

**35182 : Medicinal Chemistry - IV : T-8.2**

P. Pages : 1

Time : Three Hours

**AW - 2326**

Max. Marks : 60

- Notes :
1. All question carry equal marks.
  2. Answer **five** question.
  3. Discuss the reaction, mechanism wherever necessary.
  4. Use of pen Blue/Black ink/refill only for writing the answer book.

1. Explain in detail rational drug design. Give its advantages over conventional drug design. **12**
2. Lipophilicity, Shape and electron distribution have a major influence on drug activity. Explain the parameters that are commonly used as measure of these properties in QSAR approach to drug design. **12**
3. What is QSAR? Explain Hansch analysis and Topliss decision tree methods in QSAR approach to drug design. **12**
4. Write a detail account on structure based drug design with examples. **12**
5. Explain in detail concept of genetic engineering. Give its merits and demerits. **12**
6. What do you understand by Design of ligand? Explain ligand design based on 3D structure of receptor. **12**
7. Write a note on **any two**. **12**
  - a) Gene therapy.
  - b) Molecular mechanics.
  - c) Lead identification.

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