

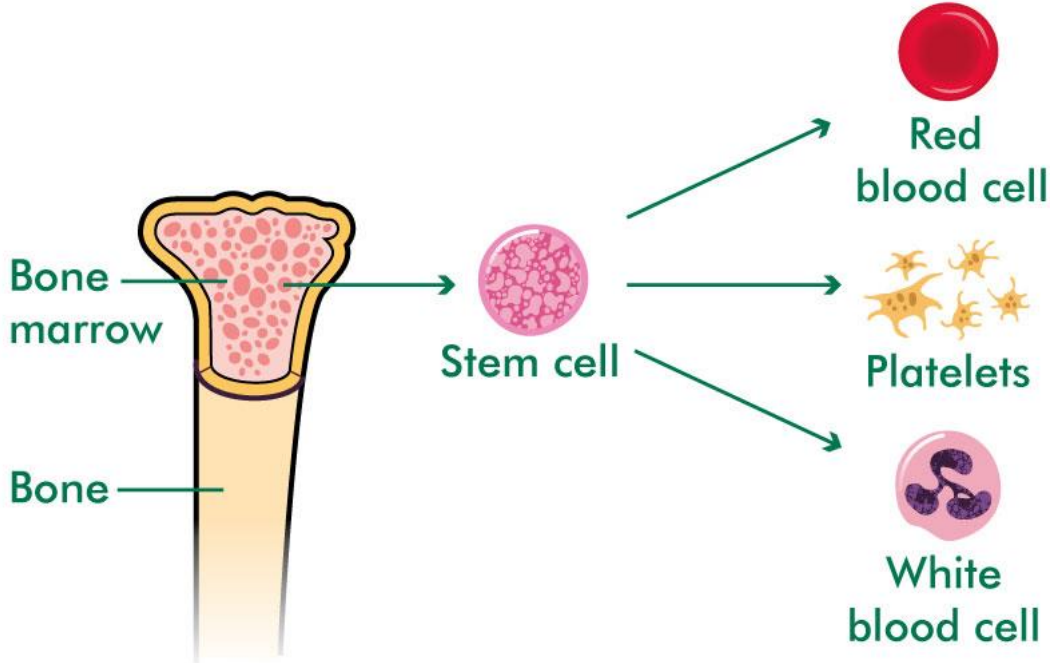
The Anatomy of a CBC

Clinical Updates Podcast: Cell lines and pathologies

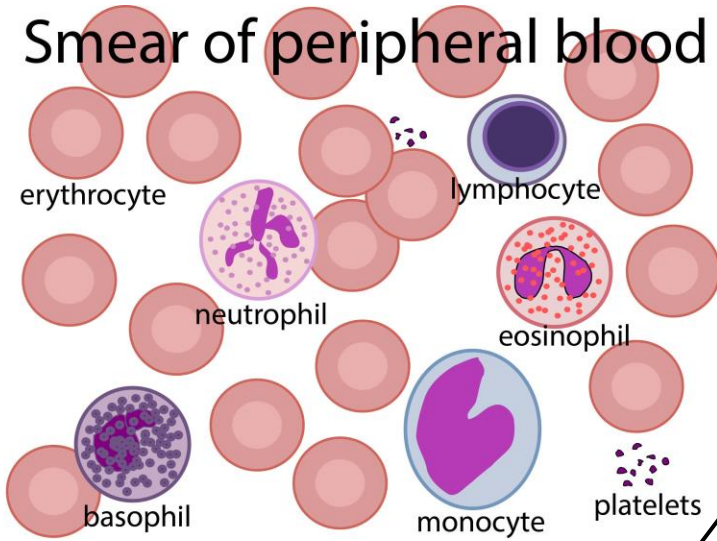
Raquel Walsh Jahnke, D.O. MBA with Lucio Minces, M.D.

