CHUKA



UNIVERSITY

UNIVERSITY EXAMINATIONS

SECOND YEAR EXAMINATION FOR THE AWARD OF BACHELOR OF SCIENCE (NURSING)

SUPPLEMENTARY EXAMINATIONS

NURS 222: HEMATOLOGY

STREAMS: BSC (NURS)

TIME: 2 HOURS

DAY/DATE: TUESDAY 10/8/2021 2.30 P.M. – 4. 30 P.M.

INSTRUCTIONS:

• Do not write anything on the question paper.

- Mobile phones and any other reference materials are NOT allowed in the examination room.
- The paper has three sections. Answer ALL questions
- All your answers for Section I (MCQs) should be on one page.
- Number ALL your answers and indicate the order of appearance in the space provided in the cover page of the examination answer booklet.

PART I: Multiple Choice Questions (20 marks). Choose the most appropriate response

- 1. A 34 year old woman sustains severe burns over most of her body surface when her propane stove exploded. What is disrupted most significantly in this patient?
 - a. Antibody production
 - b. Complement
 - c. First line defence
 - d. Phagocytosis
- 2. The evaluation in a newly diagnosed case of acute lymphoid leukemia (ALL) should routinely include all of the following EXCEPT
 - A. bone marrow biopsy
 - B. lumbar puncture
 - C. complete metabolic panel
 - D. cytogenetic testing

- 3. All the following are suggestive of iron deficiency anemia EXCEPT
 - A. koilonychia
 - B. pica
 - C. decreased serum ferritin
 - D. decreased total iron-binding capacity (TIBC)
- 4. Which of the following statements is true?
 - A. Factor VIII deficiency is characterized clinically by bleeding into soft tissues, muscles, and weight bearing joints.
 - B. Congenital factor VIII deficiency is inherited in an autosomal recessive fashion.
 - C. Factor VIII deficiency results in prolongation of the prothrombin time.
 - D. Factor VIII complexes with Hageman factor, allowing for a longer half-life.
- 5. Which of the following statements correctly describes characteristics of stem cells?
 - A. Ability to differentiate into a variety of mature cells types
 - B. Capacity for self-renewal
 - C. Generate, maintain, and repair tissue
 - D. A and C
- 6. Aplastic anemia has been associated with all of the following EXCEPT
 - a. carbamazepine therapy
 - b. methimazole therapy
 - c. non-steroidal inflammatory drugs
 - d. parvovirus infection
- 7. Which of the following is an appropriate intervention for a patient with sickle cell disease experiencing priapism that has lasted four hours?
 - a. Administration of opioid analgesics
 - b. Application of ice packs to the penis
 - c. Intravenous heparin boluses
 - d. Restriction of fluid intake
- 8. Which of the following is characteristic of pernicious anemia?
 - a. increased production of intrinsic factor
 - b. decreased absorption of vitamin B₁₂
 - c. antibodies to gastric HCl
 - d. decreased absorption of folate
- 9. Which of the following deficiencies would most likely lead to megaloblastic anemia?
 - a. vitamin E deficiency
 - b. vitamin B₆ deficiency
 - c. iron deficiency
 - d. folic acid deficiency

- 10. The peripheral blood of a patient with iron deficiency anemia will most likely show what picture?
 - a. microcytic, hypochromic red cells
 - b. microcytic, normochromic red cells
 - c. macrocytic, hypochromic red cells
 - d. normocytic, hypochromic red cel ls
- 11. With increased intravascular hemolysis which of the following will likely occur?
 - a. the test for methemalbumin will be negative
 - b. urine hemosiderin will be increased
 - c. unconjugated bilirubin levels will remain normal
 - d. the reticulocyte count will decrease
- 12. A patient has anemia, decreased RBC indices, and targets on the peripheral smear. Serum iron is normal and electrophoresis shows increased HbF and HbA₂. What is a possible diagnosis?
 - a. alpha thalassemia
 - b. sideroblastic anemia
 - c. beta thalassemia
 - d. anemia of chronic disease
- 13. The anemia of chronic disease may be caused by which of the following:
 - a. impaired iron metabolism
 - b. autoantibodies
 - c. increased EPO secretion
 - d. increased RBC lifespan
- 14. All of the following describe multiple myeloma, EXCEPT:
 - a. malignant plasma cells infiltrate the bone marrow
 - b. hypercalcemia
 - c. low levels of plasma immunoglobulin
 - d. light chains are excreted in the urine
- 15. Why is there decreased production of blood cells in the marrow in acute leukemia?
 - a. there is a stem cell defect
 - b. there is a growth factor deficiency
 - c. there is an erythropoietin deficiency
 - d. the malignant cells replace normal marrow
- 16. Which of the following coagulation factors does Antithrombin III inhibit?
 - a. factor V
 - b. factor VIII
 - c. factor IX
 - d. factor XIII

- 17. Which of the following test results would most likely be seen in Hemophilia A?
 - a. an abnormal PT
 - b. an abnormal Bleeding Time
 - c. abnormal von Willebrand factor levels
 - d. an abnormal APTT
- 18. Which of the following tests would be normal in von Willebrand disease?
 - a. a factor VIIIC assay
 - b. von Willebrand factor multimer analysis
 - c. tests for platelet aggregation
 - d. thrombin time
- 19. A five-year-old patient with sickle cell disease presents with splenic sequestration, oxygen saturation value of 95%, temperature 37.5°C (99.5°F), heart rate 148, respirations 28, and blood pressure 90/36. The patient's lab values include a white blood cell count 5,000/mm3, hemoglobin 5.2, and platelet count 175,000. What is the most important initial intervention?
 - a. Administration of intravenous antibiotics
 - b. Initiation of oxygen via nasal cannula
 - c. Ten breaths on an incentive spirometer
 - d. Transfusion of 10 ml/kg packed red blood cells
- 20. A 6-year-old boy in Kenya develops swelling of the jaw. The mass responds rapidly to chemotherapy. What is the most likely diagnosis?
 - a. Burkitt's lymphoma
 - b. Follicular lymphoma
 - c. Acute lymphoblastic leukemia
 - d. Lymphoblastic lymphoma

PART II: SHORT ANSWER QUESTIONS (40 MARKS) ANSWER ALL QUESTIONS

1. Outline the normal parameters of formed elements of blood	(10 marks)
2. Outline any five (5) clinical manifestations of Hodgkin's lymphoma	(5 marks)
3. Briefly explain the pathophysiology leading to sickle cell crises	(6 marks)
4. a. Describe the pathophysiology of acute myeloid leukemia	(4 marks)
b.State any five (5) clinical manifestations of acute myeloid leukemia.	(5 marks)
4. Describe the intrinsic pathway of coagulation	(10 marks)

PART III: LONG ANSWER QUESTIONS (40 MARKS)

- 1. Haemophillia is one of the bleeding disorders
 - a. State six (7) clinical manifestations of hemophillia (7 marks)
 - b. Outline any three (3) laboratory investigations carried out to confirm diagnosis

(3

marks)

c. Describe the management of patient with hemoplillia. (10 marks)

- 2. Anaemia is a common medical problem in resource-scarce settings.
 - a. List any four (4) types of anaemia (2 marks)
 - b. Outline ten (10) clinical features of anaemia (5 marks)
 - c. Discuss the management of a patient with a specified type of anaemia (13 marks)
