


Name:			
Enrolment No:			
<b>UPES</b> <b>End Semester Examination, May 2024</b>			
<b>Course: Medicinal Chemistry</b> <b>Program: MSc Hons 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 Chave internal choice questions.</li> <li>3. Use of Calculator is allowed.</li> </ol>			
<b>SECTION A</b> <b>(5Qx4M=20Marks)</b>			
S. No.		Marks	CO
Q1	Discuss in brief Phase I and Phase II metabolism of a drug.	4	CO1
Q2	Explain the role of the following in a pharmaceutical industry: <ol style="list-style-type: none"> <li>a. Process development.</li> <li>b. R&amp;D</li> <li>c. Quality assurance.</li> </ol>	4	CO2
Q3	What are the different routes of administration of a drug? Which is the best route considering patient compliance?	4	CO2
Q4	Justify the statement “Enzyme stimulation manifests chronic diseases” in context of inflammation and related disorders.	4	CO3
Q5	How will you differentiate the following from Lineweaver Burk Plots: <ol style="list-style-type: none"> <li>a. Competitive inhibition.</li> <li>b. Non-competitive inhibition</li> </ol>	4	CO1
<b>SECTION B</b> <b>(4Qx10M=40Marks)</b>			
Q6	Write a note on the role of Biotransformation of xenobiotics in contemporary drug discovery process.	10	CO3
Q7	Explain the following: <ol style="list-style-type: none"> <li>(i) Receptor Theory and Drug Behavior</li> <li>(ii) Types of Reversible Enzyme Inhibitors.</li> </ol>	10	CO2
Q8	Define an LD50 dose. How it is different from the IC50 and EC50 values? Explain with suitable examples.	10	CO2

Q9	What is the role of compartmental models in the quantitative evaluation of drug pharmacokinetics? Explain with examples in context of a three-compartment model.	10	CO3
	<b>OR</b>		
	Under what conditions is the 'Lipinski's Rule of 5' relaxed in a drug discovery process? Discuss in detail.	10	CO3
<b>SECTION C</b> <b>(2Qx20M=40Marks)</b>			
Q10	Schematically discuss the steps involved in a typical drug discovery process starting from the fragment-based analysis to the lead optimization. As a medicinal chemist, how will you determine the precision of the results obtained from the various <i>in silico</i> analysis during this process.	20	CO2
Q11	Write a detailed account on the various Theories of Drug Action based on the Drug-Receptor interactions.	20	CO2
	<b>OR</b>		
	Write the synthesis and uses of the following drugs: a. Salbutamol b. Penicillin c. Melphalan d. Acetaminophen	20	CO2