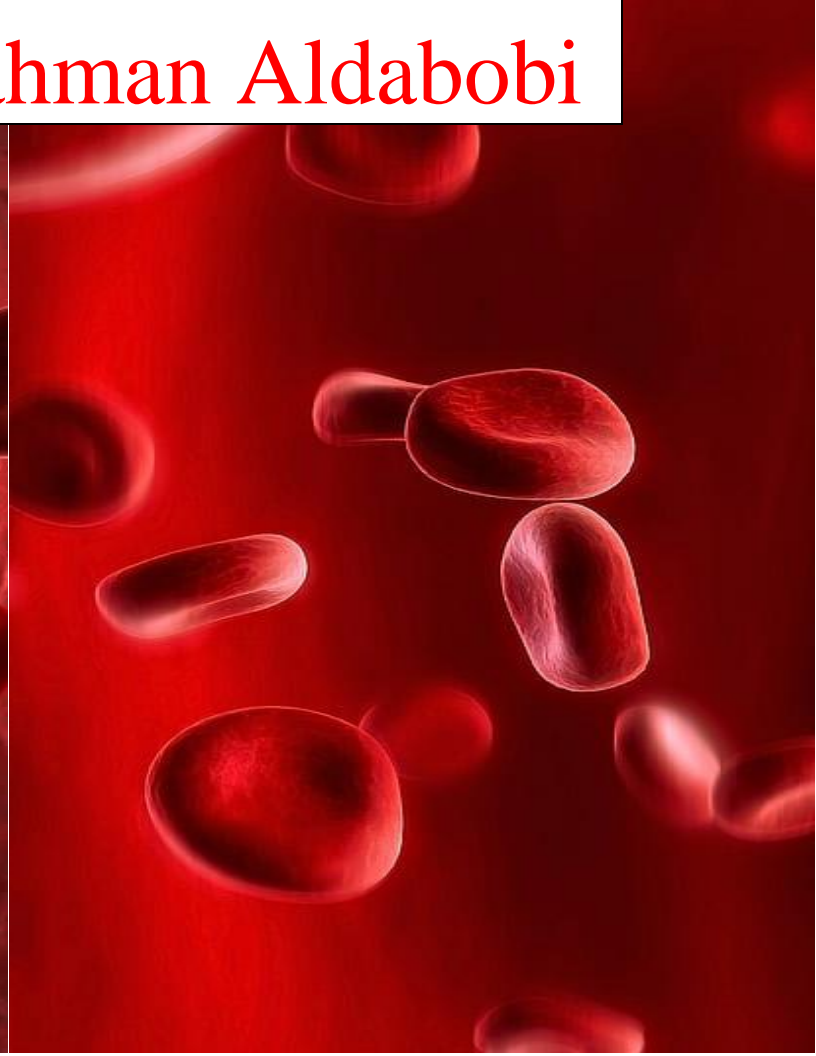
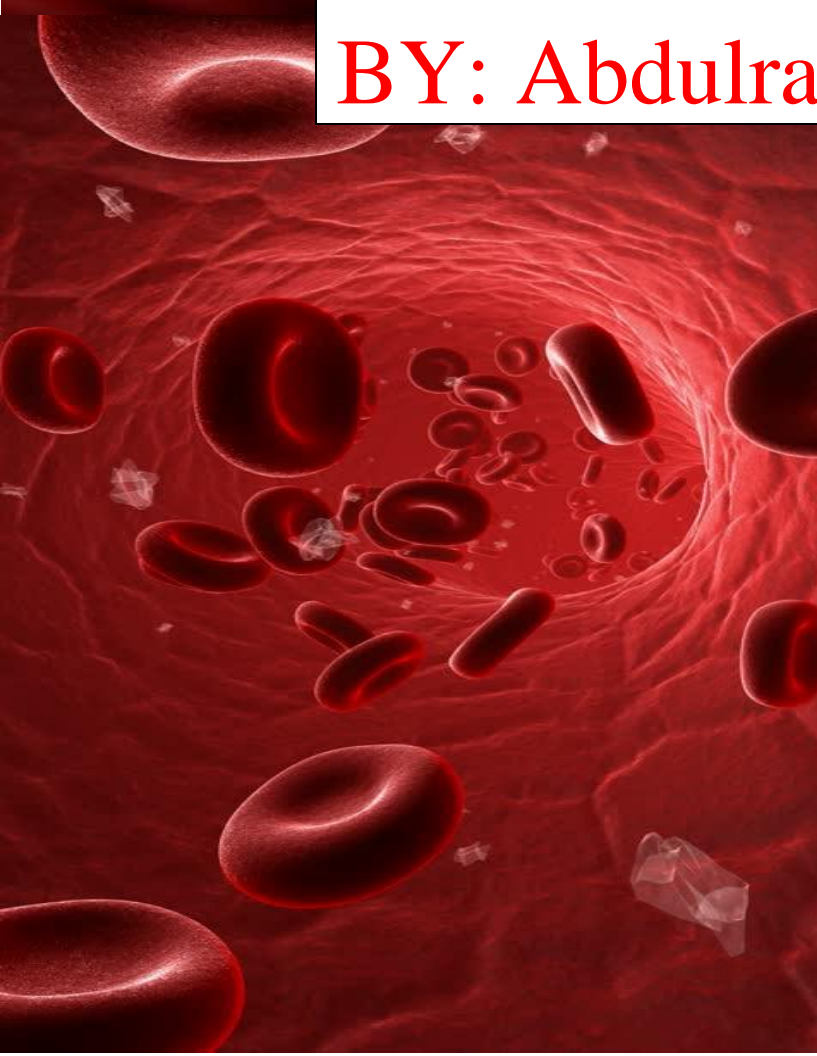


