

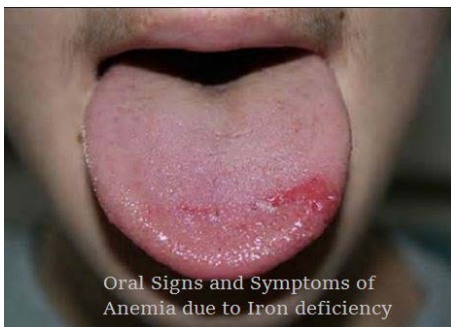
Disclaimer: This summary is meant to be for the Lab pictures only; so it contains very little details about the disease itself, to make it easier study each case and come look at the pictures after.

Lecture 1 + 2

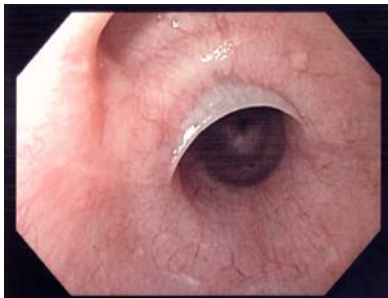
1- Iron Deficiency Anemia

***Clinical Findings:**

1. Glossitis
2. Spooning of the fingernails

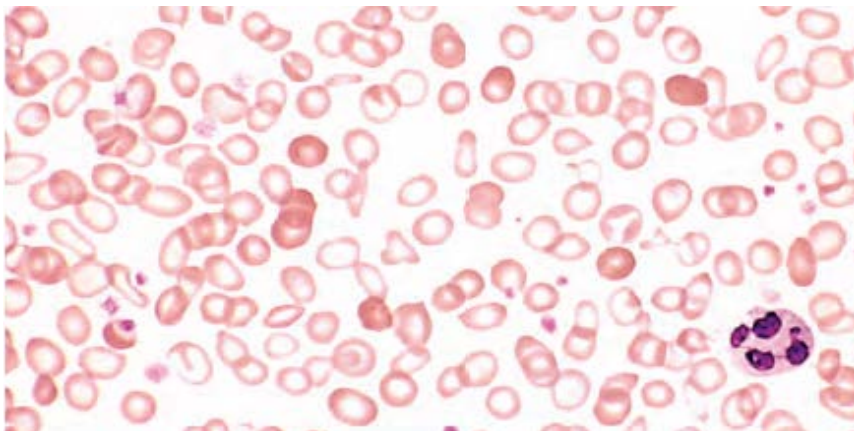


3. Esophageal web

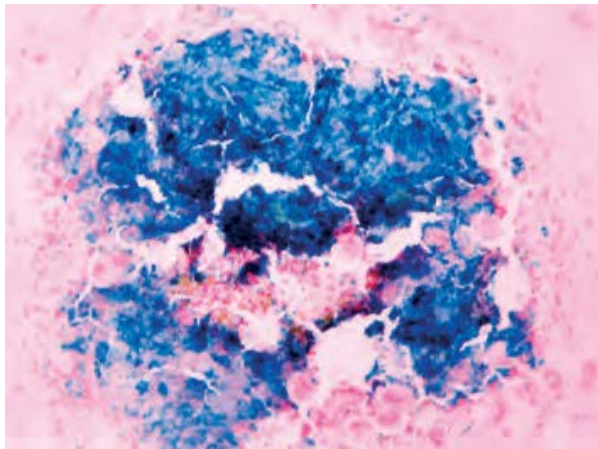
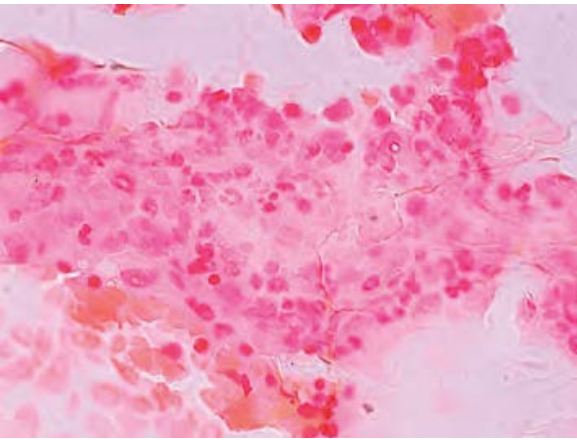


***Laboratory Findings:**

1. Microcytic (small) Hypochromic (pale from the center) RBCs



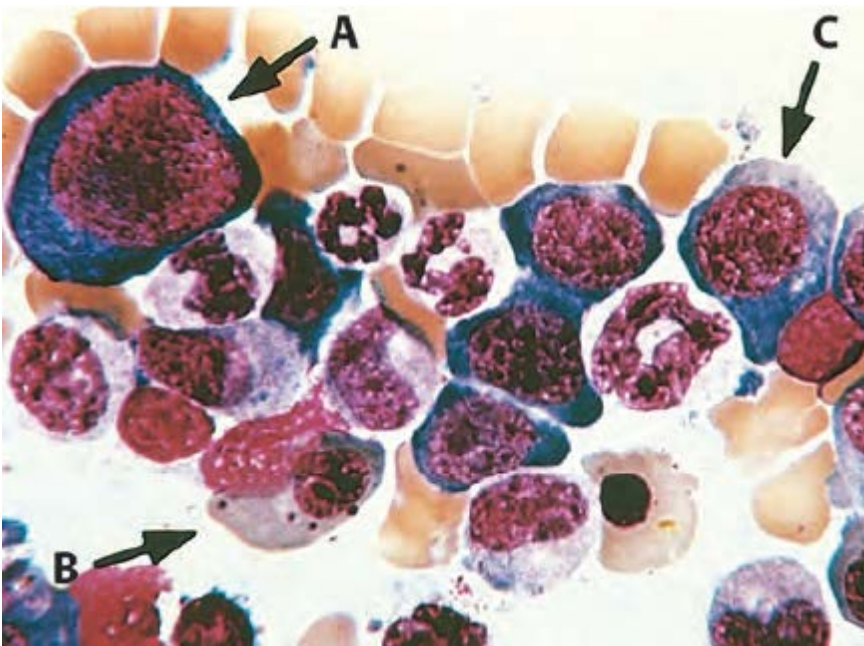
2. Low Iron levels



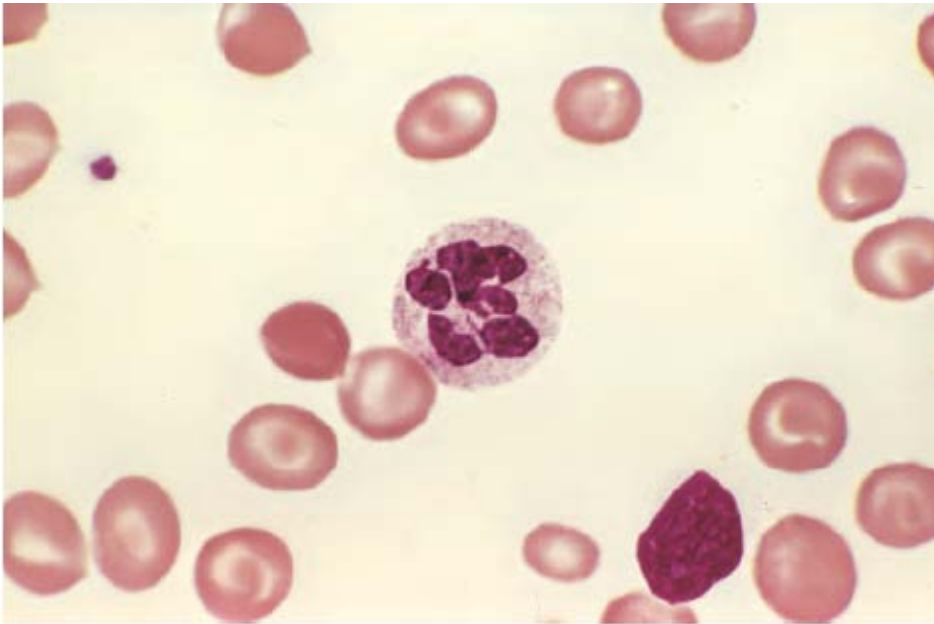
The left image shows a completely iron depleted bone marrow. The lower right image depicts a normal bone marrow with normal iron (blue).

2- Megaloblastic Anemia

1. Nuclear-cytoplasmic asynchrony; the erythroid precursor cells are larger than normal and show immature nuclei, however, the cytoplasm shows normal maturation.

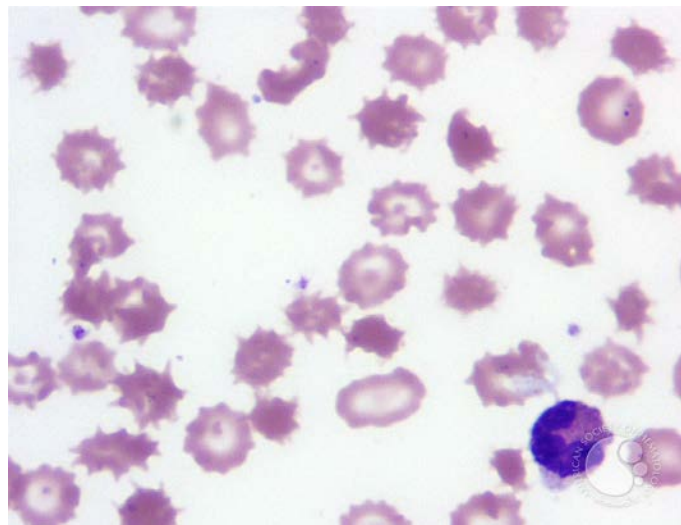
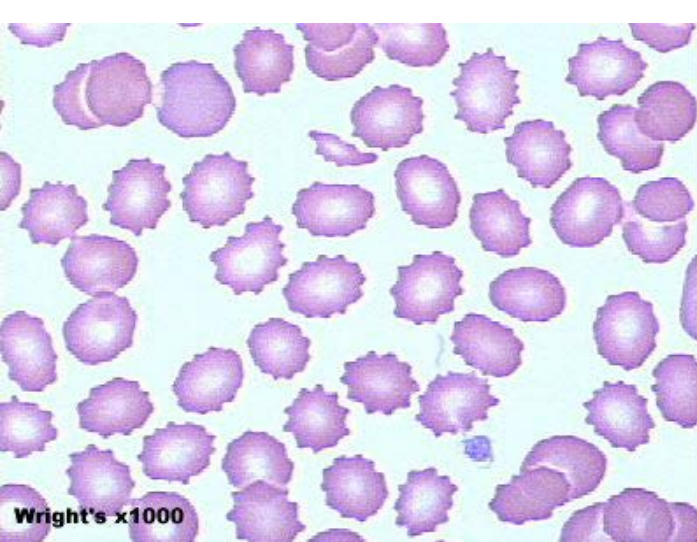


2. Hypersegmented neutrophils (neutrophils with nuclei showing more than 4 segments)



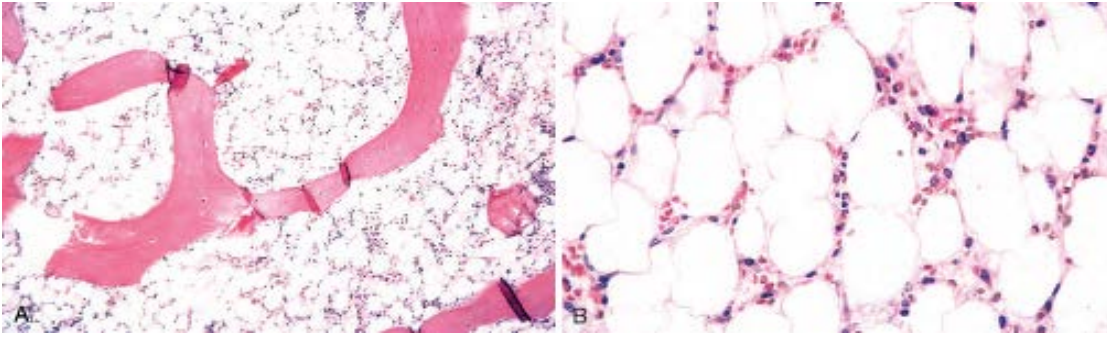
3- Anemia of Chronic Liver Disease

Characterized by presence of spur cells (also known as acanthocytes) which are large erythrocytes covered with spike-like projections that vary in width, length, and distribution.



4- Aplastic Anemia

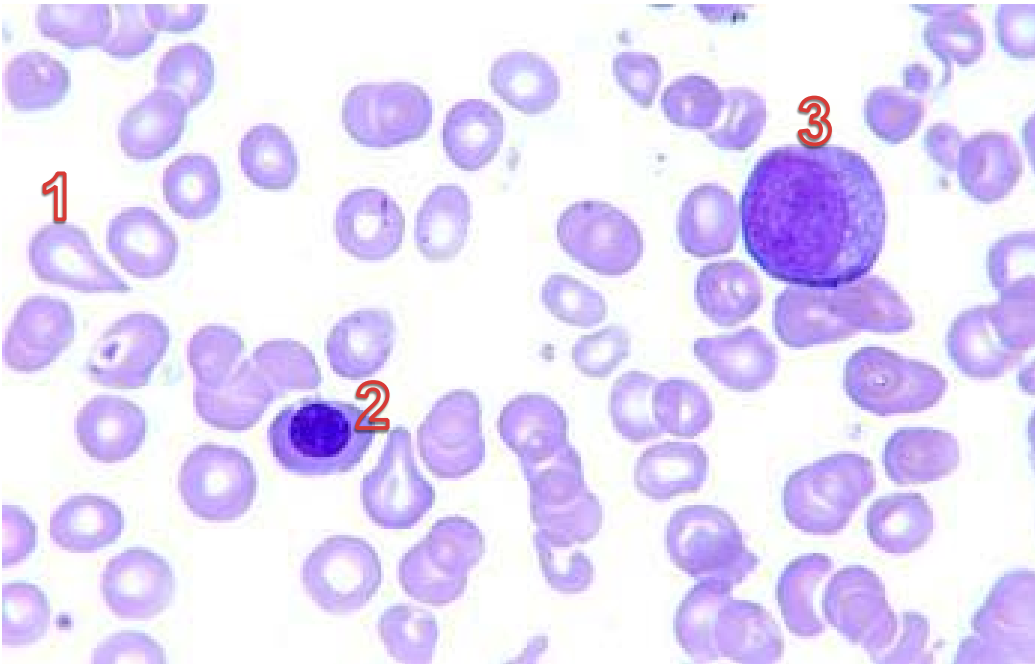
The bone marrow is completely devoid of hematopoietic cells and is filled with adipose tissue instead.



