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M. PHARM. (SEM-II)
CARRY OVER EXAMINATION 2016-17
BIO-PHARMACEUTICS & PHARMACOKINETICS

*Time : 3 Hours**Max. Marks : 100**Note : Be precise in your answer. In case of numerical problem assume data wherever not provided.*

1. **Attempt any Two parts of the following:** **2 × 10 = 20**
 - (a) Discuss the various physicochemical factors affecting drug absorption also explain flip flop model.
 - (b) How does protein binding affect distribution of drugs?
 - (c) What are the different modes of drug transport across the cellular membrane? Discuss their mechanism with examples.

2. **Attempt any Two parts of the following:** **2 × 10 = 20**
 - (a) Discuss one compartment open model (iv bolus) of administered drug.
 - (b) Give a note on non compartmental pharmacokinetics.
 - (c) Write in detail about sigma minus method and wagner nelson method.

3. **Attempt any Two parts of the following:** **2 × 10 = 20**
 - (a) What is bioavailability of drugs? Discuss various method of determining bioavailability of drugs.
 - (b) Discuss bioequivalence study design with special reference to BCS.
 - (c) Explain how plasma drug concentration data can be useful in estimating bioavailability and discuss the pharmacokinetics parameters determined from the plasma data.

4. **Attempt any Two parts of the following:** **2 × 10 = 20**
 - (a) Explain the general methods used for dosage adjustment in renal disease.
 - (b) What do you understand by individualization and optimization of dosage regimens?
 - (c) Define dose ratio? Why it is always smaller for extravascularly administered drug?

5. **Attempt any Two parts of the following:** **2 × 10 = 20**
 - (a) What are clinical trials? Discuss in detail various principles of clinical trial.
 - (b) Briefly describe about phase I and phase II clinical trials.
 - (c) Write a note on research methodology used in clinical trial study.