



Classification of anemias

What is anemia, how do you diagnose anemia, and how are the different anemias classified?



Definition of anemia

- In its broadest sense, anemia is a functional inability of the blood to supply the tissue with adequate O_2 for proper metabolic function.
- Anemia is not a disease, but rather the expression of an underlying disorder or disease.
 - A specific diagnosis is made by:



Definition of anemia

- Patient history
- Patient physical exam
- Signs and symptoms exhibited by the patient
- Hematologic lab findings
- Identification of the cause of anemia is important so that appropriate therapy is used to treat the anemia.
- Anemia is usually associated with decreased levels of hemoglobin and/or a decreased packed cell volume (hematocrit), and/or a decreased RBC count.



Definition of anemia

- Occasionally there is an abnormal hemoglobin with an increased O₂ affinity resulting in an anemia with normal or raised hemoglobin levels, hematocrit, or RBC count.
- Before making a diagnosis of anemia, one must consider:
 - Age



Definition of anemia

- Sex
- Geographic location
- Presence or absence of lung disease
- Remember that the bone marrow has the capacity to increase RBC production **5-10 times the normal production.**
 - Thus, if all necessary raw products are available, the RBC life span can decrease to about **18 days** before bone marrow compensation is inadequate and anemia develops.



Definition of anemia

- An increased production of RBCs in the bone marrow is seen in the peripheral smear as an increased **reticulocyte** count since new RBCs are released as reticulocytes.
- If the bone marrow production of RBCs remains the same or is decreased with RBCs that have a decreased survival time, anemia will rapidly develop.



Definition of anemia

- There is no mechanism for increasing RBC survival time when there is an inadequate bone marrow response, so anemia will develop rapidly.
- In summary, anemia may develop:
 - When RBC loss or destruction exceeds the maximal capacity of bone marrow RBC production or
 - When bone marrow production is impaired



Definition of anemia

- Various diseases and disorders are associated with decreased hemoglobin levels. These include:
 - Nutritional deficiencies
 - External or internal blood loss
 - Increased destruction of RBCs
 - Ineffective or decreased production of RBCs



Definition of anemia

- Abnormal hemoglobin synthesis
- Bone marrow suppression by toxins, chemicals, or radiation
- Infection
- Bone marrow replacement by malignant cells



Significance of anemia and compensatory mechanisms

- The signs and symptoms of anemia range from slight fatigue to life threatening reactions depending upon
 - Rate of onset
 - Severity
 - Ability of the body to adapt



Rate of onset and severity

- With rapid loss of blood:
 - Up to 20% may be lost without clinical signs at rest, but with mild exercise the patient may experience tachycardia (rapid heart beat).
 - Loss of 30-40% leads to circulatory collapse and shock
 - Loss of 50% means that death is imminent



Rate of onset and severity

- In slowly developing anemias, a very severe drop in hemoglobin of up to 50% may occur without the threat of shock or death.
 - This is because the body has **adaptive or compensatory mechanisms** to allow the organs to function at hemoglobin levels of 50% of normal. These include:



Adaptive or compensatory mechanisms

- An increased heart rate, increased circulation rate, and increased cardiac output.
- Preferential shunting of blood flow to the vital organs.
- Increased production of 2,3 DPG, resulting in a shift to the right in the O_2 dissociation curve, thus permitting tissues to extract more O_2 from the blood.
- Decreased O_2 in the tissues leads to anaerobic glycolysis, which leads to the production of lactic acid, which leads to a decreased pH and a shift to the right in the O_2 dissociation curve. Thus, more O_2 is delivered to the tissues per blood cell.



Diagnosis of anemia

- How does one make a clinical diagnosis of anemia?
 - Patient history
 - Dietary habits
 - Medication
 - Possible exposure to chemicals and/or toxins
 - Description and duration of symptoms



Diagnosis of anemia

- Tiredness
- Muscle fatigue and weakness
- Headache and vertigo (dizziness)
- Dyspnea (difficult or labored breathing) from exertion
- G I problems
- Overt signs of blood loss such as hematuria (blood in urine) or black stools



Diagnosis of anemia

- Physical exam
 - General findings might include
 - Hepato or splenomegaly
 - Heart abnormalities
 - Skin pallor
 - Specific findings may help to establish the underlying cause:
 - In vitamin B₁₂ deficiency there may be signs of malnutrition and neurological changes
 - In iron deficiency there may be severe pallor, a smooth tongue, and esophageal webs
 - In hemolytic anemias there may be jaundice due to the increased levels of bilirubin from increased RBC destruction



Diagnosis of anemia

- Lab investigation. A complete blood count, CBC, will include:
 - An RBC count:
 - At birth the normal range is $3.9-5.9 \times 10^6/\text{ul}$
 - The normal range for males is $4.5-5.9 \times 10^6/\text{ul}$
 - The normal range for females is $3.8-5.2 \times 10^6/\text{ul}$
 - Note that the normal ranges may vary slightly depending upon the patient population.
 - Hematocrit (Hct) or packed cell volume in % or (L/L)
 - At birth the normal range is 42-60% (.42-.60)
 - The normal range for males is 41-53% (.41-.53)
 - The normal range for females is 38-46% (.38-.46)
 - Note that the normal ranges may vary slightly depending upon the patient population.



Diagnosis of anemia

- Hemoglobin concentration in grams/deciliter - the RBCs are lysed and the hemoglobin is measured spectrophotometrically
 - At birth the normal range is 13.5-20 g/dl
 - The normal range for males is 13.5-17.5 g/dl
 - The normal range for females is 12-16 g/dl
 - Note that the normal ranges may vary slightly depending upon the patient population.
- RBC indices – these utilize results of the RBC count, hematocrit, and hemoglobin to calculate 4 parameters:
 - Mean corpuscular volume (MCV) – is the average volume/RBC in femtoliters (10^{-15} L)
 - $\text{Hct (in \%)/RBC (x } 10^{12}/\text{L)} \times 10$
 - At birth the normal range is 98-123
 - In adults the normal range is 80-100



Diagnosis of anemia

- The MCV is used to classify RBCs as:
 - Normocytic (80-100)
 - Microcytic (<80)
 - Macrocytic (>100)
- Mean corpuscular hemoglobin concentration (MCHC) – is the average concentration of hemoglobin in g/dl (or %)
 - $\text{Hgb (in g/dl)} / \text{Hct (in \%)} \times 100$
 - At birth the normal range is 30-36
 - In adults the normal range is 31-37
 - The MVHC is used to classify RBCs as:
 - Normochromic (31-37)
 - Hypochromic (<31)
 - Some RBCs are called hyperchromic, but they don't really have a higher than normal hgb concentration.

Normocytic cell

Erythrocytes - Alterations in erythrocyte size

NORMOCYTE CHRONOLAB

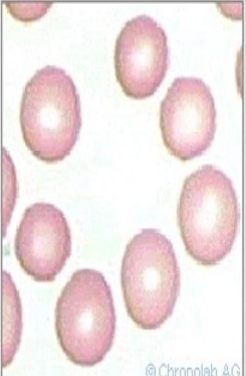
| DESCRIPTION | VARIATIONS | CONFUSIONS |
|--------------------------------------------------------------------------------------------------------------------------------|------------|------------|
| <p>Description:</p> <p>Normal size and volume, mean cell volume (MCV) 80-100 fl.</p> | | |
| <p>Shape:</p> <p>Flexible biconcave, discoid shape. Non-nucleated cell, no Golgi's complex, centriole or lysosomes.</p> | | |
| <p>Significance:</p> <p>Healthy state.</p> | | |

