

RBC disorders 3

Anemia of blood loss (1)

- Anemia of blood loss, hemorrhage
- Hemolysis
 - extrinsic
 - Immune hemolytic anemia
 - Hemolytic anemia resulting from mechanical trauma to the red cells
 - Infection

1- Other than anemia, one of the following can be seen in the setting of anemia of hemorrhage:

- A. Leukocytopenia
- B. Neutropenia
- C. Leukocytosis
- D. Lymphocytopenia
- E. Thrombocytopenia

2- Which one of the following is most helpful to differentiate between intravascular and extravascular hemolysis:

- A. LDH
- B. Haptoglobin
- C. Bilirubin
- D. Hemoglobiuria

3- Warm antibody immune hemolytic anemia is most commonly caused by:

- A. IgM
- B. IgA
- C. IgG
- D. IgD
- E. IgE

4. Which one of the following is a characteristic finding in microangiopathic hemolytic anemia:

- A. Target cells
- B. Sickle cells
- C. Spur cells
- D. Ecchinocytes
- E. Schistocytes

5. Which one of the following can cause cerebral malaria:

- A. *P.vivax*
- B. *P. ovale*
- C. *P. falciparum*
- D. *P. malarie*

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- intrinsic
 - Hereditary
 - Membranopathies-spherocytosis
 - Hemoglobinopathies-thalassemia and sickle cell disease
 - Enzymopathies-G6PD deficiency
 - Acquired
 - Paroxysmal nocturnal hemoglobinuria.

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Anemia of blood loss

- It has two types
 - Acute
 - Chronic

- Acute: resulting from external (wound) or internal (rupture aortic aneurysm) hemorrhage.
- If the blood loss is less than 20% of the total volume, healthy patients can tolerate that with a few symptoms
- If it exceeds 20% the immediate threat is from hypovolemia not anemia.

- The full effect of anemia starts to appear after 2-3 days when the fluid shifts into the intravascular space.
- Erythropoietin will be released recruiting more stem cells to proliferate and differentiate into red cells.

- Normochromic normocytic anemia
- Leukocytosis
- As Red cell production increases, it turns into slightly macrocytic anemia
- Thrombocytosis upon recovery.

- Anemia of chronic blood loss is iron deficiency anemia, discussed earlier.

