Seat N	lo.:	Enrolment No	
Sout 1		GUJARAT TECHNOLOGICAL UNIVERSITY	
В	B.PH	ARM – SEMESTER – VIII • EXAMINATION – WINTER – 2015	
•	Subject Name: Dosage Form Design II		
•		0 PM to 5.30 PM Total Marks: 80	
Instr			
		empt any five questions.	
2.	Mak	ke suitable assumptions wherever necessary.	
	_	res to the right indicate full marks.	0.6
Q.1	(a)	What do you mean by release modulation of an API and why is it necessary in suitable dosage form?	06
	(b)	What biological and physicochemical factors are responsible with their optimum range to design oral controlled DDS?	05
	(c)	Give the example of hydrophilic matrix system and drug release mechanism from it.	05
Q.2	(a)	Differentiate Novel Drug Delivery System from Conventional Dosage Form. How is it beneficiary to patient? Explain giving illustration	06
	(b)	Enlist the systems designed and developed for oral modified release and describe any one of them in detail.	05
	(c)	Which are the parenteral drug delivery systems? Give the suitable API for novel DDS in parenteral form.	05
Q.3	(a)	What do you mean by site specific and targeted drug delivery system? Give suitable example describing these systems.	06
	(b)	Differentiate transdermal from transmucosal. Write a note on buccal dosage form.	05
	(c)	Write historical aspects of ophthalmic drug delivery system. Write a note on Occuserts.	05
Q.4	(a)	How the transdermal patch is being formulated? Explain giving illustrative diagram.	06
	(b)	Differentiate transdermal patch and transdermal film. Which one is used where and when?	05
	(c)	What is GRDDS? What are the important advice for patients taking this dosage form?	05
Q.5	(a)	How will you estimate loading dose and maintenance dose in sustained release dosage form?	06
	(b) (c)	Write a note on CPOP with suitable API and its release mechanism. How will you incorporate nanostructures in suitable dosage form?	05 05
Q. 6	(a)	What do you mean by pharmacokinetic models? Discuss its importance and application of it.	06
	(b) (c)	What is apparent volume of distribution? Write an equation to calculate it. What do you mean by method of residuals? Draw an illustrative diagram for that.	05 05
Q.7	(a)	Define Clinical Pharmacokinetics and Describe scope of it. What kind of desage adjustment are needed in patients with and without renal	06 05
	(b) (c)	What kind of dosage adjustment are needed in patients with and without renal and hepatic failure? What do you mean by combination therapy? What kind of drug interactions are	05
	(0)	possible, explain giving suitable examples.	03