05/23/2024

THREE CELL LINES



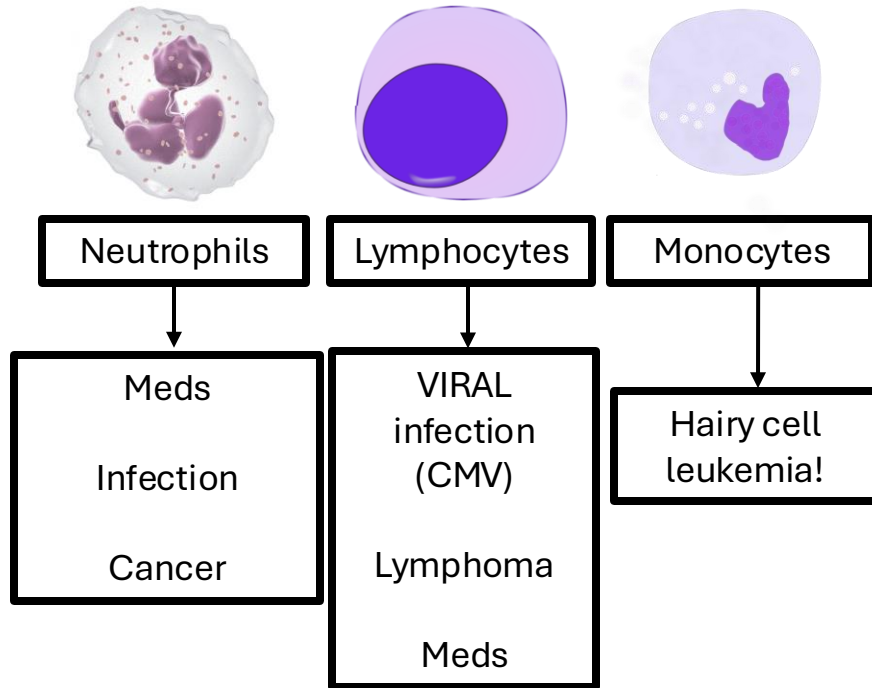
WBC



Basic Understanding Normal range 4-11 x10⁹/L

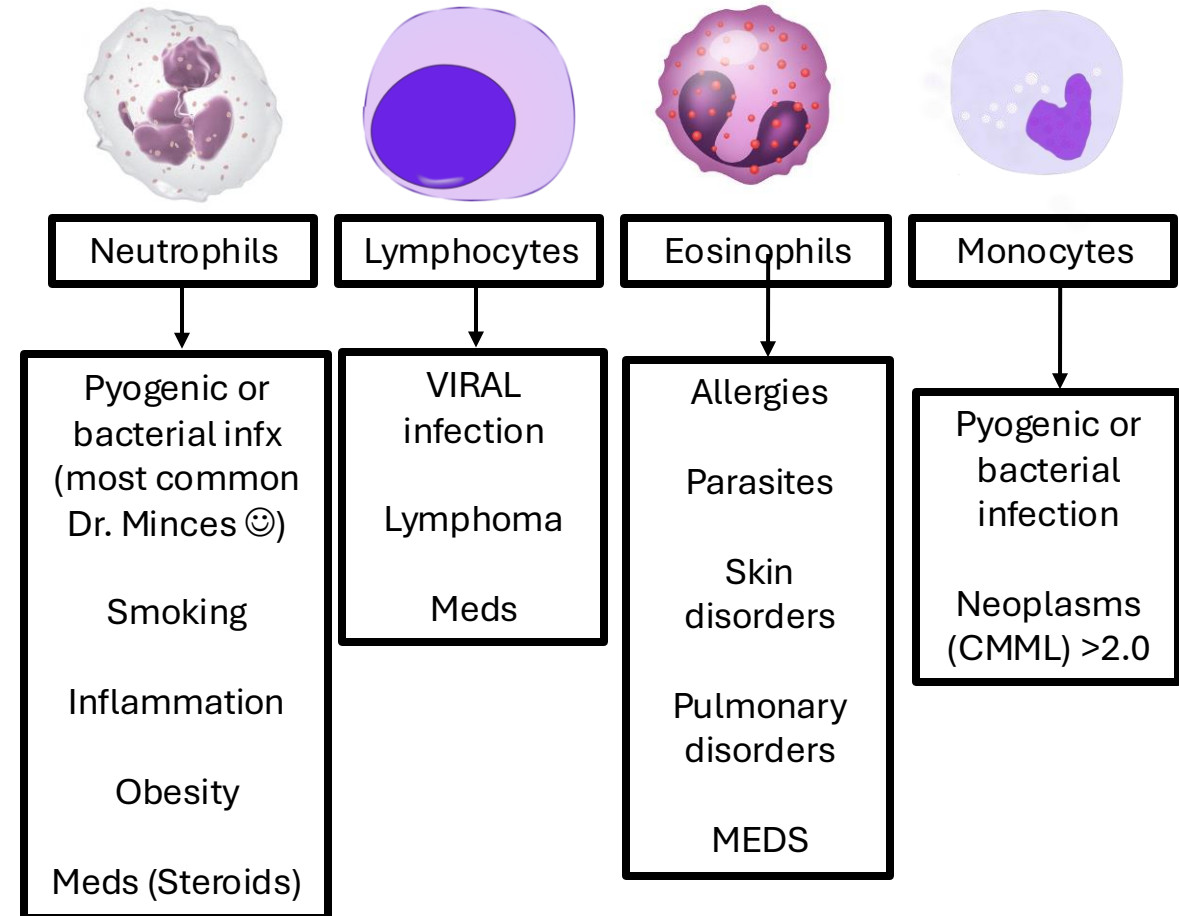
Leukopenia <4 K

- Watch Trends of ABSOLUTE counts!!
- DIFFERENTIAL



Leukocytosis > 11K

- Watch Trends of ABSOLUTE counts!!
- DIFFERENTIAL



WBC Emergencies/Urgent

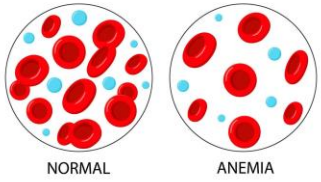
- WBC count <0.5 (risk of infections)
- WBC count >100 (sludging, bleeding)
- ANY New BLASTS detected on the differential

HGB (#)

HCT %



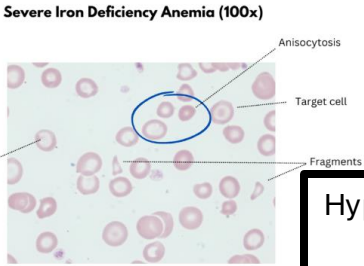
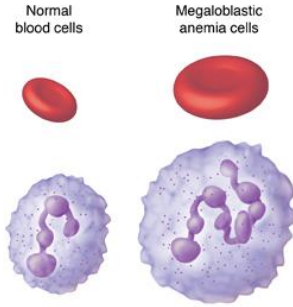
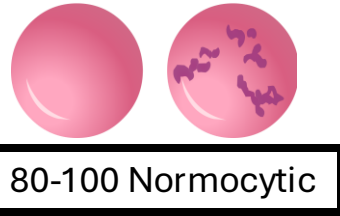
Basic Understanding Normal range ~11.5-16.5 x10⁹/L



ANEMIA <11.5

ERYTHROCYTOSIS >16.5

• Look at MCV



Hypoproliferative
Low Retic

Hyperproliferative
High retic.

