	CJ-40
Seat Number	
BP-203T	
Biochemistry	
(712203)	. 75
Total Pages: 7]	Max. Marks: 75
(Ame: 3 Hours	Sont No.
Note: (1) Do not write anything on question paper exce	ipt Seat
(2) All questions are compulsory.	
(a) Figures to right indicate full marks.	1
no supplement will be	provided.
- Wingram/cycles/pathways wherever	necessary.
should be drawn with the black ink pen or b	lack HB pencil.
	$20 \times 1 = 20$
1. Multiple Choice Questions (MCQs):	
(1) Enzymes are in nature.	
(A) Carbohydrate	
(B) Lipid	
(C) Protein	
(D) Acidic	
(2) Bile acid is synthesized in	
(A) Kidney	
(B) Liver	
(C) Intine	
(D) Stomach	

P.T.O.

(3	Which test is used for determination of ketone-bodies?
	(A) Rothera test
	(B) Shinoda test
	(C) Van Ark test
	(D) Shick test
(4)	Nucleoside is a pyrimidine or purine base :
	(A) Covalently bonded to a sugar
	(B) Ionically bonded to a sugar
	(C) Hydrogen bonded to a sugar
	(D) None of the above
(5)	Proteins are polymers of
	(A) L - α Amino Acid
	(B) L - β Amino Acid
	(C) D - α Amino Acid
	(D) D - β Amino Acid
(6)	What are the small fragments of DNA produced during replication?
	(A) Small fragments of DNA
	(B) DNA ligase
	(C) OKAZAKI fragments
0.1.46	(D) Small fragments of RNA
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(7)	Blood	urea decreases in all of the following conditions except that	n :
	(A)	Liver cirrhosis	
	(B)	Pregnancy	
	(C)	Renal Failure	
	(D)	Angina Pectoris	
(8)	Keto	one bodies are not utilized or oxidized in one of the following tiss	ues
	(A)	Heart	
	(B)	Brain	
	(C)	Liver	
	(D)	Muscles	
(9)	Ex	amples of Imino acid is :	
	(A) Arachidonic acid	
	(B) Proline	
	(C	C) Linolenic acid	
	α	O) Oleic Acid	
(10)) S	tudy of energy relationships and conversions in biological system	is is
,	t	ermed as:	
	(A) Microbiology	
		(B) Biotechnology	
		(C) Bioenergetics	
		(D) Biophysics	
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(11) One of the following is an aromatic neutral amino acid:
(A) Alanine
(B) Leucine
(C) Tyrosine
(D) Threonine
(12) The end product of oxidative phase of HMP shunt is :
(A) Ribulose-5-P
(B) Ribose-5-P
(C) Xylulose-5-P
(D) Xylose-5-P
(13) is the major site for urea synthesis in urea cycle.
(A) Mitochondria
(B) Golgi apparatus
(C) Cytosol
(D) Ribosomes
(14) Protein part of the conjugate enzyme is called:
(A) Exoenzyme
(B) Endoenzyme
(C) Apoenzyme
(D) Holoenzyme
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	(15)	Allosteric enzyme have allosteric site for	*
		(A) Activation	
		(B) Inhibition	
	• 1	(C) Moderation	
		(D) Both A and B	
	(16)	Hypercholesterolemia is observed in the di	sorder :
		(A) Hypothyroidism	
		(B) Diabetes mellitus	
		(C) Nephrotic Sydrome	
1		(D) All of the above	
	(17)	Cellular respiration is an example of:	
		(A) Endergonic Reaction	
		(B) Oxidation Reaction	
		(C) Exergonic Reaction	
ימנ		(D) None of the above	
•	(18)	During RNA Replication hydrogen bond is	breaked by :
		(A) DNA ligase	
		(B) DNA helicase	
		(C) DNA gyrase	
	CJ-40	(D) DNA polymerase	
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	(19)	Site of gluconeogenesis is:
		(A) Skeletal muscle
		(B) Kidneys
		(C) Liver
		(D) Both (B) and (C)
	(20)	The blocking of enzyme action by blocking its active site is called as:
		(A) Allosteric inhibition
		(B) Feedback inhibition
		(C) Competitive inhibition
		(D) Non-competitive inhibition
2.	Long	Answers (Answer 2 out of 3): 2×10=20
	(1)	Explain in detail glycolysis with its significances.
	(2)	Describe biosynthesis of purine and pyrimidine nucleotides.
	(3)	Explain in detail B oxidation of fatty acid.
3.	Shor	t Answers (Answer 7 out of 9) 7×5=35
	(1)	Explain in detail Citric Acid Cycle with its significances.
	(2)	Give applications of enzymes and add short note on enzyme inhibition.
	(3)	Write a note on DNA Replication.
	(4)	Elaborate Electron Transport Chain with oxidative phosphorylation.
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- (5) Write a brief note on Pentose Phosphate Pathway with its significances.
- (5) Write a brief note on 1 street on and excretion of ketone bodies.
 (6) Write a note on formation, utilization and excretion of ketone bodies.
- (6) Write a note on formation, Deamination and Decarboxylation,
 (7) Enlighten the term Transamination, Deamination and Decarboxylation,
- (8) Give therapeutic and diagnostic applications of enzymes and isoenzymes.
- (9) Explain the glycogen storage diseases (GSD).

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