GUJARAT TECHNOLOGICAL UNIVERSITY M. Pharm. – SEMESTER – III • EXAMINATION – WINTER • 2014

Subject Code: 930104Date: 06-12-2014Subject Name: Validation and Product Development			
Time: 10:30 am - 01:30 pm Total Marks: 80 Instructions:			
IIISU	1. 2.	Attempt any five questions. Make suitable assumptions wherever necessary. Figures to the right indicate full marks.	
Q.1	(a) (b) (c)	Define validation. Discuss types and advantages of Validation. Discuss design and installation qualification in process validation. Describe calibration of pH meter.	06 05 05
Q.2	(a) (b)	Enumerate in process controls employed in manufacturing process design of ophthalmic & parenteral preparations. What are limitations of sterility test? Explain techniques of rapid or real time	06 05
	(c)	sterility test. Write note on packing material for parenteral product.	05
Q.3	(a) (b)	Discuss the validation process for fluid bed and tray dryer. What are the objectives of cleaning validation? Describe the methods of cleaning validation.	06 05
	(c)	Describe validation of water system for pharmaceuticals.	05
Q.4	(a) (b)	Explain the terms Validation Protocol and Validation Report. Give Protocol for validation of tablet compression. What is Validation Master Plan? Explain its importance in setting up of a new manufacturing facility.	08 08
Q.5	(a)	Define D-value and F-value. Describe OQ and PQ steps involved in the validation of autoclave.	08
	(b)	Describe validation of HVAC system in sterile product manufacturing area.	08
Q. 6	(a)	Discuss the requirements of scale-up and post-approval changes in immediate release dosage form (SUPAC-IR).	08
	(b)	Write a note on validation of integrated lines by media fill test.	08
Q.7	(a)	Tabulate USP validation requirements for each of four categories of analytical procedures. Why and how is robustness and Reproducibility of assay method determined?	08
	(b)	Describe validation of UV-VIS spectrophotometer.	08