# HLS-Histopathology

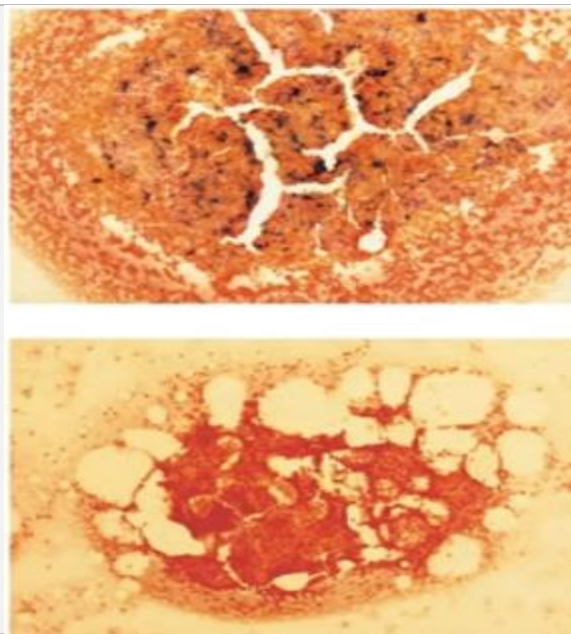
BY: Abdulrahman Aldabobi



**Image**

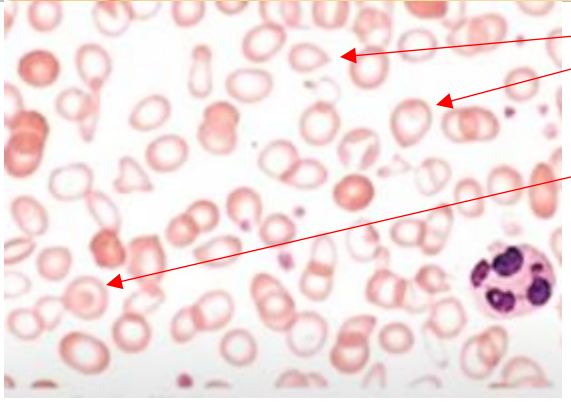
**Description**

**Diagnosis + Notes**



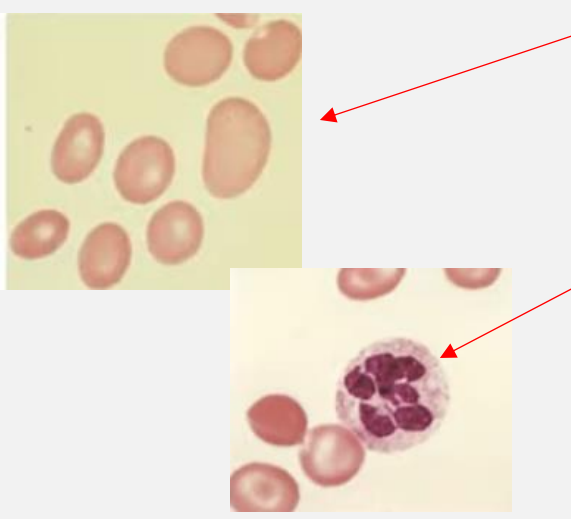
Bone marrow with decreased iron storage

1-Iron deficiency  
 2-The upper image indicates normal BM with iron represented in the blue dots  
 3-The lower image is from patient with Iron deficiency



