National Board of Examinations

Question Paper Name :	DrNB CLINICAL HAEMATOLOGY Paper2	
Subject Name :	DrNB CLINICAL HAEMATOLOGY Paper2	
Creation Date :	2023-10-15 14:13:29	
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Share Answer Key With Delivery Engine :	No	
Actual Answer Key :	No	

DrNB CLINICAL HAEMATOLOGY Paper2

Group Number : 1	
Group Id : 327187	72623
Group Maximum Duration : 0	
Group Minimum Duration : 180	
Show Attended Group?: No	
Edit Attended Group?: No	
Group Marks: 100	
Is this Group for Examiner? : No	
Examiner permission : Cant V	iew
Show Progress Bar? : No	

DrNB CLINICAL HAEMATOLOGY Paper2

Section Id :	3271872626
Section Number :	1
Section type :	Offline

Mandatory or Optional :	Mandatory
Number of Questions to be attempted :	10
Section Marks :	100
Enable Mark as Answered Mark for Review and Clear Response :	Yes
Maximum Instruction Time :	0
Sub-Section Number :	1
Sub-Section Id :	3271872630
Question Shuffling Allowed :	No
Is Section Default? :	null

Question Number : 1 Question Id : 32718726914 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

Please write your answers in the answer booklet within the allotted pages as follows:-

Question Number	Answer to be attempted within	Question Number	Answer to be attempted within
Q. 1	Page 1-5	Q. 6	Page 26-30
Q. 2	Page 6-10	Q. 7	Page 31-35
Q. 3	Page 11-15	Q. 8	Page 36-40
Q. 4	Page 16-20	Q. 9	Page 41-45
Q. 5	Page 21-25	Q. 10	Page 46-50

1. a) Describe risk factors and clinical presentation of a patient with hemophagocytic

lymphohistiocytosis (HLH). [4]

b) Describe management strategies in an infant with congenital HLH. [4]

c) How do you manage a patient of SLE with macrophage activation syndrome. [2]

Question Number : 2 Question Id : 32718726915 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) Enumerate various causes of inherited bone marrow failure syndromes. [4]

b) What is telomeropathy and its clinical manifestations. [3]

c) How do you manage a child with fanconi anemia. [3]

Question Number : 3 Question Id : 32718726916 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) Describe principles of leucodepletion and irradiation of blood products. [4]

b) Describe the clinical utilities and various methods for leucodepletion and irradiation of blood products. [3]

c) Describe the pathophysiology & clinical manifestations of transfusion associated GvHD. [3]

Question Number : 4 Question Id : 32718726917 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) Describe steps to be taken for granulocyte collection and transfusion. [4]

b) Describe evidence based practice regarding granulocyte transfusion in a patient with

neutropenic sepsis. [3]

c) Enumerate risks associated with granulocyte transfusions. [3]

Question Number : 5 Question Id : 32718726918 Question Type : SUBJECTIVE Consider As

Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) Describe structural organisation of lymph node. [3]

b) Classify and enumerate causes of lymph node enlargement based on structural organisation of lymph node. [4]

c) Write a note on Kikuchi disease. [3]

Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) Describe monoclonal gammopathy of renal significance. [5]

b) Describe neurological manifestations associated with plasma cell dyscrasia and their management. [5]

Question Number : 7 Question Id : 32718726920 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) How do you approach the diagnosis of a child with prolonged isolated neutropenia. [6]b) Describe management strategies in a child with congenital severe neutropenia/ Kostman syndrome. [4]

Question Number : 8 Question Id : 32718726921 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) Describe the pathophysiology of various clinical syndromes associated with paroxysmal nocturnal hemoglobinuria (PNH). [4]

b) Enumerate various treatment options & their mechanism of action in a patient with PNH. [3] c) Describe indications, conditioning regimens and outcomes of matched sibling allogenic stem cell transplantation in a patient with PNH. [3]

Question Number : 9 Question Id : 32718726922 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

- a) Describe various secondary causes of monocytosis. [3]
- b) Describe diagnostic approach to a patient with prolonged monocytosis. [4]
- c) Describe the treament approach for a young patient with chronic myelomonocytic leukemia

Question Number : 10 Question Id : 32718726923 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) Describe diagnostic approach to isolated thrombocytopenia in an elderly individual. [6]

b) Describe role of thrombopoietin analogues in management of secondary thrombocytopenias. [4]