

Seat No.: _____

Enrolment No. _____

GUJARAT TECHNOLOGICAL UNIVERSITY
B.Ph. - SEMESTER– VIII EXAMINATION – WINTER -2020**Subject Code: 2280001****Date: 01/01/2021****Subject Name: Dosage Form Design - II****Time: 02:00PM TO 04:00PM****Total Marks: 54****Instructions:**

1. Attempt any **THREE** questions from Q-1 to Q-6.
2. Q.7 is compulsory to attempt.
3. Make suitable assumptions wherever necessary.
4. Figures to the right indicate full marks.

- Q.1** (a) Write a note on pharmacokinetic drug interactions. **06**
(b) What is clinical pharmacokinetics? Explain dosage adjustment in patients with hepatic failure. **05**
(c) Define pharmacokinetics. Explain scope and significance of plasma drug concentration measurement. **05**
- Q.2** (a) What are pharmacokinetic models? Explain first order absorption rate constant using Wagner-Nelson methods. **06**
(b) Write a note on determination of pharmacokinetic parameters from plasma after oral administration of drug. **05**
(c) Describe method of residuals for determination of absorption rate constant. **05**
- Q.3** (a) Explain formulation and evaluation of a transdermal patch. **06**
(b) Write a detail note osmotic pressure-controlled system. **05**
(c) Explain fabrication of parenteral drug delivery system with special reference to parenteral suspension and emulsion. **05**
- Q.4** (a) Explain rationale for Gastro Retentive Drug Delivery. Enlist different approaches of Gastro Retentive Drug Delivery System. **06**
(b) Define Colon Targeted Drug Delivery Systems. Describe Advantages and limitations of it. **05**
(c) Mention the parameters and methods used for evaluation of nanoparticles. **05**
- Q.5** (a) Write a note on factors influencing design and performance of modified drug delivery systems. **06**
(b) How can a loading and maintenance dose be estimated? **05**
(c) Explain Ocuser[®] and Lacrisert[®] **05**
- Q. 6** (a) Explain merits and demerits of controlled drug delivery systems. **06**
(b) Explain effect of porosity and tortuosity for dissolution-controlled system. **05**
(c) Write a note on biological and physicochemical factors affecting design of oral sustained release systems. **05**
- Q.7** (a) Explain hydrophilic matrix system with suitable example and drug release mechanism from it. **06**

OR

- (a) Discuss apparent volume of distribution. Write an equation to calculate it. **06**

OR

- (a) Write a note on non-linear pharmacokinetics using Michaelis-Menten equation. **06**