Staining method:
May-Grünwald/Giemsa

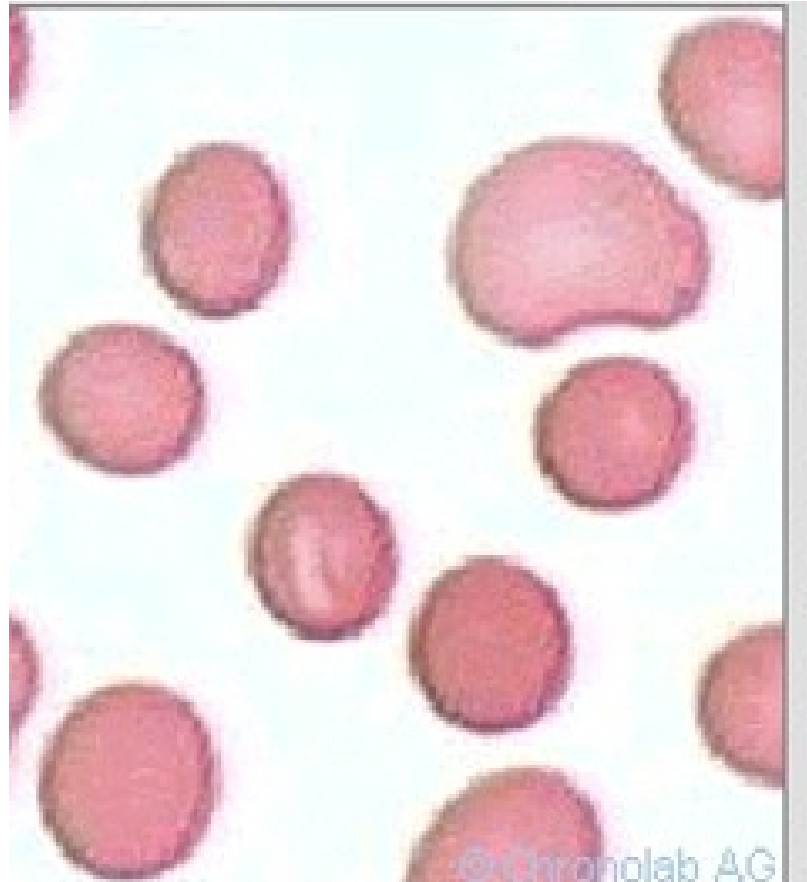
Microscope: Light

Magnification: 1:1000

Size:
7 - 8,5 μm



Microcytic cell




Macrocytic cell

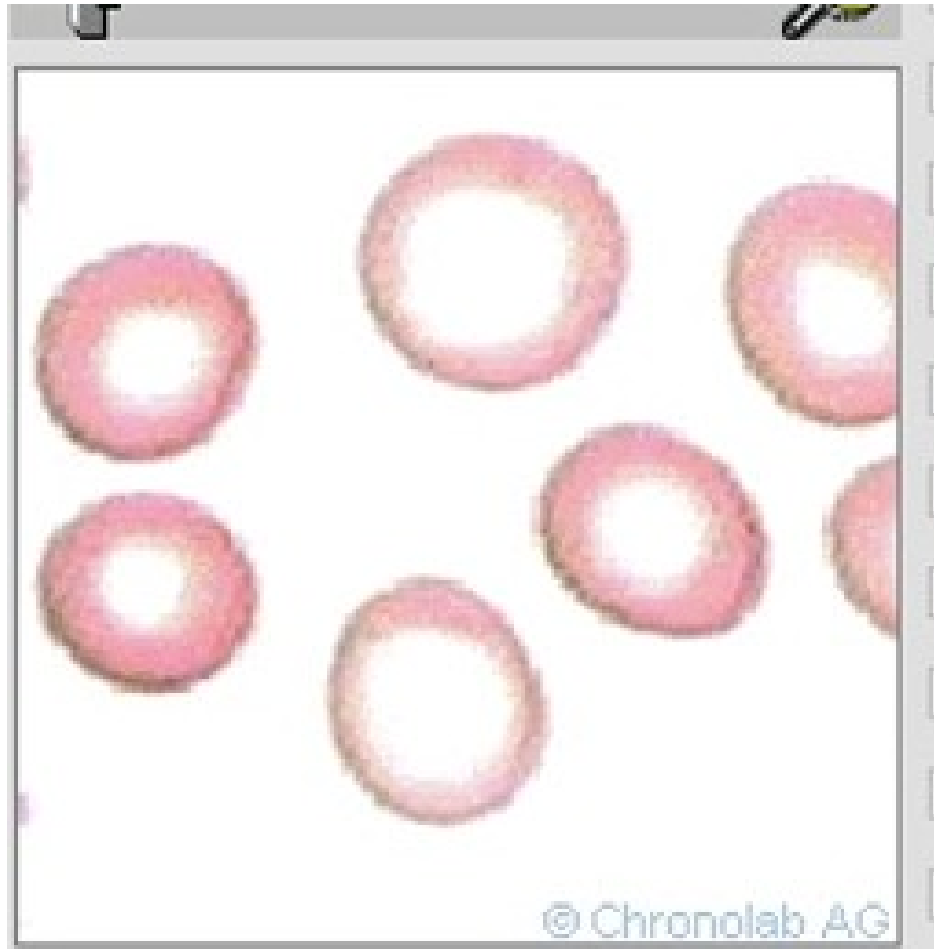


Normochromic cell

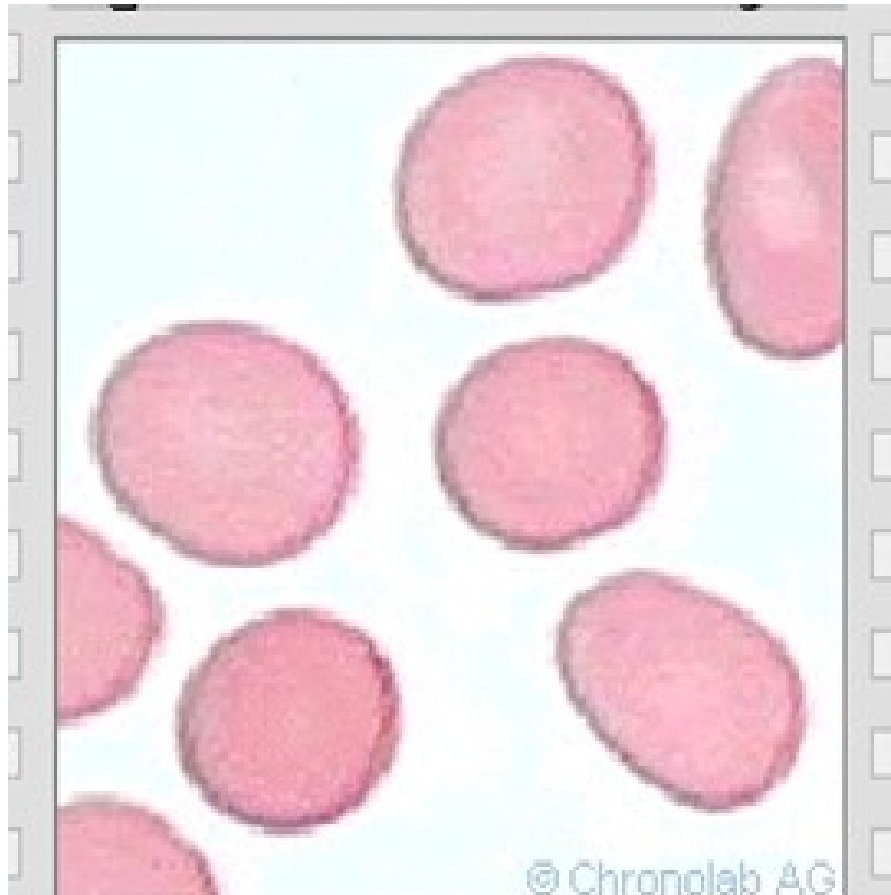
NORMOCHROMIC ERYTHROCYTE
CHRONOLAB

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|-----------------------------------------------------------------------------------------------------------------------------|------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <p>Description: Normally colored.</p> | |  <p>© Chronolab AG</p> |
| <p>Shape: Flexible biconcave, discoid shape. Non-nucleated cell, no Golgi's complex, centriole or lysosomes.</p> | | <p>Staining method: <input type="checkbox"/> May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> <p>Size: 7 - 8,5 μm</p> |

Hypochromic cell



Hyperchromic cell

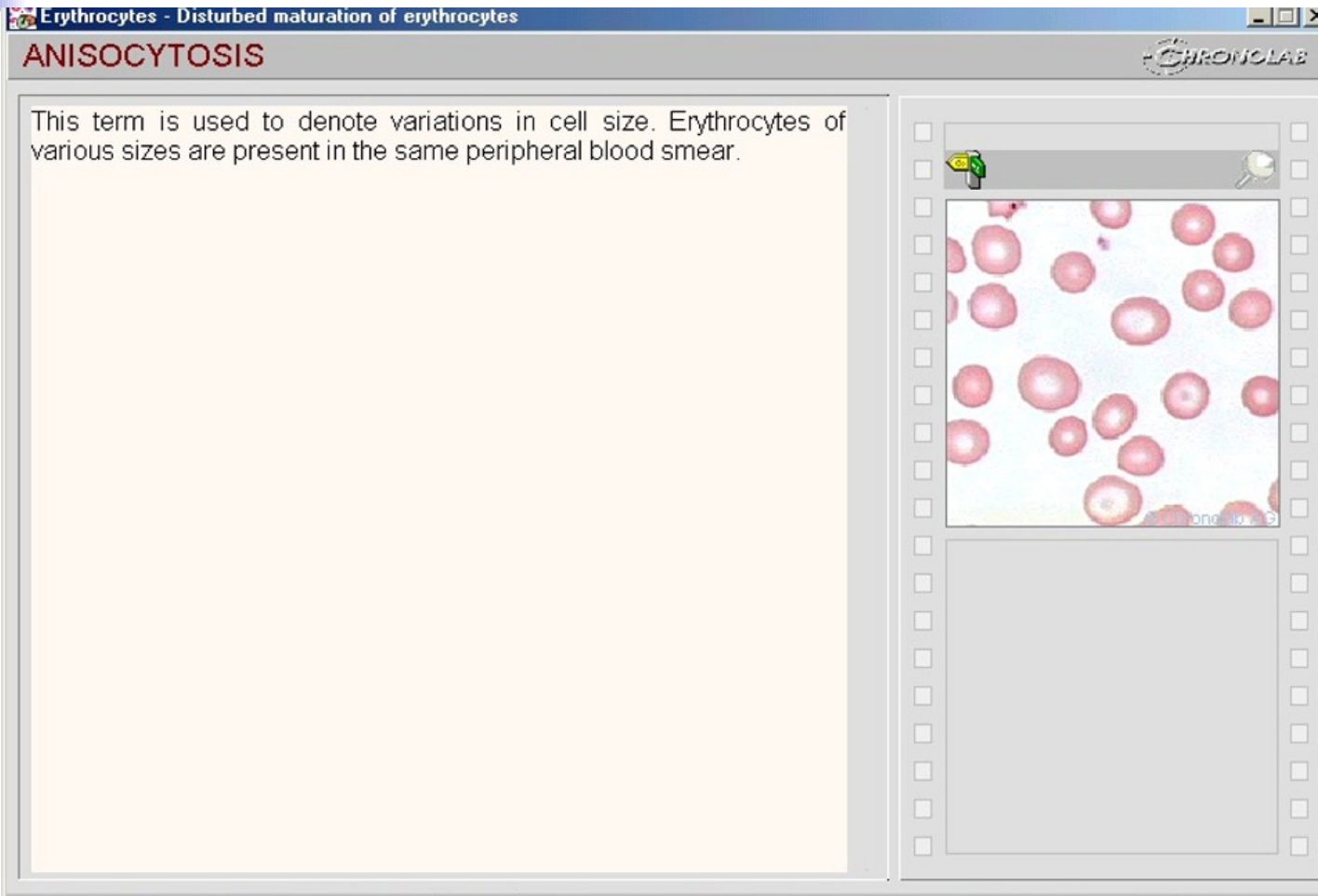




Diagnosis of anemia

- Mean corpuscular hemoglobin (MCH) – is the average weight of hemoglobin/cell in picograms ($\text{pg} = 10^{-12} \text{ g}$)
 - $\text{Hgb (in g/dl)} / \text{RBC} (\times 10^{12} / \text{L}) \times 10$
 - At birth the normal range is 31-37
 - In adults the normal range is 26-34
 - This is not used much anymore because it does not take into account the size of the cell.
- Red cell distribution width (RDW) – is a measurement of the variation in RBC cell size
 - $\text{Standard deviation} / \text{mean MCV} \times 100$
 - The range for normal values is 11.5-14.5%
 - A value > 14.5 means that there is increased variation in cell size above the normal amount (anisocytosis)
 - A value < 11.5 means that the RBC population is more uniform in size than normal.