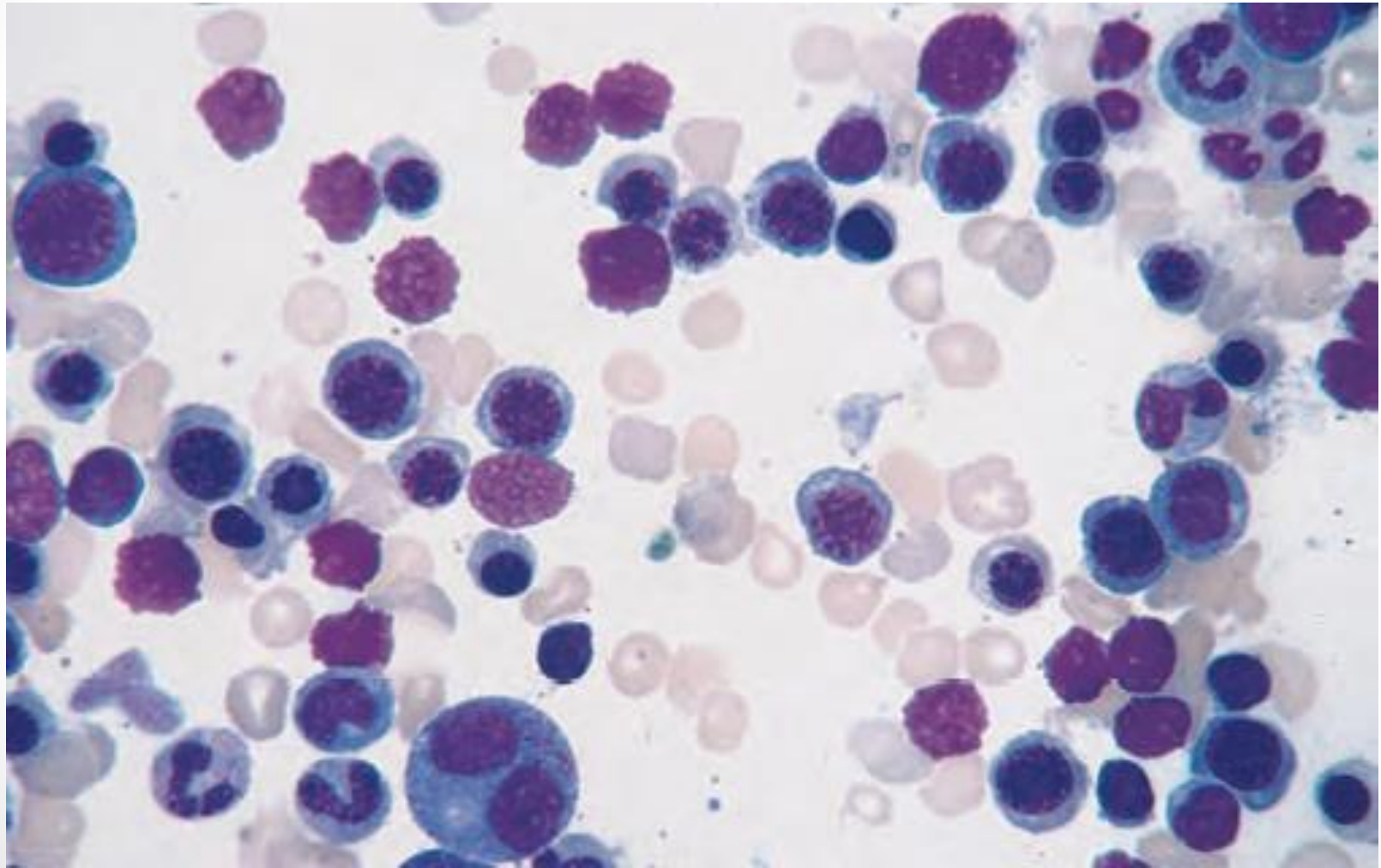
Hemolysis- general

- Increase destruction of red cells resulting in decreased RBC survival.
- Elevated erythropoietin levels
- Accumulation of hemoglobin degradation products, bilirubin and iron.
- Erythroid precursor hyperplasia in the bone marrow and reticulocyte counts in the blood.

- Extravascular or intravascular.
 - Extravascular: spleen or liver
 - No hemoglobinuria or hemoglobinemia
 - Low haptoglobin
 - High LDH
 - Splenomegaly.
 - Jaundice; high bilirubin and possibly gallbladder stones.

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Immune hemolytic anemia

*Hemolytic anemias in this category are caused **by antibodies that bind to red cells**, leading to their premature destruction.*

Warm Antibody Type

Primary (idiopathic)

Secondary: B cell neoplasms (e.g., chronic lymphocytic leukemia), autoimmune disorders (e.g., systemic lupus erythematosus), drugs (e.g., α -methyldopa, penicillin, quinidine)

