2 O FEB 2020



Faculty of Medicine, University of Ruhuna
Medical Laboratory Science Degree Programme
Year-End Examination-Year 3
6th Batch, July 2017

Clinical Biochemistry (MLS 3102) - Theory

25th July 2017

9.00 a.m. - 11.00 a.m.

Duration: Two hours

Answer all four questions.

Index number:.....

(22)

Question 1

- 1.1 A 32 year-old male admitted due to a road traffic accident develops symptoms of acute kidney injury after 24 hours.
 - 1.1.1 List the biochemical investigations that may be requested for this patient and comment on the requirements for sample collection. (20 marks)
 - 1.1.2 State the biochemical changes that you would expect in the above investigations.

 (10 marks)
 - 1.1.3 List the three categories of acute kidney injury and predict the category to which the above patient may belong. (10 marks)
- 1.2 Briefly state the pathological basis of the following clinical conditions;
 - 1.2.1 Glycosuria in diabetes mellitus
 - 1.2.2 Oedema in nephrotic syndrome
 - 1.2.3 Polyuria in diabetes insipidus

(30 marks)

- 1.3
- 1.3.1 List the properties of a substance that can be used for renal clearance test and briefly discuss how clearance tests can be used to assess kidney function

 (10 marks)
- 1.3.2 Describe the sample collection requirements for 24 hour urine creatinine clearance test and discuss the pre-analytical factors that might affect the test results

 (15 marks)
- 1.3.3 "A normal plasma creatinine concentration does not necessarily imply normal renal function". Comment on the above statement. (05 marks)

Question 2

2.1 Briefly describe what is proficiency testing.

(20 marks)

- 2.2 Describe how you would reduce the cost of quality management in the biochemistry section of your laboratory. (20 marks)
- 2.3 Describe the factors that you would consider when purchasing a new biochemistry analyser for your laboratory. (30 marks)
- 2.4 Explain how you would control the analytical quality of fasting plasma glucose test using patient data, individual and multiple patient test results. (30 marks)

Question 3

3.1 What is the difference between osmolality and osmolarity?

- 3.2 Freezing point depression is the most common laboratory method for the determination of osmolality in biological fluids.
 - Describe the principle behind freezing point depression osmometer using the 3.2.1 standard freezing curve.

The freezing point of a patient sample received to the Clinical Pathology laboratory was - 0.53 °C. Calculate the osmolality of the above patient (1 mole of dissolved molecules per kg depresses the freezing point of water by 1.86 °C).

(21) marks)

- 3.3 Discuss the variation of the osmolal gap in following conditions.
 - 3.3.1 Alcoholism
 - 3.3.2 Dehydration

(20 marks)

Question 4

- 4.1 Classify chromatographic techniques according to the method of separation and explain how separation is achieved in two of the mentioned methods. (30 marks)
- 4.2 Draw a schematic diagram of a gas chromatographic system and label the components.

(15 marks)

- 4.3 Briefly comment on the following
 - 4.3.1 Temperature programming in Gas Chromatography.
 - 4.3.2 Isocratic elution in Liquid Chromatography.
 - 4.3.3 Features of an ideal detector.

(15 marks)

- 4.4 Separation and identification of growth hormone variants were performed using a reverse phase Liquid Chromatography-Mass Spectrometry system.
 - 4.4.1 Explain what is reverse phase chromatography and why it is preferred for separation of biomolecules. (10 marks)
 - 4.4.2 List factors that would affect band broadening in the above separation.

(10 marks)

4.4.3. Explain how to minimize band broadening using the Van Deemter equation.

(20 marks)

