

GUJARAT TECHNOLOGICAL UNIVERSITY
B. Pharm - SEMESTER- VI EXAMINATION – WINTER 2021

Subject Code: 2260003**Date: 25/11/2021****Subject Name: Pharmaceutical Analysis IV****Time: 02:30PM TO 05:30PM****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** (a) Explain the principle and instrumentation of Super Fluid Chromatography. **06**
(b) Describe Flame Ionization Detector. **05**
(c) Explain the principle and applications of RIA. Explain Enzyme immuno assay-ELISA. **05**
- Q.2** (a) Explain different types of IPR and write the steps for patent filing. **06**
(b) Write a note on GLP. **05**
(c) What is analytical method validation and explain various validation parameters as per ICH guideline. **05**
- Q.3** (a) Explain Bragg's law equation for diffraction of X-rays by crystals and list applications of X-ray diffraction. **06**
(b) Explain technique of Affinity chromatography. **05**
(c) Explain principle and applications of ion exchange chromatography. **05**
- Q.4** (a) Explain hyphenation technique: LC-MS. **06**
(b) Explain various stationary phases used in gas chromatography. **05**
(c) Write a note on mobile phase used in HPLC. **05**
- Q.5** (a) Differentiate:
1. Reverse and Normal phase chromatography **06**
2. HPTLC and HPLC
3. Capillary and Open tubular column
(b) Write a note on Turbidimetry. **05**
(c) Write a brief note on Size exclusion chromatography. **05**
- Q.6** (a) List ideal requirements of detector for HPLC and explain UV-Visible detector in HPLC. **06**
(b) What are the various steps for chromatographic development in HPTLC. **05**
(c) Explain isotope dilution analysis and liquid scintillation counter. **05**
- Q.7** (a) Differentiate: HPLC and GC. **06**
(b) Give an overview of GC-MS. **05**
(c) Explain objective and advantages of ISO 9001 standards. Also explain quality management system in pharmaceutical industry. **05**
