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Roll No. Tota	I No. of Pages : 01
Total No. of Questions : 06	
M.Pharmacy (Pharmaceutical Quality Assurance) (2 (Sem1)	(017 & Onwards)
PRODUCT DEVELOPMENT & TECHNOLOGY Subject Code: MQA-104T M.Code: 74702	TRANSFER
Time: 3 Hrs.	Max. Marks: 75
INSTRUCTIONS TO CANDIDATES: 1. Attempt any FIVE questions out of SIX questions. 2. Each question carries FIFTEEN marks.	
Q1. Define Qualitative and quantitative technology models.	(15)
Q2. Describe manufacturing techniques of semisolids and parenteral	dosage form. (15)
Q3. Explain the role of documentation in technology transfer.	(15)
Q4. Define:	
 a) Product registration guidelines according to USFDA. 	(8)
b) SUPAC.	(7)
Q5. Define Pharmaceutical Dosage from and their packing requirement	ents? (15)
Q6. Write note on :	
 a) Pre-formulation protocol. 	(7)
 b) Stability testing during product development. 	(8)

NOTE: Disclosure of Identity by writing Mobile No. or Making of passing request on any page of Answer Sheet will lead to UMC against the Student.

1 M-74702 (S31)-1427

