Time: Three Hours Max. Marks: 100 Marks

Bio-Pharmaceutics and Pharmacokinetics (Revised Scheme 4) Q.P. CODE: 9338

Your answers should be specific to the questions asked. Draw neat, labeled diagrams wherever necessary. Answer any ten questions.

LONG ESSAY (Answer any TEN)

10 X 10 = 100 Marks

- 1. Explain in detail various non per-oral routes of drug absorption.
- 2. Write the objectives of bioavailability studies. Explain the methods for measurement of bioavailability.
- 3. Define drug distribution. Explain various physiological barriers for drug distribution.
- 4. Define the equivalence terms. Write the equation for similarity and dissimilarity factor and explain.
- 5. Explain Michael's Menten equation. How do you estimate Km and Vmax?
- 6. Explain the pharmacokinetics of a drug given by extravascular route which follows one compartment open model.
- 7. What is protein binding? Explain the kinetics and significance of protein binding.
- 8. Classify pharmacokinetic models and explain briefly about each.
- 9. Explain pH partition hypothesis and its limitations.
- 10. What are the pharmacokinetic considerations in designing a dosage regimen?
- 11. Write a note on Phase-II biotransformation reactions.

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