

# Hematology

## Examination



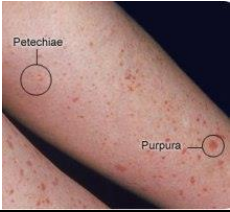
Examination	
Subject steps	Pics / Notes
<b>A-Preparation: Pre-exam Checklist: WIPE</b>	
1- Wash your hands	<p><b>Very important.</b> Ask the patient about his racial origin (e.g. Thalassemia)</p>
2- Introduce yourself to the patient, confirm patient's ID, explain the examination & take consent.	
3- Positioning of the patient " <b>lying flat</b> " and insure his/her Privacy.	
4- Exposure. Full exposure of the trunk.	
<b>B-General appearance: ABC2DEVs</b>	
1- Appearance: young, middle aged, or old, and looks generally ill or well.	Observe the patient's general appearance (age, state of health, nutritional status and any other obvious signs e.g. jaundice, cyanosis, dyspnea)
2- Body built: normal, thin, or obese	<p>♣ Begin by observing the patient's general health from the end of the bed.</p> <p>♣ The patient looks well (not cachectic), overweight (don't say obese), not connected to IV line nor O2 mask, no obvious <b>pallor or cyanosis</b>, no respiratory or pain distress (not tachypnic). Oriented to time, place and person.</p> <p>♣ Also look for syndromes that associated with cardiac disease (Marfan, Down and Turner syndromes).</p>
3- Connections: such as nasal cannula (mention the medications), nasogastric tube, oxygen mask, canals or nebulizer, Holter monitor, I.V. line or cannula (mention the medications).	
4- Color: jaundiced, pale, or cyanosed.	
5- Distress: in pain, respiratory (using accessory muscles), or neurological (abnormal movements) distress	
6- Else: mental functions: consciousness, alertness, and orientation.	
7- Vital signs: 1) Pulse rate* 2) Blood pressure (BP) 3) Temperature 4) Respiratory rate	<p>Take the patient's radial pulse (Determine the Rate, Rhythm and the Character of the pulse). Take his/her blood pressure (Lying and standing or sitting-postural hypotension). <b>Rate:</b> counting over 30 seconds, normally 60-100. <b>Rhythm:</b> regular or irregular. Synchronization by comparing with the other side (radio radial or radio femoral delay). <b>Character and volume:</b> determined from the carotid. <b>Blood pressure (BP)</b> Normal BP defined as a systolic reading less the 140, and diastolic reading less than 90. <b>Temperature:</b> Normal body temperature ranges from 36.6-37.20C. <b>Respiratory rate:</b> It is traditional to count it while taking the pulse. The normal rate at rest should not exceed 25 beat per minute (range 16-25).</p>
<p><b>*Tachycardia</b> in anemic patient: Increased cardiac output due to reduced oxygen-carrying capacity of their blood.</p>	

## Hematological Examination



- 1- Physical examination – inspection
- 2- Abdominal examination especially splenomegaly
- 3- Lymphadenopathy
- 4- Special hematological exams

### 1-Physical examination – inspection


#### A-Skin and its appendices





Paleness	anemia
Plethoric aspect	polyglobulia
Jaundice	haemolytic anaemia, pernicious anaemia(vit B12 deficiency)
Thrombocytopenic purpura <sup>1</sup> (petechia, ecchymoses, suffusion)  1 sort of bruising due to hemorrhage into the skin. petechiae: small bruising , ecchymoses: large bruising. Systemic vasculitis: petechiae are raised (palpable purpura), painful.	Thrombocytopenia  
Skin infections	neutropenia
dry skin, brittle hair and nail, hair loss, itching	Iron deficiency

#### B-Oral cavity, Mucous membranes

plummer-vinson syndrome	mucous membrane atrophy in iron deficient anaemia
Hunter's glossitis	vitamin B12 deficiency
Petechia	thrombocytopenia
Gingiva hypertrophy	Leukaemia 
Aphthosus stomatitis, angina	agranulocytosis, leukaemia 
Confluent tonsillitis	mononucleosis

