

MANIPAL UNIVERSITY

FIRST MBBS DEGREE EXAMINATION – JUNE/JULY 2015

SUBJECT: BIOCHEMISTRY – PAPER I (ESSAY)

Thursday, July 02, 2015

Time: 10:20 – 13:00 Hrs.

Maximum Marks: 80

✍ **Answer ALL the questions.**

1. A three week old infant was brought to pediatric OPD with complaints of excessive crying, refusal to feeds and lethargy. The mother also gave history of mousy odour of the urine and laboratory tests revealed phenylketonuria.

1A. What is the biochemical defect?

1B. What are the products accumulated and excreted in urine?

1C. Discuss the metabolism of phenylalanine.

1D. Add a note on the biochemically significant products formed in the metabolism of phenylalanine

(1+1+5+3 = 10 marks)

2. Illustrate the Krebs's cycle. Justify the anaplerotic and amphibolic role of the cycle with TWO examples each.

(6+2+2 = 10 marks)

3A. Classify lipoproteins and write their functions.

(4 marks)

3B. Describe competitive enzyme inhibition with TWO clinical applications.

(4 marks)

3C. Name the enzyme defect associated with:

- | | |
|--------------------------------------|--------------------------|
| i) Tay Sach's disease | ii) von Gierke's disease |
| iii) Hereditary fructose intolerance | iv) Albinism |

(1 mark × 4 = 4 marks)

3D. Write the components of the ETC in sequence indicating the ATP generating sites.

(3+1 = 4 marks)

3E. Give ONE significance of:

- | | |
|----------------------|-----------|
| i) BAL | ii) CK-MB |
| iii) Glucuronic acid | iv) GSH |

(1 mark × 4 = 4 marks)

3F. Classify amino acids based on their chemical properties.

(4 marks)

3G. Write the steps of ketogenesis. Add a note on Ketosis.

(3+1 = 4 marks)

3H. Give the normal serum/plasma values of:

- i) Postprandial glucose
- ii) Urea
- iii) Cholesterol
- iv) Glycated haemoglobin

(1 mark × 4 = 4 marks)

3I. Discuss enzyme specificity with suitable examples.

(4 marks)

3J. Explain the significance of:

- i) Galactose
- ii) Ribose

(2 marks × 2 = 4 marks)

3K. An obese male with high triglycerides was also found to have hepatomegaly. What is your probable diagnosis? Discuss the other causes for this condition and name the factors that help prevent it.

(1+2+1 = 4 marks)

3L. Discuss the metabolism of methionine.

(4 marks)

3M. Explain the steps of β -oxidation of palmitic acid in mitochondria with a note on its energetics.

(3+1 = 4 marks)

3N. **Write briefly on:**

- i) Acute phase proteins
- ii) Use of radioisotopes in medicine

(2 marks × 2 = 4 marks)

3O. Give biochemical reasons for the following:

- i) PUFAs can prevent atherosclerosis
- ii) When heated, proteins lose their biological activity
- iii) The ETC is located in the inner mitochondrial membrane
- iv) Deficiency of glucose-6-phosphate dehydrogenase can lead to haemolysis

(1 mark × 4 = 4 marks)



MANIPAL UNIVERSITY**FIRST MBBS DEGREE EXAMINATION – JUNE/JULY 2015****SUBJECT: BIOCHEMISTRY– PAPER II (ESSAY)**

Friday, July 03, 2015

Time: 10:20 – 13:00 Hrs.

Maximum Marks: 80

✍ **Answer ALL the questions.**

✍ **Long answer questions.**

1. A 48 year old male patient presented with symptoms of yellowish discoloration of skin, sclera. After the clinical investigations and ultrasonography he was found to have carcinoma of head of the pancreas. His biochemistry laboratory report showed the following:

Total bilirubin level: 20 mg/dl, conjugated bilirubin: 18mg/dl, Alkaline phosphatase (ALP):100 KAU/dl

(Normal level: 3-13 KAU units/dl), Alanine transaminase (ALT): 55 IU/l (Normal level: 5-40IU/l)

Answer the following questions:

1A. Diagnose the type of jaundice. Explain biochemical basis for diagnosis

1B. Explain the formation and metabolism of bilirubin

1C. What happens to prothrombin time in this patient and why?

((1+2)+(2+3)+2 = 10 marks)

2. With the help of diagrams explain translation under the following aspects:

2A. Activation of amino acids

2B. Initiation

2C. Elongation

2D. Termination

(2+3+3+2 = 10 marks)

3. **Short answer questions:**

3A. Explain the formation of active vitamin D from its provitamin form. Add a note on its role in serum calcium homeostasis

(2+2 = 4 marks)

3B. Describe recombinant DNA technology with the help of diagrams. Write two applications.

(3+1 = 4 marks)

3C. Write notes on:

i) HbS

ii) Westernblot technique

(2+2 = 4 marks)

- 3D. Write **one** coenzyme reaction catalyzed by and deficiency manifestations of each of the following vitamins:
 i) Vitamin C ii) Niacin
 (2+2 = 4 marks)
- 3E. Enumerate the steps of heme biosynthesis indicating enzymes and coenzymes.
 (4 marks)
- 3F. Write on the causes and alterations in the acid base parameters in respiratory acidosis.
 (2+2 = 4 marks)
- 3G. Write the formation of PRPP (5-phosphoribosyl pyrophosphate). Add a note on purine salvage.
 (1+3 = 4 marks)
- 3H. Explain the absorption of iron. What are the causes of iron deficiency anemia?
 (2+2 = 4 marks)
- 3I. A 55 year old, 80Kg weighing male corporate executive had his daily dietary intake of 535 gm carbohydrate, 140 gm protein, 100 gm fat everyday and 50gm alcohol
 i) Calculate his caloric intake per day
 ii) What dietary advise would you recommend regarding his diet composition?
 iii) Define balanced diet
 (1+2+1 = 4 marks)
- 3J. Write notes on:
 i) RFLP ii) Tumor markers
 (2+2 = 4 marks)
- 3K. Write notes on protein energy malnutrition disorders
 (4 marks)
- 3L. Explain:
 i) Causes and deficiency manifestations of vitamin B₁₂
 ii) Anion gap
 (2+2 = 4 marks)
- 3M. Describe Wald's visual cycle. List **two** ocular manifestations observed in vitamin A deficiency
 (3+1 = 4 marks)
- 3N. Write notes on:
 i) Creatinine clearance test ii) Biochemical significance of copper
 (2+2 = 4 marks)
- 3O. Give biochemical basis for the following:
 i) Folate deficiency causes megaloblastic anemia
 ii) 61 tRNA's need not be there to recognize 61 codons during translation
 iii) B₁₂ deficiency results in neurological symptoms
 iv) Genetic code is described as degenerate and unambiguous
 (1 mark × 4 = 4 marks)

