

GUJARAT TECHNOLOGICAL UNIVERSITY
B.PHARM - SEMESTER- 8 EXAMINATION – WINTER -2019

Subject Code: 2280001**Date: 14-11-2019****Subject Name: Dosage form Design II****Time: 02:30 PM TO 05:30 PM****Total Marks: 80****Instructions:**

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

- Q.1** (a) Discuss controlled release formulation with suitable example. **06**
(b) Explain Hixson and Crowell's cube root law of dissolution. **05**
(c) Give the difference between conventional and controlled release system, describe the evaluation of oral controlled drug delivery system. **05**
- Q.2** (a) Discuss formulation of osmotic based drug delivery system. **06**
(b) Enlist parameters to be considered in design of CRDDS. Discuss approaches for oral delayed release DDS. **05**
(c) Give account of polymers used in formulation of colon drug delivery system. **05**
- Q.3** (a) Discuss Penetration enhancer in TDDS. **06**
(b) Discuss in detail formulation of transdermal drug delivery system. **05**
(c) Discuss in vitro drug release and permeation studies for TDDS. **05**
- Q.4** (a) Discuss method of preparation of nano particles. **06**
(b) Discuss over view of anatomy and physiology of eye and explain absorption pathway for drug in eye. **05**
(c) Give an account of approaches for designing of gastro retentive dosage form. **05**
- Q.5** (a) Discuss on Wagner nelson and Loo-Riegelman method. **06**
(b) Note on volume of distributions. **05**
(c) Discuss one compartment open model, I.V. infusion model and discuss the effect of loading IV injection dose. Describe the derivation of various pharmacokinetic parameters for the model. **05**
- Q. 6** (a) Explain Michaelis Menten kinetics and method to determine K_m and V_{max} . **06**
(b) What are pharmacokinetic model? Explain in detail compartment model? **05**
(c) Define clearance, total body clearance and organ clearance .what is extraction ratio? **05**
- Q.7** (a) Explain dosage adjustment in patients with renal and hepatic failure. **06**
(b) What are the various ways of reducing risk of drug interaction? **05**
(c) Define clinical pharmacokinetics and describe its scope. **05**