#### C-Hands

Koilonychias (Dry, brittle, ridged, spoon-shaped nails)	Sever iron deficiency anemia. Occasionally due to fungal infection 
Digital infarction	Abnormal globulins

Pallor of palmar creases	Hemoglobin level is less than 8.1g/dl.
Gouty tophi	
Arthropathy	Hemophilia
<b>D-Face</b>	
<b>Eyes:</b> -jaundice, hemorrhage, injection - conjunctival pallor: <b>more reliable</b> than nail beds or palmar creases for diagnosing anemia	
<b>Mouth:</b>	
Hypertrophy of the gum	Leukemia
Gum bleeding	
Ulceration, infection and hemorrhage of buccal and pharyngeal mucosae	
Atrophic glossitis	Megaloblastic anemia, iron deficiency anemia 
Waldeyer's ring enlargement Waldeyer's ring is a circle of lymphatic tissue in the posterior part of the oropharynx and nasopharynx, and includes the tonsils and adenoids.	Non-Hodgkin lymphoma
<b>E-Legs</b>	
bruising	
pigmentation	
Palpable pleura	Henoch-Schönlein purpura
Scratch marks	
Leg ulcer	Hemolytic anemia (including sickle cell anaemia and hereditary spherocytosis), probably as a result of tissue infarction due to abnormal blood viscosity.
Neurological abnormalities	vitamin B12 deficiency

## 1-Abdominal Examination

- Hepatomegaly.
- Splenomegaly.
- Para-aortic adenopathy: lymphoma, lymphatic leukemia.

### • Splenomegaly:

- Palpation, percussion, auscultation
- Size, tenderness, pain
- Soft, thicken, hard
- "Hypersplenism" – pancytopenia due to sequestration

**Ludwig Traube (spleen triangle):** superiorly by the sixth rib, laterally by the mid-axillary line and inferiorly by the left costal margin that is normally resonant to percussion.

### Causes:

- Portal hypertension
  - Liver cirrhosis
  - Hepatic, portal or splenic vein thrombosis
- Storage disorders – Gaucher, Niemann-Pick
- Systemic diseases – Sarcoidosis, amyloidosis, RA
- Infections
  - Acute
  - Sepsis, IE, typhus abdominalis, Mononucleosis sy.
  - Chronic
  - Tuberculosis, brucellosis, syphilis, malaria, leishmaniasis, schistosomiasis

### Haematological disorders

- Haemolysis (RES hyperplasia): thalassaemia major and intermedia, sickle cell anaemia, any type of haemolytic anaemia
- Malignancies
  - Lymphoid: CLL, HCL, Lymphoma, ALL
  - Myeloid: CML, MF, PRV, AML
  - (Metastases of solid tumors)

### What are the causes of splenomegaly? (HICCUPS)

- Hematological: hemolytic anemia, sickle cell disease, thalassemia major.
- Infection: malaria, kala azar, CMV, HIV, schistosomiasis.
- Congestion: congestive heart failure, portal or splenic vein thrombosis.
- Connective tissue disease: SLE.
- Unknown etiology.
- Primary malignancy: chronic myeloid leukemia (CML), lymphoma, multiple myeloma.
- Storage disease: glycogen storage disease.

## 1-Lymphadenopathy

- Palpable – non-palpable
- Regional – generalized
- Size (soliter, conglomerate), speed of development
- Tenderness, pain
- Soft, thicken, hard
- Relation to the surrounding tissues (fixed or mobile)
- Fluctuation, abscess/fistula formation
- Mass effect (Tracheal/bronchial obstruction, bowel obstruction, DVT)

