

"Investing in Africa's Future"

COLLEGE OF HEALTH, AGRICULTURE & NATURAL SCIENCES

DEPARTMENT OF HEALTH SCIENCES

SLS 408 HAEMATOLOGY THEORY

END OF SECOND SEMESTER SUPPLEMENTARY EXAMINATIONS

APRIL/MAY 2019

LECTURER: DR MABOREKE

DURATION: 3 HOURS

INSTRUCTIONS

Write your candidate number on the space provided on top of each page Answer **all** questions in sections A on the question paper.

Answer **all** questions in section B on separate answer sheets provided.

Answer any **6** questions in section C on separate answer sheets provided Credit will be given for logical, systematic and neat presentations in sections B and C

Candidate number	

Sections A(40 marks)

Answer all questions by indicating T for TRUE or F for FALSE in front of each of the statements (a) to (d) in all the questions

1. Regarding FBC analyser technology

- a) Forward light scatter is proportional to cell size
- b) Side light scatter is proportional to cell complexity
- c) Cell complexity refers to degree of lobularity and granularity of the cell
- d) A lymphocyte is a complex cell

2. Which of the following conditions usually shows a high RDW?

- a) Sickle cell anaemia
- b) Iron deficiency anaemia
- c) Megaloblastic anaemia
- d) Thalassaemia trait

3. Which of the following are causes of microcytic hypochronic anaemia

- a) Iron deficiency
- b) Thalassaemia trait
- c) Early anaemia of chronic disease
- d) Vitamin B12 deficiency

4. Which test can be used to confirm iron deficiency?

- a) Serum ferritin
- b) Serum iron and Transferin
- c) Zinc protoporphyrin
- d) Serum iron alone is a reliable test

5. Regarding Myeloproliferative neoplasms,

- a) JAK2 mutation is positive in 100% of cases of polythemia Vera
- b) JAK2 mutation is positive in just over 50% of cases of Essential thrombocythemia and just over 50% of primary myelofibrosis
- c) Leukocythroblastic picture with teardrop poikilocytosis is a typical finding on blood film of a patient with primary myelofibrosis.
- d) CALR and MPL mutations are found in a small proportion of patients with Essential thrombocythaemia and primary myelofibrosis

6. Regarding accurate diagnosis of acute Leukaemia,

- a) Auer rods confirm diagnosis of Acute myeloid leukaemia
- b) Flow cytometric immunophenotype is most useful test
- c) Cytogenetics can be diagnostic and prognosticative
- d) t(8:21) t(15:17) and inV16 are good prognostis cytogenetic features

7. Regarding neutropenia

- a) Severe neutropenia refers to absolute neutrophil count of less than 0.5x109/L
- b) Prolonged severe neutropenia is that which lasts for more than seven days
- c) Patients with severe neutropenia are prone to serious infections
- d) Clinical features of neutropenia include recurrent infections and recurrent mouth ulcers

8. Regarding Acute Leukaemia,

- a) Onset of illness is insidious
- b) Predominant cells are blasts on blood or bone marrow aspirate film examination.
- c) Acute lymphoblastic Leukaemia (ALL) is more common in children than adults
- d) Acute myeloid leukaemia (AML) is more common in children than adults

9. About chronic myeloid leukaemia,

- a) On blood or marrow aspirate film exam
 - -all neutrophil precursors are represented but with predominant cells being mature neutrophils and myelocytes
- b) The cytogenetic marker is t(9:22) resulting a fusion gene.
- c) The fusion gene is BCR/ABL
- d) The diagnostic fusion gene is detected and quantified by polymerase chain reaction (PCR)

	d)	Platelet count	
11.	A t	wo year old boy has the following coagulation test results	
	•	PT13 secs (10-14)	
	•	APTT 50 secs (27-43)	
	•	FBC normal	
	•	aPTT after 50/50 mix with normal plasma = 35 secs	
	Which factor assays do you do to detect the deficiency?		
	a)	FVIII	
	b)	FVII	
	c)	FV	
	d)	FIX	
12.	Sub	oclassification of anaemia based on MCV include:	
	a)	Macrocytic hypochromic	
	b)	Microcytic hyperchromic	
	c)	Normocytic hypochromic	
	d)	Microcytic hypochromic	
13.	Wŀ	nich are the features of low grade non-Hodgkin's lymphoma?	
	a)	Slow onset of presentation	
	b)	Cells are usually small on microscopic examination	
	c)	Cells are slowly dividing	
	d)	Usually curable	
14	RD.	W is a quantification of:	
14.	a)	Microcytosis	
	a) b)	Poikilocytosis	
	c)	Anisocytosis	
	d)	Red cell dilution with water	
	۷)	ned sen dilution with water	

