

35181 : Pharmaceutics - VI : T-8.1

P. Pages : 1

Time : Three Hours

**AW - 2325**

Max. Marks : 60

- Notes :
1. All question carry equal marks.
 2. Answer **any five**.
 3. Illustrate your answer necessary with the help of neat sketches.
 4. Use of pen Blue/Black ink/refill only for writing the answer book.

1. Discuss in detail various approaches used to prepare parenteral controlled release drug delivery system & add a note on its evaluation. **12**
2.
 - i) Discuss in detail the need & importance of stabilization of pharmaceutical product. **6**
 - ii) Explain in detail stability testing protocol for liquid oral & semisolid dosage form. **6**
3. Discuss in detail about preparation of microcapsules by coacervation phase separation method & multi orifice centrifugation methods. **12**
4. Define & classify validation method. Explain in detail process validation involved in production of tablets. **12**
5.
 - i) Explain in detail various types of transdermal drug delivery systems. **6**
 - ii) Define & classify liposomes. Enlist various methods to prepare liposome & describe ethanol injection method in detail. **6**
6.
 - a) Explain in detail about resealed erythrocytes & add a note on limitation of it. **6**
 - b) Add a note on "Evaluation of microcapsules". **6**
7. Write notes on **any two**. **12**
 - a) Applications of polymers in pharmaceutical formulations.
 - b) Retrospective validation
 - c) Nano Spheres