Anisocytosis





Diagnosis of anemia

- Reticulocyte count gives an indication of the level of the bone marrow activity.
 - Done by staining a peripheral blood smear with new methylene blue to help visualize remaining ribosomes and ER. The number of reticulocytes/1000 RBC is counted and reported as a %.
 - At birth the normal range is 1.8-8%
 - The normal range in an adult (i.e. in an individual with no anemia) is .5-1.5%. **Note that this % is not normal for anemia where the bone marrow should be working harder and throwing out more reticulocytes per day. In anemia the reticulocyte count should be elevated above the normal values.**

Reticulocytes

Description:

The development begins with the extrusion of the normoblast nucleus and ends when the reticulocyte has lost its organelles.

Cytoplasm:

In panoptic stains the cytoplasm is blue-pink. When stained by brilliant-cresyl blue, the cytoplasm shows characteristic blue-purple fibers, remnants of the endoplasmatic reticulum.

Nucleus:

Absent.

Nucleoli:

Not visible.

Significance:

By determining the number of reticulocytes in the peripheral blood, the erythropoietic activity of the bone marrow can be estimated. ▼



Staining method:

Supravital stain brilliant cresyl blue

Microscope: Light

Magnification: 1:1000

Size:

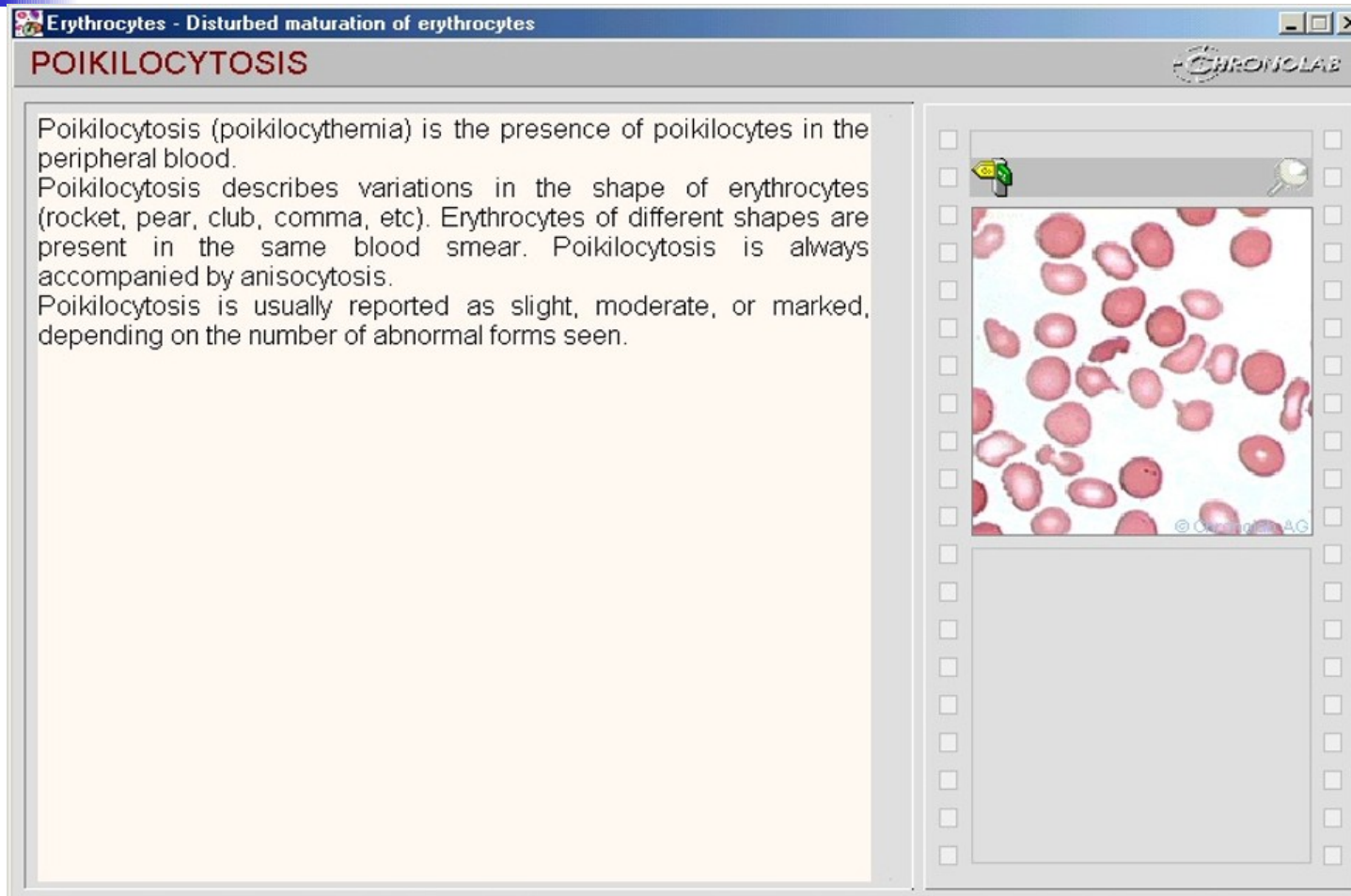
7-9 μm




Diagnosis of anemia

- The numbers reported above are only **relative values**. To get a better indication of what is really going on, a corrected reticulocyte count (patients Hct/.45 (a normal Hct) x the reticulocyte count) or an absolute count (% reticulocytes x RBC count) should be done.
- As an anemia gets more severe, younger cells that take longer than 24 hours to mature, are thrown out into the peripheral blood (shift reticulocyte). This may also be corrected for to give the **reticulocyte production index (RPI)** which is a truer indication of the real bone marrow activity.
- Blood smear examination using a Wright's or Giemsa stain. The smear should be evaluated for the following:
 - **Poikilocytosis** – describes a variation in the shape of the RBCs. It is normal to have some variation in shape, but some shapes are characteristic of a hematologic disorder or malignancy.

Poikilocytosis



Spherocytes



Erythrocytes - Alterations in erythrocyte shape

SPHEROCYTE

DESCRIPTION

VARIATIONS

CONFUSIONS

Description:

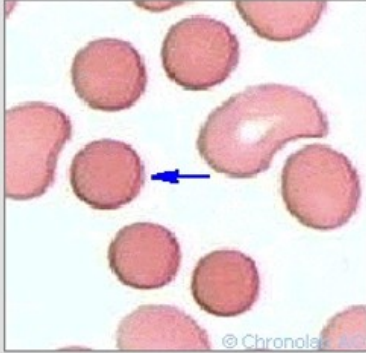
Small erythrocytes containing an increased concentration of hemoglobin, resulting from a loss of the red cell membrane. Their life span in the circulation is extremely short, and osmotic resistance is decreased to hypotonic solution.

Shape:

Spherical cells with very increased thickness (about 3 μm) and reduced diameter (about 6 μm), normal volume and biconcave shape.

Significance:

Spheroid cells occur in hereditary spherocytosis and hemolytic anemia.



© Chronolab

Staining method:
May-Grünwald/Giemsa

Microscope: Light

Magnification: 1:1000


Size:

about 6 μm


Ovalocytes (elliptocytes)

Erythrocytes - Alterations in erythrocyte shape

OVALOCYTE (ELLIPTOCYTE, PENCIL CELL, CIGAR CELL)

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|----------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Results from a defect in the red cell membrane. The life span in the circulation is shortened, and the osmotic resistance is decreased in most cases. Hemolysis is not as marked as in spherocytosis.</p> | |  <p>© Chronolab AG</p> |
| <p>Shape:</p> <p>Elliptic shape is apparent after the reticulocyte stage.</p> | | <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> |
| <p>Significance:</p> <p>Found in hereditary elliptocytosis. Can also be found in iron deficiency anemia, sickle diseases, thalassemia, and myelofibrosis.</p> | | <p>Size: 7 - 8,5 µm</p> |

Leptocyte



Erythrocytes - Alterations in erythrocyte shape

LEPTOCYTE

DESCRIPTION

VARIATIONS

CONFUSIONS

Description:

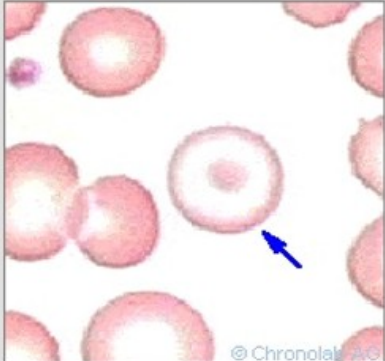
Erythrocytes of normal volume and very reduced thickness. In the smears they appear like anulocytes or "target" cells. Anulocyte has a more or less large, pale area in the center due to the lowered hemoglobin content. "Target" cells ("Mexican-hat cell") have ring zones on the periphery as well, but there is an increase of hemoglobin in the central part. An area of pallor between peripheral and central parts is visible.

