Seat No.:	Enrolment No.

GUJARAT TECHNOLOGICAL UNIVERSITY M. Pharm. – SEMESTER – I • EXAMINATION – WINTER 2013

Subject Code: 910208 Date: 26-12-2013 **Subject Name: Industrial Pharmacy - II** Time: 10.30 am - 01.30 pm **Total Marks: 80 Instructions:** 1. Attempt any five questions. 2. Make suitable assumptions wherever necessary. 3. Figures to the right indicate full marks. **Q.1** Write significance of pilot plant scale up study. Discuss about large scale (a) 06 manufacturing techniques of injectables. Describe general layout and facilities requirements of pilot plant. 05 **(b)** Write note on technology transfer: Pilot plant to production. (c) 05 **Q.2** 06 Discuss the cost control elements in production management. (a) Write a short note on documentation importance in maintenance. 05 **(b) (c)** Discuss vendor development capacity assessment for raw materials to be used in 05 semisolids. **Q.3** (a) Write in detail note on various techniques used in inventory management. 06 Define EOQ. Discuss advantages and limitation of this system. 05 **(b)** (c) Explain production scheduling and forecasting. 05 0.4 Discuss the management of scrap disposal in pharmaceutical plant. 06 (a) Classify types of management and discuss material management in detail. 05 **(b)** Classify maintenance and discuss in detail preventive maintenance. (c) 05 Explain COD. Discuss monitor and controls for COD of industrial waste water. **Q.5** 06 (a) **(b)** Discuss the role of personnel in production management. 05 Discuss environmental factors influencing on production management. 05 (c) Classify the industrial hazards and discuss fundamentals of accident prevention. 06 Q. 6 (a) **(b)** Write a detail note on sources & prevention of electrical hazards. 05 Discuss on safety measures and pollution control measures applicable in 05 (c) pharmaceutical industries. **Q.7** Justify: Pilot plant and industrial scale equipments does not create any problem (a) 06 with the equipment design. Importance of stability parameter in scale up techniques. 05 **(b)** (c) Discuss product uniformity parameter of dosage form during scale up techniques 05
