

UNIVERSITY OF JAFFNA, SRI LANKA
FACULTY OF ALLIED HEALTH SCIENCES
SECOND YEAR SECOND SEMESTER EXAMINATION IN BSc HONS (MLS)- 2019
MLSHE 2225 HAEMATOLOGY II
PAPER II

Date: 31.01.2022

Time: 2 hours

Answer all Eight Questions.

Answer Part A and B in Separate Answer Books.

Part A

1. Haemostasis is a complex process.
 - 1.1. List the major components of normal haemostasis. (10 Marks)
 - 1.2. Briefly explain the mechanism of platelet activation and aggregation in primary hemostasis. (40 Marks)
 - 1.3. Schematically enumerate the coagulation cascades of normal hemostasis (50 Marks)

2.
 - 2.1. Briefly explain the principles of the following tests
 - 2.1.1. Prothrombin Time (20 Marks)
 - 2.1.2. Activated Partial Thromboplastin Time (20 Marks)
 - 2.1.3. Thrombin Time (20 Marks)
 - 2.1.4. Fibrinogen assay (Clauss Technique) (20 Marks)
 - 2.2. List five (5) preanalytical errors which cause abnormal results in the above tests. (20 Marks)

3.
 - 3.1. Explain the role of von Willebrand factor in haemostasis (30 Marks)
 - 3.2. Tabulate the classification of von Willebrand disease based on multimer analysis (30 Marks)
 - 3.3. Write a note on PFA-100 system (40 Mark)

4.

4.1. Define "Leukemia" (15 Marks)

4.2. Explain how Philadelphia chromosome contributes to leukemogenesis (60 Marks)

4.3. List five (5) laboratory techniques which can detect Philadelphia chromosome (25 Marks)

5.

5.1. List four (4) coagulation inhibitors present in blood (20 Marks)

5.2. Briefly explain the mechanism of action of coagulation inhibitors mentioned in 5.1 (40 Marks)

5.3. Explain the significance and the mechanism of action of fibrinolytic system. (40 Marks)

Part B

6. A patient was admitted to a medical ward with suspected haemolytic anaemia.

6.1. Define haemolytic anaemia. (10 Marks)

6.2. List four different blood tests you would receive at laboratory from the ward which are useful to diagnose haemolytic anaemia and indicate expected findings in each test in haemolytic anaemia. (30 Marks)

6.3. Outline how you would classify haemolytic anaemia based on site of red cell destruction and how you would differentiate each. (30 Marks)

6.4. Outline the pathological basis for increased urobilinogen in urine in haemolytic anaemia. (30 Marks)

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7. Defects of haemoglobins cause anaemia.
 - 7.1. State the confirmatory test of beta thalassaemia major, its principle, quality control, and sample collected. (35 Marks)
 - 7.2. State the inheritance of beta thalassaemia and expected findings of the test mentioned in question 7.1, in parents of a beta thalassaemia major baby. (15 Marks)
 - 7.3. Compare and contrast alpha thalassaemia and beta thalassaemia. (35 Marks)
 - 7.4. State the principle of sickling test and how you would ensure quality of sickling test (15 Marks)
8. You are appointed as a member of the team assigned to design a haematology laboratory in a small private hospital in a major city. The management insist optimal quality and at the same time cost saving as well. After initial workload survey, it is estimated to receive approximately 25-50 FBC, 20 ESR, 4 Retic counts, and one (1) PT/INR daily and 1-2 APTT per week.
 - 8.1. Outline the key equipment/items you would suggest to run this laboratory giving reasons. (25 Marks)
 - 8.2. Outline basic safety requirements needed in a haematology laboratory. (25 Marks)
 - 8.3. Discuss what specifications are needed when selecting equipment required for FBC testing. (25 Marks)
 - 8.4. Discuss quality management processes you would implement in this laboratory for FBC testing. (25 Marks)