# [Total No. of Questions - 7] [Total No. of Printed Pages - 2] (2064)

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## M. Pharmacy 2nd Semester Examination Industrial Pharmacy

**MP-122** 

Time: 3 Hours Max. Marks: 90

The candidates shall limit their answers precisely within the answerbook (40 pages) issued to them and no supplementary/continuation sheet will be issued.

Note: Section A consists of Nine questions, Section B consists of Four questions and Section C has Two questions. Attempt Seven from Section A, Three from Section B and One from Section C.

#### **SECTION - A**

- 1. (a) Explain effect of particle size, moisture content and lubrication on strength of tablets.
  - (b) Explain inventory management in industrial pharmacy.
  - (c) Discuss layout and design of pharmaceutical industries.
  - (d) Explain regulatory guidelines for sustained/controlled release products.
  - (e) What is pilot plant scale up? Explain pilot study of capsules in detail.
  - (f) Explain finished product specification in pilot plant scale up techniques.
  - (g) Write a short note on industrial hazards alongwith their preventive measures.

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- (h) Explain USFDA requirements for animal drugs.
- (i) Explain role of TQM (Total Quality Management) in production of dosage funds. (7×5=35)

#### **SECTION - B**

- 2. Write a note on formulation aspects of sublingual tablets.
- 3. Explain the structure of pharmaceutical industry with special reference to packaging division.
- 4. Explain the role of documentation in production management with suitable examples.
- 5. Explain ISO 9000 series.

 $(10 \times 3 = 30)$ 

### **SECTION - C**

- 6. Discuss process validation for tablets in detail.
- 7. Define GMP. Explain GMP considerations in production management in detail. (25×1=25)