



## Sessional I (Even) Semester Examination March 2025

Name of the Course and semester: B.Pharm 4<sup>th</sup> Semester

Roll no.....

Name of the Paper: Medicinal Chemistry- I

Paper Code: BP- 402 T

Time: 1.5 hours

MM: 30

Note: This question paper contains three sections.

All sections are compulsory.

### Section- A

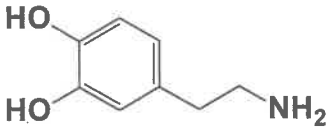
Multiple choice questions (attempt all questions)

(10 x 1=10 Marks)

<ol style="list-style-type: none"><li>The process of bioisosterism in drug design is mainly used to:<ol style="list-style-type: none"><li>Enhance the solubility of a drug in water</li><li>Modify the pharmacokinetic properties of a drug</li><li>Replace a functional group with another to optimize the drug's pharmacological activity</li><li>Increase the degree of ionization of a drug molecule</li></ol></li><li>Which of the following reactions is typically associated with Phase I drug metabolism?<ol style="list-style-type: none"><li>Conjugation with glucuronic acid</li><li>Hydrolysis</li><li>Reduction</li><li>Methylation</li></ol></li><li>What is the main characteristic of Phase II drug metabolism?<ol style="list-style-type: none"><li>Involves the addition of functional groups to the drug molecule</li><li>Includes reactions like oxidation and reduction</li><li>Conjugates drug molecules</li><li>Primarily performed by cytochrome P450 enzymes</li></ol></li><li>Which of the following factors would most likely increase the solubility of a drug in water?<ol style="list-style-type: none"><li>An increase in the molecular weight of the drug</li><li>A decrease in the ionization of the drug</li><li>A higher degree of ionization at the physiological pH</li><li>A higher partition coefficient</li></ol></li><li>What is the main effect of increasing a drug's partition coefficient (log P) value?<ol style="list-style-type: none"><li>Increased solubility in water</li><li>Increased ability to cross cell membranes</li><li>Increased ionization at physiological pH</li><li>Decreased interaction with receptors</li></ol></li></ol>	CO1
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<p>6. Which among are the precursor for the biosynthesis of catecholamine</p> <ul style="list-style-type: none"><li>a) Phenylalanine and L-tyrosine</li><li>b) Dopamine and L-tyrosine</li><li>c) Phenyl epinephrine and D-tyrosine</li><li>d) Epinephrine and D- tyrosine</li></ul> <p>7. Identify the structure</p> <div style="text-align: center;"></div> <ul style="list-style-type: none"><li>a) Adrenaline</li><li>b) Nor-Adrenaline</li><li>c) Catechol</li><li>d) Dopamine</li></ul> <p>8. Which of the following drugs is used for the treatment of asthma and acts as a selective beta-2 adrenergic agonist?</p> <ul style="list-style-type: none"><li>a) Dobutamine</li><li>b) Salbutamol</li><li>c) Epinephrine</li><li>d) Naphazoline</li></ul> <p>9. Which drug is direct acting sympathomimetic agent</p> <ul style="list-style-type: none"><li>a) Epinephrine</li><li>b) Pseudoephedrine</li><li>c) Hydroxyamphetamine</li><li>d) Propylhexedrine</li></ul> <p>10. Enzyme responsible for the termination of action catecholamine</p> <ul style="list-style-type: none"><li>a) MAO</li><li>b) COMT</li><li>c) Isomerase</li><li>d) Both a) and b)</li></ul>	<p>CO2</p>
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**Section- B**

**Short Answer type questions (attempt any two questions)**

**(2 x 5=10 Marks)**

1. Write a note on drug metabolism?	CO1
2. Explain the SAR of beta-blocker?	CO2
3. Give the synthesis and therapeutic uses of Propranolol?	CO2



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**Section- C**

**Long Answer type questions (attempt any one question)**

**(1 x 10=10 Marks)**

1. Discuss the history of Medicinal and development of Drug in detail?	CO1
2. Detail the biosynthesis of catecholamine?	CO2