

Seat No.: _____

Enrolment No. _____

GUJARAT TECHNOLOGICAL UNIVERSITY

M. Pharm. IST Semester Examination – June- 2011

Subject code: 910102

Subject Name: Pharmaceutical Formulation Development & Biopharmaceutics

Date: 22/06/2011

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)	Frame the preformulation study for preparation of oral tablets.	06
	(b)	Give the details of dissolution equipments in pharmacy.	05
	(c)	Give the Importance of half life in stability study.	05
Q.2	(a)	Give the importance of particle size and particle size distribution in preformulation study.	06
	(b)	Comment on: accelerated stability study is only temperature affected stability study.	05
	(c)	Give the methods for studying bioavailability.	05
Q.3	(a)	Why OVIs are considered in preformulation study?	06
	(b)	How the stability of formulation is lost by environmental factors?	05
	(c)	Give the importance of Vd, Cl in pharmacokinetic.	05
Q.4	(a)	Techniques for improvement of solubility of poorly soluble drugs.	06
	(b)	Give the regulatory aspects related to stability study.	05
	(c)	Give the importance and classification of compartment models.	05
Q.5	(a)	Define solubility terms and give their applications.	06
	(b)	How the salt form of the drug affected the drug absorption?	05
	(c)	Detail the methods for establishing IVIVC.	05
Q. 6	(a)	How the dissolution media and conditions are selected in dissolution study?	06
	(b)	How the gastric emptying time affected the drug absorption?	05
	(c)	Give the factors affecting IVIVC.	05
Q.7	(a)	Give the importance of BCS in dosage form development.	06
	(b)	Importance of CACO2 monolayer cell line for drug absorption.	05
	(c)	Give the formulation aspect of herbal dental powder in detail.	05
