

## NF-02011264

Seat No.

## M. Pharm. (Sem. I) (CBCS) Examination

January - 2017

## Pharmaceutical Preformulation: Paper - I

(Multidisciplinary/Elective)

Time: 3 Hours] [Total Marks: 80

Instructions:

- (1) Answer any three from each section except question 1 and 5 are compulsory.
- (2) Figure to the right indicates marks.
- (3) Draw neat and clean diagrams as required.

## **SECTION - I**

- 1 Answer any seven out of given 10 questions:
- $2 \times 7 = 14$
- (a) Enumerate the methods for particle size reduction.
- (b) What do you mean by BCS Classification system.
- (c) Define Polymorphism.
- (d) Give difference between Amorphous and crystalline forms.
- (e) Define Partition coefficient.
- (f) What do you mean by pka.
- (g) What is drug exicipient compatibilities studies.
- (h) Define: dissolution.
- (i) What is bulk density.
- (j) What do you mean by ICH Guidelines.
- 2 Answer the following:
  - A. Discuss in detail about the methods to increase the solubility 7 of drug.
  - B. Write a note on Polymorphism.

6

3	Answer the following:		
	A.	Defin, objective, goals and criterias of preformulation studies.	7
	B.	Discuss in detail about pharmaceutical solid polymorphism.	6
4	Ans	wer the following: Discuss the various methods to determine partition coefficient of drug.	7
	B.	Write a note on pharmaceutical salt form.	Ć
		SECTION - II	
5	Ans	wer any two out of three : $7 \times 2 =$	14
	(a)	Write a note on methods for evaluation of solid state.	
	(b)	Discuss characterization of Phytomedicine.	
	(c)	Describe in detail about solid state properties.	
6	Answer the following		
	A.	Write a note on phase solubility analysis.	7
	B.	Discuss partitioning of Acid and Base in Biological membrane.	6
7	Answer the following		
	A.	Discuss the influence of particle size in the formulation of	7
		dosage forms.	
	B.	Write a note in brief regarding ionization and lipophilicity of	6
		drug.	
8	Answer the following:		
	A.	Discuss in brief about factors affecting dissolution Rate.	7
	B.	Write a note on freeze dried products.	6

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