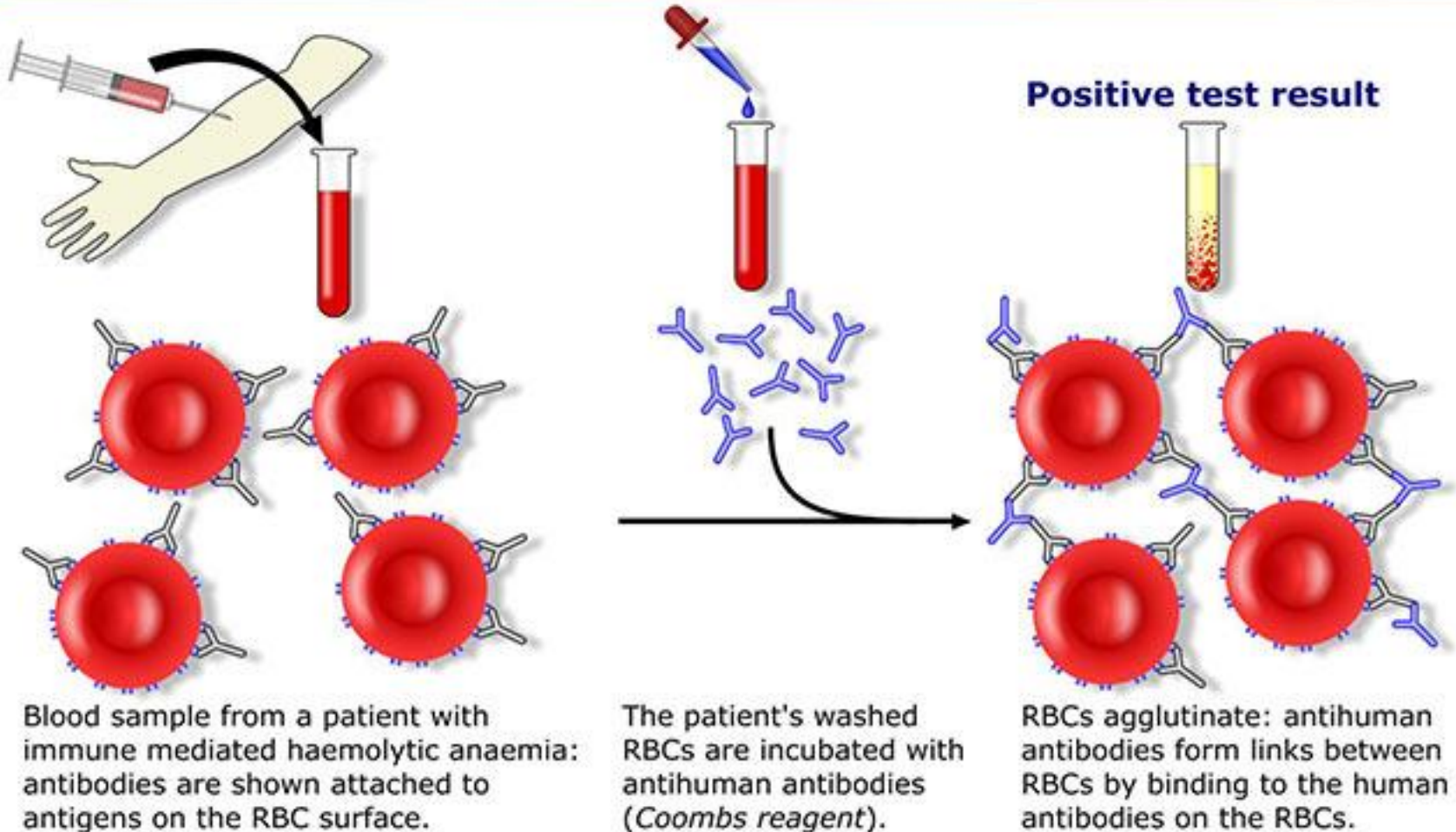
Cold Antibody Type

Acute: *Mycoplasma* infection, infectious mononucleosis

Chronic: idiopathic, B cell lymphoid neoplasms (e.g., lymphoplasmacytic lymphoma)

