

### **ENZYMES**

1.	The catalytic activity of an enzyme is restricted to its small portion call									
	(A)	Intermediate	<b>(B)</b>	Above all						
	(C)	Passive site	<b>(D)</b>	Active site						
2.	An a	ctivated enzyme made of poly	peptio	de chain and a co-factor is:						
	(A)	Co-enzyme	<b>(B)</b>	Substrate						
	<b>(C)</b>	Apoenzyme	<b>(D)</b>	Holoenzyme						
3.	The	rate of reaction of enzyme dire	ectly o	depends upon:						
	(A)	Maximum pH level								
	(B)	Amount of enzyme present at a specific time at unlimited sub concentration								
	(C)	Nature of substrate								
	<b>(D)</b>	Low temperature								
4.	Kosl	nland in 1959 proposed the mo	dified	l form of:						
	(A)	Unit membrane model	<b>(B)</b>	Reflective index model						
	(C)	Fluid mosaic model	(D)	Induce fit model						
5.	The	reversible inhibitors usually co	onstit	ute:						
	(A)	No linkage with enzyme	<b>(B)</b>	Weak linkage with enzyme						
	<b>(C)</b>	Medium linkage	<b>(D)</b>	Strong linkage with enzyme						
6.	The	detachable co-factor of an enz	yme (	if it is an inorganic ion) is called as:						
	(A)	Activator	<b>(B)</b>	Catalytic						
	<b>(C)</b>	Aqueous medium	<b>(D)</b>	Apoenzyme						

7.	If the non-protein part is covalently bonded to the protein part of an oit is called as:			
	(A)	Prosthetic group	<b>(B)</b>	Catalytic
	(C)	Activator	<b>(D)</b>	Optimum
8.	An o	enzyme with its coenzyme or p	rosth	etic group, removed is called as:
	(A)	Apoenzyme	<b>(B)</b>	Aqueous medium
	(C)	Activator	<b>(D)</b>	Prosthetic group
9.	The	active site of an enzyme is con	npose	d of binding site and:
	(A)	Apoenzyme	<b>(B)</b>	Catalytic site
	(C)	Prosthetic site	(D)	Substrate site
10.	Med	lium required for Enzymes vig	gorou	s activity:
	(A)	Colloidal	<b>(B)</b>	Transparent
	(C)	Aqueous	(D)	Gel
11.	Eve	ry enzyme functions effectively	at:	
	(A)	9.00 pH	<b>(B)</b>	Optimum pH
	(C)	2.00 pH	<b>(D)</b>	7.00 pH
12.	Alm	ost all enzymes are:		
	(A)	Fibrous proteins	<b>(B)</b>	Globular proteins
	(C)	Triangular proteins	<b>(D)</b>	All of the above
13.	The	optimum pH value for pancre	atic l	ipase is:
	(A)	8.00	<b>(B)</b>	10.00
	(C)	7.00	<b>(D)</b>	9.00
14.	The	enzymes involved in the cellul	ar re	spiration in eukaryotes are found in:
	(A)	Chloroplast	<b>(B)</b>	Aqueous medium
	(C)	Mitochondria	<b>(D)</b>	Nucleoplasm
15.	Eve	n traces of enzymes can bring	abou	change in large amount of:
	(A)	Catalytic	<b>(B)</b>	Activator
	(C)	Substrate	<b>(D)</b>	Optimum
16.	Co-c	enzymes:		
	(A)	Globular protein	<b>(B)</b>	Oral cavity
	(C)	Vitamin	<b>(D)</b>	Co-factors