Shape:

Center and the periphery are colored and separated by a clear ring. So, it has the appearance of a target (bull's-eye).

Significance:

Found most frequently in iron deficiency - sideropenic anemia, as a result of disturbances in the hemoglobin synthesis. They can be found in hypochromic anemia, homozygous and heterozygous forms of C, D, E hemoglobinopathies. The presence of leptocytes in β -thalassemia major is a pathognomonic.


© Chronolab AG

Staining method:

May-Grünwald/Giemsa

Microscope: Light

Magnification: 1:1000

Size:

Changeable μm

Acanthocyte

Erythrocytes - Alterations in erythrocyte shape

ACANTHOCYTE (SPUR CELL)

DESCRIPTION

Description:

Erythrocytes that have lost their discoid shape and which have 8-10 spicules of uneven length irregularly distributed over the red cell surface. They have a star-like shape. Lack a central area of pallor.

Shape:

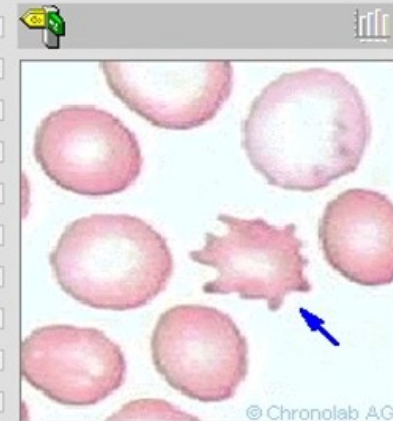
Acanthocytes appear smaller than discoid cells because they have spheroidal shape. Characteristically, the spicules are distributed irregularly over the surface, and individual spicules differ from each other. It resembles head burdock.

Significance:

They can be found in patients with an inherited decrease or complete deficiency of β -lipoproteins (abetalipoproteinemia) and in patients with some neurological diseases. They can sometimes be found in azotemia, stomach cancer, bleeding stomach ulcer, and vitamin E deficiency.

VARIATIONS

CONFUSIONS



Staining method:

May-Grünwald/Giemsa

Microscope: Light

Magnification: 1:1000


Size:

Changeable μm

Stomatocyte

Erythrocytes - Alterations in erythrocyte shape

STOMATOCYTE (MOUTH CELL, CUP FORM, MUSHROOM CAP)


| DESCRIPTION | VARIATIONS | CONFUSIONS |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|----------------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Erythrocytes with a linear central zone, mouth-shaped area of pallor as a result of the disturbance of erythrocyte membrane.</p> | |  |
| <p>Shape:</p> <p>May have a form of fish mouth.</p> | | <p>Staining method: May-Grünwald/Giemsa</p> <p>Microscope: Light</p> <p>Magnification: 1:1000</p> |
| <p>Significance:</p> <p>Present in a rare type of congenital hemolytic anemia, when intracellular potassium is decreased and sodium increased (hereditary stomatocytosis), and in alcoholism.</p> | | <p>Size:</p> <p>7 - 8,5 μm</p> |

Schistocyte

Erythrocytes - Alterations in erythrocyte shape

SCHIZOCYTE (SCHISTOCYTE, FRAGMENTED CELL)

CHRONOLAB

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|-------------------------------------------------------------------------------------------------------------------------------------|------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Erythrocyte fragment, 2-4 μm in diameter, due to localized membrane damage.</p> | | <div><p>© Chronolab A.C.</p></div> <p>Staining method: May-Grünwald/Giemsa</p> <p>Microscope: Light</p> <p>Magnification: 1:1000</p> <p>Size: 2 - 4 μm</p> |

Dacrocyte

DACRYOCYTE (TEAR DROP, TENNIS RACQUET CELL)

DESCRIPTION

Description:

Dacryocytes or "tear drop" cells are in fact discocytes drawn into a spicule on one end. They can be of normal, reduced or increased size.

Shape:

Pear shaped with a more or less extended tail that sometimes ends in a swelling.

Significance:

They can be found in patients with myelofibrosis with myeloid metaplasia and metastatic cancer to the bone marrow.

VARIATIONS

CONFUSIONS



Staining method:

May-Grünwald/Giemsa

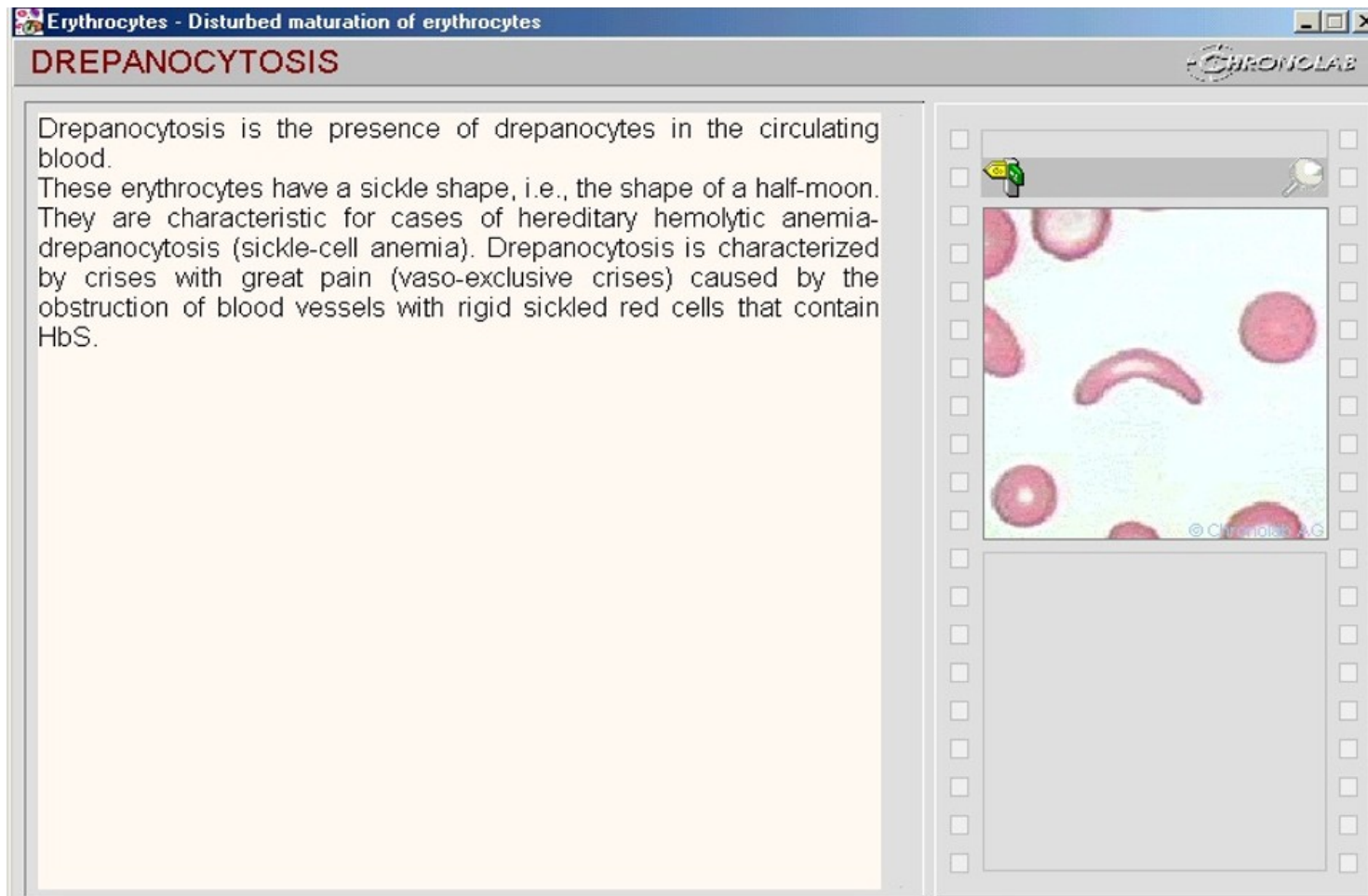
Microscope: Light

Magnification: 1:1000

Size:

Changeable μm

Sickle cells (depranocytes)




