(16) SARDAR PATEL UNIVERSITY

T.Y.B. Sc. (Biochemistry) – Fifth Semester Examination (CBCS) Monday, 18th November 2013

10:30 a.m. to 1:30 p.m.

US05CBCH03: Metabolism - I **Total Marks: 70** Note: (1) Figures to the right indicate marks. (2) Draw a neat and labeled diagram, wherever necessary. 0.1 Choose the most appropriate answer from the four alternatives given: [10]i. Which one of the following catalyzes irreversible reaction during glycolysis? (a) Hexokinase (b) Enolase (c) Aldolase (d) Phosphotriose isomerase ii. Cori's cycle is also known as (a) Glucose alanine cycle (b) Lactic acid cycle (c) Citric acid cycle (d) TCA cycle iii. How many ATP molecules are generated per molecules of glucose under aerobic conditions in glycolysis? (b) 8 (a) 10 (c) 4(d) 2 iv. β (beta)) oxidation of fatty acid take place in _____. (a) Cytosol (b) Mitochondria (c) Endoplasmic reticulum (d) None of the above v. Cholesterol contain carbons. (a) Thrirty seven (b) Twenty seven (c) Fifty Seven (d) Seventeen vi. Which of the following catalyzes irreversible and rate limiting reaction during cholesterol biosynthesis? (a) HMG CoA reductase (b) Decarboxylase (c) Isomerase (d) HMG CoA synthase vii. Glucose 6 phosphatase enzyme deficiency causes _____ disease. (a) Cori's (b) Pompe's (c) Hers's (d) Mc Ardle's viii. Branching enzyme deficient in disease. (b) Amylopectinosis (c) Both (a) and (b) (a) Andersen's (d) Forbe's ix. Niemann Pick's disease arise due to accumulation of _____. (a) Sphingol (b) Sphingomyelin (c) Glucocerebroside (d) Ganglioside x. The most important phosphosphingolipid is ____ (a) Sphingomyelin (b) Choline (c) Ganglioside (d) Ceramide Answer any TEN from the following: **Q.2** [20] i. Write names of enzymes associated with TCA cycle. What is substrate level phosphorylation? Write at least 1 such reaction. iii. Differentiate between glucokinase and hexokinase. P.T.O.

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	iv.	. What is PUFA. Write additional enzymes used for the Oxidation of PUFA.	
	v.	Differentiate between alpha and omega oxidation.	
	vi.	Give diagrammatic representation of carnitine shuttle.	
	vii.	Define pentosuria.	
	viii.	What is Cori's disease.	
	ix.	Explain amylopectinosis.	
	x.	What is kerasin? Where it is present?	
	xi.	What is Tay Sach's disease?	
	xii.	Write salient features of Niemann Pick's disease.	
Q.3		Explain the following in detail:	
	(a)	Gluconeogenesis	[5]
	(b)	PDH complex	[5]
		<u>OR</u>	
Q.3		Describe the following:	
	(a)	Significance of HMP shunt pathway	[5]
	(b)	Regulation of glycolysis	[5]
Q.4	(a)	Discuss the role of adipose tissues in lipid metabolism.	[5]
	(b)	Explain oxidation of odd chain fatty acids.	[5]
		<u>OR</u>	
Q.4		Write a detail notes on the following:	
	(a)	Fatty acid synthase complex	[5]
	(b)	Energetics and regulation of β (beta)) oxidation	[5]
Q.5		Write detail notes on the following (ANY TWO):	[10]
	(a)	Pompe's disease	
	(b)	Galactosaemia	
	(c)	Limit dextrinosis	
	(d)	Glycosuria	
Q. 6		Discuss Refsum's disease and Gaucher's disease.	[10]
		<u>OR</u>	
Q. 6		What is ketosis. Discuss carnitine / carnitine acyl transferase deficiency.	[10]
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Q.4

Q.4

Q.5

Q. 6

Q. 6