



## Biochemistry Unit V

Practice MCQ For Govt Pharmacist Exam, in this article we will solve, Practice MCQ on the UNIT V under the subject Biochemistry of second semester. Read following article for your reference.

[Enzymes » PHARMACAREERS](#)

### 1. Enzymes are biological catalysts that:

- (a) Increase the rate of a reaction without being consumed
- (b) Decrease the rate of a reaction
- (c) Initiate new chemical reactions
- (d) Are used up in the reaction they catalyze

### 2. Most enzymes are:

- (a) Lipids
- (b) Carbohydrates
- (c) Proteins (with some exceptions like ribozymes)
- (d) Nucleic acids

### 3. Enzymes are highly specific and act on a specific molecule called the:

- (a) Effector
- (b) Inhibitor
- (c) Substrate
- (d) Cofactor

### 4. Enzymes can be denatured by factors such as:

- (a) Extreme temperatures
- (b) Extreme pH
- (c) Heavy metals
- (d) All of the above

### 5. Competitive inhibitors bind to the:

- (a) Active site of the enzyme
- (b) Allosteric site of the enzyme

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- (c) Substrate binding site
- (d) Both (a) and (c)

**6. Enzyme names often end with the suffix “-ase” and indicate the:**

- (a) Source of the enzyme
- (b) Type of reaction catalyzed
- (c) Function of the enzyme
- (d) Substrate of the enzyme

**7. The International Union of Biochemistry and Molecular Biology (IUB) classifies enzymes based on the type of reaction they catalyze. Which class includes enzymes that transfer functional groups between molecules?**

- (a) Oxidoreductases
- (b) Transferases
- (c) Hydrolases
- (d) Lyases

**8. Each enzyme in the IUB classification system is assigned a unique Enzyme Commission (EC) number. This number consists of:**

- (a) A single digit
- (b) Two digits separated by a period
- (c) Four digits separated by periods
- (d) A letter and a number

**9. Hexokinase (EC 2.7.1.1) belongs to subclass 7 of class 2 in the IUB classification. What type of reaction does it catalyze?**

- (a) Oxidation-reduction
- (b) Transfer of a phosphoryl group
- (c) Hydrolysis of a bond
- (d) Cleavage of C-C, C-O, or C-N bonds

**10. Enzymes with similar EC numbers are likely to have similar:**

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- (a) Substrate specificities
- (b) Reaction mechanisms
- (c) Both (a) and (b)
- (d) Neither (a) nor (b)

**11. The Michaelis constant ( $K_m$ ) represents the substrate concentration at which the reaction rate reaches:**

- (a) 0%
- (b) 50%
- (c) 100%
- (d) It depends on the specific enzyme

**12. A higher  $K_m$  value indicates:**

- (a) Lower affinity of the enzyme for the substrate
- (b) Higher affinity of the enzyme for the substrate
- (c) Slower rate of the reaction
- (d) All of the above

**13. The Michaelis-Menten equation relates the reaction rate ( $v$ ) to the substrate concentration ( $[S]$ ). The Lineweaver-Burke plot is a graphical representation of this equation and allows for the determination of:**

- (a)  $K_m$  and  $V_{max}$  (maximum reaction velocity)
- (b)  $K_m$  only
- (c)  $V_{max}$  only
- (d) Neither  $K_m$  nor  $V_{max}$

**14. In a Lineweaver-Burke plot, the y-intercept is equal to:**

- (a)  $1/V_{max}$
- (b)  $K_m$
- (c)  $V_{max}$
- (d)  $-K_m$

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**15. A competitive inhibitor will:**

- (a) Increase the  $K_m$  value on a Lineweaver-Burke plot
- (b) Decrease the  $K_m$  value on a Lineweaver-Burke plot
- (c) Have no effect on the  $K_m$  value on a Lineweaver-Burke plot
- (d) It depends on the specific inhibitor

**16. Enzyme inhibitors are molecules that decrease the activity of an enzyme. They can be classified into two main types. Competitive inhibitors bind to the:**

- (a) Active site of the enzyme
- (b) Allosteric site of the enzyme
- (c) Neither, they bind a different site entirely
- (d) Substrate binding site, but not the active site