## Lymph nodes examination

Site	Palpable nodes may be localized to one region (e.g. local infection, early lymphoma) or generalized (e.g. late lymphoma )  The palpable lymph nodes areas are: <ul style="list-style-type: none"> <li>- <b>Epitrochlear</b></li> <li>- <b>Axillary</b></li> <li>- <b>Cervical</b></li> <li>- <b>Supraclavicular</b></li> <li>- <b>Para-aortic (rarely palpable)</b></li> <li>- <b>Inguinal</b></li> <li>- <b>Popliteal</b></li> </ul>
Size	Large nodes are usually abnormal (greater than 1 cm)
consistency	<b>Hard</b> nodes suggest carcinoma deposits; <b>soft</b> nodes may be normal; and <b>rubbery</b> nodes may be due to lymphoma
Tenderness	This implies infection or acute inflammation
Fixation	Nodes that are fixed to underlying structure are more likely to be infiltrated by carcinoma than mobile nodes
Overlying skin	Inflammation of the overlying skin suggests infection, and tethering to the overlying skin suggest carcinoma

## DDX of lymphadenopathy

### I. Infectious

- Viruses: mononucleosis sy.(EBV, CMV, HIV), hepatitis infectiosa, herpes simplex, rubella
- Bacterias: Streptococcus, Staphylococcus, Brucella, Francisella tularensis, Treponema pallidum, Chlamydia, mycobacterias
- Fungi: histoplasmosis, coccidiomycosis
- Parazites: toxoplasmosis, leishmaniasis, trypanosomiasis

### II. Malignant disorders

- Primary hematological disorders: lymphomas; myeloid disorders (acute and chronic myelomonocytic leukaemia)
- Solid tumor metastases

### III. Disorders with immunpatomechanism

- Autoimmun disorders: SLE, RA, MCTD, Sjögren-sy., vasculitis
- Sarcoidosis

### IV. Storage diseases Gaucher, Niemann–Pick-, Fabry-, Tangier-disease

### V. Endocrin disorders (lymphoid hyperplasia) Hyperthyreosis

### VI. Other rare disorders - Castleman disease - Kikuchi disease - Histiocytosis - Dermatopathic lymphadenitis - Mucocutan lymphnode sy.(Kawasaki disease)

## 1-Special hematological exams

Useful laboratory tests:

#### General

- CBC, reticulocyte
- Peripheral blood smear MGG: cytomorphology

- ESR, CRP
- EPO

Iron deficiency

- SeFe, Tf, Sat, SolTfR, Ferritin

Megaloblastic anaemia

- B12, folsav

Haemolysis

- SeBi, LDH
- Direct Coombs, irregular antibodies
- Haptoglobin, plasma free haemoglobin • HgbELFO • RBC enzyme activity

Blood loss

- Stool benzidin test, urine sediment

Plasma cell dyscrasia

- Total protein, ELFO, immunelfo, quantitative Ig

Useful haematopathological exams

- Bone marrow aspiration, biopsy & Lymphnode biopsy
  - Cytomorphology, histology
  - Immunohistochemistry
  - Flow cytometry
  - Cytogenetics (metaphase analysis, FISH)
  - Molecular genetic

FNAB of lymphnodes is not recommended!

Sometimes other organs (skin, stomach, bowel, spleen) for these examinations

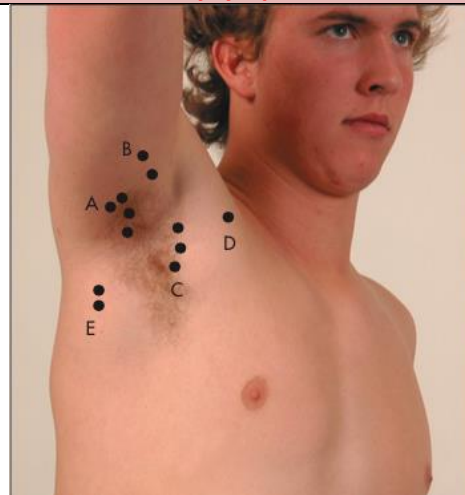
Imaging techniques

- Xray (bone laesions of MM)
- US
- CT
- MRI
- Gallium scan
- PET/CT

**Epitrochlear lymph nodes**



**Axillary lymph nodes**



The main groups of axillary lymph nodes A = central; B = lateral; C = pectoral; D = infraclavicular; E = subscapular.