(adipose tissue in the bone marrow, again)

5- Myelophthitic Anemia

Characterized by misshapen red cells, some resembling teardrops, immature granulocytic and erythrocytic precursors also may be present too.



1-tear drop RBC

2-immature erythroid precursor cell (the one with the nucleus)

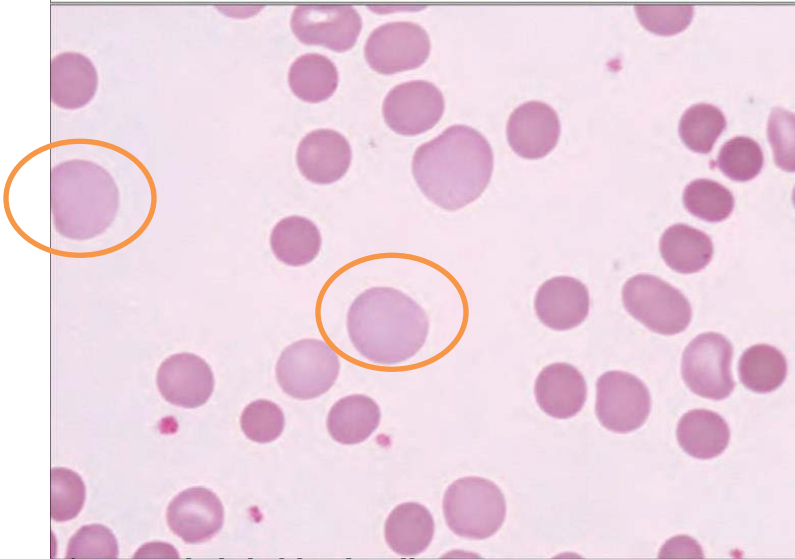
3-immature myeloid cell

Lecture 3

1- Hemolysis

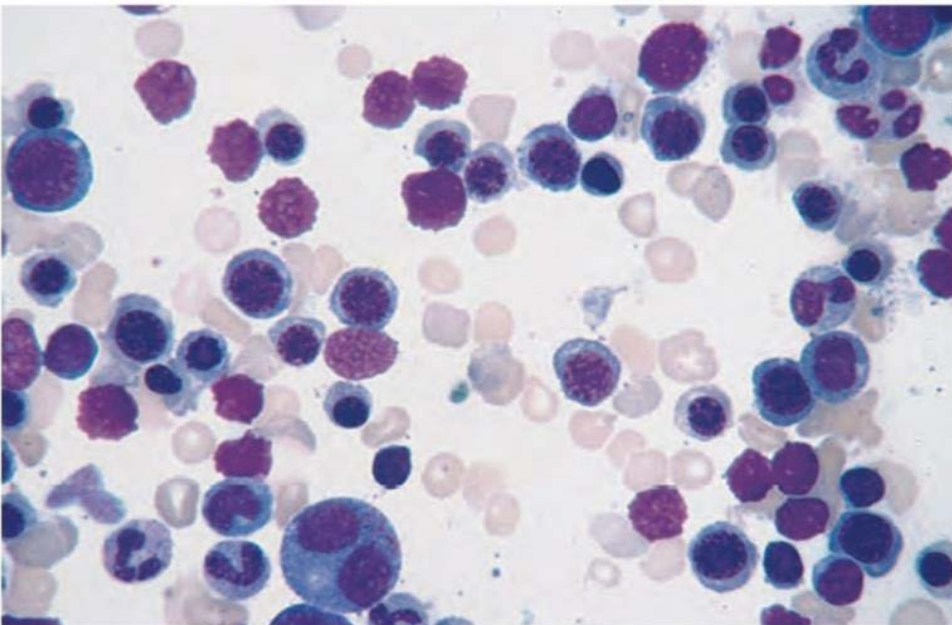
Characterized by:

1. High reticulocyte count

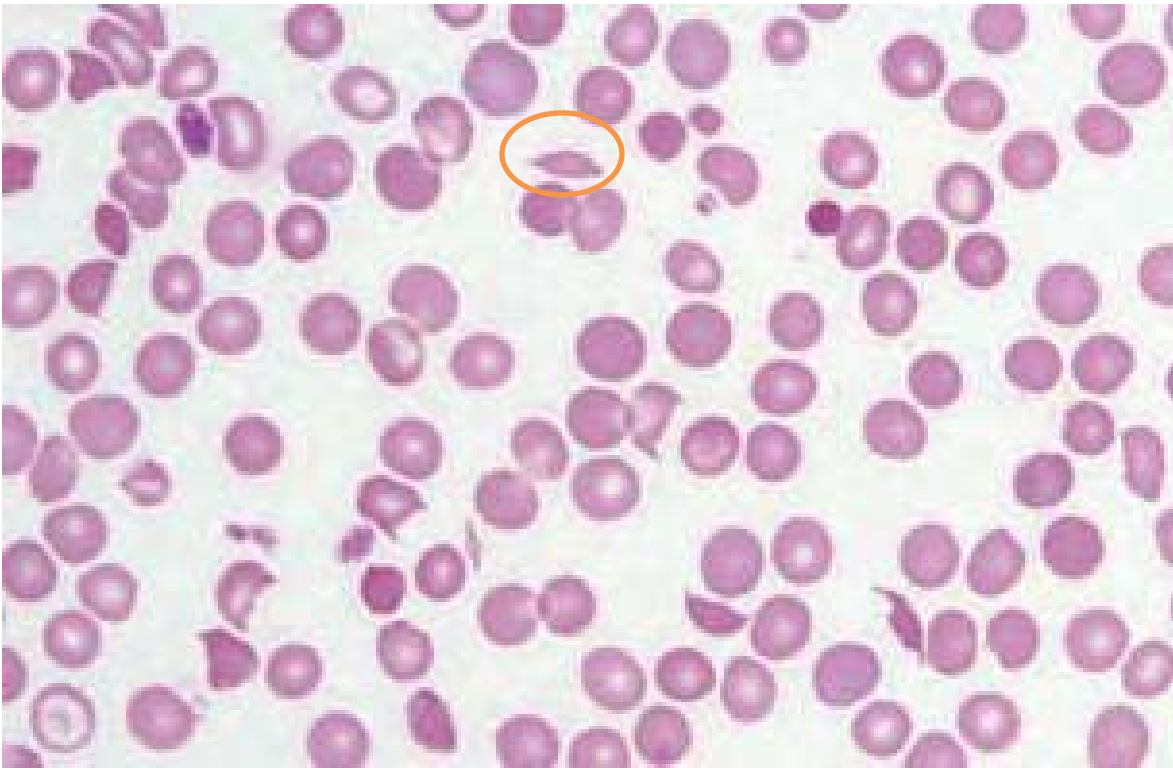


*Large, slightly bluish cells

2. Erythroid precursor cells hyperplasia

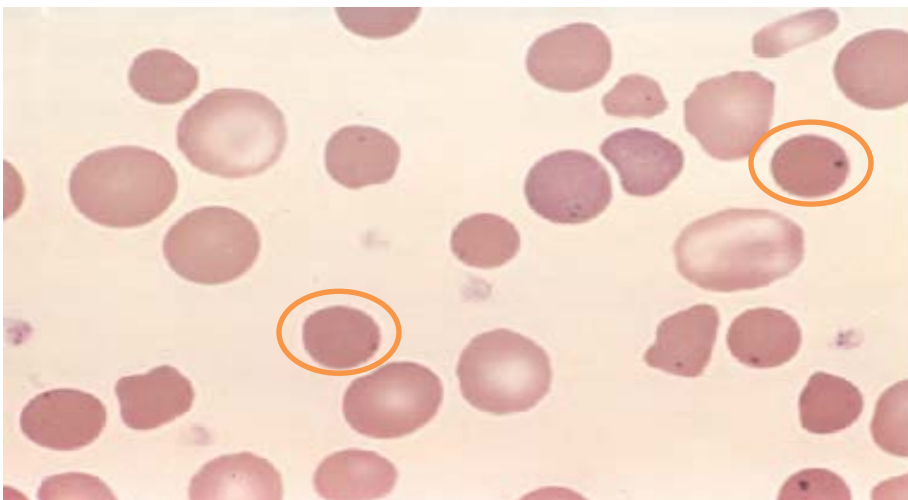


→ Schistocytes is a torn RBCs with a helmet shape, associated with some parasitic infections such as Malaria and babesia. (note from the sheet: Schistocyte is a marker of intravascular hemolysis and is not specific for a single cause of hemolysis)



Lecture 4

1- Spherocytosis



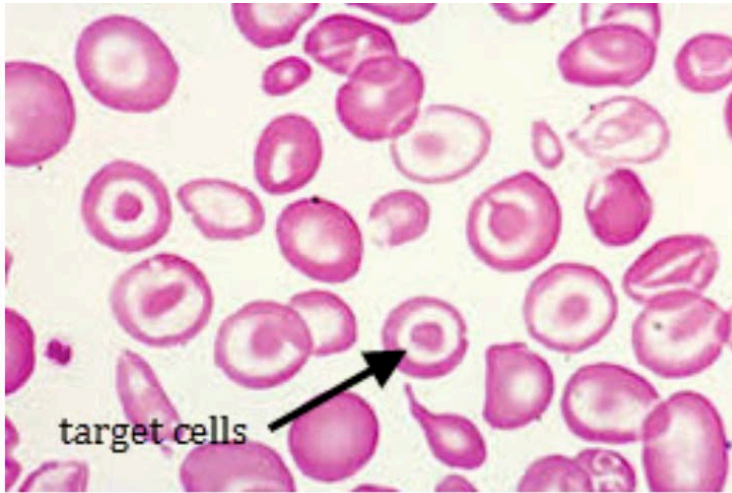
2- Sickle cell anemia (RBCs are sickle shaped)



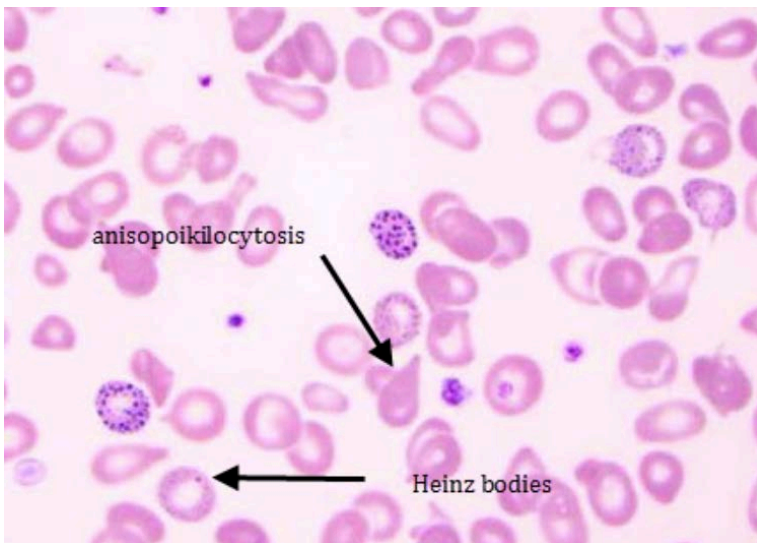
3- B-Thalassemia Major

Morphology:

1. Target Cells

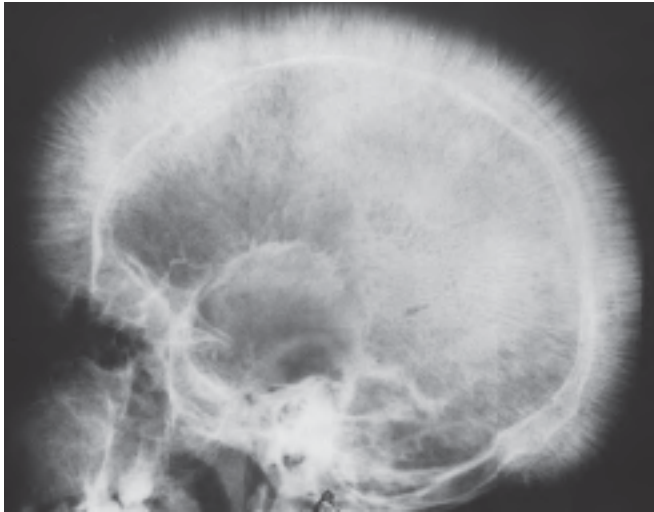


2. Heinz Bodies (inclusions within red blood cells composed of denatured hemoglobin)



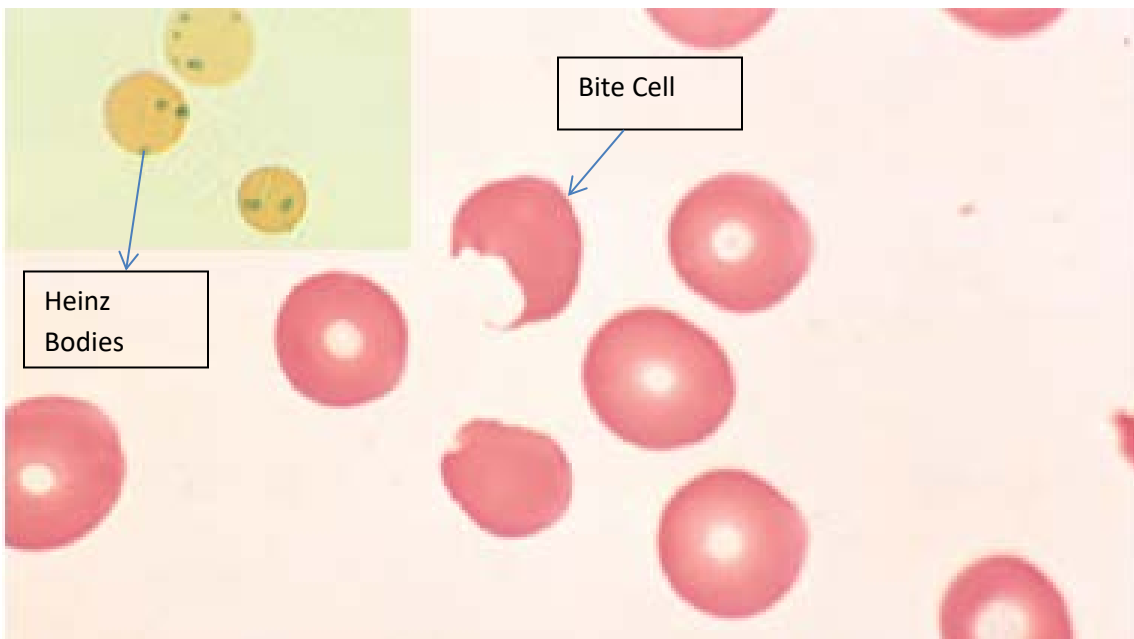
3. X-ray of the skull shows perpendicular radiations resembling a crew-cut).