Poikilocytosis  
 Target cells  
 Microcytic hypochromic RBCs

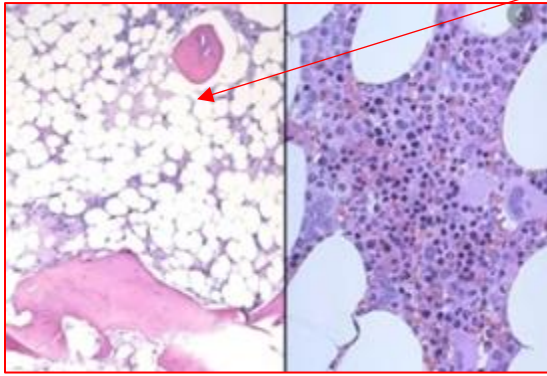
1-**Iron deficiency Anemia**  
 2-Poikilocytosis → RBCs with different shape  
 3-Target cells are RBC with a dark area in the middle of its pale center (not specific)



Macro ovalocytes  
 Hyper – segmented neutrophil

1-**Megaloblastic anemia**  
 2-May causes of macrocytes but only megaloblastic anemia causes the oval shape



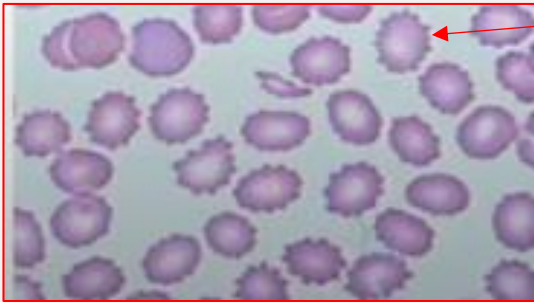


**Fatty Bone Marrow**

1- **Aplastic anemia** is characterized by Destruction of hematopoietic cells and BM is replaced by fat (note the white spots in the image to the left)

2- To the right is the normal BM to compare

3- Normoctic to Macrocytic + Pancytopenia

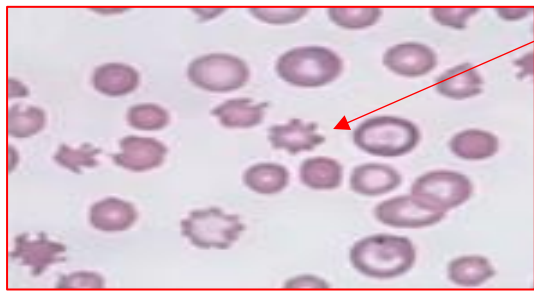


**Echinocytes (Burr Cells)**  
الخلايا الشوكية

1- **Anemia of Renal Disease**

2- The shape is caused by Uremia

3- Uremia can also impair platelet aggregation → Bleeding



**Acanthocytes (Spur Cells)**

1- **Anemia of Liver Disease**

2- The shape is thought to be due to impaired liver function

3- Longer and larger than Echinocytes



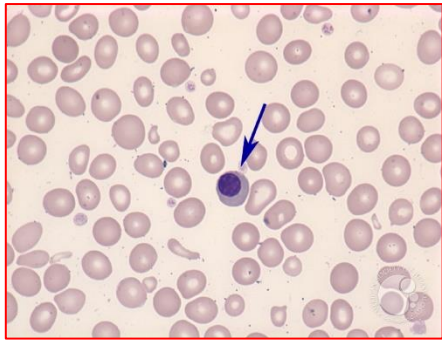
**Basophilic striplings**

**Target Cells**

1- **Thalassemia**

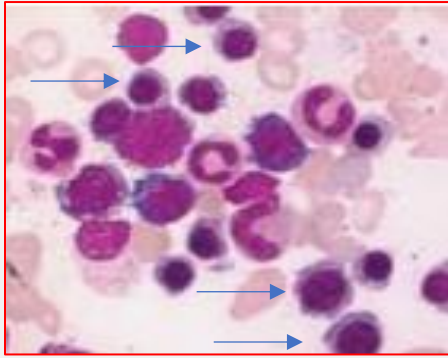
2- Target cell are due to impaired hemoglobinization (not specific)

3- Basophilic Striplings are ribosomal remnants (Also seen in sideroblastic anemia)



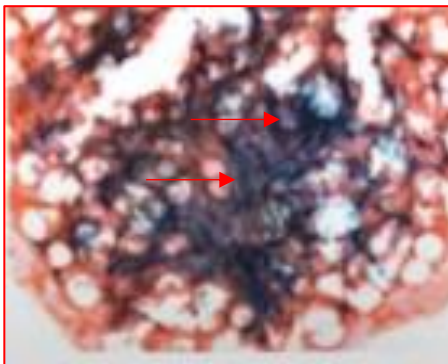
Nucleated RBC in peripheral Smear

**Beta Thalassemia Major**



Increased Normoblasts in the Bone Marrow

1-**Beta Thalassemia Major**  
 2-Normoblasts are nucleated precursors of RBCs found in the bone marrow  
 3-Normoblasts are usually 1/3 to 1/4 of cells in the bone marrow only



