C.U.SHAH UNIVERSITY Summer Examination-2017

Subject Name : Pharmaceutical Dosage Form Design I

Subject Code : 4PS07DFD1		Branch :B.Pharm	
Semester : 7	Date : 23/03/2017	Time : 02:30 To 05:30	Marks : 70
Instructions:			
(1) Use of Pro	grammable calculator & an	y other electronic instrument	is prohibited.
(2) Instruction	s written on main answer be	ook are strictly to be obeyed.	-
(3) Draw neat	diagrams and figures (if ne	cessary) at right places.	
(4) Assume su	itable data if needed.		

Q-1 Attempt the following questions:

Define the following terms:

- a) Habit
- **b**) Amorphous
- c) Monotropic
- d) Energy of Activation
- e) Arrhenius Factor
- **f**) Area under curve
- g) Dissolution
- **h**) Osmosis
- i) Disintegrant
- j) Overages
- **k**) Sink condition
- l) Absorption
- **m**) Elimination
- n) Pharmacodynamic

Attempt any four questions from Q-2 to Q-8

Q-2	a)	Attempt all questions Define Preformulation? Explain the physicochemical properties related to solubility analysis in Preformulation.	(14) (7)
	b)	Describe the various measures to prevent hydrolysis decomposition.	(7)
Q-3	a) b)	Attempt all questions Explain the effect of salt formation and racemization in preformulation. Write a short note on anti-oxidants.	(14) (7) (7)

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(14)

Q-4		Attempt all questions	(14)
	a)	Define prodrugs. Discuss pharmaceutical application of prodrugs.	(7)
	b)	Describe matrixing and bracketing in stability study.	(7)
Q-5		Attempt all questions	(14)
	a)	Discuss plasma protein drug binding.	(7)
	b)	What is BCS? Classify and give the significance of this system.	(7)
Q-6		Attempt all questions	(14)
C	a)	Differentiate absolute and relative bioavailability. Discuss the pharmacokinetic methods for the bioavailability measurement.	(7)
	b)	What is gastric emptying? Describe its role in drug absorption.	(7)
Q-7		Attempt all questions	(14)
Ľ	a)	Write a short note on drug distribution to blood brain barrier.	(7)
	b)	Explain AUC. What is its significance? How will you measure it?	(7)
Q-8		Attempt all questions	(14)
τ°	a)	Explain any one dissolution test apparatus as per I.P.	(7)
	b)	Write a note on stability testing guideline as per ICH for pharmaceutical drug product.	(7)