>100 Macrocytic

<80 Microcytic

CKD
Liver dz
Inflammation
Bone marrow disease

Hemolysis
Bleeding

B12/folate def
Meds
EtOH
Hypothyroid
MDS (if persistent)

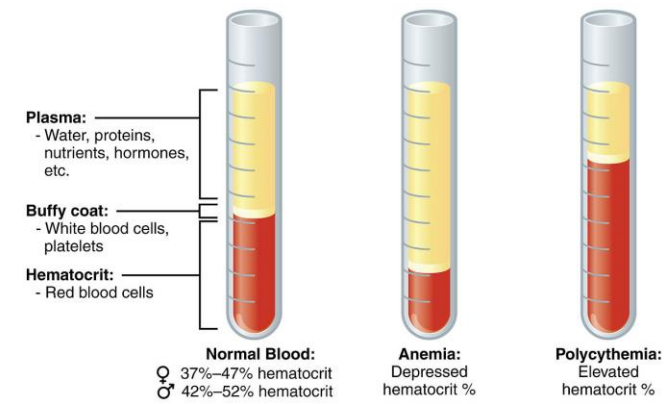
Iron deficiency, iron deficiency, iron deficiency
Thalassemia. Clue (Elevated RBC)

Primary
Low or Nm EPO

Secondary
High EPO

Congenital
Polycythemia vera
Get a JAK2 mutational study!

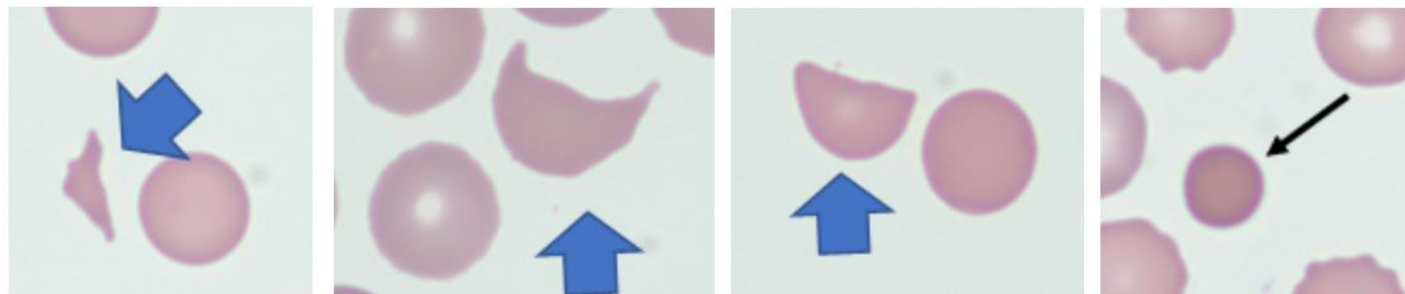
Congenital
Acquired (chronic hypoxemia)
• Renal dz
• Lung dz
• Smoking (most common)
• Meds (testosterone)
• Sleep Apnea



HGB Emergencies/Urgent

- Acute blood loss with hemodynamic instability
- Hematocrit >50 with symptoms of sludging (chest pain, visual loss, severe headache, etc)
- Hemolysis (Look for **Schistocytes or spherocytes** on blood smear, low haptoglobin, elevated LDH, elevated bilirubin (WHY?))