**17. Non-competitive inhibitors bind to a site other than the active site, but still cause inhibition. An example of a non-competitive inhibitor is:**

- (a) Malonate, which competes with succinate in the citric acid cycle
- (b) Allopurinol, which inhibits xanthine oxidase in gout treatment
- (c) Aspirin, which inhibits cyclooxygenase and reduces inflammation
- (d) Penicillin, which inhibits bacterial cell wall synthesis

**18. Irreversible inhibitors covalently modify the enzyme, permanently inactivating it. An example of an irreversible inhibitor is:**

- (a) Methotrexate, which inhibits folate synthesis in cancer treatment
- (b) Diuretics, which increase urine output by inhibiting water reabsorption
- (c) Statins, which lower cholesterol by inhibiting HMG-CoA reductase
- (d) Acarbose, which delays carbohydrate breakdown by inhibiting intestinal glucosidase

**19. Enzyme induction is a regulatory mechanism where the synthesis of an enzyme is increased in response to the presence of its substrate. This is commonly seen in the regulation of:**

- (a) Digestive enzymes
- (b) Detoxification enzymes
- (c) Both (a) and (b)

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(d) Neither (a) nor (b)

**20. Enzyme repression is the opposite of induction, where the synthesis of an enzyme is decreased in response to high levels of its product. This helps to maintain:**

- (a) Cellular homeostasis
- (b) Substrate availability
- (c) Energy balance
- (d) All of the above

**21. Feedback inhibition is a specific type of enzyme repression where the end product of a metabolic pathway inhibits an earlier enzyme in the pathway. This provides a mechanism for:**

- (a) Coordinated regulation of metabolic pathways
- (b) Prevention of substrate depletion
- (c) Product formation only when needed
- (d) All of the above

**22. Allosteric enzymes are enzymes that have one or more regulatory sites distinct from the active site. These regulatory sites can bind to:**

- (a) Allosteric activators, increasing enzyme activity
- (b) Allosteric inhibitors, decreasing enzyme activity
- (c) Both (a) and (b)
- (d) Neither (a) nor (b)

**23. Binding of an allosteric activator to an allosteric enzyme can induce a conformational change that:**

- (a) Increases the affinity of the enzyme for its substrate
- (b) Decreases the affinity of the enzyme for its substrate
- (c) Increases the  $V_{max}$  of the enzyme
- (d) All of the above

**24. An example of an allosteric enzyme is phosphofructokinase (PFK-1), a key regulatory enzyme in glycolysis. Citrate, a product of a later step in the pathway, acts as an allosteric:**

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- (a) Activator, indicating high ATP levels and slowing glycolysis
- (b) Inhibitor, indicating high ATP levels and slowing glycolysis
- (c) Activator, indicating low ATP levels and stimulating glycolysis
- (d) Inhibitor, indicating low ATP levels and stimulating glycolysis

**25. The regulatory properties of allosteric enzymes allow for a more:**

- (a) Flexible and responsive metabolic control
- (b) Simple and linear reaction rate
- (c) Increased enzyme production
- (d) Decreased substrate availability

**26. Enzyme replacement therapy is a treatment approach used for some genetic disorders caused by:**

- (a) Deficiencies in specific enzymes
- (b) Mutations in enzyme structure leading to reduced activity
- (c) Both (a) and (b)
- (d) Excessive enzyme activity

**27. Thrombolytic drugs like streptokinase are enzymes used to dissolve blood clots by:**

- (a) Degrading fibrin, a major component of clots
- (b) Inhibiting platelet aggregation
- (c) Reducing blood viscosity
- (d) All of the above

**28. Digestive enzymes like lactase can be used as dietary supplements to help individuals with:**

- (a) Lactose intolerance
- (b) Celiac disease
- (c) Crohn's disease
- (d) Ulcerative colitis



**29. Measuring the activity of certain enzymes in the blood can be used to diagnose diseases like:**

- (a) Liver damage (elevated ALT and AST)
- (b) Myocardial infarction (elevated cardiac troponin)
- (c) Prostate cancer (elevated PSA)
- (d) All of the above

**30. Isoenzymes are enzymes with slight variations in structure that can be found in different tissues. Measuring specific isoenzymes can help pinpoint the:**