10. Which of the following tests are part of the clotting screen?

a) PT

c) aPTT

b) FVIII assay

15. Regarding Hodgkin's Lymphoma (HL),

- a) It is of B cell origin
- b) The diagnostic neoplastic cells is the Reed-Sternberg cell in classical Hodgkin lymphoma
- c) Blood film examination is useful in making a diagnosis
- d) Immunohistochemistry is useful for accurate diagnosis and distinguish between classical HL and nodular lymphocyte predominant

16. Thrombin time (TT) is used mainly to evaluate the level and function of

- a) Fibrinogens
- b) Heparin contamination of blood sample where it is prolonged but reptilase time is not prolonged
- c) FVIII
- d) None of the above

17. Which of the following are features of High grade non-Hodgkin's lymphoma?

- a) Cells are usually large
- b) Cells are rapidly dividing
- c) High lactic dehydrogenate (LDH) indicates high disease bulk
- d) Potentially curable.

18. Causes of eosinophilia include the following,

- a) Parasitic infestations
- b) Atopic disease
- c) Viral infection
- d) Bleeding

19. The components of good Quality Management System (Quality Assurance) include

- a) Inspections from the international standards Organization (ISO)
- b) External quality assessment
- c) Internal Quality Control activities
- d) Continuous Quality Improvement Activities

20. Which of the following blood film features are suggestive of infection?

a) Increased rouleaux

Candidate number..... b) High ESR or CRP c) Neutrophilia with left shift and toxic granulations, vacuolations and Dohle bodies d) Peudo Pelger Huet neutrophils SECTION B (20 marks) **Answer all questions** Each question carries 5 marks 1. A 6 year old boy presents with yellow eyes. FBC shows WBC 9.4 Hb 6.5 MCV 101 Plt 170 Reticulocyte count 6% Blood film shows spherocytosis. Total bilirubin 48 (2-22) a) List 3 possible causes of spherocytosis. b) What single test do you do to narrow the diagnostic posibilities. 2) A 7 year old girl has acute leukaemia with more than 50% blasts on blood film and bone marrow film. However the haematologist cannot on microscopic examination say whether this is acute myeoid leukaemia (AML) or acute lymphoblastic leukaemia (ALL) He requests cytochemistry. Please match (use arrow) each cytochemical stain on the left with the leukaemia it detects a) Sudan Black i) ALL

3) A 60 yr old lady comes to hospital with tiredness from anaemia. WBC 3.1 Hb 5.2

b) Myeloperoxidase

e) Acid phosphatsse

MCV 119 Plt 101

a) What are the possible causes of such an anaemia

b) What further tests do you suggest to confirm the cause

4. (a) What are the two screening tests for sickle cell disease.

d) Non specific Esterase

c) PAS

ii) Monocytic AML

iii) AML

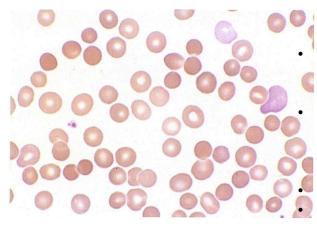
iv) T-ALL

(b) List the tests used to confirm the phenotype of sickle cell disease.

SECTION C (60 marks)

Answer any six cases on separate answer sheets provided. Each case carries 10 marks.

Case 1



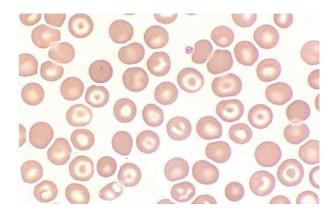
A 22 Year old female presents with history of

A 45 year old male found collapse on the flour
• FBC: Hb 13,4 g/dl, MCH 91fL, MCH 34,6 pg,
and 3 bottles of chibuku next to him.
MCHC 37,1 g/dl, RDW-CV 14,8%, WBC 8,4 x10

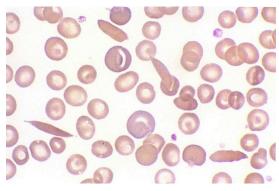
9/L Plt. 377 x10.9/L FBC: Hb 15,1 g/dl, MCV 99,8fL, MCH 34,4 pg, MCHC 34,3 g/dl, RDW-CV 15,5 %, WBC 11,1 X10 9/L PR 910x 10 19/Links 1994 (15) 11

