## Second Semester B. Pharm. Examination

## **BIOCHEMISTRY**

Paper - BP 203 T (USC - 35341)

P. Pag	ges :	3	
Time	: Thr	ee H	ours] [Max. Marks: 75
	Note	(	1) Answer All questions. 2) Diagrams and Chemical equations should be given wherever necessary. 3) Illustrate your answer necessary with the help of neat sketches. 4) Use pen of Blue/Black ink/refill only for writing the answer book
1.	(a)	Mul	tiple choice questions :
		(i)	Which of the following is a purine?
	,		(a) cytosine (b) Adenine (c) thymine (d) uracil
		(ii)	Fatty liver is caused due to accumulation of —
			(a) fatty acids (b) phospolipids
			(c) cholesterol (d) triacylglycerol
		(iii)	Gout is a metabolic disorder of catabolism of —
			(a) pyrimidine (b) Alanine (c) purine (d) phenylalanine
		(iv)	Citric acid cycle occurs in —
			(a) cytoplasm (b) mitochondrial matrix
			(c) endoplasmic reticulum (d) Golgi bodies
		(v)	The main site of gluconeogenesis is ——
			(a) Kidney (b) Liver (c) brain (d) muscle
		(vi)	Which of the following is not disaccharides ——
			(a) Maltose (b) Starch (c) Lactose (d) Sucrose

(vii) How many different amino acids are there -
(a) 3 (b) 100 (c) 20 (d) infinite number
(viii) The enzyme of Urea synthesis are found in
(a) Mitochondria only
(b) Cytosol only
(c) Both mitochondria and cytosol
(d) nucleus.
(ix) The sugar in RNA is ——
(a) phosphate (b) ribose (c) uracil (d) deoxyribose
(x) Which of the following are reduced coenzymes?
(a) ATP and GTP (b) Coenzyme A
(c) NADH and FADH <sub>2</sub> (d) Pyridoxial phosphate 10
(b) Solve the following questions:—
(i) Define carbohydrates. Give two examples.
(ii) Enlist disorders of urea cycle.
(iii) What is obesity?
(iv) How many ATP are generated in a Glycolysis cycle?
(v) Define co - enzymes.
Long answer questions (Answer any Two) :
(a) Describe in detail classification, chemical nature and biological role of

2.

- f carbohydrate.
- (b) Describe in detail reactions involved in  $\beta$  oxidation of saturated fatty acids.
- (c) Explain in detail reactions involved in HMP shunt with its significance.

20

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- 3. Short answer questions (solve any Seven) :--
  - (a) Explain Glycogen storage disease.
  - (b) Explain catabolism of phenylalanine and tyrosine.
  - (c) Explain Electron transport chain (ETC) and its mechanism.
  - (d) Write a note on Phenylketonuria.
  - (e) Describe structure of DNA with their functions.
  - (f) Explain formation and utilization of ketone boides.
  - (g) Explain disorders of lipid metabolism.
  - (h) Discuss therapeutic and diagnostic applications of Enyzmes.
  - (i) Describe catabolism of purine nucleotides.

35

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