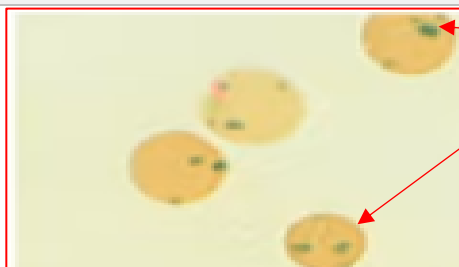
Hemosiderosis

1-**Beta Thalassemia Major**  
 2-Increased Hemosiderin in the bone Marrow  
 3-Thalassemia Major → chronic increased EPO → EPO inhibits hepcidin → unregulated iron intake → Iron deposits in the BM



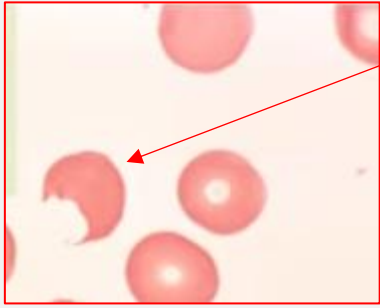
Sickle Cells  
 Target cells

1-**Sickle cell anemia**  
 2-Sickle cells are due to point mutation in the Hb  
 3-Target cells are seen in any problem with hemoglobinization (Iron deficiency, thalassemia, ....)



Heinz Bodies

1-**G6PD Deficiency**  
 2-Denaturation of Hb molecules after 2-3 days from exposure to certain oxidant



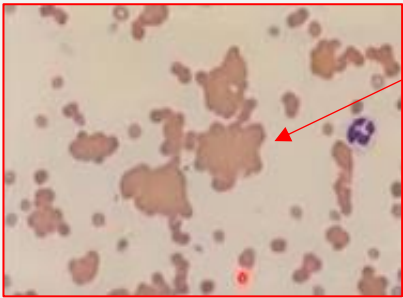
**Bite Cells  
(Degmacytes)**

1-**G6PD Deficiency**  
2-Caused by macrophages eating the preformed Heinz Bodies



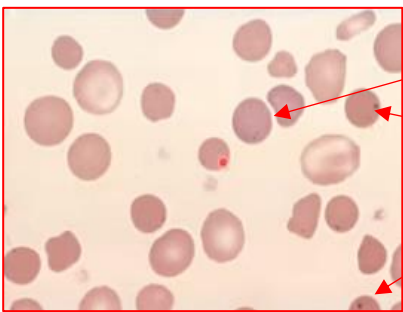
**Spherocytes**  
**Polychromasia**

1-**Warm Immune Hemolytic Anemia**  
2-Spherocytes – Round RBCS without central pallor (not specific)  
3- Polychromasia (multiple colors)  
→ The presence of reticulocytes and spherocytes result in the appearance of different colors of RBCs in the smear



**RBCs clumps**

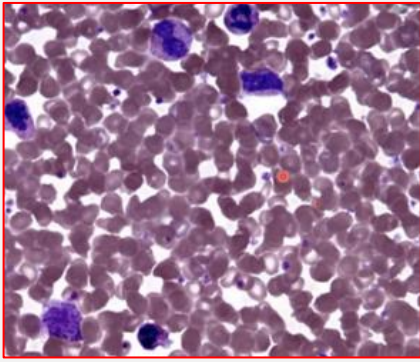
1-**Cold Immune Hemolytic Anemia**  
2-IgM antibodies bind 5 RBCs and cause agglutination  
3-very small spherocytes can also be seen



**Spherocytes**  
**Howel Jolley Bodies**

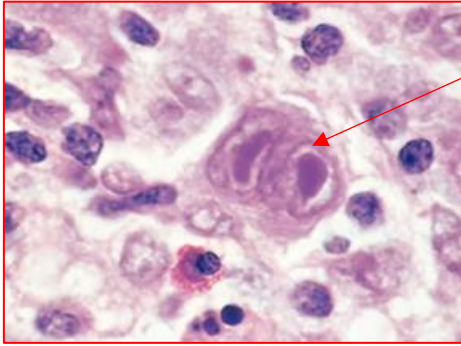
1-**Hereditary spherocytosis**  
2-Loss of some cell membrane proteins lead to the loss of biconcavity of RBCs → Spherocytes  
3-Defenitive treatment is splenectomy  
4-**Howel-Jolley bodies** are DNA remnants in the RBCs due to splenectomy (Usually these dots are removed by spleen)  
5-**Autosomal DOMINANT**





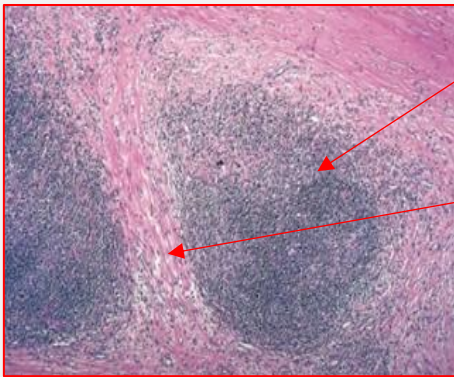
Too many RBCs are present

- 1-**Polycythemia**
- 2-Can be relative or absolute
- 3-Absolute polycythemia can be primary (Polycythemia Vera) or secondary