(الأشياء اللي زي الشعرات)



4- Glucose 6-phosphate dehydrogenase (G6PD) deficiency

1. Heinz bodies (crystal violet stain)
2. Bite Cells (forms when cells try to pluck out Heinz bodies)

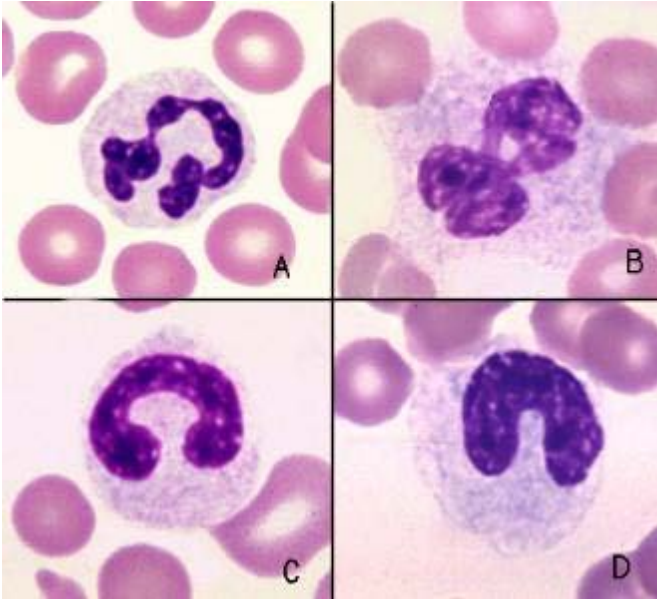


Lecture 6 (خمسة ما فيه صور)

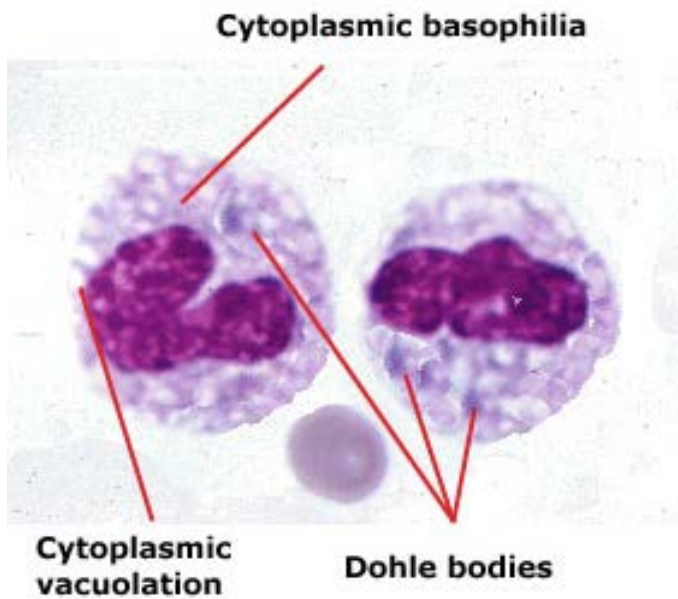
1- Neutrophilia

Morphology:

1. Toxic Granulation (cytoplasm appear granulated)

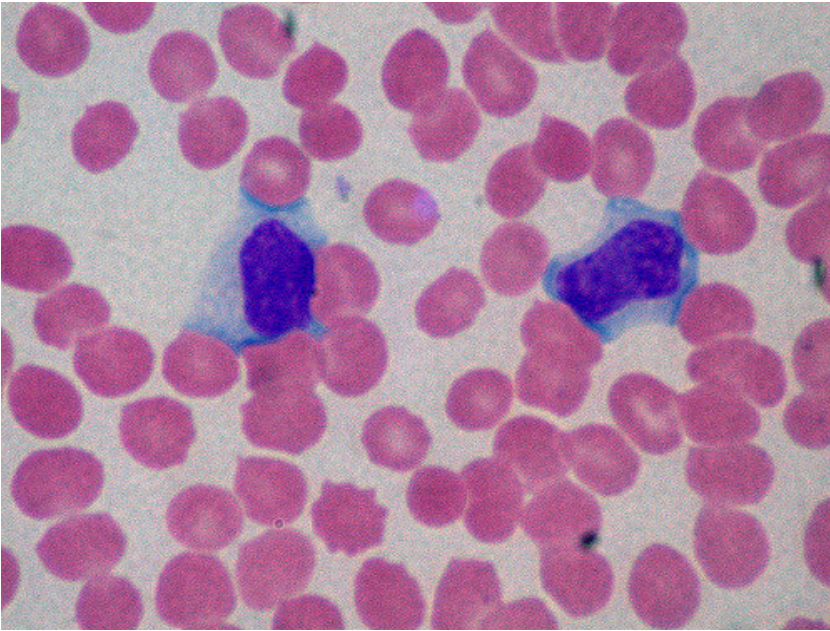


2. Cytoplasmic basophilia
3. Vacuolization of the cytoplasm
4. Döhle bodies

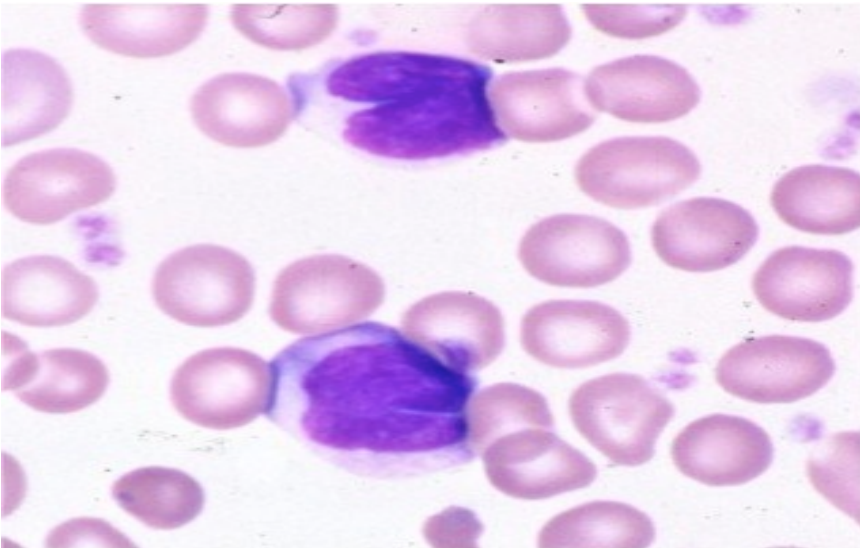


2- Lymphocytosis

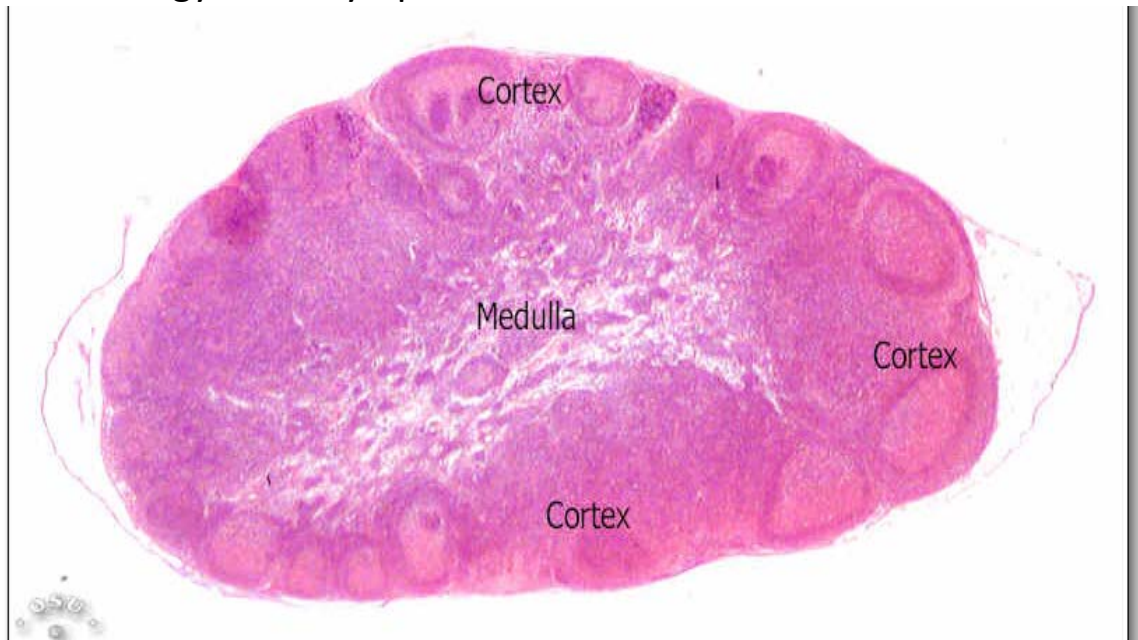
2.1 Infectious mononucleosis (atypical lymphocyte with abundant cytoplasm (somehow the cytoplasm looks like a skirt ☺))



2.2 Pertussis (whooping cough) -> a lymphocytosis; so it has almost the same look



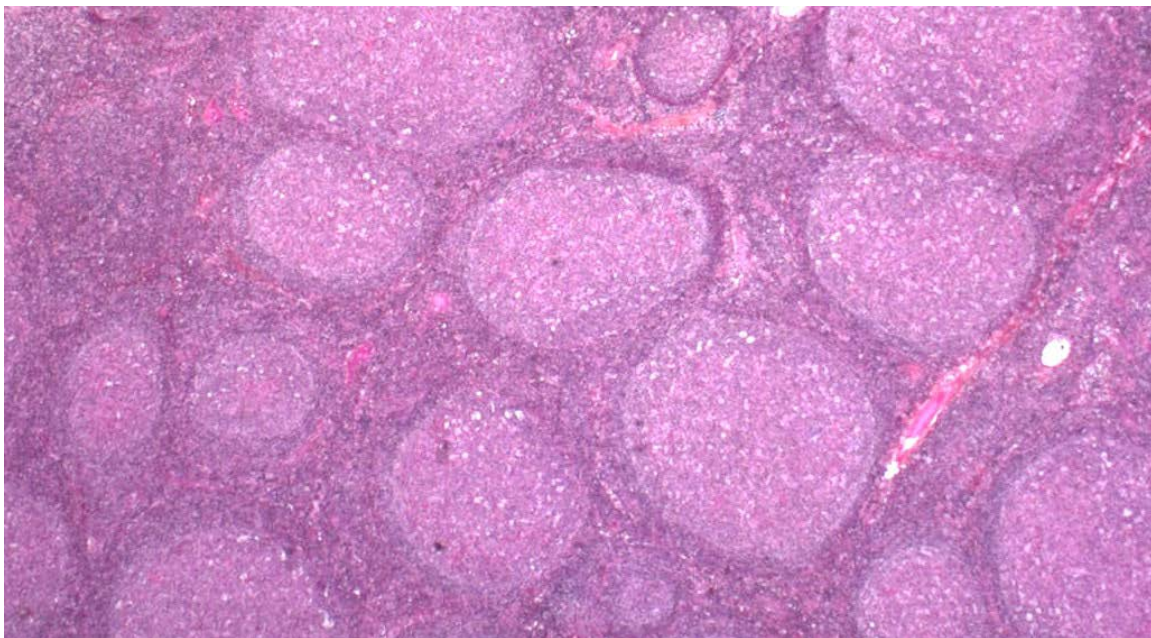
3- Histology of the lymph node



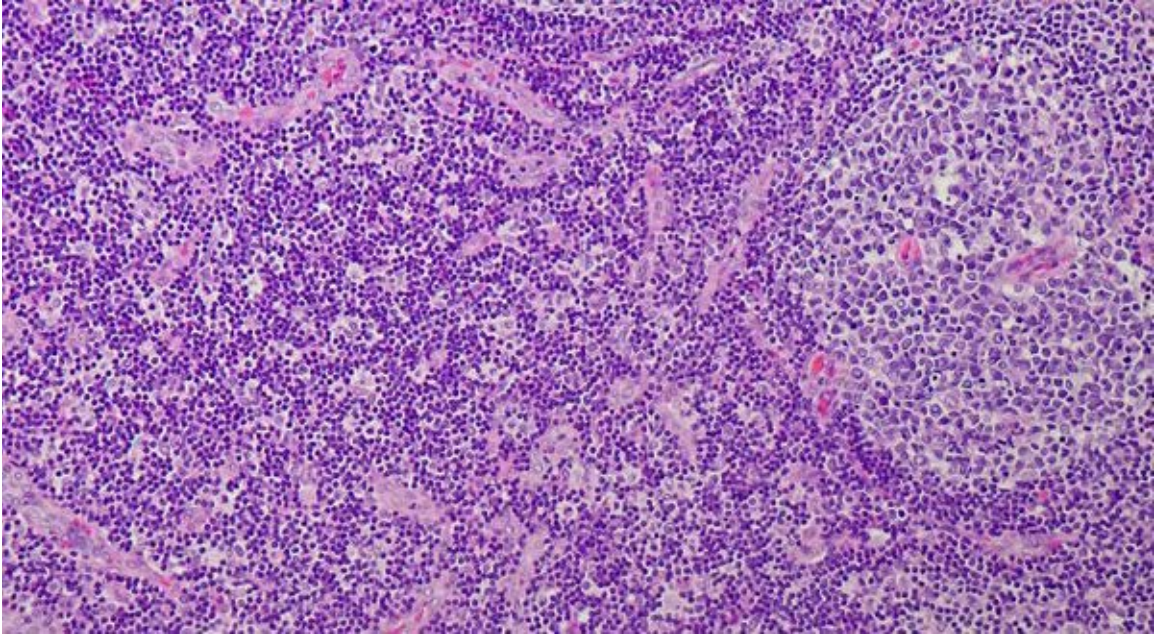
4- Chronic Nonspecific Lymphadenitis

It assumes three patterns:

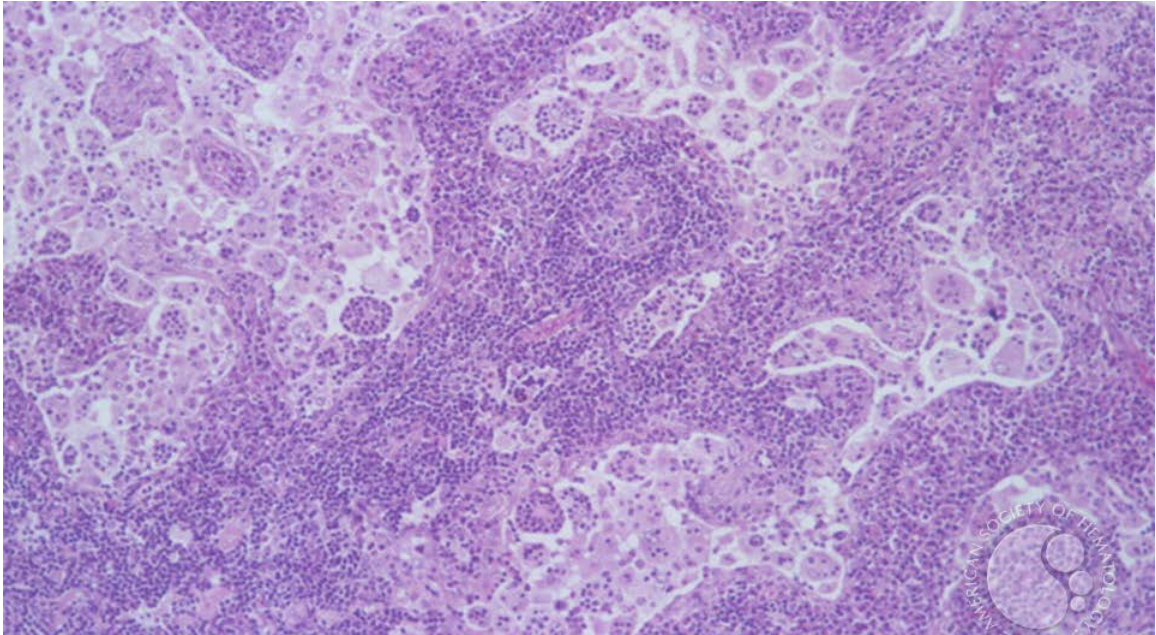
- a. Follicular hyperplasia (not to be confused with follicular lymphoma, here node architecture is preserved and the germinal centers varies in the shape and size)



b. Paracortical hyperplasia (proliferation of T-cells in the interfollicular areas)



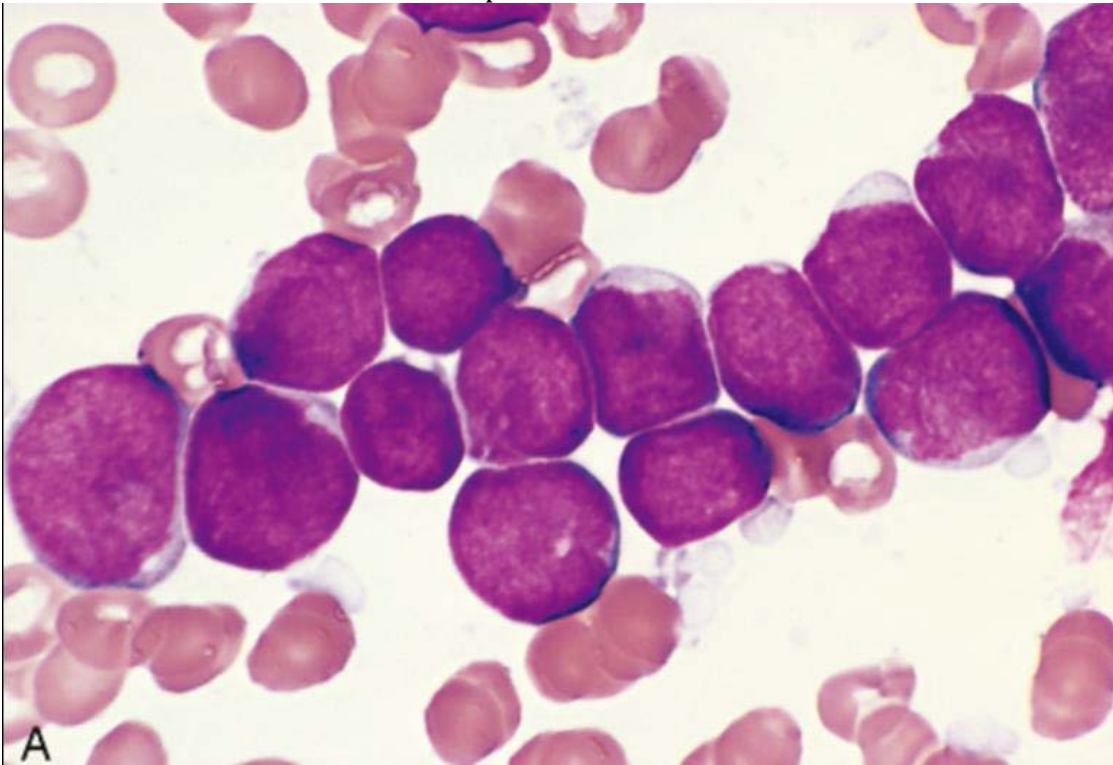
c. Sinus histiocytosis (sinuses are filled with macrophages and histiocytes.)



Lecture 7

1- Acute Lymphoblastic Leukemia

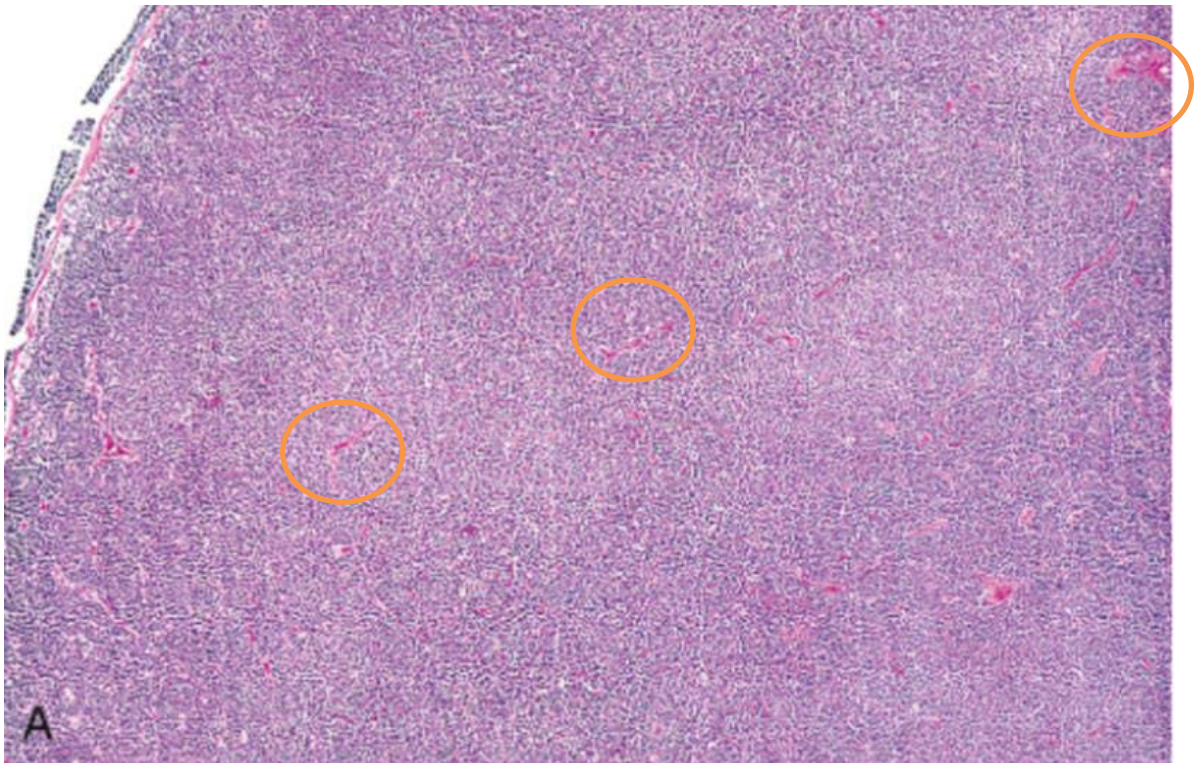
Blast Cells with fine chromatin and prominent nucleoli



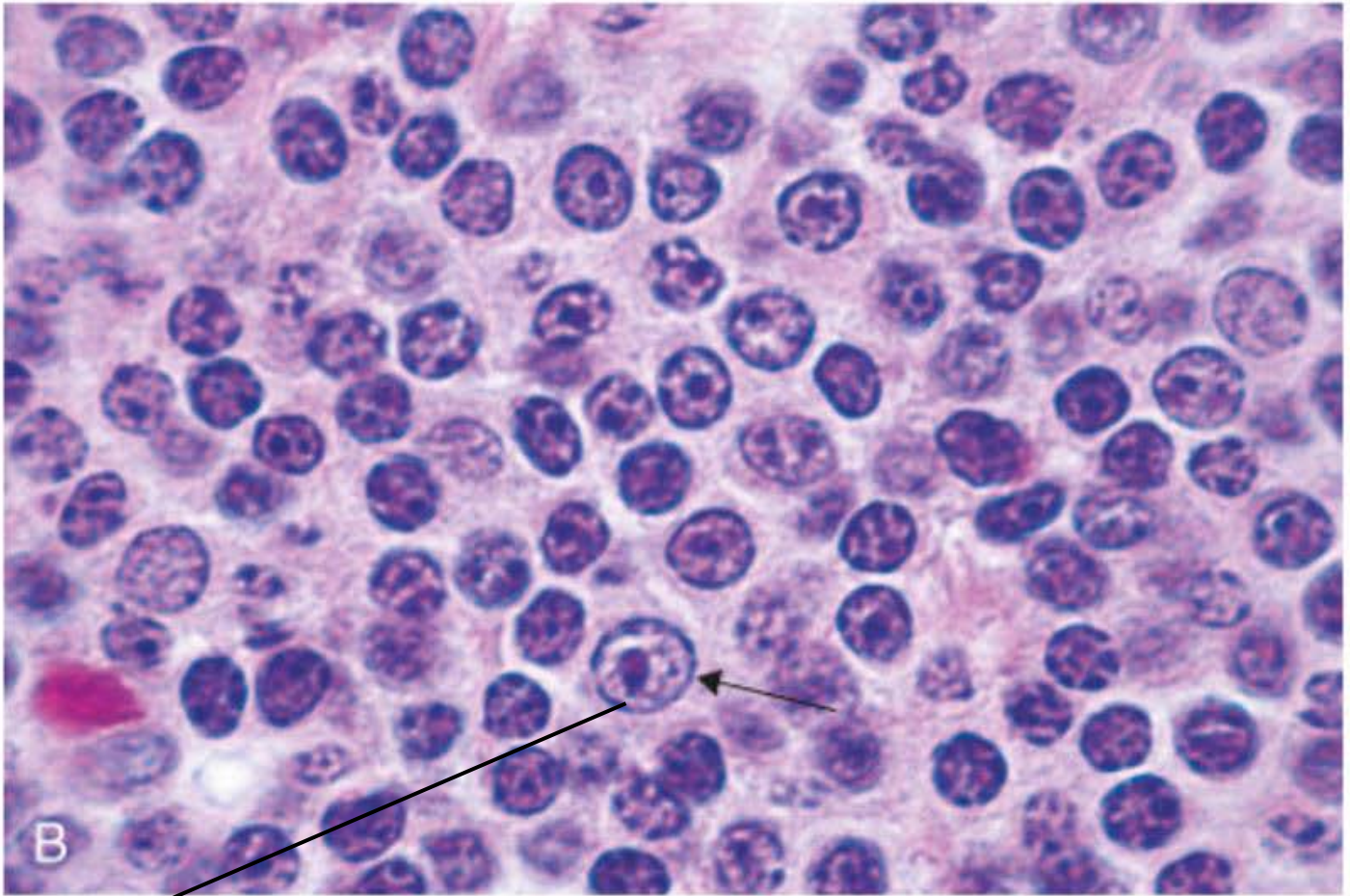
2- Chronic Lymphocytic Leukemia & SLL

Morphology:

