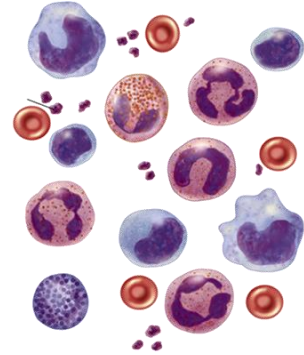


**National university Sudan**  
**Faculty of Medical Laboratory Sciences**  
**Advanced Hematology MLS –HEMA-324**

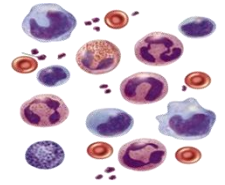
# **LEUCOCYTES BENIGN DISORDERS**



**U. Taiba Mohamed Abdalla**  
**Lecture(14)**

# LEUCOCYTES BENIGN DISORDERS

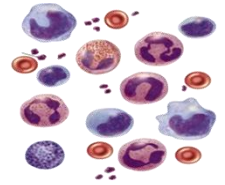
## Quantitative changes (LYMPHOCYTOSIS)



- **Infectious Mononucleosis**
  - Epstein-Barr virus
  - Saliva from infected person is the main contagion
  - Virus infect epithelial cells and B cells
  - Infection in children under the age of 10 does not cause illness and result in life long immunity
  - **Clinical features**
    - Fever, malaise, fatigue, sore throat, diagnostic red spots at the junction of soft and hard palate, splenomegaly
    - Blood picture shows leucocytosis (  $10 - 20 \times 10^9/L$ ) due to absolute increase in the number of lymphocytes
    - Diagnosis is by serological tests
    - There is no specific treatment

# LEUCOCYTES BENIGN DISORDERS

## Qualitative changes (MORPHOLOGY)

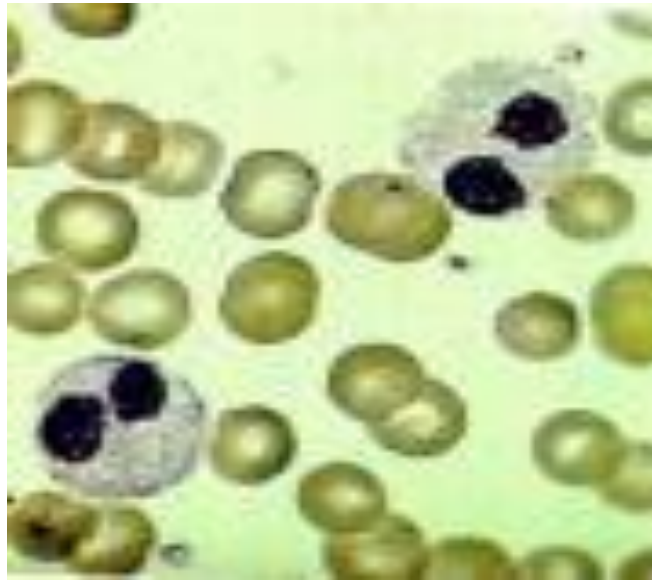


- **Congenital**

- **Pelger-Huet anomaly**

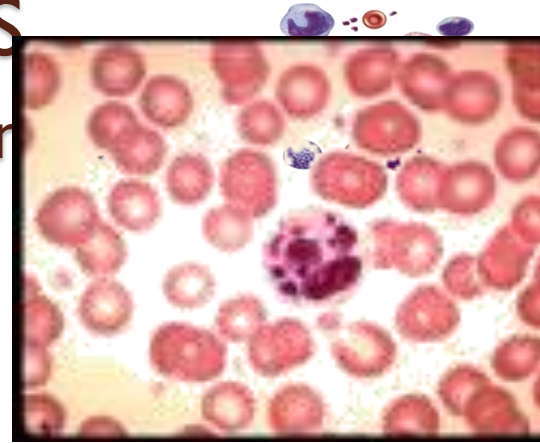
- Is a genetic disorders with an autosomal dominant inheritance pattern , neutrophils whose nucleus is hyposegmented