- 2-list 3 possible causes of the abnormality
- 1) RepWhatifurther 166st flipmou recommend to
- 2) IIBPPOSYINDUCALISES OF TIRE OBSIBILITIES TY
- 3) What further test do you recommend to narrow your diagnostic possibilities

Case 2

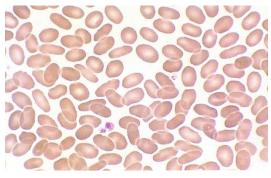


Case 3



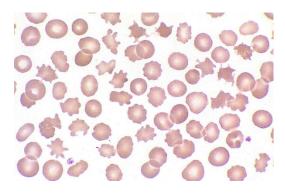
- A 47 year old male from Ghana with severe bone pain
- FBC: Hb 9,2 g/dL, MCV 89,5fL, MCH 30,2pg,RDW-CV14,9%,Retc.CT5,9%,WBC 15,2 x 10 9/L, Plt 600 x 10 9/L,NBRC 4/100 WBC, Diff: ANC 8,1 x 10 9/L
- 1) Report on the blood film?
- 2) what is your diagnosis
- 3) What are the 2 screening tests for this condition
- 4) what 4 different test are available to confirm the phenotype of this condition

Case 4



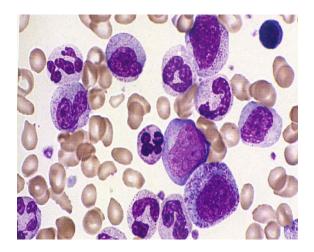
- A 37 Year old female preoperative blood test .
- FBC 12,5 g/dL,MCV
 86,3fL,MCH29,6pg,MCHC34,2g/dL,RDW-CV14,5%,WBC 11,0 x 10 9/L, Plt 204 x 10 9/L.
 Diff ANC 8,8 x 10 9/L
- 1-Report on the blood film?
- 2-What is your diagnosis?
- 3 AV99 tyleart best testale securite cand mon plassing urine
 - FBC: Hb 11,8 g/dL, MCV 92,1fL,MHC 31,0pg,MCHC 33,6g/dL,RDW-CV15,9%,WBC 17,8 x 10 9/L, Plt 138 x 10 9/L. Diff ANC 16,7 x 10 9/L





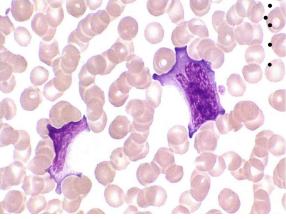
- 1-Report on the blood film?
- 2-What do you think is the diagnosis?
- 3-what biochemical test do you do to confirm your diagnosis?

Case 6



- A 64 year old male presents with lassitude anaemia and massive splenomegaly.
- FBC: Hb 10,4 WBC 170,6, PLT 338, ANC 76,8, ALC 3,4, AMC 5,1,AEC 5,1,ABC 13,6. 1-
- 1) Report on the blood film?
- 2-What do you think is the diagnosis?
- 3-what test do you recommend to confirm the diagnosis

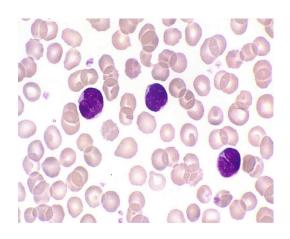
Case 7



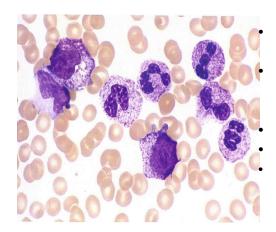
- A 6 year old boy with a cough admitted in ICU
- FBC: Hb 10.2 WBC 16.2 PLT 643 ALC 12.4 A 31 year old male with a superficial eruption of sicles on his back.

- y cause of the lymphocytosis? 3-what further test do you do to confirm the cause 2-list the virus than can cause such a picture? of caugh? 3-what test do recommend to confirm the causes?

Case 8



Case 9



A 6 month infant receiving chemotherapy for neuroblastoma.

FBC: Hb 10, MCV83,MCH 28,2, RDW-CV 15,9, WBC 38,6, PLT 69,ANC 27,8, ALC 1,9,AMC 4,2.

1-Report on the blood film?

2-What is thelikely cause of this picture?

3-what further microbiological tests do you recommend to confirm diagnosis?