Macroovalocyte

Erythrocytes - Disturbed maturation of erythrocytes

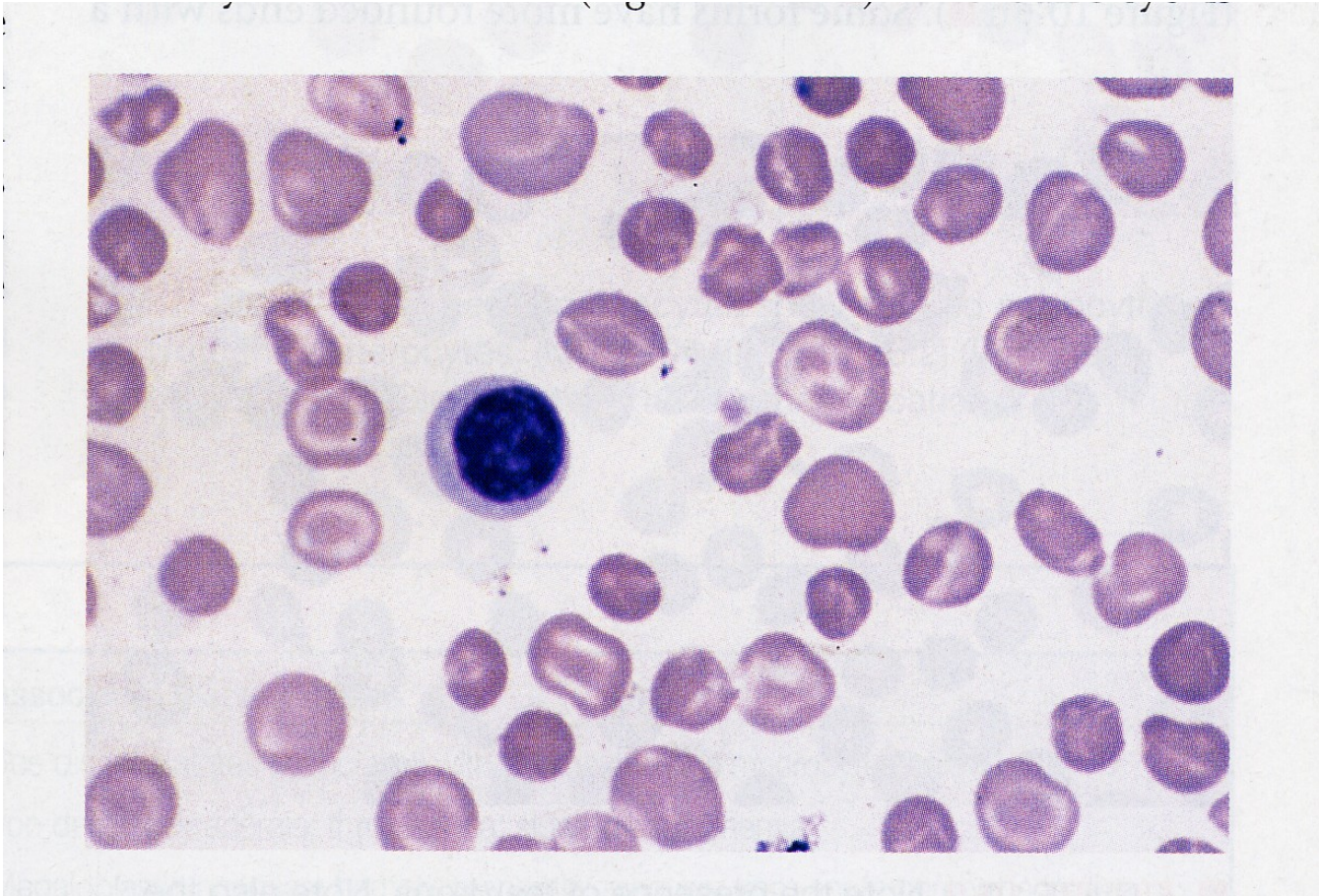
MEGALOCYTOSIS

Megalocytosis (macrocythemia) is the occurrence of unusually large numbers of macrocytes. Megalocytes have increased longitudinal diameter (oval), thickness, and volume. They can be observed in vitamin B₁₂ and folic acid deficiencies. Also called macroovalocytes.

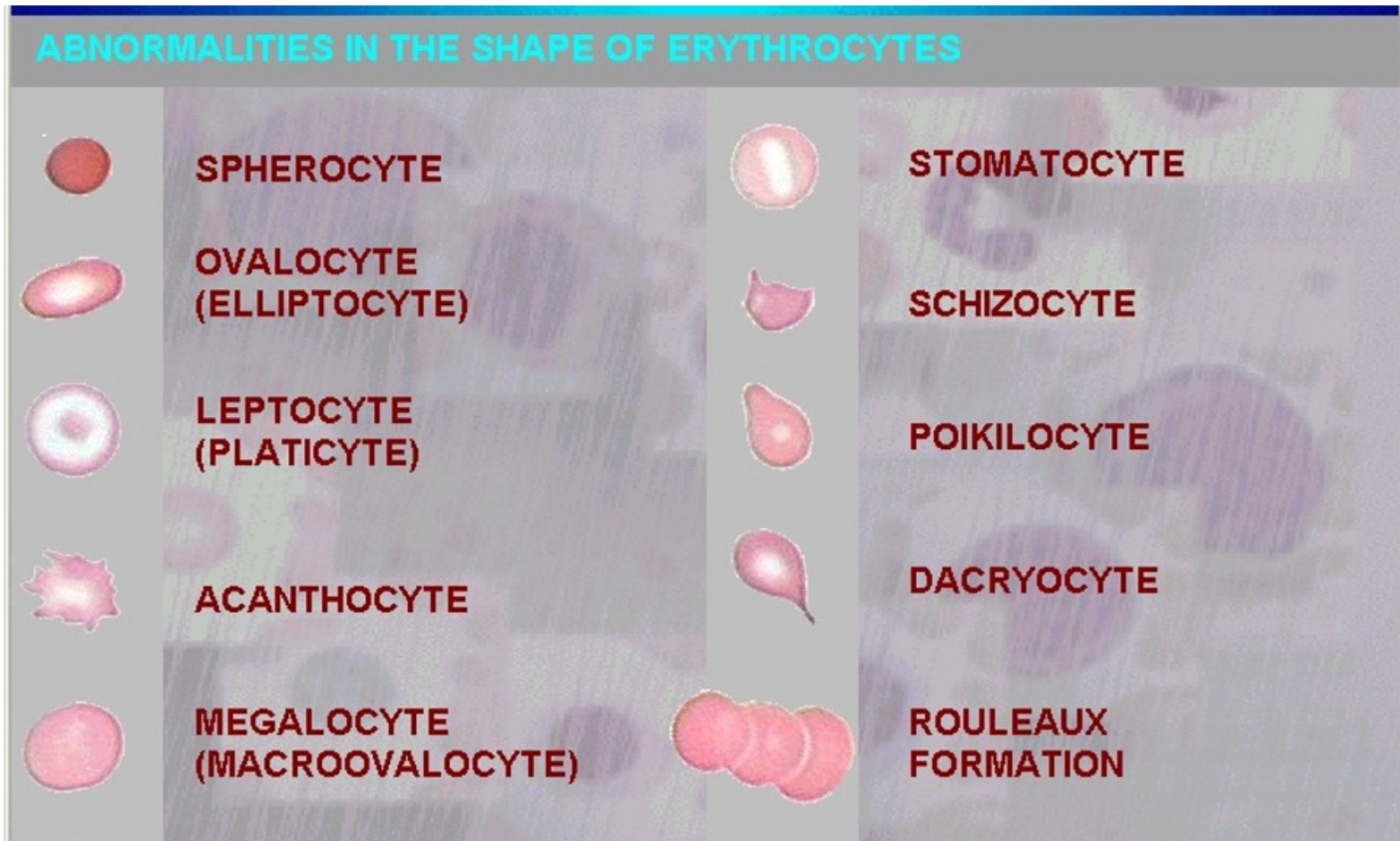


© ChronoLab AG

Target cells



Summary of variations in RBC shape (poikilocytosis)






Diagnosis of anemia

- Erythrocyte inclusions – the RBCs in the peripheral smear should also be examined for the presence of inclusions:

Cabot's rings

| CABOT'S RINGS | | CHRONOLAB |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------|
| DESCRIPTION | VARIATIONS | CONFUSIONS |
| <p>Description:</p> <p>Basophilic rings or loops, remnants of the nuclear membranes.</p> | |  <p>© Chronolab AG</p> |
| <p>Shape:</p> <p>Purplish rings, figures-of-eight, incomplete rings or similar configurations of a reddish-violet fine filament appearing in the center or near the periphery of erythrocytes.</p> | | |
| <p>Significance:</p> <p>Cabot's rings have been described in pernicious anemia, lead poisoning, leukemia, alcoholic jaundice, congenital dyserythropoietic anemia and in some forms of severe anemia. They can be seen in patients taking cytostatics.</p> | <p>Staining method: May-Grünwald/Giemsa</p> <p>Microscope: Light</p> <p>Magnification: 1:1000</p> <p>Size: Changeable μm</p> | |

Howell-Jolly bodies

HOWELL - JOLLY BODIES

CHRONOLAB

DESCRIPTION

VARIATIONS

CONFUSIONS

Description:

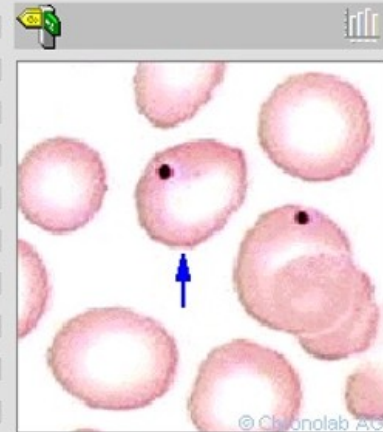
Howell-Jolly bodies are nuclear remnants, usually located eccentrically in reticulocytes and erythrocytes. They are generally derived from chromosomes during abnormal divisions. Similar small nuclear fragments may be produced by pathologic fragmentation of the nucleus (karyorrhexis).

Shape:

Round body inclusions that stain homogeneously dark purple by Romanowsky stains.

Significance:

Very rare in normal people but common after the removal of the spleen, in some hemolytic anemias and in megaloblastic anemia. They may occur in very severe anemias, especially megaloblastic anemia.



