

## **IMPORTANT QUESTIONS**

For

**B. Pharmacy Third Year II-Semester** 

Subject:

**Biopharmaceutics & Pharmacokinetics (BPPK)** 

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### Section-I: Very Short Answer Questions.

- 1. Write Noyes Whitney equation. Explain the terms?
- 2. Describe the absorption of a drug on rectal administration.
- 3. Define apparent volume of distribution.
- 4. Define Biopharmaceutics & Pharmacokinetics.
- 5. Mention factors affecting absorption.
- 6. Differentiate passive transport and active transport.
- 7. What is protein binding? How it affects bioavailability?
- 8. What is Lipinskis rule of five?
- 9. Differentiate between plasma drug binding and tissue drug binding.

### Section-II: Short Answer Questions

- 1. Discuss about pH-partition hypothesis.
- 2. Describe the absorption of drugs from extravascular routes.
- 3. Explain briefly about kinetics of protein binding.
- 4. How do you determine absorption rate constant, Ka by Wagner nelson method.
- 5. Write a note on carrier mediated transport?
- 6. Describe about the physiological barriers to the distribution of drugs.
- 7. Write about gastric emptying rate and volume of distribution.
- 8. Discuss the mechanism of active diffusion in absorption of drugs?
- 9. How the organ size and perfusion rate influence the drug distribution.

### Section-III: Long Answer Questions

- 1. Define absorption. Write in detail about mechanism of drug absorption with diagram.
- 2. Discuss protein binding and various factors affecting drug protein binding.

### UNIT-II

### Section-I: Very Short Answer Questions.

- 1. Write a note on excretion of drugs through skin.
- 2. Define absolute bioavailability and relative bioavailability.
- 3. Describe hepatic clearance.
- 4. List the factors affecting elimination of drugs.
- 5. Define Orange book and objectives of bioavailability studies.
- 6. What are the markers used in renal clearance.
- 7. Define Creatinine and how to calculate the creatinine clearance.

### Section-II: Short Answer Questions

- 1. Explain biliary excretion of drugs.
- 2. Discuss about methods to enhance bioavailability of poorly soluble drugs.
- 3. Explain various methods for assessment of bioavailability.
- 4. Discuss in-vitro in vivo correlation.
- 5. Explain various cross over designs in bio equivalence studies.
- 6. Explain factors affecting the renal excretion of drugs.

## Section-III: Long Answer Questions

- 1. Describe the renal excretion of drugs.
- 2. Write about in-vitro drug dissolution models.
- 3. Explain in detail about Bioequivalence study protocols.

## UNIT-III

## Section-I: Very Short Answer Questions.

- 1. Define  $C_{max}$   $T_{max}$  and AUC?
- 2. Explain the term V  $_{d}$ , t  $\frac{1}{2}$ , K<sub>a</sub> and CLR.
- 3. Explain flip-flop method in extra vascular administration.
- 4. Write the equation for calculating steady state drug concentration for one compartment open model.
- 5. What is flip flop phenomenon and how is it useful in method of residual.
- 6. Define Microconstants and hybrid constants and write the relationship between them.

# Section-II: Short Answer Questions

- 1. Explain in detail about compartment models.
- 2. Explain the pharmacokinetic parameters of a drug which follows one compartment open model when given by IV bolus with relevant mathematical equations

# Section-III: Long Answer Questions

1. Problems based on compartment models pharmacokinetic parameters.

### Section-I: Very Short Answer Questions.

- 1. Write the equation for calculating loading dose
- 2. Write briefly about two compartment open model.

#### Section-II: Short Answer Questions

- 1. Derive Kinetics parameter for IV bolus administration in two compartment open model.
- 2. Explain methods of adjustment of dose and dosage regimen in patients with hepatic failure.
- 3. Write the significance of different volumes of distribution in two compartment model.

### Section-III: Long Answer Questions

1. Derive mathematical equations used to calculate pharmacokinetic parameters following IV bolus administration blood data, assuming that the drug follows two compartment open model.

### **UNIT-V**

### Section-I: Very Short Answer Questions.

- 1. What are the factors for cause of non-linear kinetic?
- 2. Write Michaelis Menten equation.

#### Section-II: Short Answer Questions

- 1. Describe estimation of  $K_m$  and  $V_{max}$  in non-linear kinetics.
- 2. Write a note on non-linear Pharmacokinetics.
- 3. Write a note on Michaelis Menten equation.

### Section-III: Long Answer Questions

1. Drive Michaelis Menten equation and how do you estimate  $K_m$  and  $V_{max}$