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**MPY - 201 (PCS)**  
**M.Pharmacy II Semester**  
 Examination, December 2015  
**Biopharmaceutics and Pharmacokinetics**  
**(Advanced Pharmaceutics - I)**

Time : Three Hours

Maximum Marks: 70

**Note:** Attempt any five questions. All questions carry equal marks.

1. a) Discuss the compartment model for a drug showing rapid distribution after IV bolus administration. How this model is useful in determination of various pharmacokinetic parameters? 7
- b) A subject received an i.v. dose of 100 mg of a drug and plasma concentration of drug were as follows: 7

Time (hr)	1	2	3	4	5	6	7
Plasma Conc. ( $\mu\text{g/ml}$ )	60.65	36.79	22.31	13.53	8.21	4.98	3.02

Assume that the drug is eliminated by an apparent first order process and  $C_0 = 100 \mu\text{g/ml}$  Calculate, Overall elimination rate constant ( $K_e$ ), Volume of distribution  $V_d$  and Elimination half-life  $t_{1/2}$ .

2. a) Describe Wagner-Nelson method for determination of absorption rate constant. 7
- b) Discuss Sigma-Minus method for determination of elimination rate constant. 7

3. a) What is linear pharmacokinetics? How it is recognised? Give the reasons for nonlinear pharmacokinetics. 7
- b) What is Michaelis-Menten kinetics? Discuss the methods for determination of  $K_m$  and  $V_m$ . 7
4. a) Discuss the concept of Steady-state plasma concentration. How it is dependent on elimination half-life of a drug? 7
- b) Discuss First order absorption kinetics in multiple dosing. 7
5. a) Discuss Physiological pharmacokinetic model. Give its application and limitations. 7
- b) Discuss concept of Statistical moment theory, MRT and MDT. 7
6. a) Define bioavailability and bioequivalence. Discuss various study designs involved in determination of bioequivalence. 7
- b) Give detail account on in-vitro dissolution and in-vivo bioavailability correlation. 7
7. Write short notes on any two of the following 14
- a) Factors affecting plasma concentration and toxicity
- b) Loading dose and maintenance dose
- c) Therapeutic index and Therapeutic window.
- d) Interrelationship between pharmacokinetic parameters and physiological variables

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