

**FINAL YEAR B.PHARM. SEMESTER-VII (2011 COURSE) :
WINTER - 2017**

SUBJECT: BIOPHARMACEUTICS AND PHARMACOKINETICS

Day : **Thursday**
Date : **16/11/2017**

W-2017-3846

Time : **02.00 PM TO 05.00 PM**
Max. Marks : **80.**

N.B.:

- 1) Q. No. 1 and Q. No. 5 are **COMPULSORY**. Out of the remaining attempt any **TWO** questions from Section-I and any **TWO** questions from Section-II.
- 2) Both the sections should be written in **SEPARATE** answer books.
- 3) Figures to the **RIGHT** indicate full marks.

SECTION-I

- Q.1** Answer the following in short (Any **Five**) **(10)**
- a) What is meant by rate determining step in the process of drug absorption?
 - b) Define volume of distribution and give its equation.
 - c) Enlist the Phase II reactions of biotransformation and give its significance.
 - d) Explain the mechanism of transport of Vit B₁₂.
 - e) Give dose adjustment in case of renal failure patients.
 - f) What is enterohepatic circulation?
- Q.2** a) What is gastric emptying? Explain the factors influencing gastric emptying. **(07)**
b) Explain the different physiological barriers to the drug distribution. **(08)**
- Q.3** a) Explain the influence of particle size on drug absorption from GIT. **(07)**
b) Give an account of drug interactions and highlight its significance. **(08)**
- Q.4** Write notes on (Any **Two**) **(15)**
- a) Passive diffusion process for drug transport
 - b) pH-partition hypothesis
 - c) Significance of protein drug binding.

SECTION-II

- Q.5** Answer the following in short (Any **Five**) **(10)**
- a) Explain trapezoidal rule.
 - b) Define clinical pharmacokinetic and pharmaceutical equivalents.
 - c) What are the advantages of urinary data over plasma data.
 - d) Define MRT. Give equation.
 - e) Give the advantages of compartmental modelling.
 - f) Give the objectives of bioavailability studies.
- Q.6** Discuss the method of residuals for determination of absorption rate constant **(15)** following oral administration of drug assuming one compartment open model.
- Q.7** a) Give an account of physiological modeling. **(07)**
b) Discuss the approaches for enhancement of bioavailability. **(08)**
- Q.8** Write notes on (Any **Two**) **(15)**
- a) Pharmacokinetic methods for determination of bioavailability
 - b) Assessment of pharmacokinetic parameters following IV bolus administration for one compartment open model
 - c) Statistical designs for BA/ BE studies.

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