17.	Sali	vary amylase:		
	(A)	Oral cavity	<b>(B)</b>	Co-factors
	(C)	Vitamin	<b>(D)</b>	Globular protein
18.	Met	al ions:		
	(A)	Vitamin	<b>(B)</b>	Oral cavity
	<b>(C)</b>	Globular protein	<b>(D)</b>	Co-factors
19.	Ami	ino acids:		
	(A)	Globular protein	<b>(B)</b>	Oral cavity
	<b>(C)</b>	Chloroplast	<b>(D)</b>	Vitamin
20.	Pho	tosynthesis:		
	(A)	Oral cavity	<b>(B)</b>	Vitamin
	(C)	Globular protein	<b>(D)</b>	Chloroplast
21.	Emi	l Fischer:		
	(A)	Cyanide	<b>(B)</b>	Lock and key model
	(C)	Temperature	<b>(D)</b>	Specific in action
22.	Inhi	bitors:		
	(A)	Temperature	<b>(B)</b>	Cyanide
	(C)	Lock and key model	<b>(D)</b>	Specific in action
23.	Acti	vation energy:		
	(A)	Temperature	<b>(B)</b>	Cyanide
	<b>(C)</b>	Lock and key model	(D)	Specific in action
24.	Enz	ymes:		
	(A)	Temperature	<b>(B)</b>	Specific in action
	(C)	Reversible inhibitors	<b>(D)</b>	Cyanide
25.	Con	npetitive:		
	(A)	Reversible inhibitors	<b>(B)</b>	Lock and key model
	(C)	Specific in action	<b>(D)</b>	Temperature
26.	The	catalytic activity of an enzyme	e is re	stricted to its small portion called:
	(A)	Active site	<b>(B)</b>	Passive site
	<b>(C)</b>	Allosteric site	(D)	All Choices are correct

27.	An a	Apoenzyme  cland in 1959 proposed:  Fluid mosaic model  Lock and key model  Chical nature of enzymes is:  Lipids  Proteinacous  (B)  Steroids  (B)  Steroids  Proteinacous  (D)  All (A), (B) and (C)  proposed "lock and key" model to study enzyme-substrate interaction?  Koshland (1959)  (B)  Wilhelm Kuhne (1878)  Fischer (1898)  (D)  None of these  Iman body the optimum temperature for enzymatic activities is:  37°C  (B)  40°C  25°C  (D)  30°C				
	<b>(A)</b>	Coenzyme	<b>(B)</b>	Substrate		
	<b>(C)</b>	Apoenzyme	<b>(D)</b>	Holoenzyme		
28.	Kos	hland in 1959 proposed:				
	<b>(A)</b>	Fluid mosaic model	<b>(B)</b>	Induce fit model		
	<b>(C)</b>	Lock and key model	<b>(D)</b>	Reflective index model		
29.	Cher	nical nature of enzymes is:				
	<b>(A)</b>	Lipids	<b>(B)</b>	Steroids		
	<b>(C)</b>	Proteinacous	<b>(D)</b>	All (A), (B) and (C)		
30.	Who	proposed "lock and key" mo	del to	study enzyme-substrate interaction?		
	<b>(A)</b>	Koshland (1959)	<b>(B)</b>	Wilhelm Kuhne (1878)		
	(C)	Fischer (1898)	<b>(D)</b>	None of these		
31.	In h	uman body the optimum temp	eratu	re for enzymatic activities is:		
	<b>(A)</b>	37°C	<b>(B)</b>	40°C		
	<b>(C)</b>	25°C	<b>(D)</b>	30°C		
32.	Opt	imum pH value for pepsin is:				
	<b>(A)</b>	5.5	<b>(B)</b>	7.4		
	<b>(C)</b>	4.1	<b>(D)</b>	1.4		
33.	Con	npetitive inhibitors stop an enz	yme	from working by:		
	<b>(A)</b>	Changing the shape of the enzy	yme			
	<b>(B)</b>	Merging with the substrate inst	tead			
	<b>(C)</b>	Blocking the active site of the	enzyn	ne		
	<b>(D)</b>	Combining with the product of	the r	eaction		
34.	The	enzymes are sensitive to:				
	<b>(A)</b>	Changes in pH	<b>(B)</b>	Changes in temperature		
	<b>(C)</b>	Both (A) and (B)	<b>(D)</b>	None of these		
35.		yme B requires Zn <sup>2+</sup> in order zinc is best identified as:	to ca	ntalyze the conversion of substrate X.		
	<b>(A)</b>	Coenzyme	<b>(B)</b>	Activator		
	<b>(C)</b>	Substrate	<b>(D)</b>	Product		