Schistocyte types

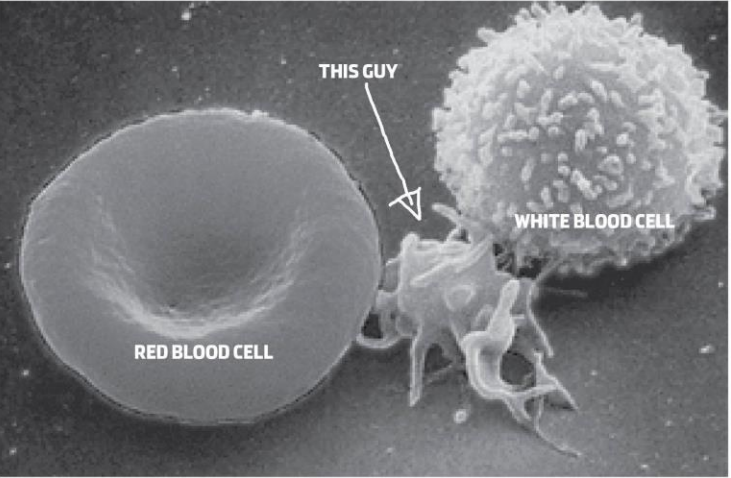


Triangular cell

Horn cell

Helmet cell

Microspherocyte

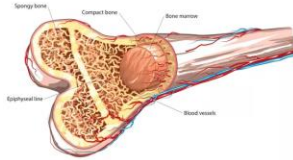
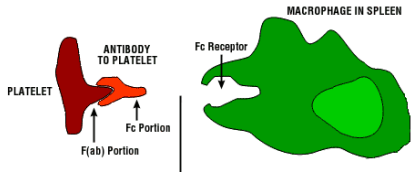


PLT

Basic Understanding Normal range 150-450

Thrombocytopenia

<150



Increased peripheral destruction

Hypersplenism

Bone marrow conditions

- ITP
- TTP (look for schistocytes)
- DIC
- HUS
- Infection

Lymphomas

Liver disease – alcohol, Hep C

Note: Liver disease leads to portal hypertension → hypersplenism → decreased production of TPO (produced by liver)

Aplasia: Drugs, idiopathic

Infiltration: Leukemia, lymphoma, myeloma, MDS

B12/Folate deficiency

Thrombocytosis

>450

Primary – Chronic and slowly progressive

Secondary – More acute

JAK2, CALR and MPL mutations to screen for a chronic myeloproliferative disorder (ET, PMF)

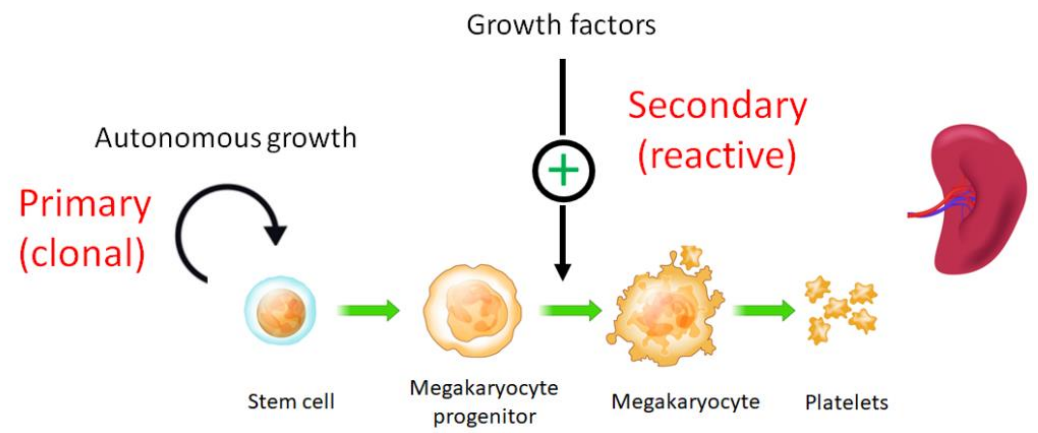
Clues: Splenomegaly, bleeding

Acute infection

Solid organ malignancy

IRON DEFICIENCY!

Inflammatory conditions (sarcoidosis, autoimmune, etc.)



Platelet Emergencies/Urgent

- Platelet count $<10,000$ (spontaneous bleeding)
- Platelet count $>1,000,000$ (in the setting of primary thrombocytosis) – risk for arterial and blood clots

CBC clues to a bone marrow condition and further evaluation.

- Pancytopenia (all cell lines affected)
- Bicytopenia (same)
- Cytopenias with lymphocytosis, lymphadenopathy or splenomegaly (lymphoma)
- Any dysplasia noted on peripheral smear (MDS)
- BLASTS on peripheral smear (MDS or AML)
- Cytopenias (most commonly anemia) with Rouleaux – need to exclude a plasma cell process

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- Contact carla.griffin@centracare.com with questions or for a CME transcript.