- Bilobed and occasional unsegmented neutrophils
- Autosomal recessive disorder



# LEUCOCYTES BENIGN DISORDERS

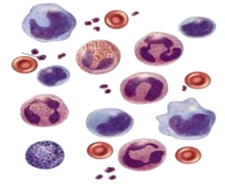
Qualitative changes (MORPHOLOGY) con



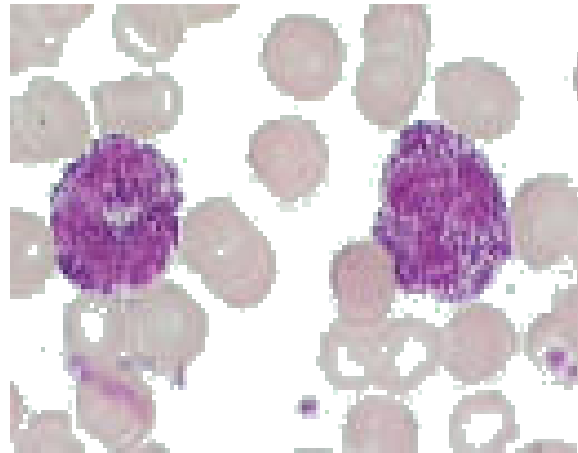
- Neutrophil hyper-segmentation
  - Rare autosomal dominant condition
  - Neutrophil function is essentially normal
- May-Hegglin anomaly
  - Also called Dohle leucocyte inclusions with giant platelets , rare genetic disorders of the blood plts that causes them to be large .
  - Neutrophils contain basophilic inclusions of RNA
  - Occasionally there is associated leucopenia
  - Thrombocytopenia and giant platelet are frequent

# LEUCOCYTES BENIGN DISORDERS

## Qualitative changes (MORPHOLOGY) contd.

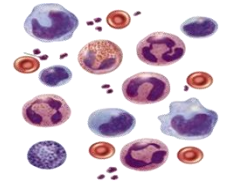


- Alder's anomaly
- Most characteristic finding is the metachromatic granules surrounded by a clear zone seen in lymphocytes.
  - Granulocytes, monocytes and lymphocytes contain granules which stain purple with Romanowsky stain
  - Granules contain mucopolysaccharides

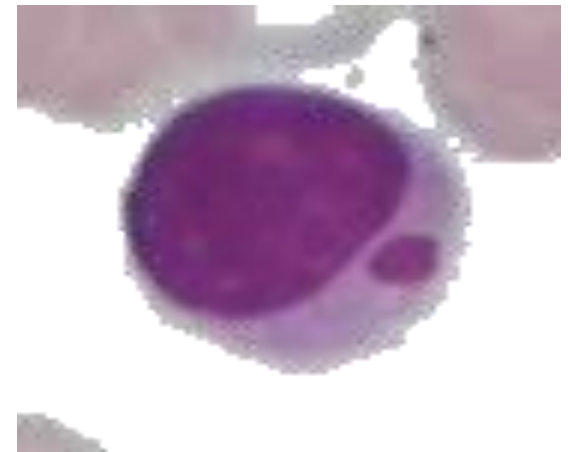


# LEUCOCYTES BENIGN DISORDERS

## Qualitative changes (MORPHOLOGY) contd.

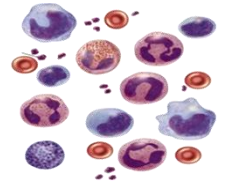


- Chediak-Higashi syndrome
- Is a rare disorders that arises from a mutation of a lysosomal trafficking regulator protein which leads a decrease in phagocytosis ,
  - Autosomal recessive disorder
  - Giant granules in granulocytes, monocytes and lymphocytes
  - Partial oculocutaneous albinism
  - Depressed migration and degranulation
  - Recurrent pyogenic infections
  - Lymphoproliferative syndrome may develop
  - Treatment is BMT



# LEUCOCYTES BENIGN DISORDERS

## Qualitative changes (MORPHOLOGY) contd.



- **Acquired**

- **Toxic granulation**

- Changes in granulocyte cells seen on examination of the blood film of patients with inflammatory conditions , dark coarse granules found in granules found particularly in neutrophils .

- **Dohle bodies**

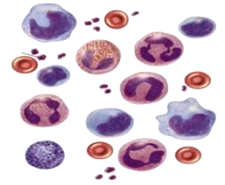
- Are light blue gray , oval , basophilic , leucocyte inclusions located in peripheral cytoplasm of the neutrophils . Seen in infections , measles , leukemia .

- **Pelger cells**

- **Hypersegmented neutrophils**

# LEUCOCYTES BENIGN DISORDERS

## Qualitative changes (FUNCTIONAL)



- **Chronic granulomatous disease**
- Also known as Bridge- Good syndrome , certain cells of the immune system have difficulty forming the reactive oxygen compounets used to kill certain ingested pathogens , this leads to the formation of granutoma in many organs . Mainly cause by the mutation in the X chromosome.