


<b>Name:</b>			
<b>Enrolment No:</b>			
<b>UPES</b>			
<b>End Semester Examination, December- 2024</b>			
<b>Course:</b> Medicinal Chemistry		<b>Semester:</b> VII	
<b>Program:</b> B.Tech (Biotechnology)		<b>Duration:</b> 3 Hours	
<b>Course Code:</b> HSBT4004		<b>Max. Marks:</b> 100	
<b>Instructions: Read all the questions carefully.</b>			
<b>S. No.</b>	<b>Section A</b>	<b>Marks</b>	<b>COs</b>
	<b>Short answer questions/ MCQ/T&amp;F</b> <b>(20Qx1.5M= 30 Marks)</b>		
<b>Q1.</b>	Define Bioisoster with an example.	1.5	CO1
<b>Q2.</b>	Carbachol differs from acetylcholine by..... a) Ester b) Amide c) Chloro group d) Hydroxyl group	1.5	CO1
<b>Q3.</b>	Generally, drugs are absorbed in which form? a) In ionized form b) In unionized form c) In both of above form d) In none of above form	1.5	CO1
<b>Q4.</b>	.....is the NSAIDs drug, which anthranilic acid derivative. a) Mefenamic acid b) Ibuprofen c) Piroxicam d) Zomepirac	1.5	CO1
<b>Q5.</b>	Choose the basic nucleus present in the sympathomimetic agents. a) Catechol nucleus b) Benzyl nucleus c) Naphthol d) Indole	1.5	CO1
<b>Q6.</b>	Draw the structure of Norepinephrine.	1.5	CO1
<b>Q7.</b>	Ultra-short-acting Barbiturates. a) Phenobarbitone b) Butobarbitone c) Pentobarbitone d) Thiopentone	1.5	CO1
<b>Q8.</b>	Enlist Phase-I reactions.	1.5	CO1
<b>Q9.</b>	The most significant protein involved in binding with drug is..... a) Albumin	1.5	CO2

	b) Glycoprotein c) Lipoprotein d) Globulin		
<b>Q10.</b>	Replacement of oxygen at C-2 position of barbituric acid by a sulfur atom ..... a) Has no change in the activity b) Increases the activity c) Decreases the activity d) Show anxiolytic activity	1.5	CO2
<b>Q11.</b>	Write the structure of Phenytoin.	1.5	CO3
<b>Q12.</b>	Write the structure of carbachol.	1.5	CO3
<b>Q13.</b>	Draw the structure of Aspirin.	1.5	CO3
<b>Q14.</b>	Pilocarpine is used for ..... a) Gout b) Glaucoma c) Urinary retention d) Infection	1.5	CO3
<b>Q15.</b>	State the examples of Narcotic antagonist.	1.5	CO4
<b>Q16.</b>	Which of the following is precursor of adrenaline synthesis.....? a) Phenylalanine b) Tyrosine c) Tryptophan d) None of the above	1.5	CO4
<b>Q17.</b>	The choline ester resistant to both true and pseudo- cholinesterase is..... a) Bethanechol b) Carbachol c) Methacholine d) Benzoylcholine	1.5	CO4
<b>Q18.</b>	Introduction of methyl group at alpha ( $\alpha$ ) position of acetylcholine forms acetyl- $\alpha$ -methyl choline which has more selectivity towards..... a) Nicotinic receptor b) Muscarinic receptor c) Both d) None of the above	1.5	CO4
<b>Q19.</b>	Dopamine is biosynthesized from..... a) L-Alanine b) L-Tyrosine c) L-Phenylalanine d) L-DOPA	1.5	CO5
<b>Q20.</b>	Choose the basic nucleus present in the Diazepam. a) Catechol nucleus b) Benzyl nucleus c) Benzodiazepine nucleus d) Tyrosine nucleus	1.5	CO5
<b>Section B</b>			

<b>(4Qx5M=20 Marks)</b>			
		<b>5</b>	
<b>Q1.</b>	What are the different metabolic pathways? Explain the Phase I and Phase II reactions.	<b>(2+3)</b>	CO1
<b>Q2.</b>	Discuss SAR and Classification of Morphine Analogs.	<b>5</b>	CO3
<b>Q3.</b>	Classify barbiturates with examples based on duration of action. Discuss in detail Mechanism of action.	<b>(2+3)</b>	CO4
<b>Q4.</b>	Discuss in detail SAR of Benzodiazepines.	<b>5</b>	CO4
<b>Section C (2Qx15M=30 Marks)</b>			
		<b>15</b>	
<b>Q1.</b>	Classify cholinergic receptors, Explain the catabolism of acetyl choline and explain the SAR of direct acting para-symphathomimetic agent.	<b>(3+4+4+4)</b>	CO2
<b>Q2.</b>	Give Chemical Classification of NSAIDS and Explain its Mechanism of Action.	<b>(3+4+4+4)</b>	CO4
<b>Section D (2Qx10M=20 Marks)</b>			
		<b>10</b>	
<b>Q1.</b>	Give the biosynthesis and metabolism of nor-adrenaline. Define, classify and write the SAR of adrenergic agents.	<b>(6+4)</b>	CO3
<b>Q2.</b>	Define sedative and hypnotics. Classify them and explain the SAR of barbiturates.	<b>(3+3+4)</b>	CO5