36.	The	enzyme minus its coenzyme is	refer	red to as the:
	(A)	Iso-enzyme	<b>(B)</b>	Metalloenzyme
	<b>(C)</b>	Apoenzyme	<b>(D)</b>	All of these
37.		"lock and key" model of endemodecule:	nzym	e action illustrates that a particular
	(A)	Forms a permanent enzyme-su	bstrat	e complex
	<b>(B)</b>	May be destroyed and resynthe	esized	several times
	(C)	Interacts with a specific type o	f subs	trate molecule
	(D)	Reacts at identical rates under	all co	nditions
38.	Con	sider this reaction. A + B ——	<b>→</b>	C + D + energy.
	(A)	This reaction is exergonic		
	<b>(B)</b>	An enzyme could still speed th	ie reac	etion
	(C)	A and B are reactants; C and D	are p	products
	<b>(D)</b>	All of these are correct		
39.		inhibitor that changes the over wn as:	erall	shape and chemistry of an enzyme is
	(A)	Auto-steric inhibitor	(B)	Competitive inhibitor
	(C)	Steric inhibitor	<b>(D)</b>	Noncompetitive inhibitor
40.	Non	-protein components of enzym	ies ar	e known as:
	(A)	Coenzvmes	<b>(B)</b>	Activators
	(C)	Cofactors	<b>(D)</b>	All (A), (B) and (C)
41.		reaction below occurs within rogen peroxide. In this reaction		cells to prevent the accumulation of alase functions as an:
		2H <sub>2</sub> O <sub>2</sub> Catal	ase → 2	$2H_2O + O_2$
	(A)	Enzyme in the breakdown of h	ydrog	en peroxide
	<b>(B)</b>	Enzyme in the synthesis of hyd	drogei	n peroxide
	(C)	Emulsifier in the digestion of h	nydrog	gen peroxide
	(D)	Indicator in the detection of hy	droge	en peroxide

42.		enzyme is generally named by	ng ——— to the end of the name	
		"-ase". coenzyme	(B)	"-ase". cell in which it is found
		"-ose". substrate	(D)	"-ase". substrate
43.	The	minimum amount of energy n	eeded	I for a process to occur is called the:
	(A)	Minimal energy theory		Process energy
	(C)	Kinetic energy	<b>(D)</b>	Activation energy
44.				est the efficiency of a certain enzyme. hange in the enzyme's efficiency?
	<b>(A)</b>	Adding an acidic solution to the	ie setu	up .
	<b>(B)</b>	Adding more substrate but not	enzyı	me
	<b>(C)</b>	Increasing temperature of solu	tion	
	<b>(D)</b>	All (A), (B) and (C) change en	zyme	's efficiency
45.	Enz	ymes function as:		
	<b>(A)</b>	Organic catalysts	<b>(B)</b>	Inorganic catalysts
	<b>(C)</b>	Inhibitors	<b>(D)</b>	All of these
46.	A ca	talyst is a chemical involved in,	but n	ot — by, a chemical reaction.
	<b>(A)</b>	Supported	<b>(B)</b>	Changed
	<b>(C)</b>	Controlled	<b>(D)</b>	All of these
47.	Mar	ny enzymes function by ———		the activation energy of reactions.
	<b>(A)</b>	Increasing	<b>(B)</b>	Promoting
	<b>(C)</b>	Lowering	<b>(D)</b>	Both (A) and (B)
48.	Anι	uncatalysed reaction requires	a:	
	(A)	Higher activation energy	<b>(B)</b>	Lower activation energy
	<b>(C)</b>	Balanced activation energy	<b>(D)</b>	All of these
49.	The	binding of the substrate to the	enzyı	me alters the structure of the enzyme:
	<b>(A)</b>	Lock and key hypothesis	<b>(B)</b>	Induced fit hypothesis
	(C)	Fischer's hypothesis	<b>(D)</b>	D.D. Wood's hypothesis
50.	The	y are non-protein organic mole	cules	bound to enzymes near the active site:
	<b>(A)</b>	Activators	<b>(B)</b>	Coenzymes
	(C)	Holoenzymes	<b>(D)</b>	All of these