Staining method:

May-Grünwald/Giemsa

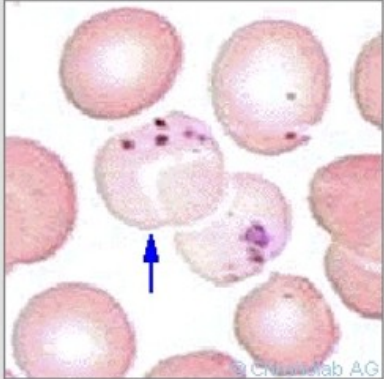
Microscope: Light

Magnification: 1:1000

Size:


1 - 1,5 μm

Nuclear dust



| NUCLEAR DUST IN ERYTHROCYTES | | CHRONOLAB |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| DESCRIPTION | VARIATIONS | CONFUSIONS |
| <p>Description:</p> <p>Fine, small, reddish-blue granules spread in erythrocytes; residues of nuclear material. They must be differentiated from basophilic granules. They occur exclusively in non-nucleated erythrocytes.</p> | |  <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> <p>Size: Changeable μm</p> |
| <p>Shape:</p> <p>Smaller than Howell-Jolly bodies and usually only just visible.</p> | | |
| <p>Significance:</p> <p>They reflect severe anemias.</p> | | |

Basophilic stippling

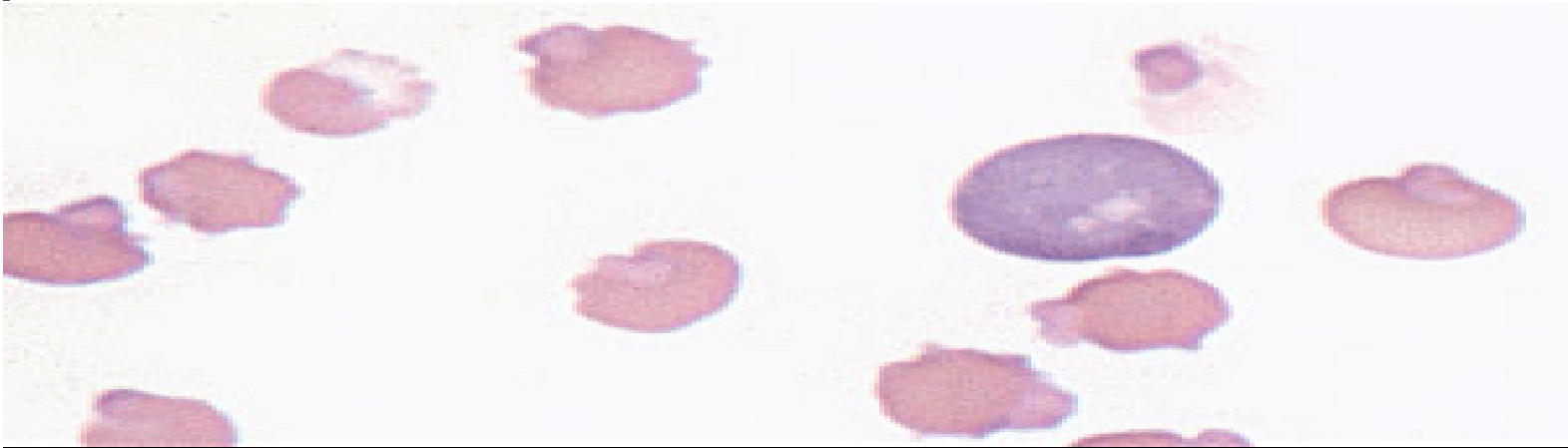
BASOPHILIC STIPPLING

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Erythrocytes with basophilic stippling are cells with bluish-black granular inclusions and usually contain 10-20 blue colored granules. Basophilic punctations are partially or completely made of RNA. They can be seen in smears stained with a Romanowsky stain. They can be counted after staining with methylazur after Manson-Schwartz method.</p> | | <div><p>© Chronolab AG</p></div> <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> <p>Size: Changeable μm</p> |
| <p>Shape:</p> <p>Basophilic punctations in erythrocytes are round or irregularly shaped granules of variable size which are stained blue by Giemsa.</p> | | |
| <p>Significance:</p> <p>They are frequent in anemias as a result of lead or other heavy metals poisoning.</p> | | |

Heinz bodies


| HEINZ BODIES | | CHRONOLAB |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| DESCRIPTION | VARIATIONS | CONFUSIONS |
| <p>Description:</p> <p>Heinz bodies are composed of denatured proteins, primarily hemoglobin. They are formed by intra-erythrocytic precipitation of hemoglobin.</p> | | <div></div> <div></div> <div></div> <div>Size: Changeable μm</div> |
| <p>Shape:</p> <p>They can be visualized by special staining procedures, e.g., acetyl-phenyl-hydrazine.</p> | | |
| <p>Significance:</p> <p>They may be found in the following conditions: anemias caused by G₆PD deficiency, phenylhydrazine therapy, sulphanilamide therapy, α-thalassemia.</p> | | |

Heinz bodies (new methylene blue stain)

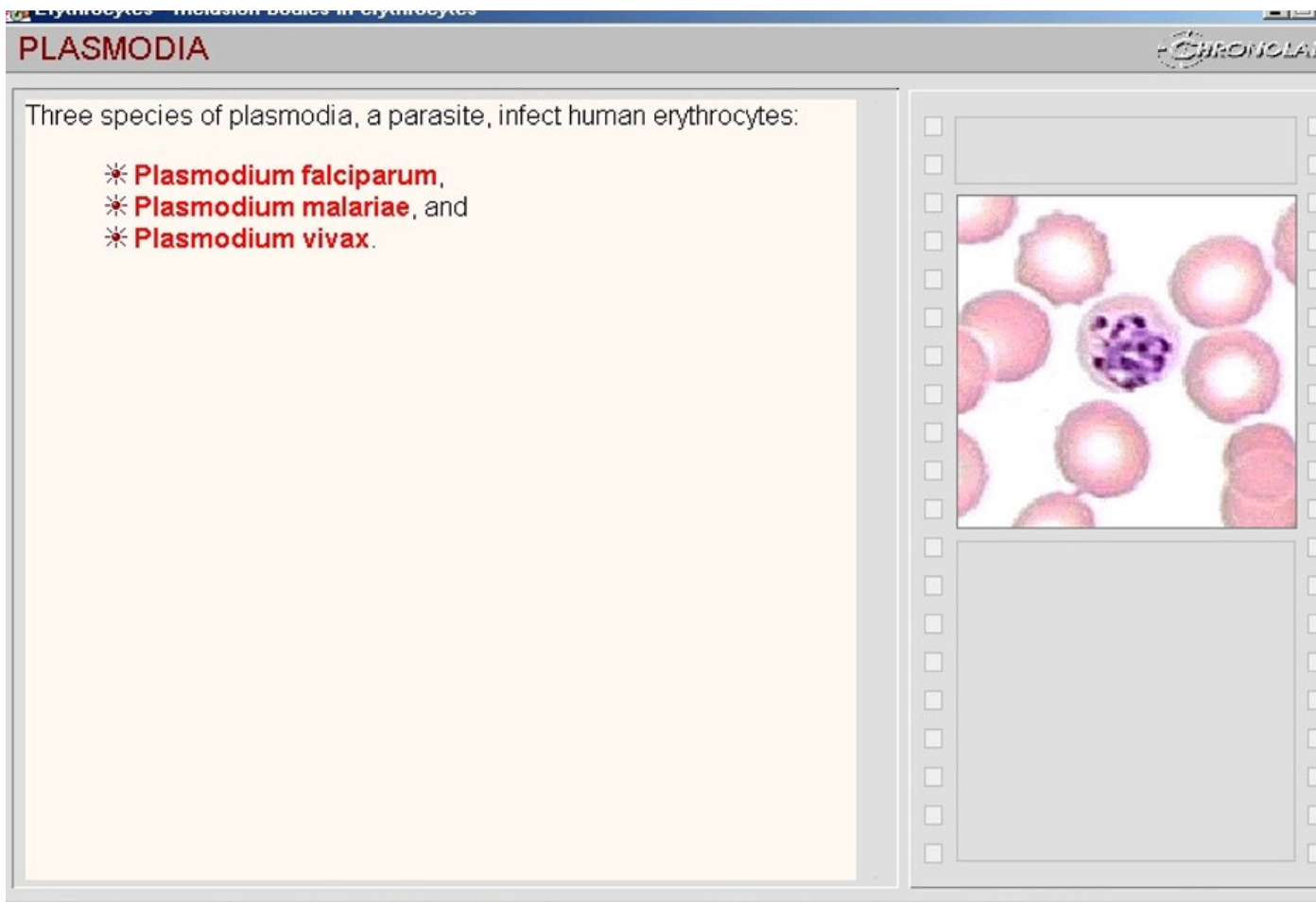


Siderocytes

SIDEROCYTES

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Erythrocytes containing one or more iron-containing granules of non-hemoglobin iron, giving a positive Prussian blue reaction. Granules appear as basophilic granules. In the bone marrow smears in healthy people, the number of sideroblasts (normoblasts containing iron granules) is 20-90%. The granule size is about 2 μm.</p> | | <div><p>© Chronolab AG</p></div> <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> <p>Size: Changeable μm</p> |
| <p>Shape:</p> <p>Granules appear as basophilic granules.</p> | | |
| <p>Significance:</p> <p>In healthy people, siderotic granules are not normally found in peripheral blood erythrocytes. They may be found in the peripheral blood in disorders associated with impaired hemoglobin synthesis, e.g., sideroblastic anemia, thalassemia and lead poisoning. They are also present in blood after splenectomy.</p> | | |

Plasmodium (malarial parasite)

