

Day: Friday
Date: 19/04/2019

S-2019-4521

Time: 02.00 P.M. TO 05.00 PM
Max. Marks: 70

N.B:

- 1) **Q. No. 1 and Q. No. 5 are COMPULSORY.** Out of the remaining attempt **ANY TWO** questions from each section.
- 2) Figures to the right indicate **FULL** marks.
- 3) Answer to the both sections should be written in **SEPARATE** answer books.

SECTION - I

- Q.1** A) Attempt **ANY FOUR** of the following: (08)
- i) Explain the influence of gastric emptying on drug absorption.
 - ii) Enlist the different steps and reactions of biotransformation.
 - iii) Define total clearance and renal clearance. Give their equations.
 - iv) Explain drug related factors affecting protein binding.
 - v) Name the specialized barriers to distribution of drug.
 - vi) Differentiate between passive and active transport mechanism.
- B) Explain the pre systemic metabolism (03)
- Q.2** Explain in details the theories of dissolution. (12)
- Q.3** a) Explain Phase I and Phase II metabolic reactions with suitable examples. (07)
- b) Give an account of physicochemical factors influencing drug distribution. (05)
- Q.4** Write short notes on **ANY THREE** of the following: (12)
- a) Concept of renal clearance
 - b) pH-partition hypothesis
 - c) Significance of protein binding
 - d) Volume of distribution

SECTION - II

- Q.5** A) Attempt **ANY FOUR** of the following: (08)
- i) What is AUC and trapezoidal rule?
 - ii) Enlist the pharmacokinetic parameters.
 - iii) What is non-linear pharmacokinetics?
 - iv) What is zero order kinetics?
 - v) What are the objectives of bioavailability studies?
 - vi) Define drug effect and drug potency.
- B) What are the various approaches to quantitative study of kinetic process of drug disposition? (03)
- Q.6** Derive the equation to obtain the pharmacokinetic parameters for i.v. loading dose followed by i.v. infusion according to one compartment open model. (12)
- Q.7** a) Differentiate between compartment modeling and physiological modeling. (07)
- b) Elaborate on the study designs employed to establish bioequivalence. (05)
- Q.8** Write short notes on **ANY THREE** of the following: (12)
- a) Sigma minus method
 - b) Method of residuals
 - c) Methods of enhancement of bioavailability
 - d) Pharmacokinetic modeling
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