51.	The first step in any reaction catalysed by an enzyme is the formation specific association between the molecules called:				
	(A)	Enzyme-product complex	<b>(B)</b>	Enzyme-intermediate complex	
	(C)	Enzyme-substrate complex	(D)	None of these	
52.	The bind	_	tors i	s defined by their ability to interact or	
	(A)	The active site of an enzyme	<b>(B)</b>	Regulatory sub-units of an enzyme	
	<b>(C)</b>	Non-competitive inhibitor	<b>(D)</b>	Enzyme cofactors	
53.		n enzyme solution is saturated iin an even faster yield of prod		n substrate, the most effective way to would be:	
	(A)	Add more of the enzymes	<b>(B)</b>	Add more substrate	
	<b>(C)</b>	Add an allosteric inhibitor	(D)	Add a non-competitive inhibitor	
54.		en the final product of a me abolic pathway it is:	tabol	ic pathway turn off the first step of	
	(A)	Positive feed back	<b>(B)</b>	Negative feed back	
	<b>(C)</b>	Competitive feed back	<b>(D)</b>	Both (A) and (C)	
55.		en the inhibitory chemical, wh Is to the enzyme other than at		oes not have to resemble the substrate, ctive site is called:	
	(A)	Noncompetitive Inhibition	<b>(B)</b>	Competitive Inhibition	
	<b>(C)</b>	Uncatalysed reaction	<b>(D)</b>	All (A), (B) and (C)	
56.	Whi	ich one is not attribute of enzy	me?		
	(A)	Specific in nature	<b>(B)</b>	Protein in chemistry	
	<b>(C)</b>	Consumed in reaction	(D)	Increases rate of reaction	
57.		ch one inactivates an enzym ve site of an enzyme?	e by	indirectly changing the shape of the	
	(A)	Non-competitive inhibitor	<b>(B)</b>	Competitive inhibitor	
	<b>(C)</b>	Coenzyme	(D)	Activator	
58.	The	enzymes are classified into:			
	(A)	Five groups	<b>(B)</b>	Three groups	
	<b>(C)</b>	Six groups	(D)	Four groups	
59.	Non	-proteinaceous part of holoena	zyme	is:	
	(A)	Prosthetic group	<b>(B)</b>	Apoenzyme	
	(C)	Tubulin	<b>(D)</b>	None of these	

60.	,	Enzymes are highly specific for a given substrate which is due to the shape of their:						
	(A)	Active site	<b>(B)</b>	Allosteric site				
	<b>(C)</b>	Non-competitive site	<b>(D)</b>	None of these				
61.	The	name enzyme was suggested in	n 187	8 by the German physiologist:				
	(A)	Wilhelm Kuhne	<b>(B)</b>	Koshland				
	(C)	Fischer	(D)	Paul Filder				
62.	Prot	teinaceous part of holoenzyme	is:					
	(A)	Prosthetic group	<b>(B)</b>	Apoenzyme				
	(C)	Lecithin	<b>(D)</b>	None of these				
63.	The	"lock and key hypothesis" attem	npts to	explain the mechanism of:				
	(A)	Vacuole formation	<b>(B)</b>	Pinocytosis				
	(C)	Sharing of electrons	<b>(D)</b>	Enzyme specificity				
64.	An enzyme that hydrolyzes protein will not act upon starch. This fact is an indication that enzymes are:							
	(A)	Hydrolytic	<b>(B)</b>	Specific				
	(C)	Catalytic	<b>(D)</b>	Synthetic				
65.	The	site where enzyme catalyzed r	eactio	on takes place is called:				
	(A)	Active site	<b>(B)</b>	Allosteric site				
	(C)	Denatures site	(D)	Dead site				
66.	Wha	at is a coractor.						
	(A)	Inorganic ions	<b>(B)</b>	Organic molecules				
	(C)	Both (A) and (B)	<b>(D)</b>	None of the above				
67.	$Mg^+$	<sup>2</sup> is an inorganic activator for	the er	nzyme:				
	<b>(A)</b>	Phosophatase	<b>(B)</b>	Carbonic anhydrase				
	<b>(C)</b>	Enterokinase	<b>(D)</b>	Amylase				
68.	Zn <sup>+2</sup>	is an inorganic activator for e	enzyn	ne:				
	(A)	Carbonic anhydrase	<b>(B)</b>	Phosophatase				
	(C)	Chymotrypsin	<b>(D)</b>	Maltase				

	ctive site of an enzyme tha	i many bacteria useu
to make cell-walls?		

