


<b>Name:</b> <b>Enrolment No:</b>			
<b>UPES</b> <b>End Semester Examination, May 2025</b>			
<b>Course: Medicinal Chemistry</b> <b>Program: MSc Chemistry</b> <b>Course Code: CHEM7055</b>		<b>Semester: II</b> <b>Time: 03 hrs.</b> <b>Max. Marks: 100</b>	
<b>Instructions:</b> <ol style="list-style-type: none"> <li>1. Read all the questions carefully and attempt questions of one section in one place.</li> <li>2. Question 9 in Section B and Question 11 in Section C have internal choice questions.</li> </ol>			
<b>SECTION A</b> <b>(5Qx4M=20Marks)</b>			
S. No.		Marks	CO
Q1	Define the bioavailability of a drug. What are the different factors that affect the bioavailability of an orally administered drug?	4	CO1
Q2	Explain the following. a. Pharmacodynamics of drug action b. Advantages of <i>in vitro</i> testing over <i>in vivo</i> testing.	4	CO2
Q3	Differentiate between a soft drug and a prodrug with suitable examples. Which of these is recommended for achieving a burst drug release?	4	CO2
Q4	Discuss drug pharmacokinetics and the various factors that determine the drug metabolism.	4	CO3
Q5	Justify the statement “Molecular docking may not provide a precise information on ligand behavior within the active site of target protein”.	4	CO1
<b>SECTION B</b> <b>(4Qx10M= 40 Marks)</b>			
Q6	Write a note on the significance of Partitioning Coefficient in medicinal chemistry. How will you determine the partitioning coefficient for a drug experimentally ?	10	CO3
Q7	Discuss competitive inhibitors and active sites directed irreversible inhibition of enzyme. How will you experimentally verify the competitive inhibition of an enzyme by a drug?	10	CO2
Q8	How will you distinguish a lead compound from a hit molecule ? Discuss the steps involved in a typical drug discovery paradigm.	10	CO2
Q9	State the importance of ‘Lipinski’s rule of five’ in medicinal chemistry. How will you determine the key physicochemical parameters associated with this rule? Give examples.	10	CO1
<b>OR</b>			

	What are drug receptors? Categorize various types of drug receptors based on the nature of drug-receptor interactions.		
<b>SECTION-C</b> <b>(2Qx20M=40 Marks)</b>			
Q10	Write a short note on the following. a. Concept of 'Bioisosterism' with examples. b. Computer programs for molecular modeling. c. Routes of drug administration and their importance.	<b>20</b>	<b>CO2</b>
Q11	Categorize anticancer drugs based on their mode of action. Also draw the structures of any one commercialized drug molecules under each class of anticancer drugs.  <div style="text-align: center;"><b>OR</b></div> Explain the drug metabolism process, covering Phase I and Phase II metabolism. Discuss their role in the pharmacological action of drugs.	<b>20</b>	<b>CO3</b>