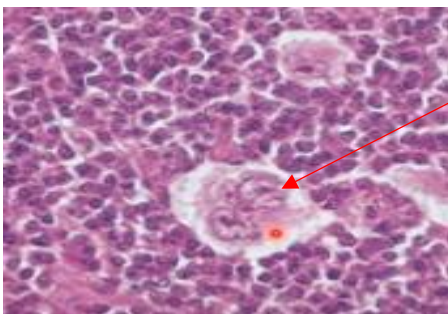
Reed-Sternberg cells

- 1-**Hodgkin Lymphoma**
- 2- Giant multinucleated, prominent nucleolus, eosinophilic lymphocyte
- 3- Hodgkin cells are the same but with one nucleus (both can be found in the lymph node)
- 3-Positive for CD30 and CD15
- 4-Negative for CD20, CD3, CD45



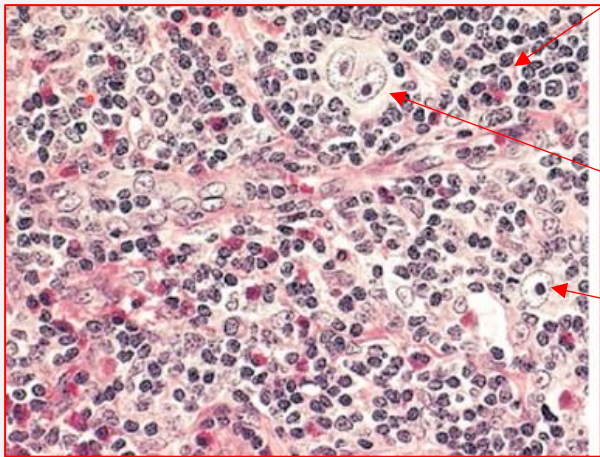
Lymphocyte nodules  
Fibrous tissue

- 1-**Nodular sclerosis**
- 2-Most common type of Hodgkin Lymphoma
- 3-Characterized by Lymphocyte nodules separated by fibrous tissue



Lacunar Cells

- 1-**Nodular Sclerosis**
- 2-Reed-Sternberg cell with peripheral cytoplasmic retractions (Empty on the periphery)



Lymphocytes nodules

RS cells

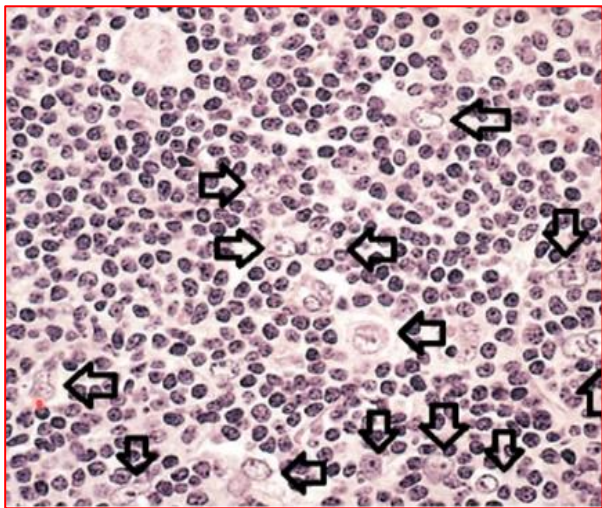
Hodgkin cells

1-**Mixed cellularity**

2-Same as nodular sclerosis but with no fibrous bands

3-More common in older patients

4-Associated with EBV



Popcorn Cells

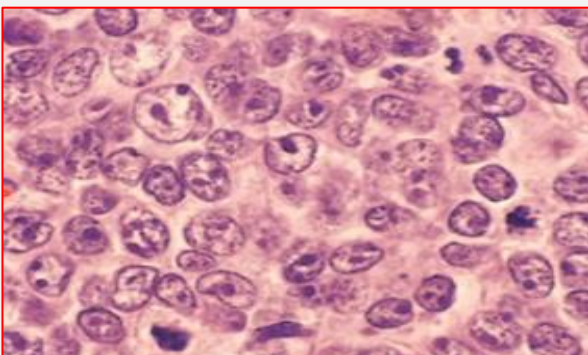
1-**Lymphocyte predominant Hodgkin Lymphoma**

2- Giant cells with Multilobated vesicular nucleus and small blue nucleoli

3- Cells were called Lymphohistiocytes.

