



**HEMATOLOGY**  
**PAPER-II**

HEM/D/17/48/II

Time : 3 hours  
Max. Marks :100

- Attempt all questions in order.
- Each question carries 10 marks.
- Read the question carefully and answer to the point neatly and legibly.
- Do not leave any blank pages between two answers.
- Indicate the question number correctly for the answer in the margin space
- Answer all the parts of a single question together.
- Start the question to a question on a fresh page or leave adequate space between two answers.
- Draw table/diagrams/flowcharts wherever appropriate.

Write short notes on:

1. a) Neutrophil function tests 4+3+3  
b) Chediak Higashi syndromes  
c) Granulocyte transfusions
2. a) Management of idiopathic hypereosinophilic syndrome 3+4+3  
b) Role of PDGFR $\alpha$  in hypereosinophilia  
c) Acquired eosinophilia
3. a) Diagnosis of systemic mastocytosis 4+3+3  
b) Granules of basophils  
c) Causes of basophilia in peripheral blood.
4. a) Diagnosis of chronic myelomonocytic leukemia 3+2+3+2  
(CMML).  
b) How does it differ from MDS with excess blasts?  
c) Management of CMML  
d) Infections causing monocytosis.
5. Histomonophagocytic Syndrome: 3+4+3  
a) What are the causes?  
b) Criteria for diagnosis  
c) Langerhan's cell histiocytosis.
6. Chronic lymphocytic leukemia: 2+1+3+4  
a) Diagnosis of CLL  
b) How to distinguish it from monoclonal B lymphocytosis.  
c) The genetic landscape of CLL and the implications in prognosis.  
d) Current management of CLL

7. Plasma cells: 4+4+2
- a) What are the imaging techniques for the diagnosis of bone disease in multiple myeloma and their advantages/disadvantages?
  - b) Criteria for the diagnosis of multiple myeloma, smoldering multiple myeloma and MGUS.
  - c) How to differentiate between multiple myeloma and lymphoplasmacytic lymphoma?
8. a) What is hemovigilance? 5+5
- b) How will you implement this in blood bank?
9. a) Risk stratification in MDS 5+5
- b) Treatment options in low risk MDS
10. a) Acquired causes of thrombocytosis 2+4+4
- b) JAK 2 pathway and its inhibitors in myeloproliferative neoplasms
- c) Management of essential thrombocythemia

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