[Max. Marks: 100] [Time: 3 Hours]

Preformulation and Production Management (Revised Scheme 4)

Q.P. CODE: 9337

Your answers should be specific to the questions asked. Draw neat, labeled diagrams wherever necessary.

LONG ESSAY (Answer any TWO)

2 X 20 = 40 Marks

- Explain the validation of manufacturing process for sterile and non-sterile products including 1. brief protocols and reports.
- 2. Explain the various techniques to study the crystal properties and polymorphism.
- 3. Write a note on compression and consolidation of powders. Explain the influence of compression force on the properties of tablets.

SHORT ESSAY (Answer any FIVE)

5 X 10 = 50 Marks

- List-out and explain preliminary evaluation tests required before preformulation studies. 4.
- 5. Give the difference between ICH and WHO guidelines for stability studies.
- Give all the general considerations required for setting up a pilot-plant. 6.
- 7. Explain classical optimization and simplex method.
- 8. Give the detailed procedure to obtain a patent.
- 9. Classify material handling equipments and explain in detail.

Explain briefly the various levels of inventory.

Mechanical hazards

***** SHORT NOTES 2 X 5 = 10 Marks

- 10.
- 11.

