Enrolment No

GUJARAT TECHNOLOGICAL UNIVERSITY B. PHARM (SEMESTER-8) SUMMER-2015

B. PHARM (SEMESTER-8) SUMMER-2015						
•	Subject code: 280001 Date: 28/04/201					
•	Subject Name: Dosage Form Design - II					
Time: 10:30 am to 1:30 pmTotal Marks: 80						
	ctions:	, no ,o				
		pt any five questions.				
 Make suitable assumptions wherever necessary. Figures to the right indicate full marks. 						
5.	Figure	is to the right mulcate full marks.				
Q.1	(a)	Mention the parameters and methods used for evaluation of	06			
C		microspheres.				
	(b)	Describe in detail physicochemical factors affecting design of oral	05			
		sustained release dosage forms.				
	(c)	What are the important characteristics to be evaluated for design of	05			
		parenteral suspensions?				
Q.2	(a)	Describe solvent emulsification evaporation method for preparation	06			
		of nanoparticles. Mention advantages and disadvantages of the method.				
	(b)	Write in detail formulation design of transdermal drug delivery	05			
	(0)	system.	05			
	(c)	Classify the polymers used for preparation of matrix tablets. Give	05			
		two names of each class.				
Q.3	(a)	Enlist different pharmacokinetic models. What is compartment	06			
		model? Mention advantages and disadvantages of the same.				
	(b)	Discuss the properties of drug that can be enhanced by drug carrier	05			
		delivery system for drug-targeting.	05			
	(c)	What are the merits and demerits of using urinary excretion data for pharmacokinetic parameters?	05			
		pharmacoknetic parameters:				
Q.4	(a)	What properties are required for the drug to be a candidate for	06			
C		transdermal drug delivery system? Write evaluation method for				
		adhesive properties of TDDS				
	(b)	A physician administering a loading dose and simultaneously IV	05			
		infusion to get C_{ss} (steady state concentration) of $2mg/L$ if drug has				
		elimination rate constant $K = 0.2hr^{-1}$.Vd=15L. Calculate the required				
		loading dose and infusion rate.	0.5			
	(c)	Explain the term "renal clearance". Describe graphical method for determination of renal clearance.	05			
Q.5	(a)	Describe in detail preparation of HBS for gastric retention of oral	06			
Q	(u)	dosage forms. What is the method of evaluation of such dosage	vu			
		forms for gastric retention time?				
	(b)	A long acting antibiotic has a half life of 14 days, how long will it	05			
		take for the drug concentration in the blood to drop to 90% of its				
		initial value?				
	(c)	Write a note on osmotic ocular inserts. Mention the components of	05			
		each part.	of			

Q. 6	(a) (b) (c)	Write a note on Wagner Nelson method for pharmacokinetics of drug absorption. Mention the advantages and limitations of colon targeted drug delivery system What is the composition of liposome? Mention the therapeutic applications of liposome.	06 05 05
Q.7	(a) (b)	What type of drugs can benefit from using gastric retentive devices? What are the limitations of these devices? Explain non-linear pharmacokinetics using Michaelis Menten equation	06 05
	(c)	A new antihypertensive drug was given to a patient in single I.V. dose of 200mg to a 60Kg male. After 6hrs the blood concentration of drug was found to be 2.0mg/100ml of blood. If the Vd is 10% of body weight, compute the total amount of drug in body fluids after 6hrs. What is the half-life of the drug?	05