1. Sheets of lymphocytes within the lymph node

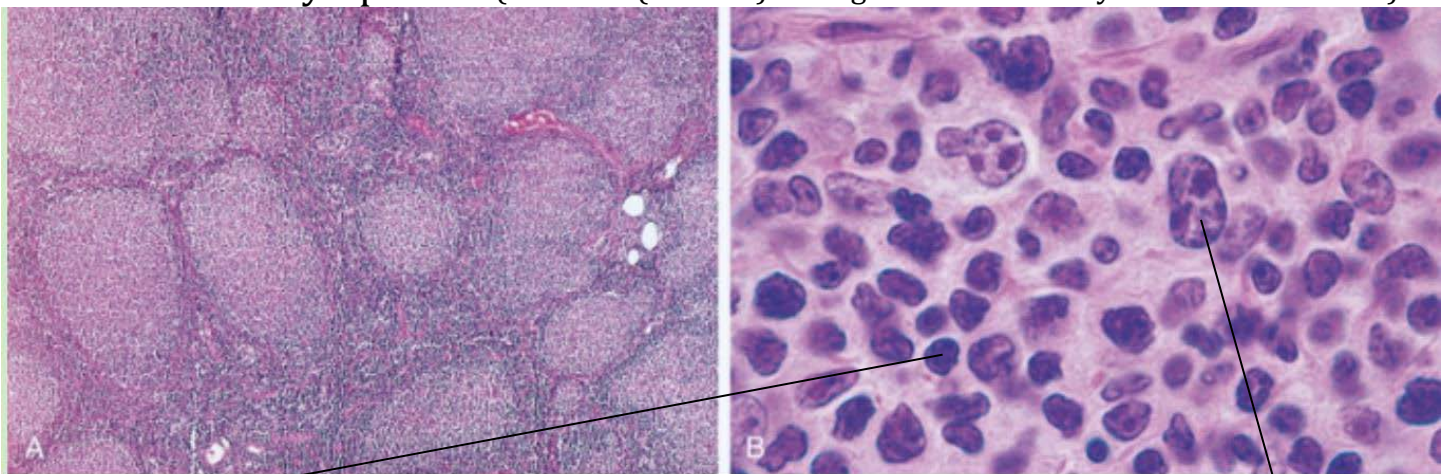


2. Small mature lymphocytes that appear like normal lymphocytes with dense chromatin and small nucleoli



A small percentage of lymphocytes show prominent nucleoli, called **Prolymphocytes**.

3- Follicular Lymphoma (follicular (nodular) arrangement of centrocytes and centroblasts)



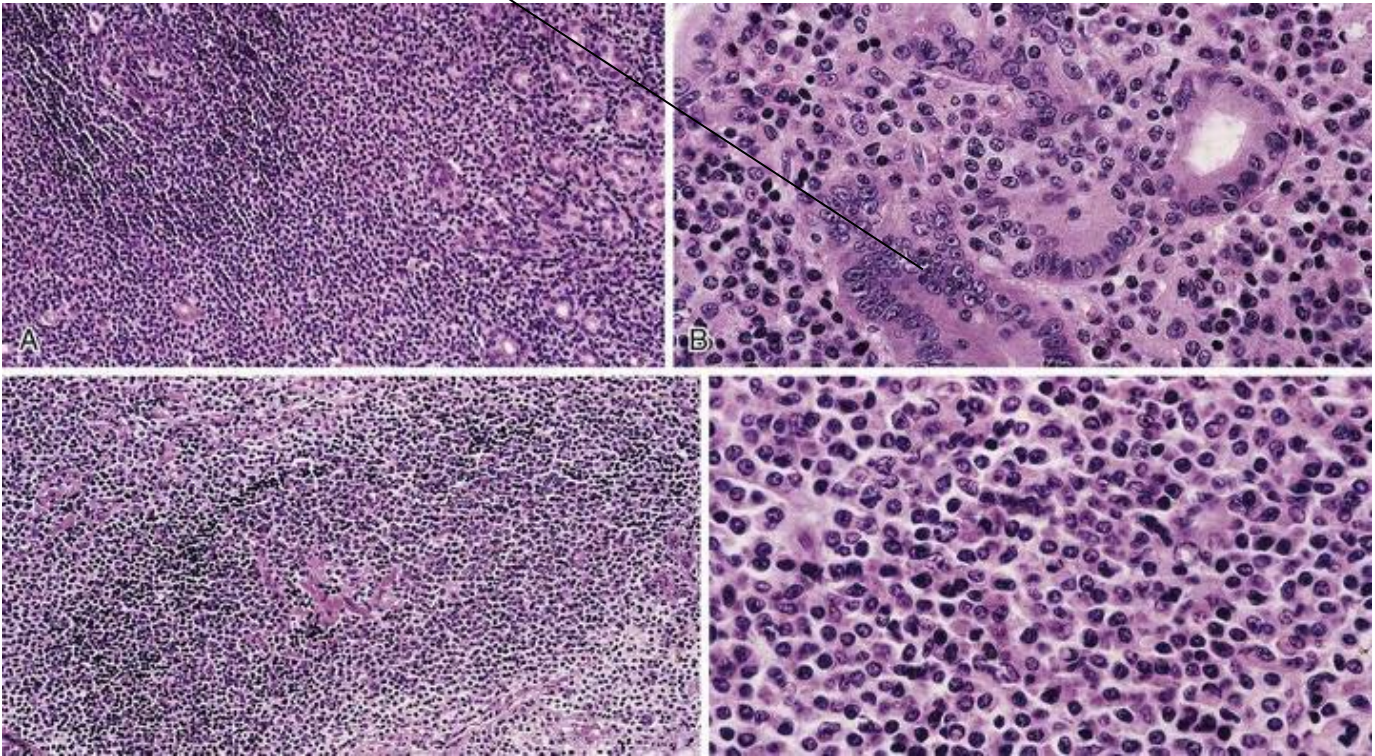
Centrocyte

centroblast

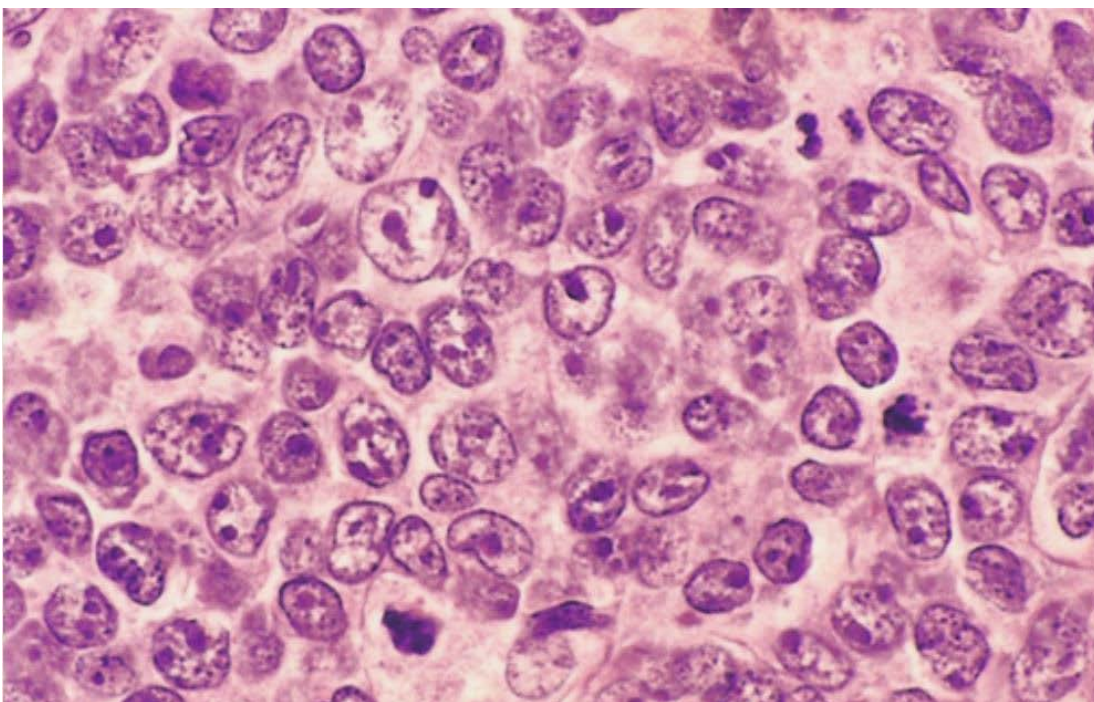
4- Extranodal Marginal Zone Lymphoma

Morphology:

1. lymphocytes are small to medium in size with variable cytoplasm
2. Lymphoepithelial lesions



5- Diffuse large cell lymphoma (presents as large cells with at least double the size of a normal lymphocyte and diffused arrangement)



Lecture 8

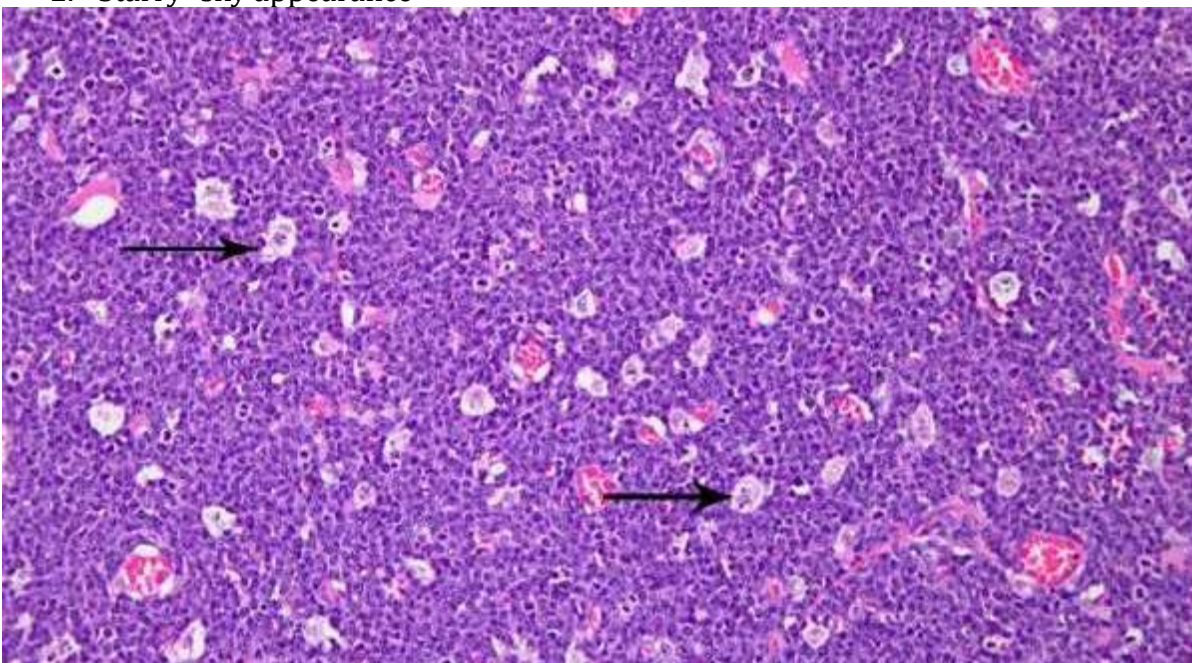
1- Burkitt Lymphoma

Clinical Presentation: (fastest growing tumor in humans)

