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UNIVERSITY OF JAFFNA, SRI LANKA BACHELOR OF PHARMACY SECOND YEAR SECOND SEMESTER EXAMINATION – FEBRUARY 2017 PHACE 2234 PHARMACEUTICS III

Date: 13.02.2017 Time: 02 Hours

ANSWER ALL EIGHT QUESTIONS.

1.1 Define "pharmaceutical dissolution".	(10 marks)	
1.2 Explain the	¥.	
1.2.1 process of dissolution of solids using Noyes whitney equation.	(40 marks)	
1.2.2 factors that affect the dissolution rate of solids.	(50 marks)	
, , , , , , , , , , , , , , , , , , , ,	(60 marks)	
2.2 Enumerate the pharmaceutical application of colloids.	(40 marks)	
3.1 Explain the effect of the following on the stability of suspension		
	(30 marks)	
	(25 marks)	
3.2		
3.2.1 Classify emulsifying agents.	(15 marks)	
3.2.2 Explain how above mentioned emulsifying agents stabilize the emulsions?		
	(30 marks)	
4.1 List the trues of wheelesical systems	(15 montes)	
**	(15 marks)	
	(30 marks)	
	(30 marks)	
100	()	
4.3.1 suspensions.	(15 marks)	
4.3.2 ophthalmic solutions.	(10 marks)	
	 1.2 Explain the 1.2.1 process of dissolution of solids using Noyes whitney equation. 1.2.2 factors that affect the dissolution rate of solids. 2.1 Classify and differentiate different types of colloids. 2.2 Enumerate the pharmaceutical application of colloids. 3.1 Explain the effect of the following on the stability of suspension. 3.1.1 Electrolytes 3.1.2 Polymers. 3.2 3.2.1 Classify emulsifying agents. 3.2.2 Explain how above mentioned emulsifying agents stabilize the emuls. 4.1 List the types of rheological systems. 4.2 Explain the mechanism of 4.2.1 shear thickening system. 4.2.2 thixotropy. 4.3 Explain the importance of rheology in 4.3.1 suspensions. 	

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	5.1 What is the importance of interfacial phenomena in the development of	
	pharmaceutical formulations?	(20 marks)
	5.2 Classify surfactants with examples.	(20 marks)
	5.3 Write an account on	
	5.3.1 Hydrophilic Lipophilic Balance value of surfactants.	(20 marks)
	5.3.2 application of surfactants in pharmaceutical formulations.	(40 marks)
6.		
	Explain the methods that improve the flow property of powders?	(30 marks)
	6.2 Briefly explain how drugs can be protected from	
	6.2.1 hydrolysis.	(40 marks)
	6.2.2 oxidation.	(30 marks)
7.	Straight for	
	7.1 Explain the influence of water/lipid solubility of drugs in the passive tra	ansport
	across biological membrane.	(30 marks)
	7.2 Briefly describe the accelerated stability testing of drugs.	(70 marks)
0	with the following out in an analysis of the second of the	
8.	Write an account on	
	8.1 significance of protein-drug binding.	(30 marks)
	8.2 structural features of gels.	(30 marks)
	8.3 β-cyclodextrins.	(40 marks)

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