

POSTGRADUATE INSTITUTE OF MEDICINE
UNIVERSITY OF COLOMBO

POSTGRADUATE DIPLOMA IN CLINICAL HAEMATOLOGY
EXAMINATION – MAY 2022

Date :- 31st May 2022

Time:- 1.00 p.m. – 4.00 p.m.

Answer **four (04)** questions only.

Answer each question **in a separate book**.

All questions carry equal marks.

PAPER I - ESSAY

1. Discuss the causes, their natural history and relevant diagnostic investigations of neutropaenia in infancy and childhood. (100 marks)
2.
 - 2.1. Describe the WHO 2016 Classification of Myelodysplastic Syndromes (MDS). (40 marks)
 - 2.2. Discuss the pitfalls in the diagnosis of MDS and how they can be overcome. (60 marks)
3.
 - 3.1. Discuss the diagnostic approach to a 25-year-old woman with long standing menorrhagia who is suspected to have von Willebrand disease (vWD). (60 marks)
 - 3.2. List, giving reasons the factors which will affect the von Willebrand factor level. (40 marks)
4. Write short notes on the following:
 - 4.1. Factor V Leiden mutation. (30 marks)
 - 4.2. Acquired factor VIII inhibitors. (40 marks)
 - 4.3. Acquired disorders of platelet function. (30 marks)
5.
 - 5.1. Describe the pathogenesis of haemolysis in paroxysmal nocturnal haemoglobinuria (PNH). (30 marks)
 - 5.2. Discuss the diagnostic approach to PNH and outline the management. (70 marks)

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POSTGRADUATE DIPLOMA IN CLINICAL HAEMATOLOGY
EXAMINATION – MAY 2022

Date :- 1st June 2022

Time:- 9.00 a.m. - 12.00 noon

Answer **six (06)** questions only.

Answer each question **in a separate book**.

All questions carry equal marks.

PAPER II
STRUCTURED ESSAY QUESTIONS (SEQ)

1.
 - 1.1. List the inherited and acquired causes of ring sideroblasts in the bone marrow. (30 marks)
 - 1.2. Describe the molecular basis and pathophysiology of one (01) of the inherited causes that you listed in 1.1. (30 marks)
 - 1.3. Outline the clinical and laboratory findings in the condition described in 1.2. (20 marks)
 - 1.4. Briefly outline the management of the condition mentioned in 1.2. (20 marks)
2. A previously healthy 25-year-old woman presented to the emergency unit with fever of two days, a purpuric rash over both lower limbs and confusion.

Investigations revealed:

Haemoglobin	10 g/dL	
Platelet count	$2 \times 10^9/L$	
Serum creatinine	1.0 mg/dL	(0.9-1.2)
Coagulation profile	Normal	
Urine pregnancy test	Negative	

The blood film confirmed thrombocytopaenia and marked red cell fragmentation.

- 2.1. State the most likely diagnosis. (10 marks)
- 2.2. Describe the pathophysiology of the condition mentioned in 2.1 (40 marks)
- 2.3. Outline further investigations on this patient. (30 marks)
- 2.4. Outline the management of this patient. (20 marks)

3. A 35-year-old man presented with tiredness and easy bruising of one month's duration.

His full blood count revealed:

Haemoglobin	5.6 g/dL
WBC count	$16.7 \times 10^9/L$
Platelet count	$23 \times 10^9/L$
Blood film	Blasts cells were noted

- 3.1. Briefly outline the investigations to confirm an **acute myeloid leukaemia (AML)** in this patient. (40 marks)
- 3.2. How would you subclassify AML in this patient according to the 2016 WHO classification of AML. (40 marks)
- 3.3. Outline the prognostic factors of AML. (20 marks)

4. A 72-year-old woman was found to have a paraproteinaemia of 17g/L.

The full blood count revealed:

Haemoglobin	9.8 g/dL
MCV	107 fL
WBC total count	$21 \times 10^9/L$
Neutrophils	$1.5 \times 10^9/L$
Lymphocytes	$18 \times 10^9/L$
Platelet count	$212 \times 10^9/L$

The patient was suspected to have Waldenstrom macroglobulinaemia.

- 4.1. Outline the important further investigations. (45 marks)
- 4.2. Briefly describe how you would differentiate monoclonal gammopathy of undetermined significance (MGUS) from Waldenstrom macroglobulinaemia. (30 marks)
- 4.3. Discuss the principles of management of Waldenstrom macroglobulinaemia (25 marks)

Contd...../3-

5. A 3-day-old term neonate was admitted with an episode of convulsions. The baby was afebrile and had bruises over the legs and abdomen.

The full blood count revealed:

Haemoglobin	18 g/dL
WBC count	$9 \times 10^9/\text{L}$
Platelet count	$12 \times 10^9/\text{L}$

- 5.1. List the five (05) most likely causes for this clinical presentation. (20 marks)
- 5.2. Describe the investigations you would do to arrive at a diagnosis. (40 marks)
- 5.3. Outline the management of this baby. (20 marks)
- 5.4. The parents wish to know the possibility of thrombocytopaenia in the next baby. What is your answer? (20 marks)
- 6.
- 6.1. Describe how **patients' results** are used for quality control of a full blood count analyzer. (30 marks)
- 6.2. Outline the process of verifying the performance of a newly installed coagulometer in your laboratory for basic coagulation tests (PT, APTT, TT, Fibrinogen). (40 marks)
- 6.3. List the sources of error in automated coagulation based FVIII assay following sample acceptance in the laboratory. (30 marks)
- 7.
- 7.1. What are the definitions of "leucodepletion" and "leucoreduction" of blood components. (10 marks)
- 7.2. Discuss indications for "leucodepleted and leucoreduced" blood components. (50 marks)
- 7.3. Briefly discuss the methods available for "leucodepletion and leucoreduction" giving their advantages and disadvantages. (40 marks)