



MIDTERM ASSESSMENT
2nd YEAR MBBS - MCQs

ENCIRCLE one best answer.

Any cutting or overwriting will not be accepted and no marks will be given even if the answer is correct.

1 Which of the following is primary bile salt (a) Fumaric acid (b) Pyruvic acid (c) Deoxycholic acid (d) Chenodeoxycholic acid	2 Salivary amylase is inactivated in stomach by (a) HCl (b) Pepsin (c) Pepsinogen (d) Renin
3 Increased level of serum amylase is observed in (a) Prostate cancer (b) Liver disease (c) Pancreatitis (d) Myocardial infarction	4 All of the following are components of electron transport chain except (a) FMN (b) FAD (c) CoQ (d) CDP
5 NADH+H ⁺ by oxidative phosphorylation gives (a) More ATP than FADH ₂ (b) Equal ATP to FADH ₂ (c) Less ATP than FADH ₂ (d) It gives 3.5 ATP	6 During starvation the first nutrient to be depleted is (a) Proteins (b) Glycogen (c) Triacylglycerol (d) Cholesterol
7 Propionyl-S-CoA which is liberated during the oxidation of odd chain fatty acids can be used for (a) Synthesis of cholesterol (b) Synthesis of thyroid hormones (c) Synthesis of epinephrine (d) Gluconeogenesis	8 All of the following statements about ketone bodies are true except (a) Their synthesis increases in diabetes mellitus (b) These are synthesized in liver mitochondria (c) These can deplete alkali reserve (d) These can be oxidized in liver to provide energy
9 Insulin stimulates (a) Hepatic glycogenolysis (b) Hepatic glycogenesis (c) Gluconeogenesis (d) Lipolysis	10 Which of the following is an intestinal enzyme (a) Maltase (b) Ribonuclease (c) Amylase (d) Pepsin
11 Glucose-1-PO ₄ is liberated from glycogen by the action of (a) Glucokinase (b) Hexokinase (c) Glycogen synthase (d) Phosphorylase	12 Before pyruvic acid enters into the citric acid cycle it must be converted to (a) Lactate (b) Acetyl-CoA (c) Fructose-1-PO ₄ (d) Citrate
13 Which of the following vitamin is the component of electron transport chain? (a) Vitamin E (b) Thiamine (c) Niacin (d) Vitamin D	14 2,4 dinitrophenol is an (a) Inhibitor of electron transport chain (b) Activator of electron transport chain (c) uncoupler of oxidative phosphorylation (d) Activator of glycogenolysis

<p>15 Rotenone a fish poison is</p> <p>(a) Uncoupler of oxidative phosphorylation (b) Activator of oxidative phosphorylation (c) Inhibitor of electron transport chain (d) Activator of electron transport chain</p>	<p>16 Clay coloured stools are due to the absence of</p> <p>(a) Vitamin D (b) Calcium carbonate (c) Bile salts (d) Bile pigments</p>
<p>17 Biosynthesis of secondary bile acids occurs in</p> <p>(a) Liver (b) Lungs (c) Intestine (d) Stomach</p>	<p>18 Chronic exposure to drugs or toxins induce increased synthesis of</p> <p>(a) Glucronate (b) Citrate (c) Pyruvate (d) Malate</p>
<p>19 Deficiency of Dopamine in brain may lead to</p> <p>(a) Maple syrup urine disease (b) Albinism (c) Parkinsonism (d) Phenylketonuria</p>	<p>20 Hormone which activates the absorption of glucose from intestine is</p> <p>(a) Insulin (b) Oxytocin (c) Thyroid hormone (d) Parathyroid hormone</p>
<p>21 Serine is the precursor of all of the following except</p> <p>(a) Glycine (b) Choline (c) Ethanolamine (d) Methionine</p>	<p>22 Which of the following is the major transport form of NH_3 in the blood?</p> <p>(a) Glutamate (b) Glutamine (c) Transaminase (d) Alpha ketoglutarate</p>
<p>23 In Transamination reaction</p> <p>(a) Urea is formed (b) NH_3 is formed (c) Amino group of one amino acid is transferred to a ketoacid (d) Amino group from a ketoacid is transferred to amino acid</p>	<p>24 In the muscles NH_3 is transported as</p> <p>(a) Asparagine (b) Glutamate (c) Glutamine (d) Alanine</p>
<p>25 End product of protein metabolism</p> <p>(a) NH_3 (b) Urea (c) Glutamine (d) Alanine</p>	<p>26 Glutamate dehydrogenase enzyme is present in</p> <p>(a) Cytoplasm of every cell (b) Cytoplasm of liver (c) Mitochondria of liver (d) In mitochondria and cytoplasm of liver</p>
<p>27 In urea one nitrogen comes from NH_3 and second comes from</p> <p>(a) Glutamine (b) Alanine (c) Glutamate (d) Aspartate</p>	<p>28 Which amino acid becomes Essential in Phenylketonuria?</p> <p>(a) Tryptophan (b) Aspartate (c) Tyrosine (d) Alanine</p>
<p>29 Urea cycle starts in</p> <p>(a) Cytoplasm of cells (b) Cytoplasm of liver (c) Kidney (d) Mitochondria of liver</p>	<p>30 Urine become black on standing in which disease?</p> <p>(a) Phenylketonuria (b) Albinism (c) Maple syrup urine disease (MSUD) (d) Alkaptonuria</p>