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Paper Code :DMS-221 Roll No.

MBA-14(Pharmaceutical Marketing) 2nd Year Examination, Academic Batch 2016-17 Pharmaceutical Formulation Design and Development

Time: 3 Hours [Max. Marks: 100

Note. Attempt any *Five* questions. All questions carry equal marks.

- Q1. Define Preformulation studies. Explain the role of crystallinity and polymorphism study in the development of a new drug formulation in solid state.
- Q2. Define and classify polymers. Write their synthesis and properties..
- Q3. Explain the following terms:
- a) BCS system of classification of drugs. b) Similarity factor.
- Q4. Explain various factors influencing *IVIVC*
- Q5. Explain various additives used in liquid dosage forms.
- Q.6. Explain pharmaceutical ingredients and excipients-definition and types
- Q.7. Discuss various additives used in semisolid dosage forms.
- Q.8. Write a note on accelerated stability testing of pharmaceutical products.