Seat No.: _____

Enrolment No.	Enrol	Iment	No.	
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GUJARAT TECHNOLOGICAL UNIVERSITY

B. Pharm-Semester-VIII Summer-2012 Examination

Subject code: 280001

Subject Name: Dosage Form Design-II

Time: 10:30am to 1:30pm

Date:16 -05-2012 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)	Enlist various approaches for pharmacokinetic analysis of experimental data and explain compartmental models in detail.	06
	(b)	Enlist the methods for determination of absorption rate constant. Explain any one method in detail.	05
	(C)	What process of drug ADME are known to show non linearity Explain giving suitable examples	05
Q.2	(a) (b)	Explain in detail: Ideal requirements for sustained release formulations. Why loading dose is required in sustained release formulations? Give the equations for loading dose and maintenance dose.	06 05
	(C)	Explain osmotic pressure controlled drug delivery system.	05
Q.3	(a)	Define "Drug interaction". Explain pharmacokinetic drug interactions giving suitable examples	06
	(b) (c)	Explain dosage adjustment in patients with renal and hepatic failure. Explain volume of distribution and distribution coefficient	05 05
Q.4	(a)	Define clearance, total body clearance and organ clearance. What is extraction ratio?	06
	(b)	Name different methods for determination of Ke from urinary excretion data. Give the criteria for obtaining valid urinary excretion method.	05
()	(C)	Write a note on Wagner Nelson method.	05
Q.5	(a) (b) (c)	Explain targeted drug delivery systems giving examples. Discuss formulation of diffusion layer controlled drug delivery systems. Write a note on matrix based controlled drug delivery system.	06 05 05
Q. 6	(a) (b) (c)	Discuss in detail formulation of transdermal drug delivery system. Explain gastro retentive drug delivery systems. Write a note on buccal drug delivery systems.	06 05 05
Q.7	(a)	Enlist different approaches for formulation of colon targeted drug delivery system and explain any two of them.	06
	(b) (c)	Write a note on ophthalmic controlled release systems Differentiate between microspheres and micro capsules	05 05