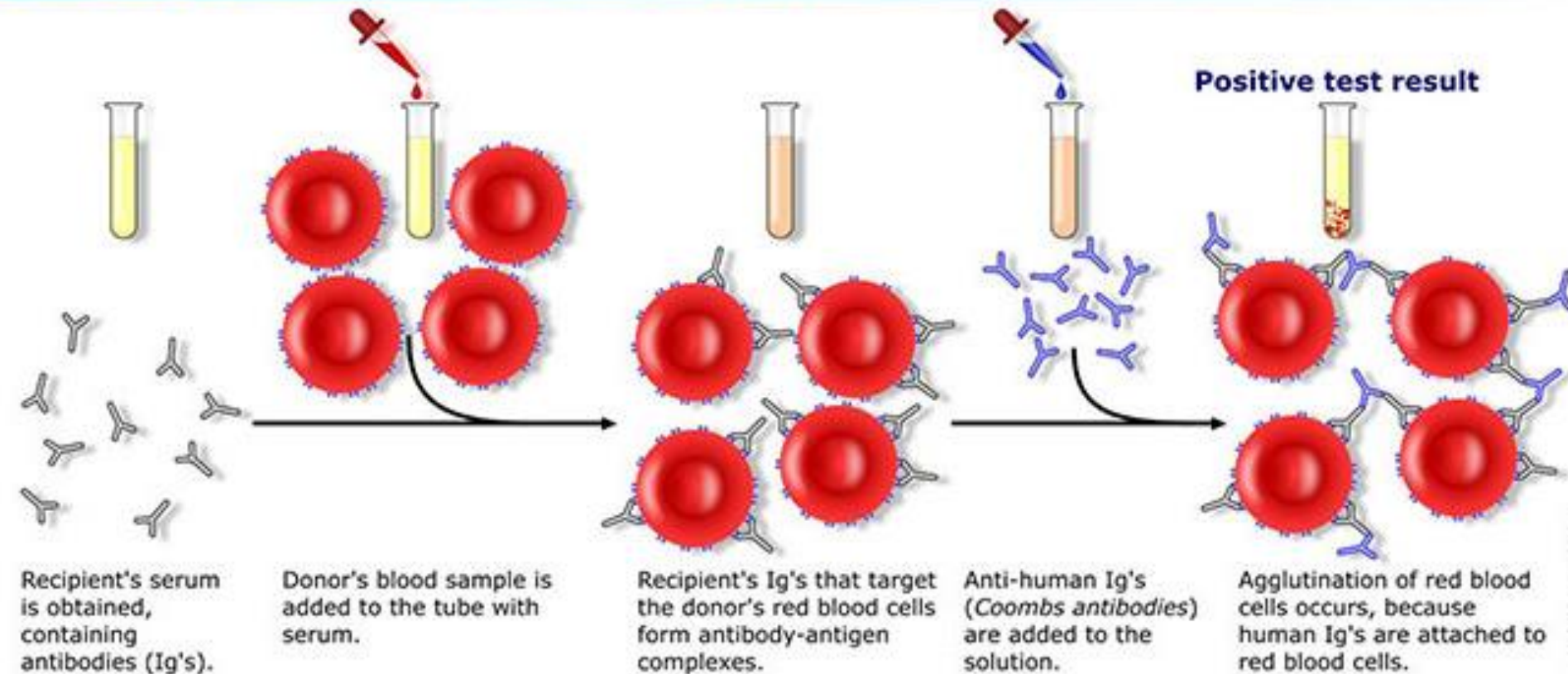
Coombs test-direct

Direct Coombs test / Direct antiglobulin test



Coombs test-indirect

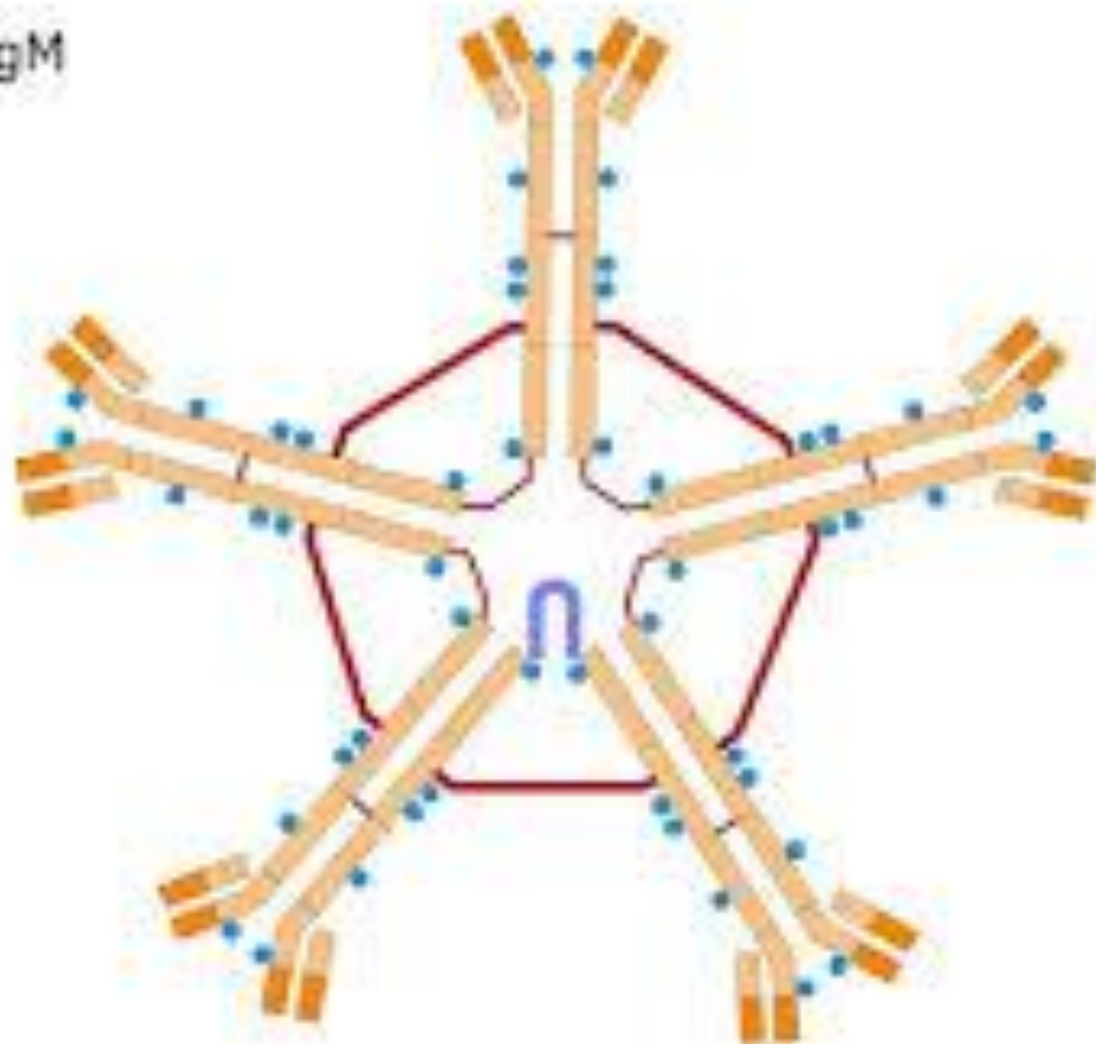
Indirect Coombs test / Indirect antiglobulin test



- Warm immunohemolytic anemia:
 - IgG and rarely IgA
 - 37 C°
 - Over 60% idiopathic.
 - Remaining minority is caused by autoimmune diseases (SLE), B cell neoplasms (CLL), or drugs (methyl-dopa, penicillin).
 - Most patients have mild anemia with splenomegaly and require no treatment.

- Cold immunohemolytic anemia:
 - IgM
 - Low temperatures in cold weather
 - Extravascular hemolysis.
 - Could be idiopathic
 - Secondary to mycoplasma, infectious mononucleosis, B cell neoplasms (lymphoplasma cytic lymphoma)
 - Usually mild without clinical significance.

