

Seat No.: \_\_\_\_\_

Enrolment No. \_\_\_\_\_

**GUJARAT TECHNOLOGICAL UNIVERSITY**  
**B.PHARM - SEMESTER- 8 EXAMINATION – SUMMER -2019**

**Subject Code: 2280001****Date: 04-05-2019****Subject Name: Dosage form Design II****Time: 10:30 AM TO 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.

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|-------------|-----|--|-----------|
| <b>Q.1</b>  | (a) | Explain biological factors affecting the design of oral sustained release drug delivery system.                          | <b>06</b> |
|             | (b) | Write note on Bio erodible controlled drug delivery system.  | <b>05</b> |
|             | (c) | Discuss various causes for non linearity of drug   | <b>05</b> |
| <b>Q.2</b>  | (a) | Explain effect of porosity and tortuosity on controlled drug delivery system.  | <b>06</b> |
|             | (b) | Explain “Drug interaction” Discuss ADME drug interaction with suitable example   | <b>05</b> |
|             | (c) | Define clinical pharmacokinetics. Explain methods for the calculation of creatinine from serum creatinine concentration. | <b>05</b> |
| <b>Q.3</b>  | (a) | Explain dosage adjustment in patients with renal and hepatic failure.  | <b>06</b> |
|             | (b) | Discuss in brief about PULSINCAP technology  | <b>05</b> |
|             | (c) | Write merits of non compartmental analysis. Explain AUC & AUMC plots.  | <b>05</b> |
| <b>Q.4</b>  | (a) | Give criteria for obtaining valid urinary excretion method.  | <b>06</b> |
|             | (b) | Write a note on hydrogel   | <b>05</b> |
|             | (c) | Explain the method of residuals for the calculation of absorption rate constant from oral data.                          | <b>05</b> |
| <b>Q.5</b>  | (a) | Discuss the formulation of parenteral emulsion and suspensions.  | <b>06</b> |
|             | (b) | Discuss extraction ratio and hepatic clearance in detail.  | <b>05</b> |
|             | (c) | Explain michaelis menten equation.   | <b>05</b> |
| <b>Q. 6</b> | (a) | Explain non erodible and erodible ocular control release system.   | <b>06</b> |
|             | (b) | Write note on matrix tablets.  | <b>05</b> |
|             | (c) | Write fundamental and rational of modified release drug delivery system.   | <b>05</b> |
| <b>Q.7</b>  | (a) | Give on account of approaches for designing of gasro retentive dosage form.  | <b>06</b> |
|             | (b) | Give advantages and disadvantages of compartment modeling.   | <b>05</b> |
|             | (c) | Describe floating drug delivery system.  | <b>05</b> |

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