

GUJARAT TECHNOLOGICAL UNIVERSITY
B.PHARM – SEMESTER – 8- EXAMINATION –WINTER - 2018

Subject Code: 280004

Date: 22/11/2018

Subject Name: Pharmaceutical Analysis-IV

Time: 02:30 PM TO 05:30 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 | (a) Describe generation and characteristic of X-rays. Explain principle of X-Ray Diffractometer. | [6] |
| | (b) Describe applications of X-ray diffraction. | [5] |
| | (c) Explain in brief technique of GC-MS. | [5] |
| Q-2 | (a) Write a note on stationary phases in normal and reversed phase HPLC | [6] |
| | (b) Discuss detectors used in HPLC. | [5] |
| | (c) Explain the principle, instrumentation and applications of super critical fluid chromatography. | [5] |
| Q-3 | (a) Explain the principle of Gas chromatography. Explain the terms (i) GSC (ii) GLC. | [6] |
| | (b) Enlist the detectors used in GC and discuss construction and working of any one. | [5] |
| | (c) Enumerate principles of chromatography. Discuss size exclusion chromatography. | [5] |
| Q-4 | (a) Give advantage and limitation of HPTLC. | [6] |
| | (b) Discuss purposes & techniques of chemical derivatization in HPLC& GC with illustration. | [5] |
| | (c) Write full form of TRIPS and GATT. Give requirement of TRIPS. | [5] |
| Q-5 | (a) Write a note on Good Laboratory Practice (GLP). | [6] |
| | (b) What is IPR. Explain each with example. | [5] |
| | (c) What are types of patent applications? | [5] |
| Q-6 | (a) Describe Radioimmuno assay. | [6] |
| | (b) Give a brief account on ELISA technique. | [5] |
| | (c) Write a brief note on Liquid scintillation systems. | [5] |
| Q-7 | (a) Define and explain analytical method validation as per ICH guideline. Explain: how precision of assay method are determined? | [6] |
| | (b) What is the basic difference between nephelometry and turbidimetry? Describe applications of nephelometry and turbidimetry. | [5] |
| | (c) Explain requirement of ISO 9001-2000. | [5] |