

**GUJARAT TECHNOLOGICAL UNIVERSITY**  
**B.PHARM SEMESTER VIII • EXAMINATION – WINTER-2017**

**Subject Code: 2280001**

**Date: 02/11/2017**

**Subject Name: Dosage Form Design-II**

**Time: 02:30 pm to 5:30 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) Enumerate the factors affecting the designing of oral sustained release drug delivery systems and explain any one in detail. | <b>06</b> |
|             | (b) Discuss the merits and demerits of controlled release formulation  | <b>05</b> |
|             | (c) What are loading and maintenance dose? How are they calculated?  | <b>05</b> |
| <b>Q.2</b>  | (a) Write a note on dissolution and diffusion controlled release system  | <b>06</b> |
|             | (b) Describe the key components of osmotic drug delivery system with examples.   | <b>05</b> |
|             | (c) Write a note on colon targeted drug delivery system.   | <b>05</b> |
| <b>Q.3</b>  | (a) Discuss the various formulation approaches for floating drug delivery systems.   | <b>06</b> |
|             | (b) Write about In-vitro-In-vivo evaluations of floating drug delivery systems.  | <b>05</b> |
|             | (c) Explain pH Sensitive and Prodrug approach to develop colonic Drug Delivery Systems in brief.                                 | <b>05</b> |
| <b>Q.4</b>  | (a) Write in brief about OROS-CT and EOP.  | <b>06</b> |
|             | (b) Discuss in detail about PULSINCAP Technology.  | <b>05</b> |
|             | (c) Write a note on: Microspheres.   | <b>05</b> |
| <b>Q.5</b>  | (a) Describe formulation and evaluation of transdermal drug delivery system.   | <b>06</b> |
|             | (b) Define liposomes and niosomes. Describe their evaluation.  | <b>05</b> |
|             | (c) Write a note on: Hydrogel.   | <b>05</b> |
| <b>Q. 6</b> | (a) What are pharmacokinetic models? Explain in detail one compartment model.  | <b>06</b> |
|             | (b) Describe the method of residuals for determination of absorption rate constant.  | <b>05</b> |
|             | (c) Explain Wagner nelson method in detail.  | <b>05</b> |
| <b>Q.7</b>  | (a) Explain how one can detect nonlinear pharmacokinetics? Explain Michaelis Menten equation for capacity limited process.       | <b>06</b> |
|             | (b) Define clinical pharmacokinetics. Explain dosage adjustment in patients with renal failure.                                  | <b>05</b> |
|             | (c) Explain term: Drug interaction. Discuss ADME drug interactions with suitable examples.                                       | <b>05</b> |

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