



DAO-BP604T

Seat No. _____

M. P. M. (Sem. VI) Examination

May – 2022

Biopharmaceutics & Pharmacokinetics

Time : 3 Hours]

[Total Marks : 75

Instructions :

- (1) Figure to the right indicates marks.
- (2) Draw neat and clean diagrams as required.

1 Answer the following questions : **10×2=20**

- (a) Enlist the factors affecting drug absorption from GI tract.
- (b) Define absolute bioavailability and relative bioavailability.
- (c) How do you determine renal clearance of drugs ?
- (d) Define phase I and phase II reactions of biotransformation with example.
- (e) What is a compartment model ? Discuss the various types of compartment models.
- (f) Define Distribution and enlist factors affecting distribution of drug.
- (g) Define elimination rate constant.
- (h) Define non linear pharmacokinetics.
- (i) Write the merit and demerit of the Wagner-nelsons method.
- (j) Define loading dose and maintenance dose.

2 Answer any **Two** out of the following : **2×10=20**

- (a) Discuss in detail one-compartment open model for a drug administered as IV Bolus. Give the schematic representation, graphs and equations for the same.
- (b) Discuss in detail various physico-chemical, factors affecting drug absorption.
- (c) Define biotransformation. Explain with examples phase I and phase II reactions.

3 Answer any **seven** out of the following :

7×5=35

- (a) Differentiate between one and two compartments.
- (b) Explain Michaelis Menten equation in determining non-linearity.
- (c) Explain how steady state level of the drug is achieved through I. V. infusion.
- (d) Write a short note on in-vitro-in-vivo correlations.
- (e) Explain Non renal routes of drug excretion.
- (f) Write a note an measurement of bioavailability.
- (g) Explain factors affecting protein-drug binding.
- (h) Write the importance of Compartment modeling in pharmacokinetic study.
- (i) Enlist various Mechanisms of drug absorption through GIT and explain any one in detail.