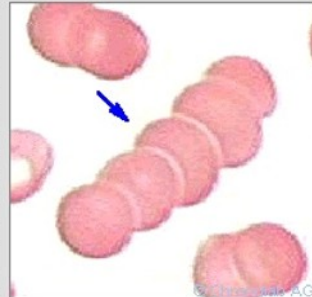


Diagnosis of anemia

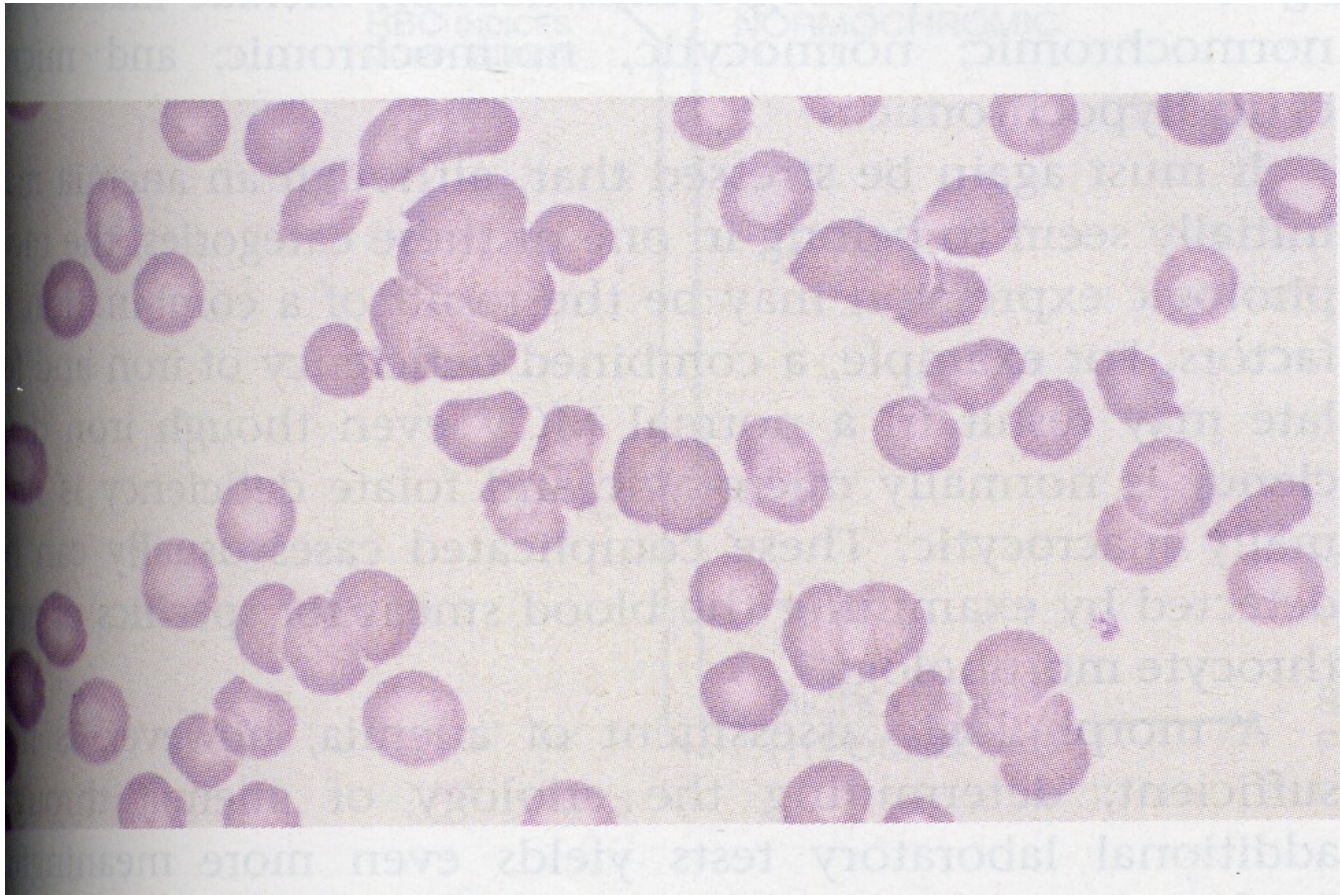
- A variation in erythrocyte distribution such as rouleaux formation or agglutination

Erythrocytes - Alterations in erythrocyte shape

ROULEAUX FORMATION

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|----------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Larger or smaller aggregation of erythrocytes.</p> | |  |
| <p>Shape:</p> <p>Erythrocytes in freshly taken blood aggregate on each other like stacked coins.</p> | | <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> |
| <p>Significance:</p> <p>Present in thick portions of smears, especially in patients with increased erythrocyte sedimentation rate. Can also be found pathologically in multiple myeloma.</p> | | <p>Size: Changeable μm</p> |

Agglutination of RBCs






Diagnosis of anemia

- A variation in size should be noted (anisocytosis) and cells should be classified as
 - Normocytic
 - Microcytic
 - Macrocytic
- A variation in hemoglobin concentration (color) should be noted and the cells should be classified as
 - Normochromic
 - Hypochromic
 - Hyperchromic
- Polychromasia (pinkish-blue color due to an increased % of reticulocytes) should be noted

Normocytic RBC


| Erythrocytes - Alterations in erythrocyte size | | |
|--------------------------------------------------------------------------------------------------------------------------------|------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| NORMOCYTE | | |
| DESCRIPTION | VARIATIONS | CONFUSIONS |
| <p>Description:</p> <p>Normal size and volume, mean cell volume (MCV) 80-100 fl.</p> | |  <p>© Chronolab AG</p> <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> <p>Size: 7 - 8,5 µm</p> |
| <p>Shape:</p> <p>Flexible biconcave, discoid shape. Non-nucleated cell, no Golgi's complex, centriole or lysosomes.</p> | | |
| <p>Significance:</p> <p>Healthy state.</p> | | |

Microcytic RBC

Erythrocytes - Alterations in erythrocyte size

MICROCYTE


CHRONOLAB

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|-------------------------------------------------------------------------------------------------------------------------------------------------|------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Smaller diameter and volume than normal, round or slightly oval red cells, normal thickness (MCV < 70 fl).</p> | |  <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> <p>Size: < 6 µm</p> |
| <p>Shape:</p> <p>Flexible biconcave, discoid shape.</p> | | |
| <p>Significance:</p> <p>Iron deficiency, thalassemias, sideroblastic anemia.</p> | | |


Macrocytic RBC

Erythrocytes - Alterations in erythrocyte size

MACROCYTE

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|----------------------------------------------------------------------------------------------------------------------------------|------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Larger volume than normal (MCV > 100 fl).</p> | |  <p>A microscopic image showing several large, pink, spherical red blood cells (macrocytes) against a light blue background. A blue arrow points to one of the cells. The cells are significantly larger than normal red blood cells.</p> |
| <p>Shape:</p> <p>Discoid shape, a complete mature cell filled with hemoglobin.</p> | | <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> |
| <p>Significance:</p> <p>Liver disease, drug induced anemia, Vitamin B₁₂ deficiency, folic acid deficiency.</p> | | <p>Size: 9 - 10 μm</p> |

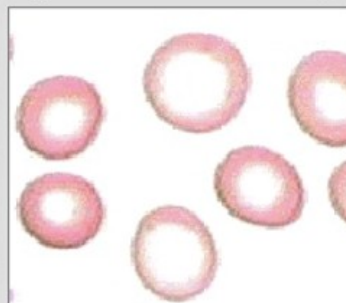
Normochromic RBC

| NORMOCHROMIC ERYTHROCYTE | | CHRONOLAB |
|--------------------------------------------------------------------------------------------------------------------------------|------------|-------------------------------------------------------------------------------------------------------------------------------|
| DESCRIPTION | VARIATIONS | CONFUSIONS |
| <p>Description:</p> <p>Normally colored.</p> | |  <p>© Chronolab AG</p> |
| <p>Shape:</p> <p>Flexible biconcave, discoid shape. Non-nucleated cell, no Golgi's complex, centriole or lysosomes.</p> | | |
| | | <p>Staining method:</p> <p>May-Grünwald/Giemsa</p> <p>Microscope: Light</p> <p>Magnification: 1:1000</p> |
| | | <p>Size:</p> <p>7 - 8,5 μm</p> |

Hypochromic RBC

Erythrocytes - Alterations in erythrocyte coloration

HYPOCHROMIC ERYTHROCYTE

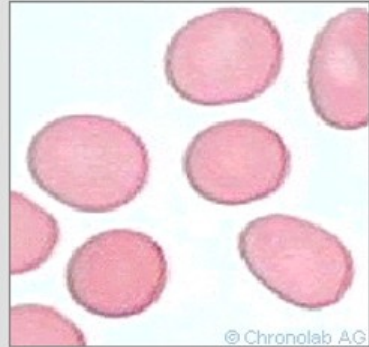
| DESCRIPTION | VARIATIONS | CONFUSIONS |
|------------------------------------------------------------------------------------------------------|------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Increased area of pallor to $>1/3$ diameter of cell.</p> | |  <p>© Chronolab AG</p> <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> <p>Size: 7 - 8,5 μm</p> |

Hyperchromic RBC

Erythrocytes - Alterations in erythrocyte coloration

HYPERCHROMIC ERYTHROCYTE

CHRONOLAB

| DESCRIPTION | VARIATIONS | CONFUSIONS |
|---------------------------------------------------------------------------------------------------------------|------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <p>Description:</p> <p>Intensive coloration, "stuck-on" appearance. No central area of pallor.</p> | |  <p>© Chronolab AG</p> <p>Staining method: May-Grünwald/Giemsa Microscope: Light Magnification: 1:1000</p> <p>Size: 7 - 8,5 μm</p> |

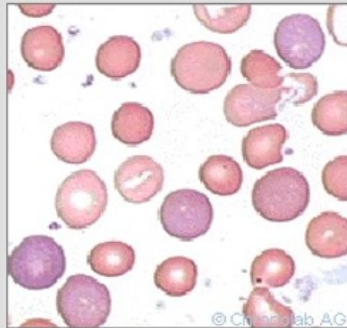
Polychromasia

POLYCHROMIA (POLYCHROMASIA)

Erythrocytes stain bluish. This is the result of the persistence of RNA and is a sign of cell immaturity.







