Seat No.:	Enrolment No.

GUJARAT TECHNOLOGICAL UNIVERSITY M.PHARM – SEMESTER -1 - EXAMINATION – WINTER - 2018

Date: 05/01/2019 Subject Code: MPC103T **Subject Name: Advanced Medicinal Chemistry** 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. Enumerate stages of drug discovery and explain methods of lead discovery. 06 **Q.1** (a) Explain the theories involved in drug receptor interaction. **(b)** 05 Why we required lead optimization? Explain methods involved in lead 05 (c) optimization. **Q.2** Add a note on rational behind prodrug design. **06** (a) Explain strategies to combat drug resistance in antibiotics therapy. 05 **(b)** Why is cancer so difficult to cure? Explain with drug resistance. 05 (c) **Q.3** Why analog design important in drug design? Explain any one strategy 06 (a) involved in analog design. **(b)** Write recent advancement in anticancer agents. 05 Why stereochemistry is important in drug action? 05 (c) 0.4 Explain the strategies involved in development antiviral agents. 06 (a) Explain enzyme kinetics & principles of enzyme inhibitors. 05 **(b)** Add a note on enzyme inhibitors in basic research. 05 (c) 0.5 Explain design of peptidomimetics. 06 (a) Explain chemistry of prostaglandins, leukotrienes and thromboxanes. 05 **(b)** Add a note on identification, validation and diversity of drug targets. (c) 05 Q. 6 (a) Explain latest development of antihypertensive medication. 06 Add a note on carrier linked prodrugs. **(b)** 05 Explain simplification approach in lead optimization (c) 05 Explain non-covalently and covalently binding enzyme inhibitors. **Q.7** 06 (a) Add a note on COX-1 & COX-2 inhibitors. **(b)** 05 Explain H1 & H2 receptor antagonists. (c) 05