- (a) Overall level of enzyme activity
- (b) Tissue origin of a disease process
- (c) Specific genetic mutation causing an enzyme deficiency
- (d) Effectiveness of enzyme replacement therapy

**31. Creatine kinase (CK) has multiple isoenzymes. CK-MB, primarily found in heart muscle, is elevated in the blood following a heart attack. This is an example of using isoenzymes for:**

- (a) Diagnosis of tissue-specific damage
- (b) Monitoring enzyme replacement therapy
- (c) Studying enzyme structure-function relationships
- (d) Identifying genetic polymorphisms

**32. Coenzymes are small organic molecules that function as:**

- (a) Catalytic components of enzymes
- (b) Regulatory molecules for enzyme activity
- (c) Building blocks for macromolecules
- (d) Energy carriers in cellular metabolism

**33. Coenzymes often participate in reactions by accepting or donating:**

- (a) Functional groups like phosphates or methyl groups
- (b) Electrons
- (c) Both (a) and (b)
- (d) Neither (a) nor (b)

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**34. Nicotinamide adenine dinucleotide (NAD<sup>+</sup>) is a coenzyme involved in numerous oxidation-reduction reactions. The reduced form of NAD<sup>+</sup> is:**

- (a) NADH
- (b) NADP<sup>+</sup>
- (c) NADPH
- (d) FAD

**35. Flavin adenine dinucleotide (FAD) is another important coenzyme that functions as an electron carrier in cellular respiration. The reduced form of FAD is:**

- (a) NADH
- (b) NADP<sup>+</sup>
- (c) NADPH
- (d) FADH<sub>2</sub>

**36. Coenzyme A (CoA) is a crucial coenzyme involved in:**

- (a) Fatty acid metabolism
- (b) Amino acid metabolism
- (c) Carbohydrate metabolism
- (d) All of the above

**37. Biotin is a B vitamin that acts as a coenzyme for enzymes involved in:**

- (a) Gluconeogenesis and fatty acid synthesis
- (b) Transamination reactions of amino acid metabolism
- (c) Decarboxylation reactions
- (d) Activation of fatty acids for metabolism

**38. Thiamine pyrophosphate (TPP) is a coenzyme essential for the activity of enzymes in the:**

- (a) Citric acid cycle
- (b) Pentose phosphate pathway
- (c) Electron transport chain
- (d) Urea cycle

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**39. A deficiency in vitamin B6 (pyridoxine) can lead to symptoms like:**

- (a) Peripheral neuropathy
- (b) Dermatitis
- (c) Anemia
- (d) All of the above

**40. Coenzyme deficiencies can disrupt various metabolic pathways, leading to a range of diseases. Understanding coenzyme function is crucial for:**

- (a) Development of targeted therapies for metabolic disorders
- (b) Design of personalized nutrition plans
- (c) Both (a) and (b)
- (d) Neither (a) nor (b)

#### Answers

1. Enzymes are biological catalysts that: **(a) Increase the rate of a reaction without being consumed**
2. Most enzymes are: **(c) Proteins (with some exceptions like ribozymes)**
3. Enzymes are highly specific and act on a specific molecule called the: **(c) Substrate**
4. Enzymes can be denatured by factors such as: **(d) All of the above**
5. Competitive inhibitors bind to the: **(a) Active site of the enzyme**
6. Enzyme names often end with the suffix “-ase” and indicate the: **(b) Type of reaction catalyzed**
7. The International Union of Biochemistry and Molecular Biology (IUB) classifies enzymes based on the type of reaction they catalyze. Which class includes enzymes that transfer functional groups between molecules? **(b) Transferases**
8. Each enzyme in the IUB classification system is assigned a unique Enzyme Commission (EC) number. This number consists of: **(c) Four digits separated by periods**
9. Hexokinase (EC 2.7.1.1) belongs to subclass 7 of class 2 in the IUB classification. What type of reaction does it catalyze? **(b) Transfer of a phosphoryl group**
10. Enzymes with similar EC numbers are likely to have similar: **(c) Both (a) and (b)**
11. The Michaelis constant ( $K_m$ ) represents the substrate concentration at which the reaction rate reaches: **(b) 50%**
12. A higher  $K_m$  value indicates: **(a) Lower affinity of the enzyme for the substrate**
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