IgM

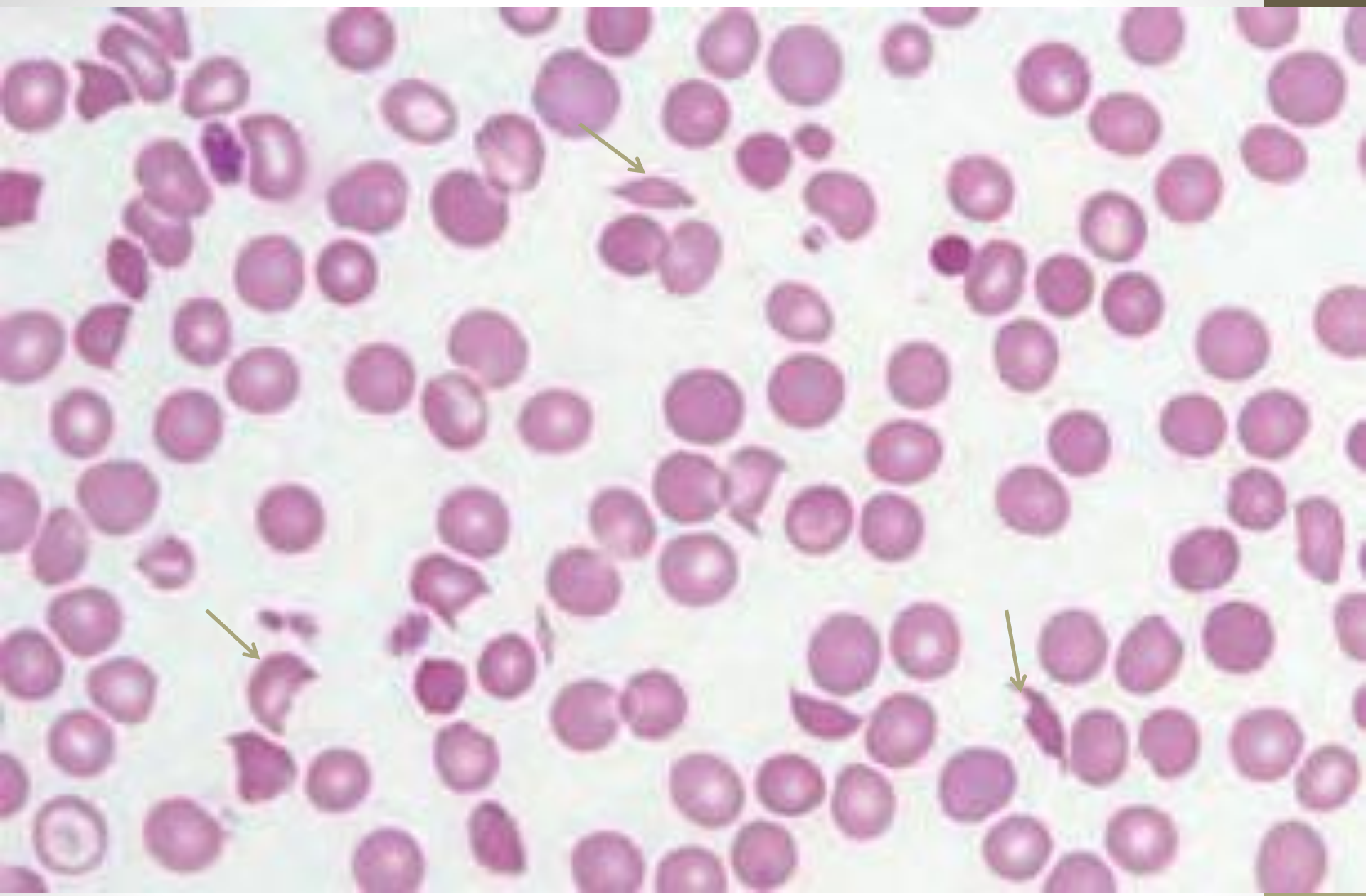




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Hemolytic anemia resulting from mechanical trauma to the red cells

- Repeated physical activity
- Cardiac valves
- Microangiopathic hemolytic anemia
 - DIC, most commonly
 - Malignant hypertension
 - SLE
 - Thrombotic thrombocytopenic purpura
 - Hemolytic uremic syndrome
 - Disseminated cancer



schistocytes

- Microangiopathic hemolytic anemia is not, by itself, a serious disease, but it points to a serious underlying disorders.

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infection

- Malaria:
 - Parasites within the RBCs, rupture resulting in hemolysis and episodic symptoms
 - Hematin released from the RBCs results in brown pigmentation of the spleen, liver and bone marrow
 - Massive splenomegaly and occasional hepatomegaly.
 - Falciparum can cause cerebral malaria which can be fatal.

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THANK YOU