4- Arranged in nodules with no fibrous bands

5- express B-cell markers → CD45 and CD20 and negative for CD30 and CD15



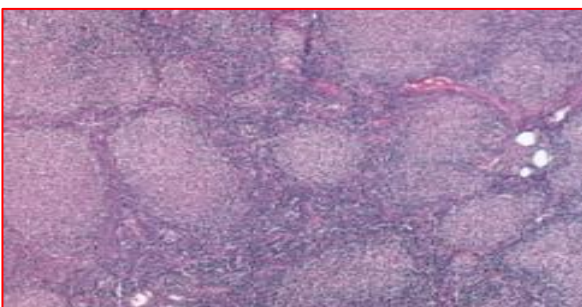
Large B-cells

1-**Large diffuse B-Cell Lymphoma**

2-No nodules at lower magnification

3-At higher magnification → Large cell with irregular nuclei and small nucleoli

4-CD20 Positive

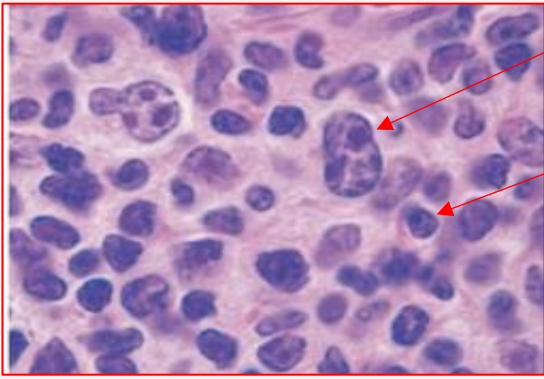


Follicles within the lymph nodes

1-**Follicular Lymphoma**

2-Lymph nodes are filled with follicles affecting its architecture



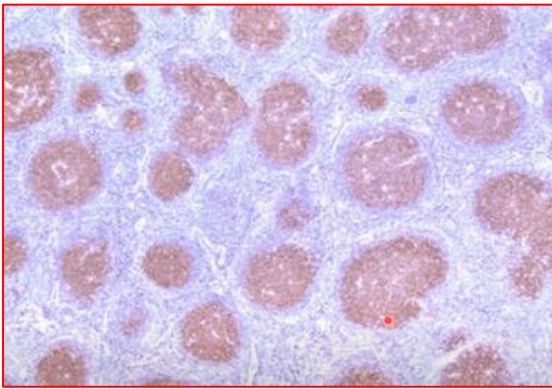


Centroblasts

Centrocytes

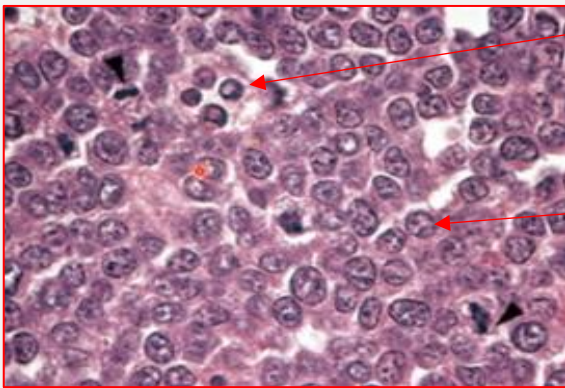
1-**Follicular Lymphoma**

2-Centrocytes are small and differentiated → Predominant → Low grade lymphoma  
 2-Centroblasts are large and undifferentiated → Predominant → High Grade Lymphoma



Follicular Lymphoma with immunohistochemistry testing for Bcl2

Used to differentiate between Follicular Lymphoma and Reactive follicular hyperplasia



Tingible bodies

Intermediate lymphocytes

1-**Burkitt Lymphoma**

2-High rates of mitosis → Nuclear debris engulfed by the macrophages  
 3-B-Lymphocytes are intermediate in size and monomorphic (The same shape) and usually round  
 4-Starry like appearance → Pale areas resemble the sky and crowded areas resemble the stars



