

GUJARAT TECHNOLOGICAL UNIVERSITY**M.PHARM- SEM-II-EXAMINATION – JULY 2012****Subject code: 2920107****Date: 06/07/2012****Subject Name: Pharmaceutical Analysis 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	(a)	Discuss the principle, instrumentation and application of the GC-MS method used in pharmaceutical analysis.	06
	(b)	Describe and discuss: counter current chromatography	05
	(c)	Describe and discuss: Size exclusion chromatography	05
Q.2	(a)	Discuss principle and application of supercritical fluid chromatographic technique for analysis and isolation.	06
	(b)	Discuss the principle and applications of electrophoretic techniques	05
	(c)	Write a short note on the methods of automated analysis.	05
Q.3	(a)	Describe and discuss with respect to analysis of analysis of protein: isoelectric focusing and tryptic mapping.	06
	(b)	Write a short note on amino acid sequence analysis.	05
	(c)	Discuss methods of assay for anti hypertensive drugs and drugs acting on CV system.	05
Q.4	(a)	Discuss methods of solid state analysis of drugs.	06
	(b)	Discuss principles and procedures involved in the analysis of sedatives and tranquilizers.	05
	(c)	Discuss principle and applications of electrophoretic techniques.	05
Q.5	(a)	Discuss importance of Reporting threshold, Identification threshold and qualification threshold as discussed in the ICH Q3 guidelines.	06
	(b)	Discuss methods of determination of total solids in asava/aristha.	05
	(c)	Describe methods for the determination of iodine value in herbal formulations.	05
Q. 6	(a)	Discuss specific sample preparation techniques from biological matrices.	06
	(b)	Discuss UV and fluorescent analysis of powdered crude drugs and their applications	05
	(c)	Write a note on proximate analysis of crude drugs	05
Q.7	(a)	Enlist and discuss different techniques and methods applied in the extraction and identification of phytochemicals.	06
	(b)	Write a short note on pharmaceutical sample preparation.	05
	(c)	Discuss WHO guideline about coarseness of powder material and the sieves to be used.	05