

GUJARAT TECHNOLOGICAL UNIVERSITY
B.PHARM - SEMESTER-6 EXAMINATION – WINTER -2023

Subject Code: BP604TT**Date: 12/12/2023****Subject Name: Biopharmaceutics and Pharmacokinetics****Time: 02.30 p.m. to 5.30 p.m.****Total Marks: 80****Instructions:**

1. Attempt any five questions.
2. Make suitable assumptions wherever necessary.
3. Figures to the right indicate full marks.

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| Q.1 | (a) Discuss the objectives and factors affecting drug absorption through oral route. 06 |
| | (b) What is apparent volume of distribution. Discuss the influence of protein binding on apparent volume of distribution. 05 |
| | (c) Discuss Michaelis-Menten Equation. 05 |
| Q.2 | (a) Define following terms: Absolute bioavailability, Relative Bioavailability, Bioequivalence. 06 |
| | (b) Write a short note on kinetics of protein-drug binding. 05 |
| | (c) Briefly describe factors affecting distribution of drug. 05 |
| Q.3 | (a) Introduce the concept of IVIVC. Discuss BCS based biowaivers in IVIVC briefly. 06 |
| | (b) Write a short note on non-renal excretion of drugs. 05 |
| | (c) Define high soluble and high permeable as per BCS classification. Enumerate various methods for improvement of dissolution rates of poorly soluble drug. 05 |
| Q.4 | (a) Explain Latin crossover design in bioequivalence studies. 06 |
| | (b) Explain briefly what is compartmental model. Explain Catenary and Mammillary compartment models in detail 05 |
| | (c) Describe the method of residuals for determination of absorption rate constant. 05 |
| Q.5 | (a) Write a note on PBPK models. 06 |
| | (b) Enumerate USP dissolution apparatus along with examples of relevant dosage forms evaluated in each dissolution apparatus. 05 |
| | (c) Explain Wagner nelson method in detail. 05 |
| Q. 6 | (a) Discuss briefly the concept of loading dose and maintenance dose. 06 |
| | (b) Write a brief note on two compartment open model. 05 |
| | (c) Write a note on factors affecting protein binding with suitable examples. 05 |
| Q.7 | (a) Explain the factors causing non linearity in pharmacokinetics in drugs. 06 |
| | (b) Explain merits and demerits of non-compartmental pharmacokinetics. 05 |
| | (c) Differentiate between plasma-protein drug binding and tissue-drug binding. 05 |