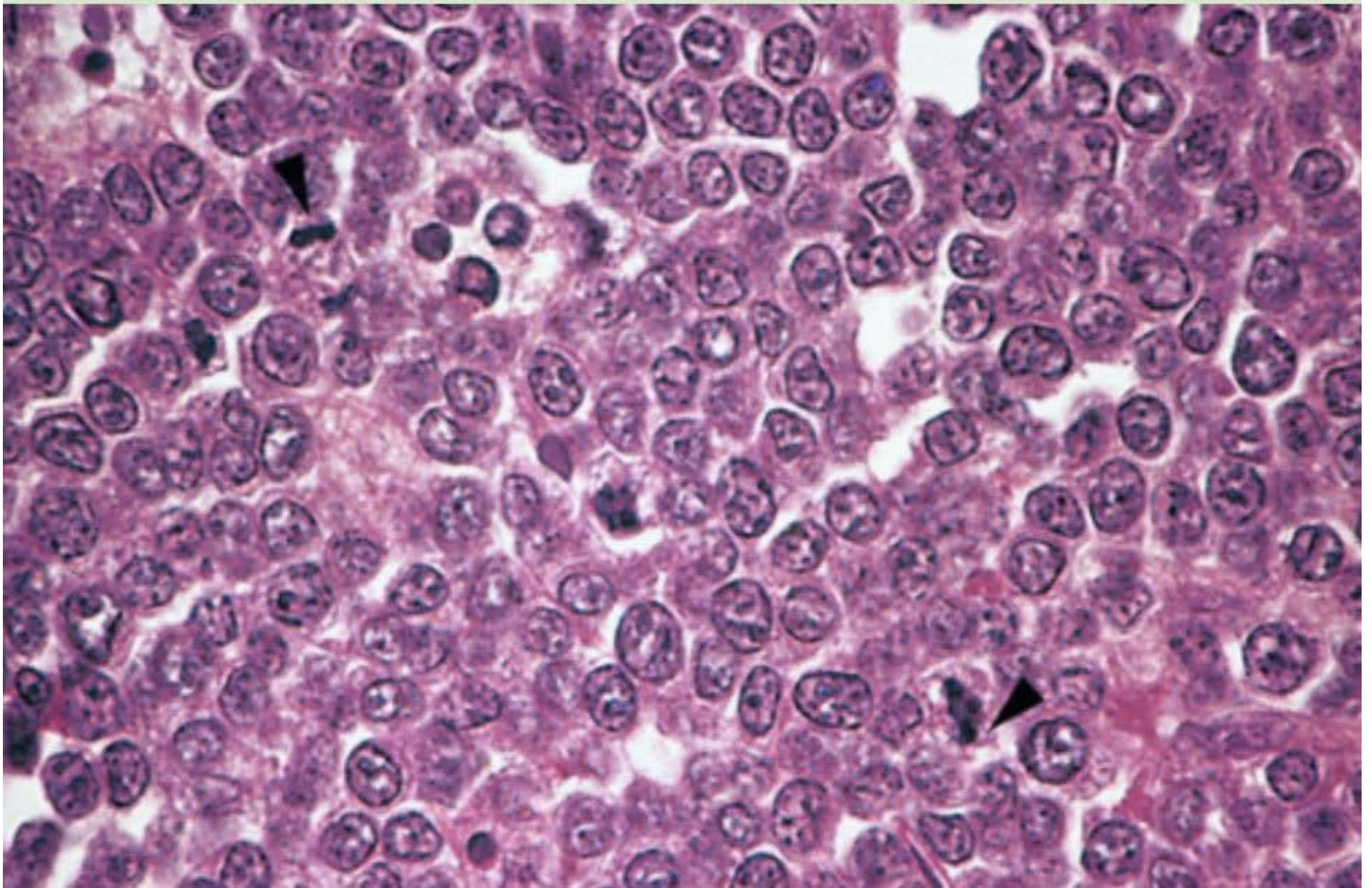


Morphology:

1. Sheets of medium sized cells
2. Starry -sky appearance



3. Variable cytoplasm, several nucleoli
4. Frequent mitosis



2- Plasma cell myeloma (multiple myeloma)

Morphology: (شكرا لصاحب الشيت)



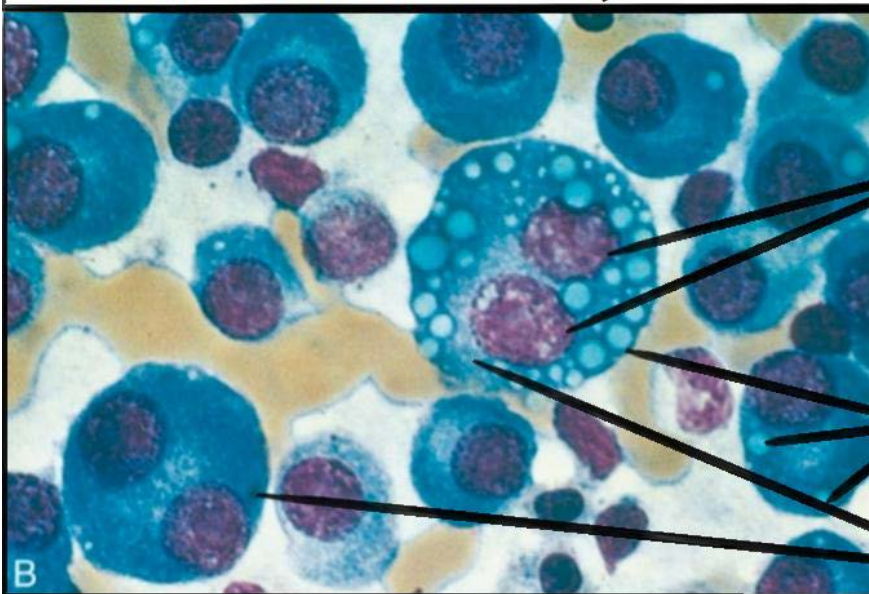
Lytic lesions:

most common in central axial bones such as the skull and the vertebra (still it can happen anywhere). caused by increase osteoclastic activity and decreased osteoplastic activity (increased RANKL activity).



Plasma cells:
have abundant
basophilic
cytoplasm,
prominent
perineuclear hof,
round eccentric
nucleus with
coarse chromatin
(clock-face)

Multiple myeloma cells may have normal plasma cell appearance and may have an abnormal appearance (binucleation, large size and vacuoles - which might be cytoplasmic or nuclear)

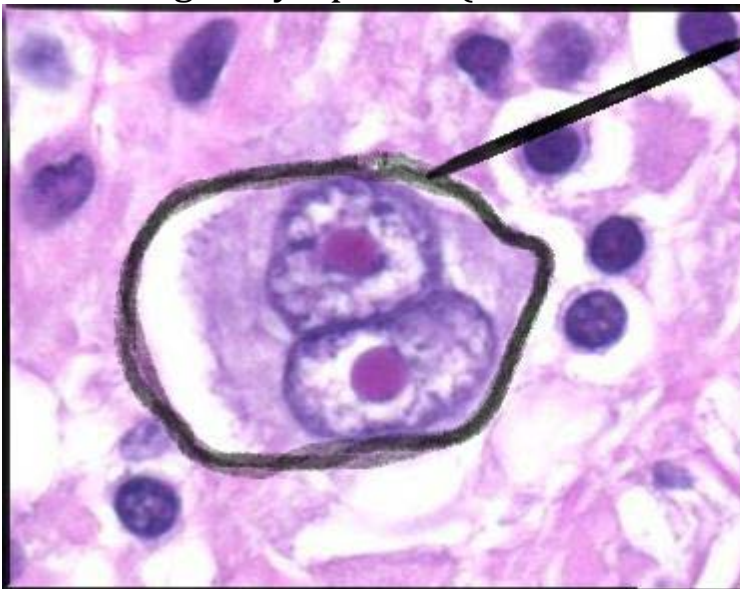


Binucleation

Vacuoles

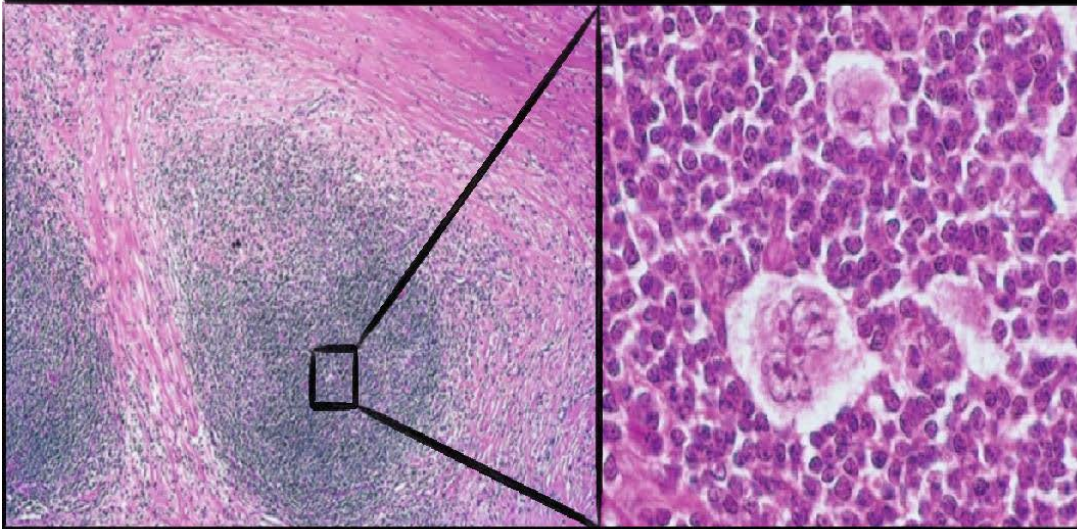
Large size

3- Hodgkin lymphoma (شکرا کثیر صاحب شیت 8)



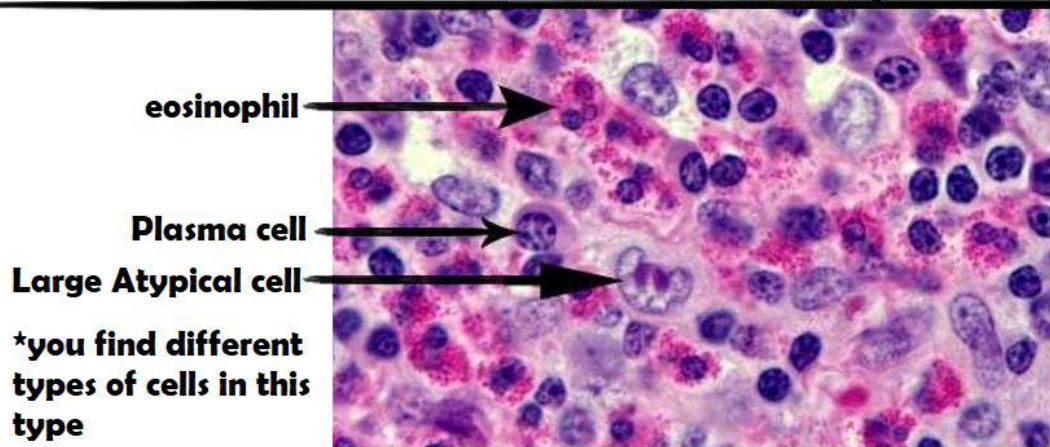
RS-cell:
***Huge cell (its nucleolus might be bigger than a normal lymphocyte).**
***two nuclei opposing each other, prominent nucleoli.**
it is hard to miss.

Classic HL: Nodular sclerosis

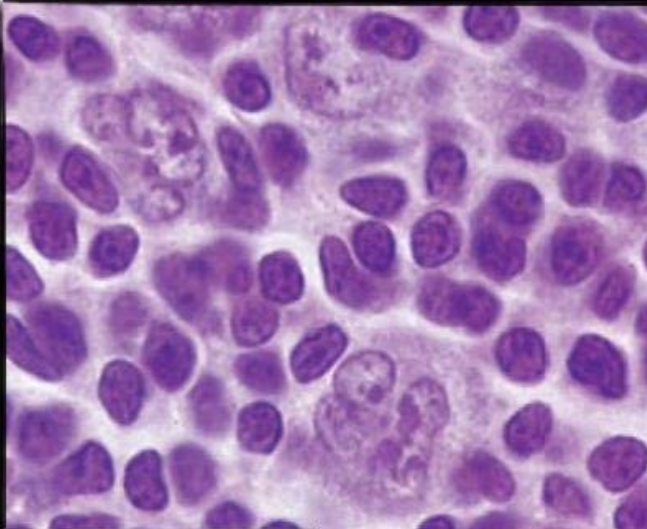


Contains nodules that are surrounded by a fibrous tissue, if we zoom in we will found RS-cells surrounded by benign reactive lymphocytes.

Classic HL: Mixed cellularity



Nodular lymphocyte predominant HL (NLP HL)



Large atypical cells called (Popcorn cells) surrounded by reactive lymphocytes and inflammatory cells

4- Mycosis fungoides

Clinical Presentation:

MF has three stages:

1- erythrodermic rash 2- plaque phase 3- Tumour phase

Patch



Plaque



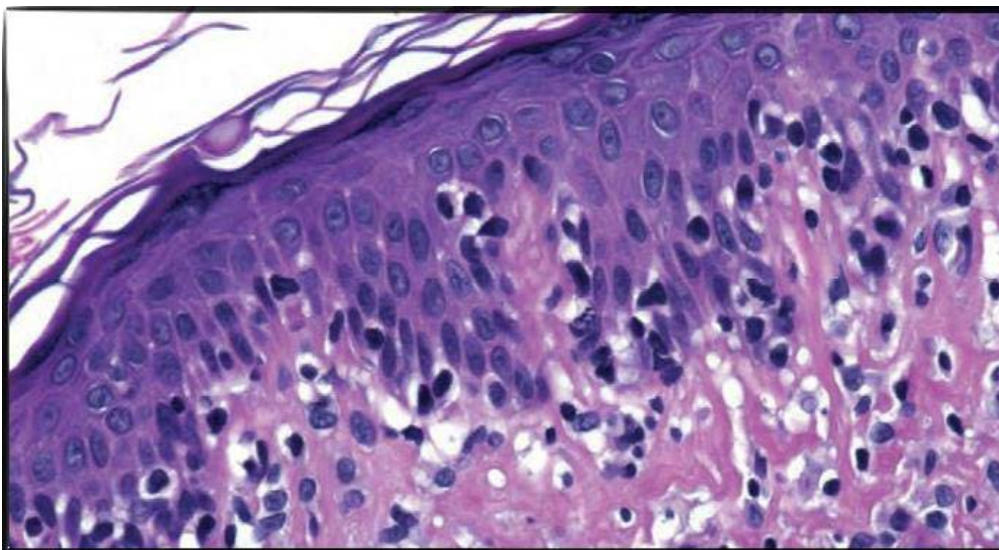
Tumor



Erythroderma

Note from the sheet: Patch: localized redness/ Plaque: can be felt (palpated)/ Tumor: causes necrosis/ Erythroderma: generalized redness.

