# **Question Paper**

Exam Date & Time: 05-May-2018 (02:00 PM - 05:00 PM)



## MANIPAL ACADEMY OF HIGHER EDUCATION

#### MANIPAL COLLEGE OF PHARMACEUTICAL SCIENCES END SEMESTER THEORY EXAMINATIONS- MAY 2018 PROGRAM: MPHARM SEMESTER 2 (PHARMACEUTICS and INDUSTRIAL PHARMACY) DATE: 05/05/2018 TIME: 2:00 PM - 5:00 PM

#### Advanced Biopharmaceutics and Pharmacokinetics [PCE-MPH202T]

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Marks: 50

Duration: 180 mins.

### Answer all the questions.

### Answer the following (5 marks x 8 = 40 marks)

1)	Explain the limitations of pH partition hypothesis.	(5)
2)	Discuss the construction and working of USP apparatus 1 for dissolution testing.	(5)
3)	Explain the effect of polymorphism of drugs on their absorption.	(5)
4)	Explain the effect of Binders and Disintegrants on the dissolution of tablets.	(5)
5)	Explain the characteristic features of active transport of drugs.	(5)
6)	Describe the Pharmacokinetics of one compartment model for an IV bolus drug administration. Support your answer with appropriate graphs and equations.	(5)
7)	What are the causes of nonlinear pharmacokinetics of a drug? Explain with suitable graphs.	(5)
8)	Write and explain the importance of BCS classification system. Give examples.	(5)
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### Answer all the questions.

### Answer the following with specific answers (2 marks x 5 = 10 marks)

<sup>9)</sup> How Clearance and Volume of Distribution related? How useful <sup>(2)</sup>
<sup>(2)</sup> this relation is?

- <sup>B)</sup> What is the loading dose required for drug A if Target concentration is 10 mg/L, Vd is 0.75 L/kg and Patient's weight is 75 kg
- <sup>C)</sup> Show in a graph the Plasma concentration of drug, when the Rate <sup>(2)</sup> of infusion of drug is equal to the rate of elimination?
- D) Why apparent volume of distribution tends to decrease for protein <sup>(2)</sup> drugs?
- E) Why the pharmacokinetic parameters calculated using plasma (2) data does not represent the actual pharmacokinetics of the drug from targeted delivery systems?

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