2012 Scheme

12. Derive the equation to determine the first order rate constant of absorption (Ka) for one compartment open model extravascular administration.

- 13. Define apparent volume of distribution and explain how the organ/tissue size and perfusion rate affects drug distribution
- 14. Explain glutathione conjugation.
- 15. Explain the official dissolution test methodology and acceptance criteria for immediate release of solid dosage forms
- 16. What are the physiological barriers to distribution of drugs.
- 17. Sigma minus method for determination of elimination rate constant for IV bolus administration.

Third Year B.Pharm Degree Supplementary Examinations November 2019 Pharmaceutics - IV

(Biopharmaceutics and Pharmacokinetics)

Time: 3 Hours

Essays

- Answer all Questions. •
- Write equations wherever necessary. •

(3x10=30)

(14x5=70)

- 1. Enumerate the various physicochemical properties of drugs affecting GI absorption. Explain each factor in detail
- 2. Describe the kinetics of one compartment open model IV bolus administration and explain how the various kinetic parameters are determined.
- 3. Explain the reasons for non-linearity in pharmacokinetics. Describe the Michaelis-Menten equation and describe how Vmax and Km are determined.

Short notes

- 4. Draw the plasma drug concentration time profile graph for one compartment open model extra vascular administration and define the various pharmacokinetic and pharmacodynamic parameters
- 5. Explain conjugation of drugs with glucuronic acid
- 6. The transport mechanism across the membrane using energy
- 7. The binding of drug to blood components
- 8. What are the various factors that affect the protein binding of drugs
- 9. Explain the entero-hepatic cycling of drugs
- 10. Oxidation of carbon-nitrogen system and explain with examples
- 11. In vitro In vivo correlation.

Reg. No:

Total Marks: 100

QP CODE: 314006