Morphology:

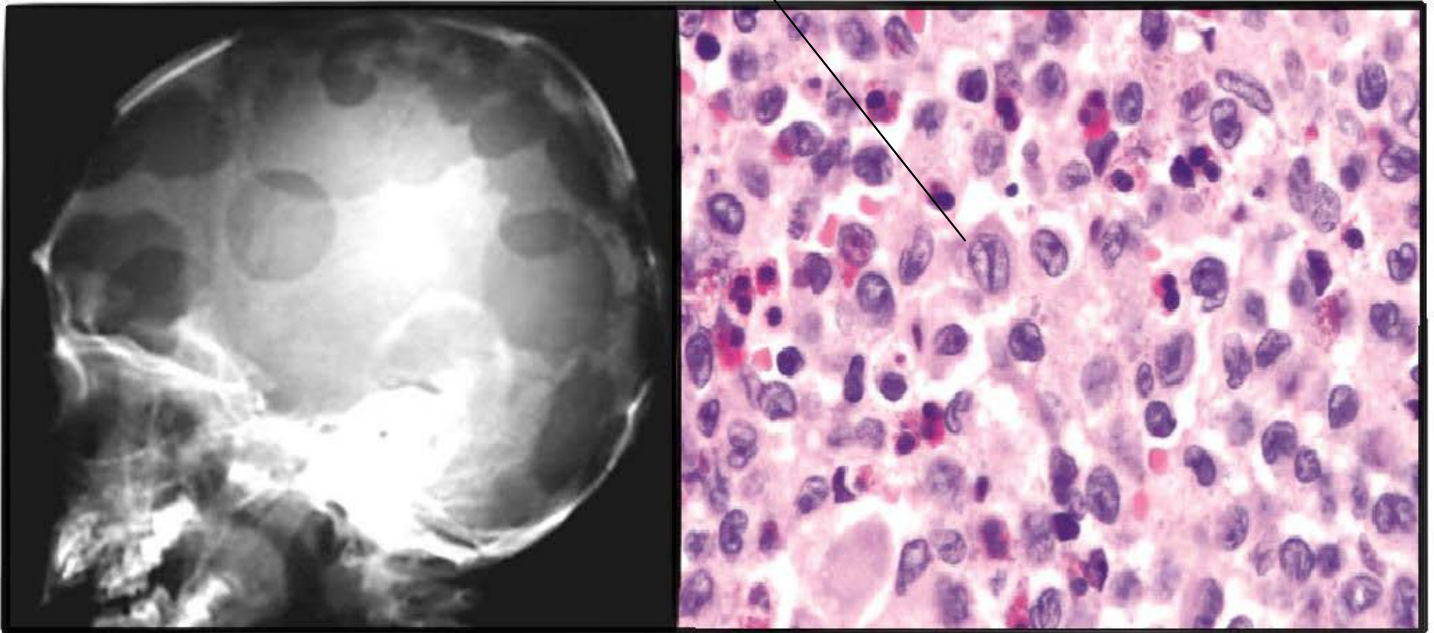


You can see the lymphocytes invading the epidermis which does not happen normally, indicates MF

5- Langerhans cell histiocytosis (A Histiocytic neoplasm)

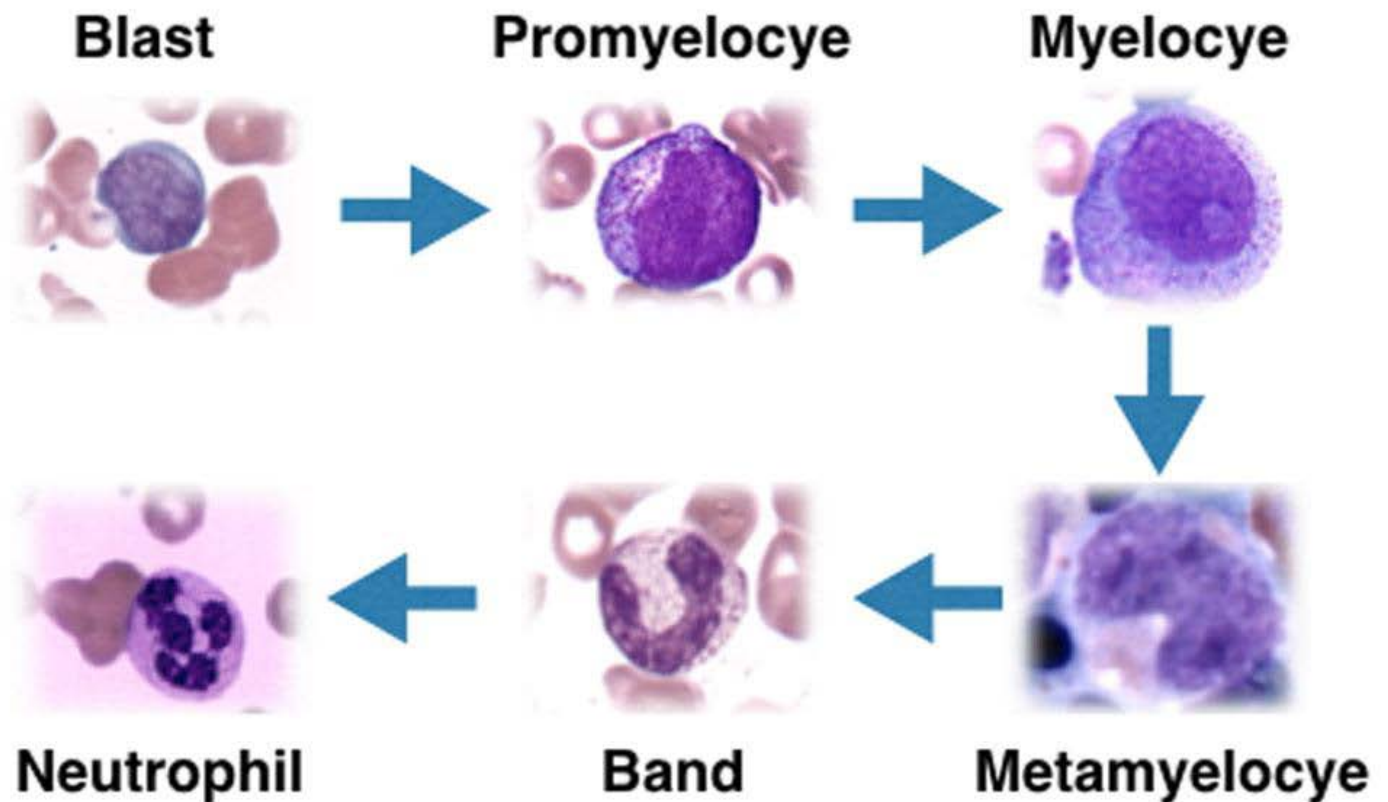
Morphology:

1. Lytic Lesions in the Skull
2. Langerhans cell histiocytosis cells have a coffee bean appearance



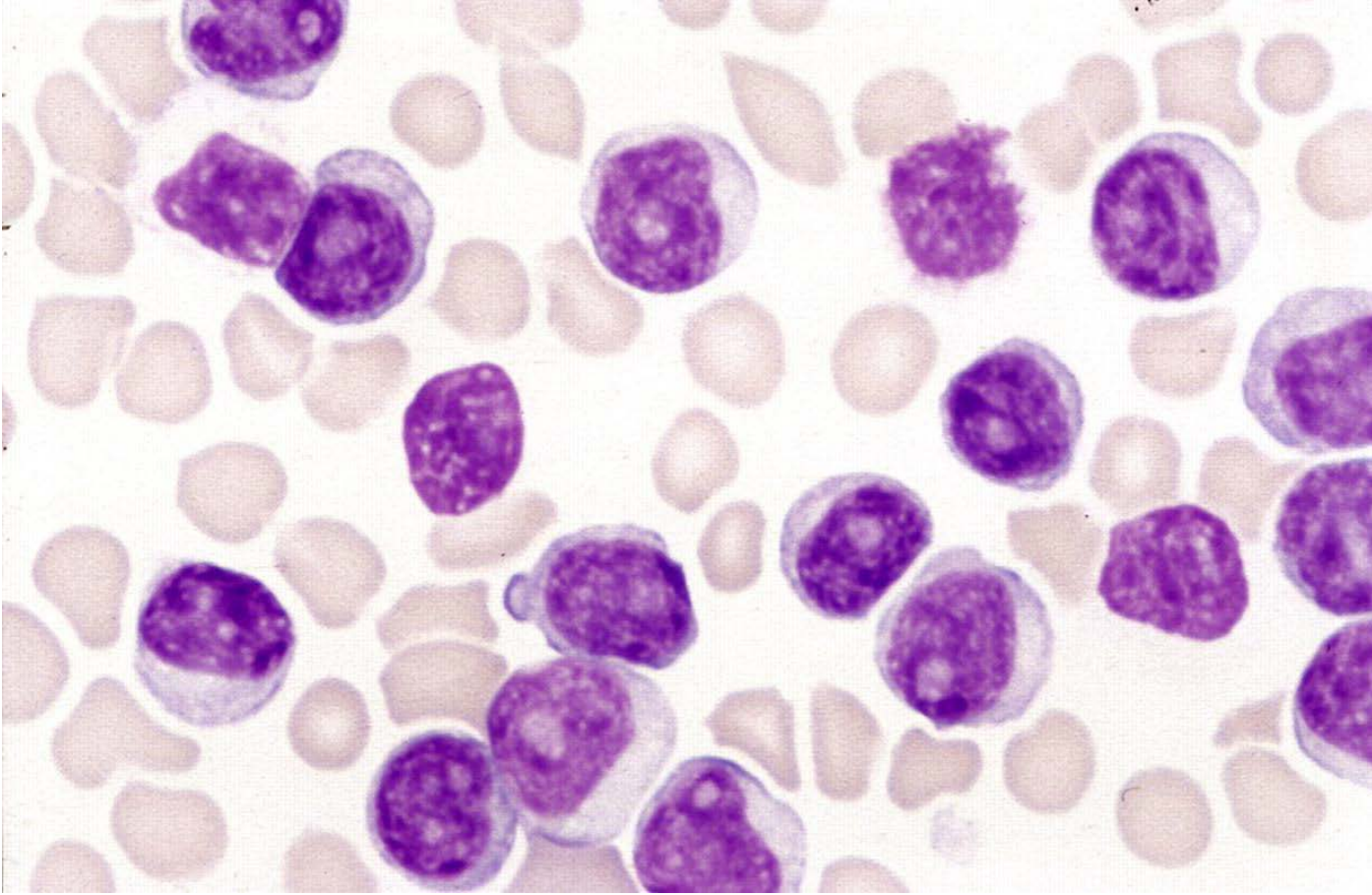
Lecture9

Neutrophil Stages of Differentiation

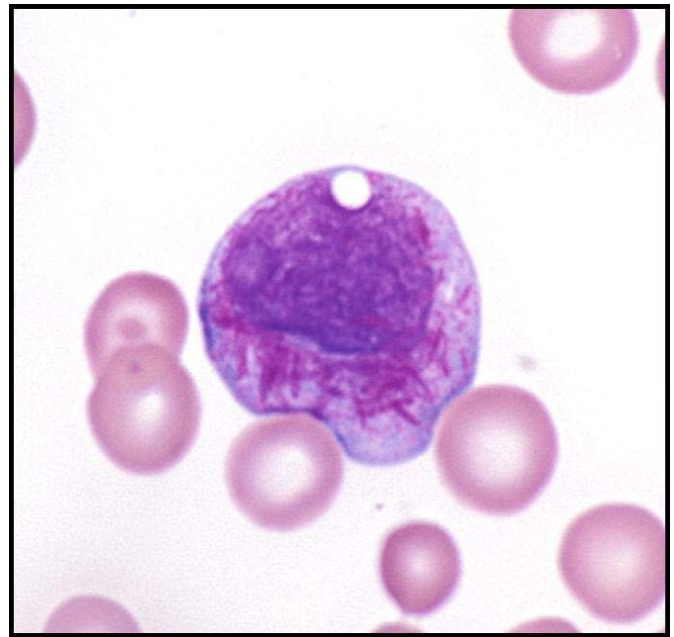
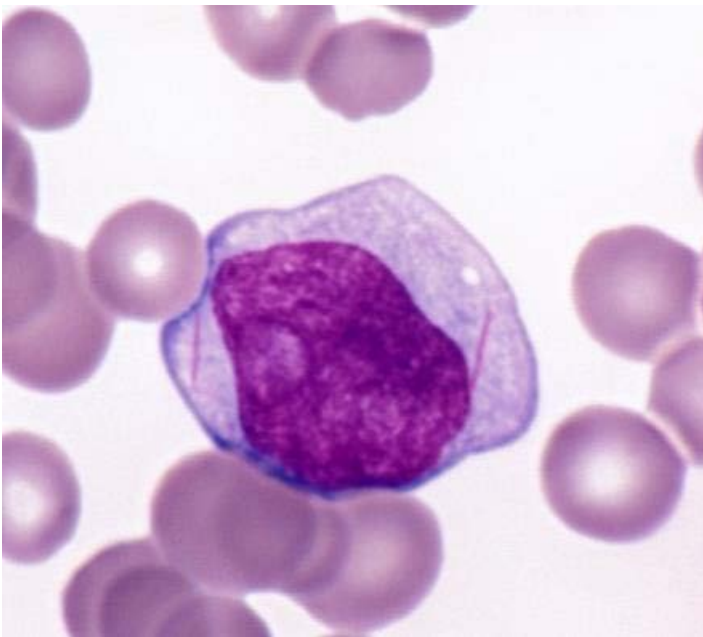


