Seat]	No.:	Enrolment No.	
	-	GUJARAT TECHNOLOGICAL UNIVERSITY	
M. Pharm. – SEMESTER – I • EXAMINATION – WINTER • 2014 Subject Code: 910104 Date: 07-01-2015			
Subject Name: Biological Evaluations and Clinical Research			
Time: 10:30 am - 01:30 pm Total Marks: 80 Instructions:			
Instr	1.	Attempt any five questions. Make suitable assumptions wherever necessary.	
Q.1	(a) (b) (c)	Describe principle of GCP as per ICH guideline. Discuss ELISA for protein hormones. Explain importance and clinical significance of Student's t-test in Bio-assay.	06 05 05
Q.2	(a) (b) (c)	What is toxicity? Discuss general methodology for toxicology. Describe Gel-Clot technique for LAL test. Write a short note on Helsinki declaration.	06 05 05
Q.3	(a) (b) (c)	Describe advantages of urinary excretion studies. Explain also the criteria for obtaining valid urinary excretion data. Describe microbial limit tests with its applications. Describe radio-immunoassay of insulin.	06 05 05
Q.4	(a) (b) (c)	Enumerate various techniques for depyrogenations. Explain any two techniques in details.What is toxicity study? Describe briefly the parameters for measuring toxic effect.Enumerate different biological methods in Pharmaceutical Analysis.	06 05 05
Q.5	(a) (b) (c)	Explain, classify and give uses of pharmacokinetic models. Describe chemical properties of pyrogens and endotoxins. Describe Indian guidelines on biomedical research and clinical trials.	06 05 05
Q. 6	(a) (b) (c)	Write a note on design, approval and execution of protocol for bioequivalence studies.Describe the significance and methods used for testing effectiveness of antimicrobial preservatives.Explain in detail about Protein precipitation method used in various studies.	06 05 05
Q.7	(a) (b) (c)	 Write a note on Module 4 and Module 5 of CTD describing non-clinical study reports and clinical study reports. Describe in detail about Objectives and consideration in bio-availability studies. Explain LD₅₀ and ED₅₀. Describe method for their determination. 	06 05 05