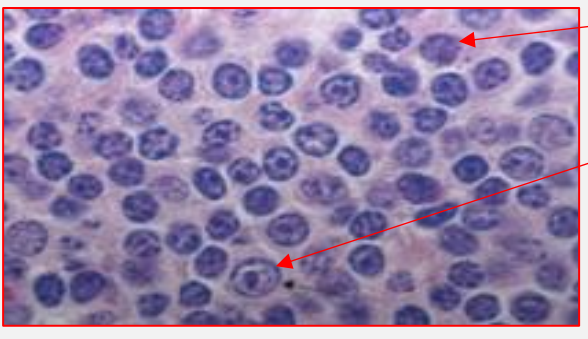

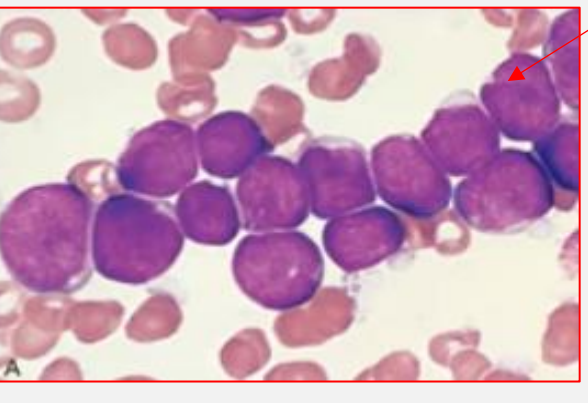

Lymph node with no apparent architecture

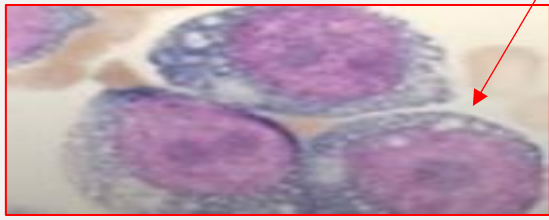
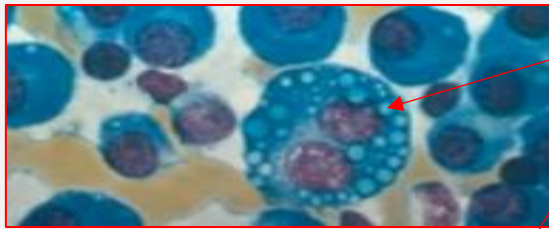
Proliferation centers (Pale)

1-**Small Lymphocyte Lymphoma**

2- LN with no architecture and pale areas called proliferation centers (prolymphocytes with larger nuclei and abundant cytoplasm → pale)

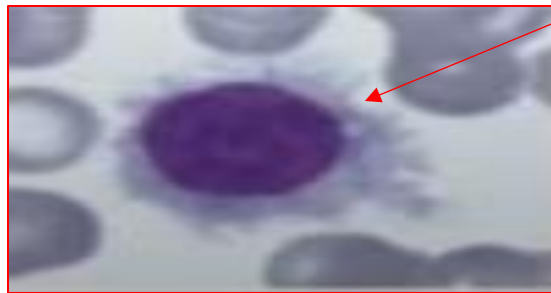


	<p>Lymphocytes</p> <p>Prolymphocytes</p>	<p>1- <b>Small Lymphocyte Lymphoma</b></p> <p>2-Lymphocytes predominant in early stages of the disease</p> <p>3-Prolymphocytes predominant in the late stages of the disease</p>
	<p>Prolymphocytes</p> <p>Smudge cells</p>	<p>1-<b>Chronic Lymphoblastic leukemia</b></p> <p>2-Smudge cells are dead lymphocytes</p>
	<p>Lymphoblasts</p>	<p>1-<b>Acute Lymphoblastic leukemia</b></p> <p>2-Lymphoblasts are large with high N/C ratio and with pale (open) chromatin</p> <p>3-Agranular (Vs. Myelogenous)</p> <p>4-Positive TdT</p> <p>5-CD22+ → B lymphoblast</p> <p>6-CD10+ → T Lymphoblast</p>
	<p>Rouleaux Formation of RBCs</p>	<p>1-<b>Multiple Myeloma</b></p> <p>2-Different from agglutination in that agglutination happens in many directions</p>



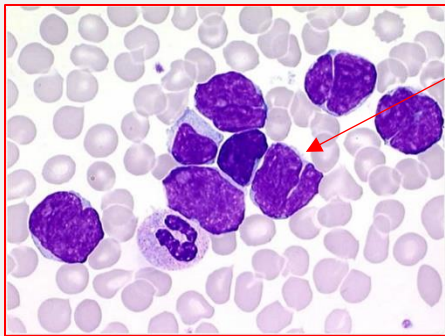
Abnormal plasma cells

1-**Mutible Myeloma**  
 2-Plasma cells with more than one nucleus and cytoplasmic vacuoles contains immunoglobulins  
 3-Have a prominent nucleolus instead of the characteristic Kary-Wheel nucleolus