1- Acute myeloid leukemia

1. At least 20% blasts by definition (Check the previous page)



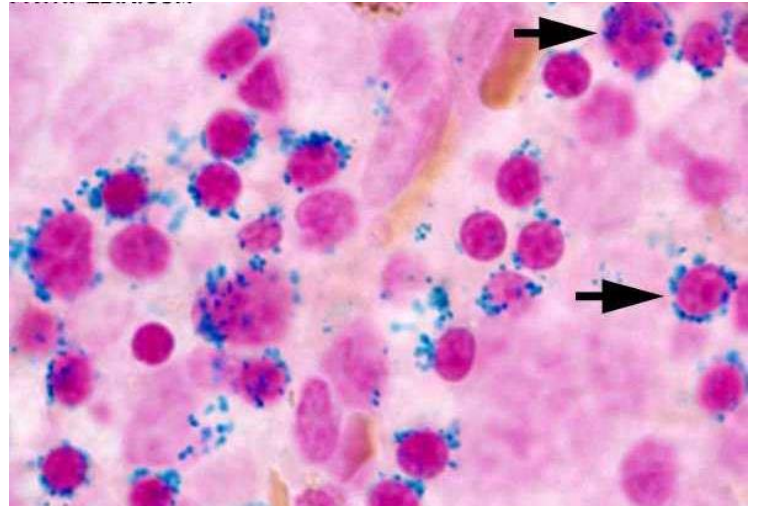
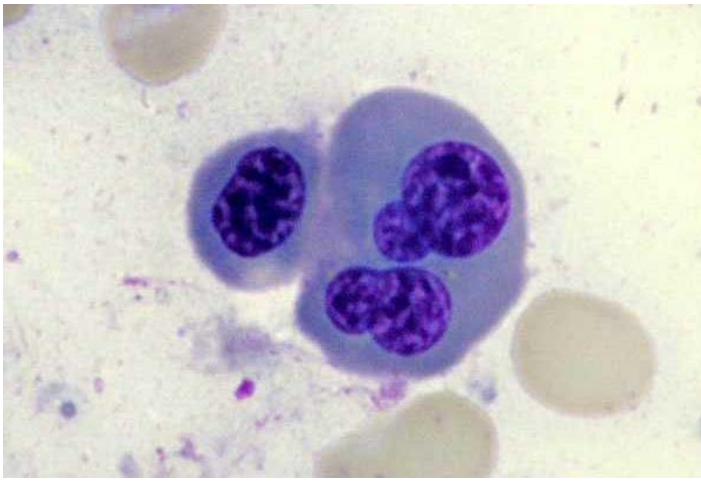
2. Auer rods (needle shaped structures in the cytoplasm)



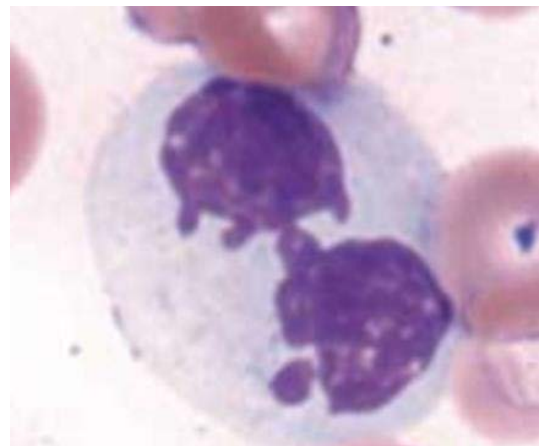
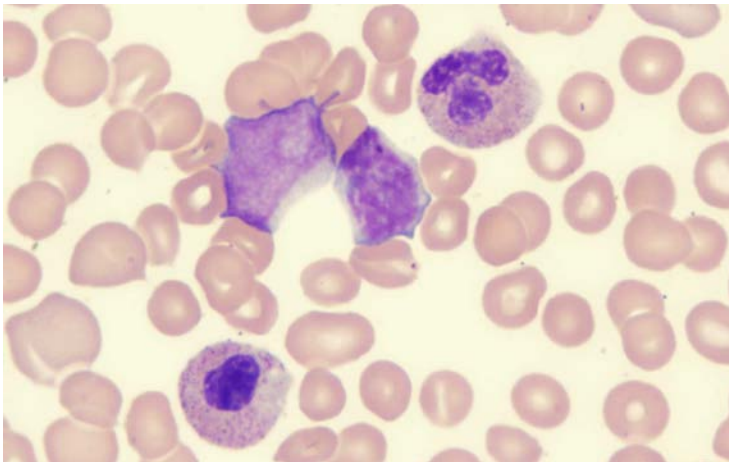
2- Myelodysplastic Syndrome

Morphology:

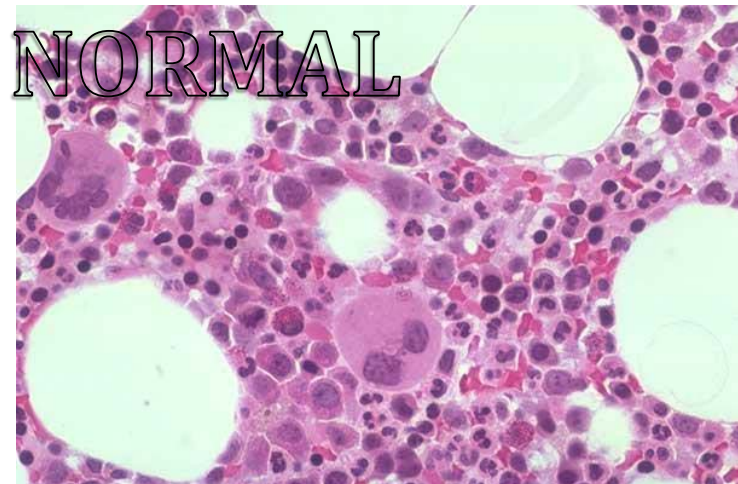
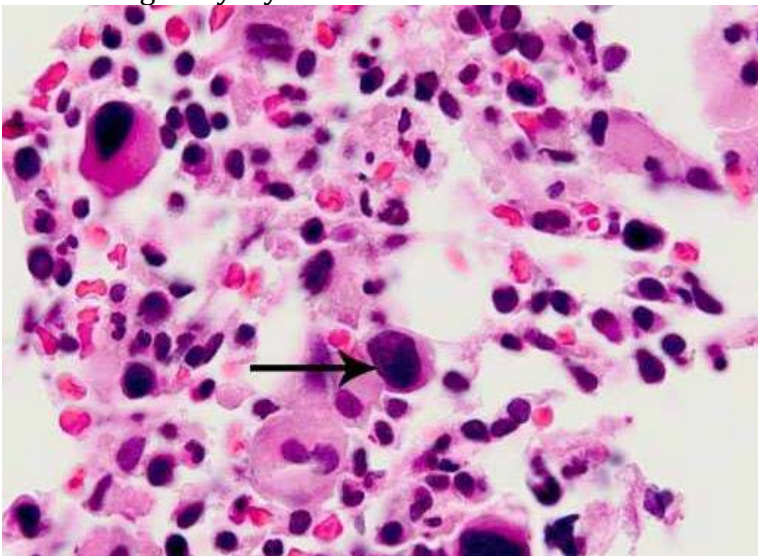
1. Erythroid: Abnormal nuclear contour and iron deposits (ring sideroblasts)



2. Myeloid: abnormal segmentation and granulation



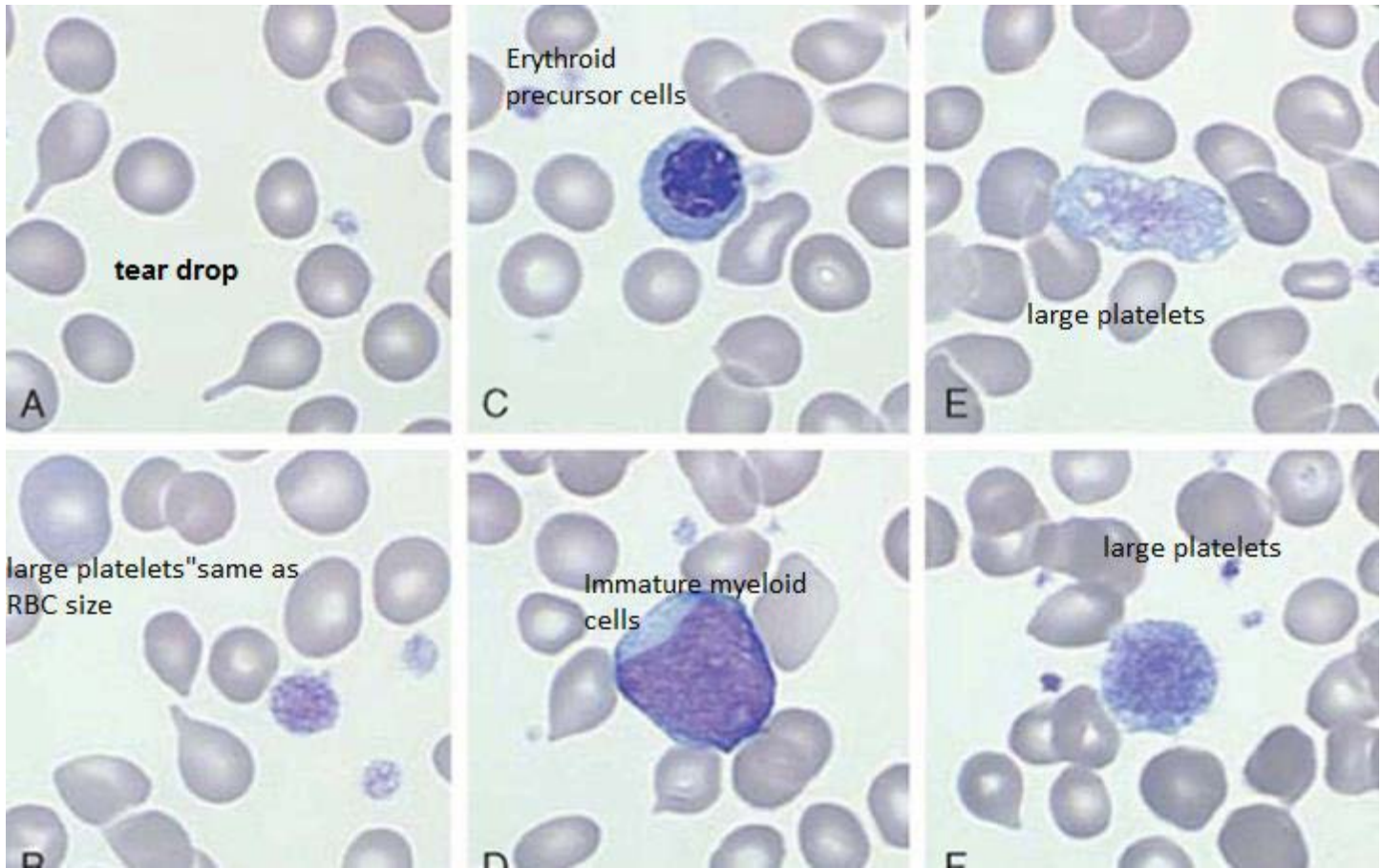
3. Megakaryocyte: small and monolobed



3- Primary myelofibrosis

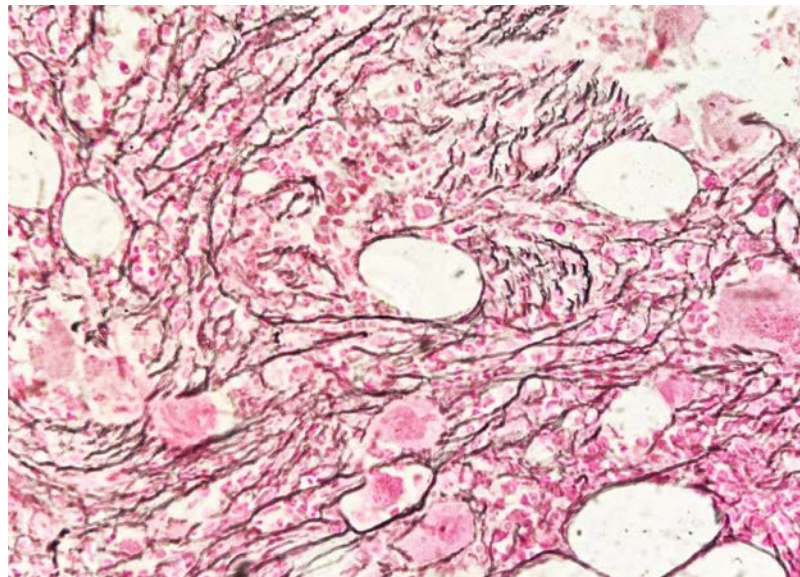
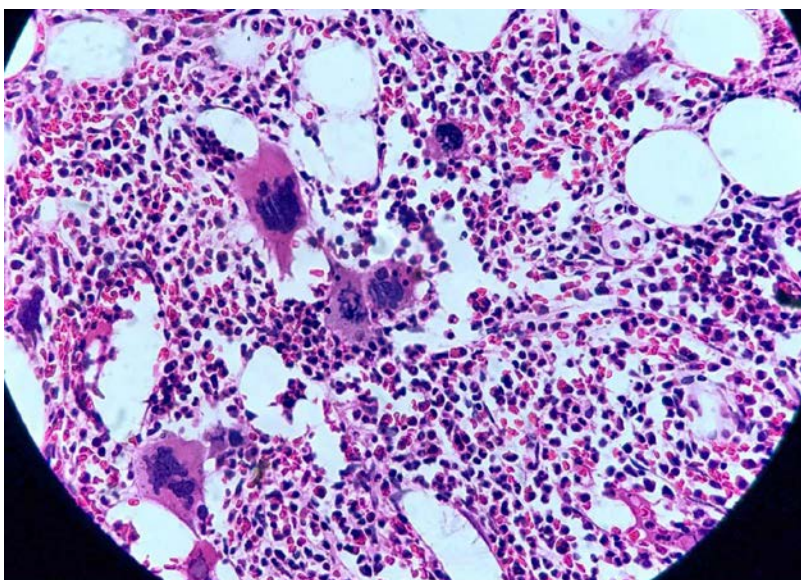
Morphology:

→ Peripheral Blood:



→ Bone Marrow:

Severe fibrosis and abnormally large and clustered megakaryocytes



Lecture 10

Sheet pictures are enough.