

UKA TARSADIA UNIVERSITY

M. Pharm. (Pharmaceutical Analysis) II Semester

040060202 - Pharmaceutical Analysis-II

SECTION I

Q.1] (a) Answer the following [7 x 1 = 7]

- 1) What is counter current chromatography?
- 2) Define super critical fluid.
- 3) What is a biological sample?
- 4) What is ultra filtration?
- 5) Define partition coefficient.
- 6) What is the principle of dialysis as a sample preparation technique?
- 7) What is peptide mapping?

(b) Attempt any four [4 x 2 = 8]

- 1) Explain the terms in SEC: Exclusion limit, Permeation limit.
- 2) What is the significance of system biocompatibility in peptide mapping?
- 3) Describe the uses of peptide mapping.
- 4) Explain column switching as a sample preparation technique.
- 5) What is the significance of post column derivatization in amino acid analysis?

Q.2]

(1) Discuss the role of SDS-PAGE in analysis of proteins and peptides. [05]

OR

(1) What is isoelectric focusing? Explain synthetic carrier ampholytes and immobilized pH gradients in IEF. [05]

(2) Discuss the significance of solid phase extraction as a sample preparation technique. [05]

OR

(2) Explain the principles and procedures for amino acid sequence analysis. [05]

Q.3] Attempt any two [2 x 5 = 10]

- (1) Explain the significance of GC-MS in quantification of drugs in biological samples with suitable examples.
- (2) What are the advantages of SFC over HPLC and GC? Enlist the components of an instrument used for SFC.
- (3) Explain how ion exchange chromatography aid in analysis of proteins and peptides. Discuss factors influencing retention in IEC.

SECTION II

Q.4] (a) Answer the following [7 x 1 = 7]

- 1) Define conformational polymorphism.
- 2) What is extractive value?
- 3) Define Crude fiber content.
- 4) What is flash chromatography?
- 5) Enlist the methods used for quantitative analysis of dosage forms containing analgesics and antipyretics.
- 6) Define Quality Control.
- 7) What do you mean by pesticidal residue?

(b) Attempt any four [4 x 2 = 8]

- 1) Residual solvent testing is important in pharmaceutical industry. Justify.
- 2) Classify analytical techniques used for solid state analysis.
- 3) How will you determine acid insoluble ash value of crude drugs?
- 4) Explain the terms: Bitterness index, foaming index.
- 5) Define Specific absorbance, Molar absorptivity.

Q.5]

(1) Define automated systems. Differentiate between types of automatic analytical systems. [05]

OR

(1) Discuss the role of chromatography in identification of plant constituents with suitable examples. [05]

(2) Discuss the principles and procedures involved in analysis of pharmaceutical dosage forms containing anti diabetics. [05]

OR

(2) Discuss the role of NMR spectroscopy in solid state analysis with suitable examples. [05]

Q.6] Attempt any two [2 x 5 = 10]

- (1) Discuss the role of titrimetric methods in analysis of antihypertensive and antihistaminic drugs.
- (2) Discuss the role of isolation techniques in degradation and impurity analysis.
- (3) Explain the principle of flow injection analysis.