


Name:			
Enrolment No:			
<b>UPES</b> <b>End Semester Examination, May 2025</b>			
<b>Course: Pharmacology</b> <b>Program: BT-Biotechnology</b> <b>Course Code: HSFN3008P</b>		<b>Semester: VI</b> <b>Time: 03 hrs.</b> <b>Max. Marks: 100</b>	
<b>Instructions: Read the question paper carefully. Attempt the questions as mentioned.</b>			
<b>S. No.</b>	<b>Section A</b>  <b>Short answer questions/ MCQ/T&amp;F</b> <b>(20Qx1.5M= 30 Marks)</b>	<b>Marks</b>	<b>COs</b>
<b>Q 1</b>	Write two aspects of pharmacology.	<b>1.5</b>	<b>CO1</b>
<b>Q 2</b>	Define ED <sub>50</sub> .	<b>1.5</b>	<b>CO2</b>
<b>Q 3</b>	Give application of plasma protein binding.	<b>1.5</b>	<b>CO2</b>
<b>Q 4</b>	Arrange the following terms in correct order – Absorption, metabolism, excretion, and distribution.	<b>1.5</b>	<b>CO2</b>
<b>Q 5</b>	Define drug action and drug effects. A. Drug action is the observable result; drug effect is molecular interaction B. Drug action is molecular interaction; drug effect is the observable result C. Both refer to side effects only D. Drug action is long-term effect; drug effect is short-term only	<b>1.5</b>	<b>CO1</b>
<b>Q 6</b>	Mention one example of a drug with a narrow therapeutic index. A. Paracetamol B. Digoxin C. Ibuprofen D. Ranitidine	<b>1.5</b>	<b>CO1</b>
<b>Q 7</b>	Define half-life (t <sub>1/2</sub> ) of a drug. A. Time required for complete elimination of the drug B. Time taken to reach maximum effect C. Time required for plasma concentration to reduce by 50% D. Time taken for drug absorption	<b>1.5</b>	<b>CO1</b>
<b>Q 8</b>	State one function of the cytochrome P450 enzyme system. A. Enhancing drug absorption B. Inhibiting drug binding C. Metabolizing drugs in the liver D. Facilitating drug excretion by kidneys	<b>1.5</b>	<b>CO1</b>

<b>Q 9</b>	Define loading dose. A. The amount of drug excreted in one hour B. The dose given to maintain drug levels C. A higher initial dose to rapidly reach therapeutic concentration D. The amount of drug required to cause toxicity	<b>1.5</b>	<b>CO2</b>
<b>Q 10</b>	Define receptor. A. A blood protein that binds drugs B. A site where a drug binds to produce an effect C. A hormone that carries signals D. An enzyme that breaks down drugs	<b>1.5</b>	<b>CO2</b>
<b>Q 11</b>	Name any one factor that affects drug absorption.	<b>1.5</b>	<b>CO2</b>
<b>Q 12</b>	Write general characteristics of sympathetic nervous system.	<b>1.5</b>	<b>CO2</b>
<b>Q 13</b>	Parasympathetic nervous system causes increased heart rate. (True/False)	<b>1.5</b>	<b>CO2</b>
<b>Q 14</b>	Define neurotransmitters.	<b>1.5</b>	<b>CO1</b>
<b>Q 15</b>	Show the effects of sedative and hypnotic on a graph.	<b>1.5</b>	<b>CO1</b>
<b>Q 16</b>	Write steps involved in general anaesthesia.	<b>1.5</b>	<b>CO2</b>
<b>Q 17</b>	Differentiate analgesic and antipyretic.	<b>1.5</b>	<b>CO1</b>
<b>Q 18</b>	Give one example of drug use in glaucoma.	<b>1.5</b>	<b>CO1</b>
<b>Q 19</b>	Write about the effect of GABA.	<b>1.5</b>	<b>CO2</b>
<b>Q 20</b>	Abbreviate the term ADME.	<b>1.5</b>	<b>CO2</b>
<b>Section B</b> <b>(4Qx5M=20 Marks)</b>			
<b>Q 1</b>	Explain the pharmacology of antidepressant drugs.	<b>5</b>	<b>CO2</b>
<b>Q 2</b>	Classify local anaesthetics and write their mechanism of actions.	<b>5</b>	<b>CO3</b>
<b>Q 3</b>	Define drug excretion. Write phases of drug metabolism.	<b>5</b>	<b>CO2</b>
<b>Q 4</b>	Define receptor. Explain their types.	<b>5</b>	<b>CO3</b>
<b>Section C</b> <b>(2Qx15M=30 Marks)</b>			
<b>Q 1</b>	Define transduction mechanisms with example. Explain any one mechanism in detail.	<b>10+5</b>	<b>CO3</b>
<b>Q 2</b>	Discuss drug distribution and its application. Explain how drug distribution and plasma protein binding affect drug action.	<b>10+5</b>	<b>CO5</b>
<b>Section D</b> <b>(2Qx10M=20 Marks)</b>			
<b>Q 1</b>	Write a note on neurotransmitters pharmacology.	<b>10</b>	<b>CO5</b>
<b>Q 2</b>	Explain the pharmacology of antiparkinsonian drugs.	<b>10</b>	<b>CO3</b>

