[Time: 3 Hours] [Max. Marks: 100]

<u>Medicinal Chemistry – I (Drug Design)</u> (Revised Scheme 4)

Q.P. CODE: 9347

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. What are the different biological targets for drug discovery? Explain with examples.
- 2. a) What is a pharmacophore? Explain a method for identification of a pharmacophore.
 - b) Write a note on Ligand-Gated Ion channels and Orphan receptors.
- 3. Discuss the various mathematical methods used for the analysis in QSAR.
- 4. Explain the application of Hansch analysis with a suitable example.
- 5. Explain the concepts of (a) Virtual screening (b) Drug likeness.
- 6. Discuss (a) In Silico De Novo design (b) Topological drug classification.
- 7. Discuss the drugs used as immuno suppressants and immuno stimulants with examples.
- 8. Describe the development of gastric proton pump enzyme inhibitors and cycloxygenase inhibitors.
- 9. Explain the significance of prodrugs in drug design with specific examples.
- 10. What is epitope mapping? Give the importance of epitope mapping in drug design.
- 11. Relate quantitative structure activity relationship with biological activity of a drug molecule.
- 12. Discuss (a) Analog approach in drug discovery (b) Applications of QSAR.
