

**M. PHARM. SEM-I (CHOICE BASED CREDIT &  
GRADE SYSTEM) : WINTER - 2017**

**SUBJECT : ADVANCED PHARMACEUTICS – I**

Day : **Monday**  
Date : **08/01/2018**

**W-2017-3857**

Time : **10.00 AM to 01.00 PM**  
Max. Marks : 60

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**N.B.**

- 1) Answer any **THREE** questions from **Section – I** any **THREE** questions from **Section – II**.
  - 2) Both the sections should be written in separate answer books.
  - 3) Figures to the right indicate **FULL** marks.
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**SECTION - I**

- Q.1** Elaborate on the significance of drug-excipient compatibility testing with the help of examples. (10)
- Q.2** Give an account of various parameters to be considered during stability testing and evaluation as per ICH guidelines. (10)
- Q.3** Explain the various kinetic models useful to describe dissolution profile. Add a note on model independent approach to compare dissolution profiles. (10)
- Q.4** Write short notes on: (10)
- a) Factorial design.
  - b) Principle and instrumentation of laser diffraction based particle size analysis

**SECTION – II**

- Q.5** What are cyclodextrins? Explain the methods of preparation and characterization of cyclodextrin complexes. (10)
- Q.6** What are bioresorbable polymers? Explain the different mechanisms of biodegradation. (10)
- Q.7** Explain the different liquid crystalline phases and the methods to identify them. (10)
- Q.8** Write notes on : (10)
- a) Glass transition temperature of polymers and its significance.
  - b) Mathematical approaches to study compressional behavior of solids.

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