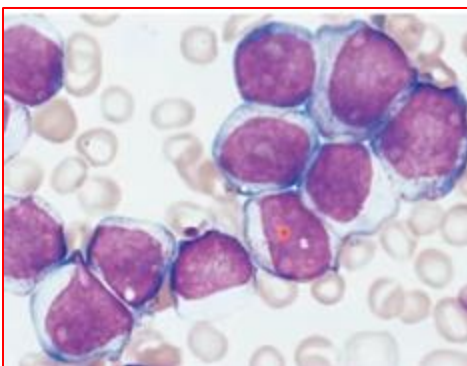
Hairy lymphocytes

1-**Hairy cell leukemia**  
 2- Few with prominent cytoplasmic projections



Neoplastic lymphocytes /  
 Sezary cells

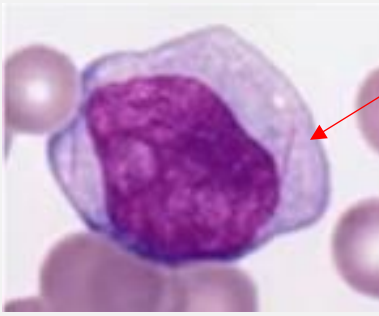
1-**Mycosis fungoides**  
 2-CD4+ cell causing erythema and may progress to plaques and tumor  
 3-Neoplastic lymphocyte with irregular nuclear membrane resembles a cerebriform  
 3- Called Sezary syndrome if causing leukemia



Granulocytes precursor  
 Aur Rods

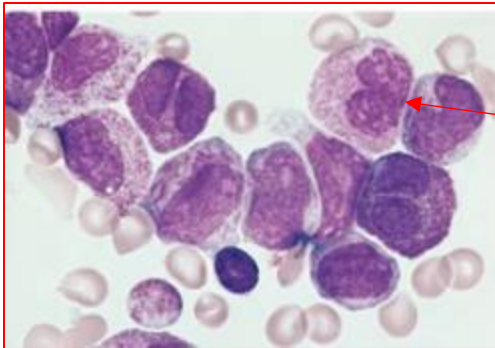
1-**Acute Myeloid Leukemia**  
 2-similar to lymphoblasts but larger and with more cytoplasm (N/C is still high) and with granules  
 3-Pale nuclei  
 4-Expresses CD34, Myeloperoxidase, CD13 and CD33  
 5-Negative for Tdt





Aur Rods

- 1- **Acute Myeloid Leukemia**
- 2- Aggregations of Myeloperoxidase enzyme produced by these cells

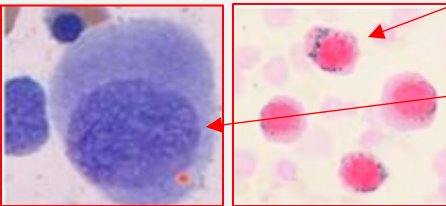


Neoplastic Promyelocytes with Cleaved nucleus

- 1- **Promyelocytic Leukemia**
- 2- To differentiate between it and AML → It is heavily granulated and CD34 negative + Nucleus resembles number 8
- 3- Aur Rods can be also present
- 4- Associated with DIC
- 5- Treated with trans



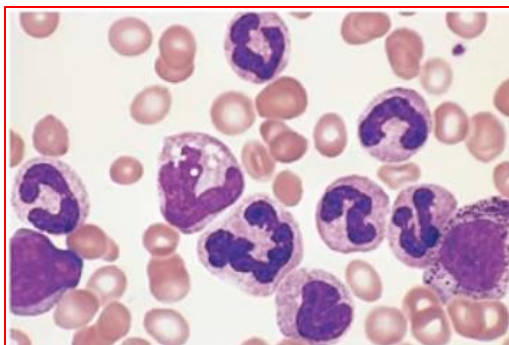
Myeloid cells



Ring Sideroblasts

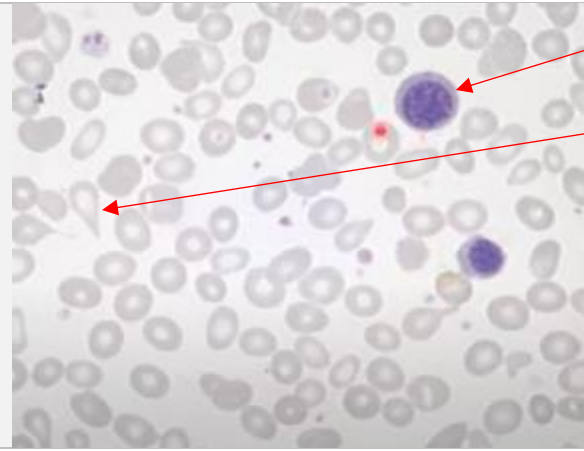
Megakaryocyte

- 1- **Myelodysplastic Syndrome**
- 2- Impaired maturation
- 3- RBCs → Ring Sideroblasts due to RNA impaired splicing
- 4- Myeloid cells → Hypogranulated and hyposegmented
- 5- Megakaryocyte → small and hypolobulated



Lymphocytosis

- 1- **Chronic Myelogenous leukemia**
- 2- Mature myelocytes in the blood
- 3- Left shift can also be seen → precursor cells in the blood



Nucleated RBC

Tear Drop RBC

1-**Primary Myelofibrosis**

2-These two with shift to left is called leucoerythroblastic anemia