(A) Amphotericin

(B) Gentamicin

(C) Penicillin

- (D) Cephalosporin
- 70. DDT and Parathion are inhibitors of key enzymes in:
  - (A) Nervous system

- (B) Respiratory system
- (C) Digestive system
- (D) Circulatory system

## 71. At high temperature the rate of enzyme action decreases because the increased heat:

- (A) Changes the pH of the system
- (B) Alters the active site of the enzyme
- (C) Neutralize acids and bases in the system
- (D) Increases the concentration of enzymes
- 72. Which of the following enzymes would digest a fat?
  - (A) Sucrase

(B) Protease

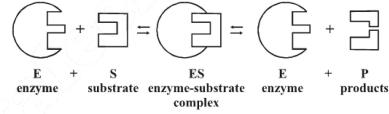
(C) Ligase

(D) Lipase

# 73. In the lock and key model of enzyme action, a part of the enzyme that recognizes the substrate is known as the:

- (A) Enzyme-substrate complex
- (B) Product
- (C) Enzyme-product comple
- **(D)** Active site

#### 74. Which model of enzyme action is represented in this diagram?



- (A) Fluid mosaic model
- (B) Induce fit model
- (C) Lock and key model
- (D) Reflective index model

## 75. A certain enzyme will hydrolyze egg white but not starch. Which statement best explains this observation?

- (A) Starch molecules are too large to be hydrolyzed
- **(B)** Enzyme molecules are specific in their actions
- (C) Egg white acts as a coenzyme for hydrolysis
- (D) Starch is composed of amino acids

76. At about 0°C, most enzymes are:

(A) Inactive

(B) Active

(C) Destroyed

(D) Replicated

77. Vitamins are essential for the survival of organisms because vitamins usually function as:

(A) Substrates

(B) Nucleic acids

(C) Co-enzymes

(D) Nucleosides

78. When a molecule binds to an area of an enzyme that is not the active site, and changes the shape of the enzyme so that it no longer can work, this is called:

(A) Denaturation

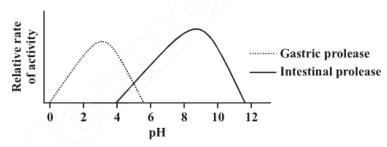
- (B) Competitive inhibition
- (C) Noncompetitive inhibition
- (D) Substrate delocation

79. What is a coenzyme?

(A) Inorganic ion

- (B) Organic molecule
- **(C)** Both (A) and (B)
- (D) None of these

80. Which statement best expresses the information represented in the graph shown?



- (A) The action of enzymes varies with pH
- **(B)** A pH of 7 provides the optimum environment for digestive enzymes
- (C) Gastric juice is active at a pH extending from 0 to 12
- (D) Acids have a pH greater than 7

81. Which type of inhibitor is shown in this diagram?



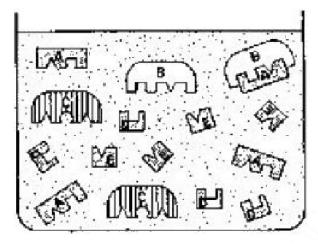
(A) Competitive

(B) Non-competitive

(C) Allosteric

**(D)** Both (B) and (C)

### 82. Which enzyme represents an enzyme functioning in this reaction?



(A) E

**(B)** C

**(C)** B

**(D)** A

Biology F.Sc. Part-I

### Answers

Sr.	Ans.								
1.	(D)	2.	(D)	3.	(B)	4.	(D)	5.	(B)
6.	(A)	7.	(A)	8.	(A)	9.	(B)	10.	(C)
11.	(B)	12.	(B)	13.	(D)	14.	(C)	15.	(C)
16.	(C)	17.	(A)	18.	(D)	19.	(A)	20.	(D)
21.	(B)	22.	(B)	23.	(A)	24.	(B)	25.	(A)
26.	(A)	27.	(D)	28.	(B)	29.	(C)	30.	(C)
31.	(A)	32.	(D)	33.	(C)	34.	(C)	35.	(B)
36.	(C)	37.	(C)	38.	(D)	39.	(D)	40.	(D)
41.	(A)	42.	(D)	43.	(D)	44.	(D)	45.	(A)
46.	(B)	47.	(C)	48.	(A)	49.	(B)	50.	(B)
51.	(C)	52.	(A)	53.	(A)	54.	(B)	55.	(A)
56.	(C)	57.	(A)	58.	(C)	59.	(A)	60.	(A)
61.	(A)	62.	(B)	63.	(D)	64.	(B)	65.	(A)
66.	(C)	67.	(A)	68.	(A)	69.	(C)	70.	(A)
71.	(B)	72.	(D)	73.	(D)	74.	(C)	75.	(B)
76.	(A)	77.	(C)	78.	(C)	79.	(B)	80.	(A)
81.	(D)	82.	(C)						