

[4]

Q.4	i.	What is enzyme inhibition? Cite with example.	3	2	1	3	1
	ii.	Explain the reversible enzyme inhibition.	7	2	1	3	1
OR	iii.	Differentiate between reversible and irreversible enzyme inhibition with suitable examples.	7	2	1	3	1
Q.5	i.	What is affinity chromatography?	4	2	1	4	1
	ii.	Write down different methods used in enzyme analysis.	6	2	1	4	1
OR	iii.	Write about site-directed mutagenesis & its role in genetic/enzyme engineering.	6	2	4	7	1
Q.6	Attempt any two:						
	i.	Write down the role of enzymes in industries.	5	2	3	5	3
	ii.	How enzymes can be used in disease diagnosis?	5	2	4	5	3
	iii.	How enzymes can be used in environmental management? Write with classic examples.	5	2	3	5	1

Total No. of Questions: 6

Total No. of Printed Pages: 4

Enrolment No.....



Faculty of Science

End Sem Examination Dec 2024

BT3SE02 Enzymology

Programme: B.Sc.

Biotechnology/Specialisation:

Biotechnology

Maximum Marks: 60

Duration: 3 Hrs.

Note: All questions are compulsory. Internal choices, if any, are indicated. Answers of Q.1 (MCQs) should be written in full instead of only a, b, c or d. Assume suitable data if necessary. Notations and symbols have their usual meaning.

Marks	BL	PO	CO	PSO
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- Q.1 i. An allosteric inhibitor of an enzyme typically-
- (a) Changes the shape of the enzyme, affecting the active site.
 - (b) Binds to the active site, directly blocking substrate binding.
 - (c) Increases the affinity of the enzyme for its substrate.
 - (d) Increases the V_{max} of the enzyme reaction.
- ii. According to the transition state theory, the transition state-
- (a) Has the same free energy as the reactants.
 - (b) Is more stable than the product.
 - (c) Represents the highest energy point on the reaction pathway.
 - (d) Is a permanent intermediate.
- iii. In the general mechanism of enzyme catalysis, the enzyme primarily lowers which of the following to increase the reaction rate?
- (a) The free energy of the products.
 - (b) The free energy of the reactants.
 - (c) The equilibrium constant.
 - (d) The activation energy.

[2]

- iv. Ubiquitination as a means to regulate enzyme levels works by- **1** 2 1 2 1
- (a) Promoting enzyme assembly into multienzyme complexes.
 - (b) Marking enzymes for degradation via the proteasome pathway.
 - (c) Facilitating enzyme activation by co-factors.
 - (d) Reducing enzyme synthesis at the transcriptional level.
- v. A suicide inhibitor is distinct from other types of inhibitors because it- **1** 2 1 3 1
- (a) Acts reversibly to reduce enzyme activity.
 - (b) Competes with the substrate for the active site.
 - (c) Inhibits the enzyme only in the presence of cofactors.
 - (d) Undergoes a reaction with the enzyme, resulting in a permanently inactivated enzyme.
- vi. In the determination of the inhibition constant (K_i) for a competitive inhibitor, a smaller K_i value indicates- **1** 2 1 3 1
- (a) Weaker binding affinity between the inhibitor and enzyme.
 - (b) Stronger binding affinity between the inhibitor and enzyme.
 - (c) That the inhibitor only binds to the enzyme substrate complex.
 - (d) A decrease in V_{max} with increasing substrate concentration.
- vii. Which of the following techniques is most commonly used as an initial step in isolating enzymes from a complex cell lysate.? **1** 2 1 4 1
- (a) Affinity chromatography
 - (b) Size-exclusion chromatography.
 - (c) Ammonium sulphate precipitation.
 - (d) Reverse-phase HPLC.

[3]

- viii. When analysing enzyme activity, a spectrophotometric assay might be chosen because- **1** 2 1 4 1
- (a) It provides a direct measure of substrate concentration.
 - (b) It allows for continuous measurement of reaction rate in real-time.
 - (c) It is the most cost-effective method regardless of enzyme type.
 - (d) It requires enzymes to be immobilized on a solid phase.
- ix. In the leather industry, proteases are employed during the "bating" process to- **1** 1 5 4 1
- (a) Soften hides by breaking down non-collagenous proteins.
 - (b) Remove natural fats and oils from hides.
 - (c) Prevent microbial growth on leather surfaces.
 - (d) Dye the leather fibers.
- x. Which of the following enzymes is widely used in the pharmaceutical industry to dissolve blood clots? **1** 1 5 4 1
- | | |
|--------------|-------------------|
| (a) Protease | (b) Lipase |
| (c) Catalase | (d) Streptokinase |
- Q.2**
- i. What is the meaning of enzyme active site? **2** 2 1 1 1
 - ii. Mention different factors affecting enzyme activities **3** 2 1 1 1
 - iii. Derive Michaelis-Menten equation. **5** 2 1 1 1
- OR**
- iv. Derive lineweaver burk plot for the determination of enzyme K_m and V_{max} . **5** 2 1 1 1
- Q.3**
- i. What is nucleophilic catalysis in enzymology? **2** 2 1 2 1
 - ii. Write down the mechanism of enzyme action. **8** 2 1 2 1
- OR**
- iii. What are collision and transition state theory in enzymology? **8** 2 1 2 1

Q. No	Answer	Answer Explanation
I	A	Changes the shape of the enzyme, affecting the active site.
II	C	Represents the highest energy point on the reaction pathway.
III	D	The activation energy
IV	B	Marking enzymes for degradation via the proteasome pathway
V	D	Undergoes a reaction with the enzyme, resulting in a permanently inactivated enzyme.
VI	B	Stronger binding affinity between the inhibitor and enzyme.
VII	C	Ammonium sulphate precipitation
VIII	B	It allows for continuous measurement of reaction rate in real-time.
IX	A	Soften hides by breaking down non-collagenous proteins
X	D	Streptokinase

Answers Of Multi-choice questions

Faculty of Science

Dept. of Biotechnology

End Sem Examination Dec 2024

Enzymology BT3SE02

- Q2
 i. Enzyme active site - 2
 ii. Factors (Name) - 3
 iii. Drive MMC - 5
 iv. Lineweaver buck - 5
- Q3
 i. who nucleophile - 2
 ii. Mechanism of EA - 8
 → Intro - 3
 - explain - 5
 iii. Transition state tho = 8

- Q4
 i. Enzyme inhibition definition - 3
 ii. Reversible enyzme inhibition -
 description 3
 Diagram 3
 iii. Differential R and Ir -
2 point 2 marks
- Q5
 i. Affinity chromatography - 4
 ii. Methods - two method - 6
 3 marks for each method
 iii. Site directed mutagenesis -
 Description 3
 Diagram 3
- Q6
 i. Role of enzyme in indu - 5
 ii. Use of enyzme in disease diagno - 5
 iii. Enzyme management - 5