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# B.Pharma (2012 to 2016) (Sem.-8) PHARMACEUTICS-IX (DOSAGE FORM DESIGN)

Subject Code : BPHM-801 M.Code : 72296

Time: 3 Hrs. Max. Marks: 80

# INSTRUCTIONS TO CANDIDATES:

- SECTION-A is COMPULSORY consisting of FIFTEEN questions carrying TWO marks each.
- SECTION-B contains FIVE questions carrying FIVE marks each and students have to attempt any FOUR questions.
- SECTION-C contains FOUR questions carrying TEN marks each and students have to attempt any THREE questions.

## SECTION-A

## Answer briefly :

- Describe the role of solubility in preformulation studies.
- Define bioequivalence.
- Define polymorphism.
- Define dissolution.
- e. Define bioavailability.
- Name any two drugs prone to hydrolysis.
- g. Define solvates.
- Name any two drugs prone to oxidation.
- Explain hydrates.
- Define prodrugs.

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- Discuss the importance of student's T test.
- 1. Name any two approaches for solubility enhancement.
- m. Define absolute and relative bioavailability.
- n. What are controlled release dosage forms?
- o. What are biodegradable polymers?

## SECTION-B

- What are the polymers used for enteric coated tablets? Discuss the in vitro testing of enteric coated tablets.
- Discuss various factors affecting the oxidation of drugs.
- What is BCS? Discuss various classes along with suitable example.
- 5. What are the prominent features of quality assurance?
- 6. Explain the rationale for retrospective validation. Also, discuss the process briefly.

#### SECTION-C

- Discuss preformulation studies. Enlist the various properties of the drugs to be considered while performing the preformulation studies.
- Discuss the ICH guidelines for accelerated stability studies.
- Classify oral controlled release drug delivery system.
- Explain the application of prodrugs in various pharmaceutical fields.

NOTE: Disclosure of Identity by writing Mobile No. or Making of passing request on any page of Answer Sheet will lead to UMC against the Student.

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