Degrees:

- * light polychromasia (+);
- * polychromasia (++);
- * marked polychromasia (+++).



The image shows a microscopic view of a blood smear. Numerous red blood cells (erythrocytes) are visible. Many of these cells exhibit a bluish or slate-colored tint, which is characteristic of polychromasia. This bluish tint is due to the presence of residual RNA within the cells, indicating they are immature. The cells vary in size and the intensity of their bluish staining, representing different degrees of polychromasia. A copyright notice '© ChronoLab AG' is visible in the bottom right corner of the image.

Summary of variations in color and size

| VARIATIONS IN ERYTHROCYTE COLOR | | ABNORMALITIES IN ERYTHROCYTES SIZE | |
|-------------------------------------------------------------------------------------|---------------------------------|---------------------------------------------------------------------------------------|------------------|
|  | NORMOCHROMIC ERYTHROCYTE |  | NORMOCYTE |
|  | HYPOCHROMIC ERYTHROCYTE |  | MICROCYTE |
|  | HYPERCHROMIC ERYTHROCYTE |  | MACROCYTE |



Diagnosis of anemia

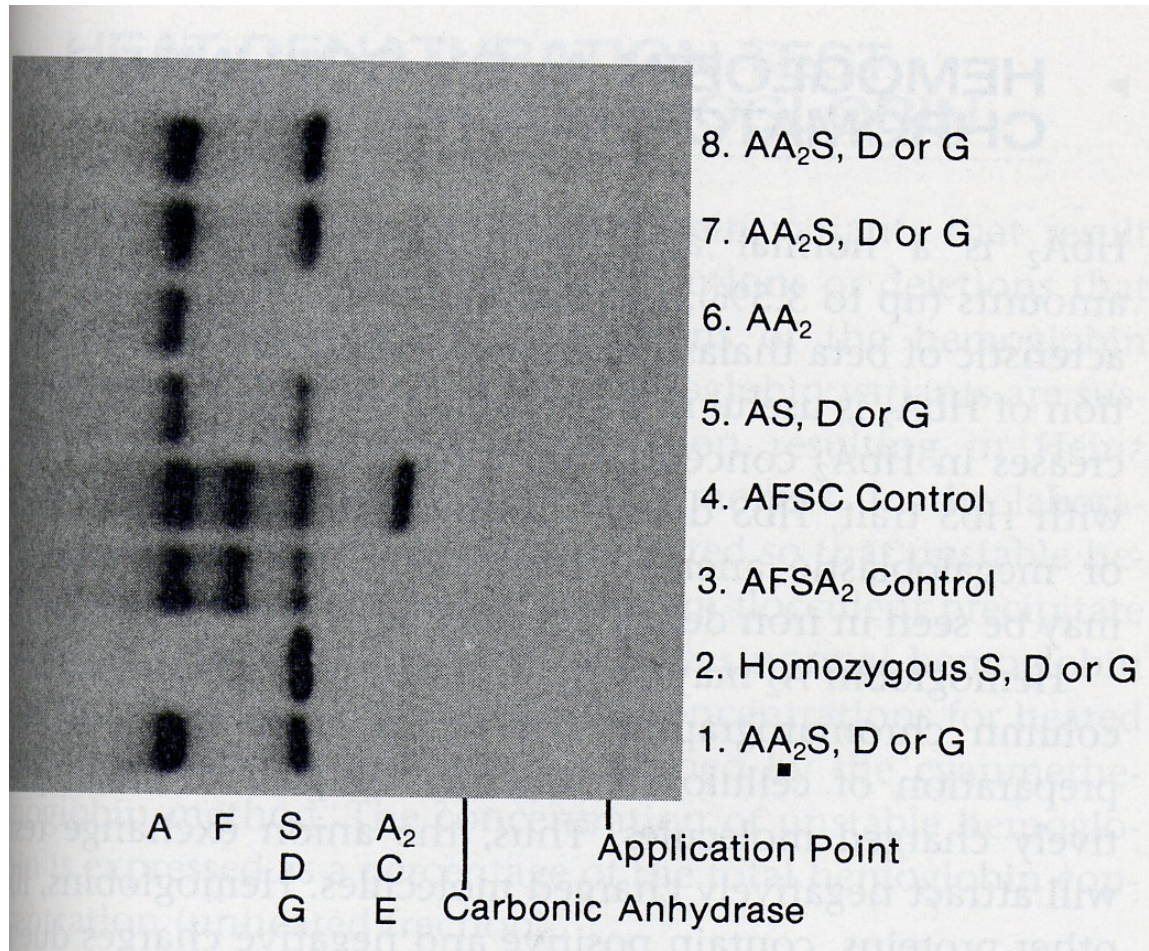
- The peripheral smear should also be examined for abnormalities in leukocytes or platelets.
 - Some nutritional deficiencies, stem cell disorders, and bone marrow abnormalities will also effect production, function, and/or morphology of platelets and/or granulocytes.
 - Finding abnormalities in the leukocytes and/or platelets may provide clues as to the cause of the anemia.
- The lab investigation may also include:
 - A bone marrow smear and biopsy
 - Used when other tests are not conclusive



Diagnosis of anemia

- In a bone marrow sample, the following things should be noted:
 - Maturation of RBC and WBC series
 - Ratio of myeloid to erythroid series
 - Abundance of iron stores (ringed sideroblasts)
 - Presence or absence of granulomas or tumor cells
 - Red to yellow ratio
 - Presence of megakaryocytes
- Hemoglobin electrophoresis – can be used to identify the presence of an abnormal hemoglobin (called hemoglobinopathies). Different hgbs will move to different regions of the gel and the type of hemoglobin may be identified by its position on the gel after electrophoresis.

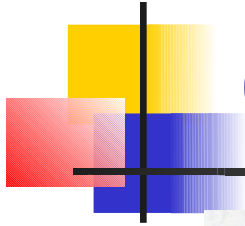
Hemoglobin electrophoresis



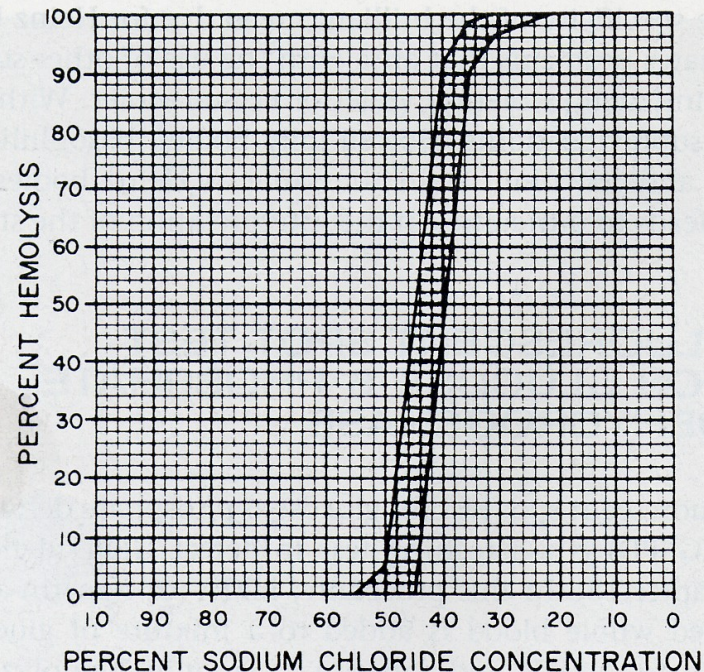


Diagnosis of anemia

- Antiglobulin testing – tests for the presence of antibody or complement on the surface of the RBC and can be used to support a diagnosis of an autoimmune hemolytic anemia.
- Osmotic fragility test – measures the RBC sensitivity to a **hypotonic** solution of saline. Saline concentrations of 0 to .9% are incubated with RBCs at room temperature and the percent of hemolysis is measured. Patients with spherocytes (missing some membrane) have increased osmotic fragility. They have a limited ability take up water in a hypotonic solution and will, therefore, lyse at a **higher** sodium concentration than will normal RBCs



Normal osmotic fragility curve



■ **FIGURE 40-4** Normal osmotic fragility curve. The osmotic fragility curve of a normal individual would fall within the area defined by the two sigmoid curves. A curve to the left of normal indicates increased fragility, and a curve to the right decreased fragility.



Diagnosis of anemia

- Sucrose hemolysis test – sucrose provides a low ionic strength that permits binding of complement to RBCs. In paroxysmal nocturnal hemoglobinuria (PNH), the RBCs are abnormally sensitive to this complement mediated hemolysis. This is used in screening for PNH.
- Acidified serum test (Ham's test) – is the definitive **diagnostic test for PNH**. In acidified serum, complement is activated by the alternate pathway, binds to RBCs, and lyses the abnormal RBCs found in PNH.

Acidified serum test

★ TABLE 40-1

Schematic Outline of Acidified Serum Test Procedure and Expected Results in PNH

| Tube | 1 | 2 | 3 | 4 | 5 | 6 | 7 |
|-------------------------|------|------|------|------|------|------|------|
| Patient serum (mL) | 0.5 | 0.5 | | | | 0.5 | 0.5 |
| Control serum (mL) | | | 0.5* | 0.5 | 0.5 | | |
| 0.2 N HCl (mL) | | 0.05 | 0.05 | | 0.05 | | 0.05 |
| Patient RBC (mL) | 0.05 | 0.05 | 0.05 | 0.05 | 0.05 | | |
| Control RBC (mL) | | | | | | 0.05 | 0.05 |
| Results observed in PNH | 0 | + | 0 | 0 | + | 0 | 0 |

PNH is suggested if hemolysis occurs in tubes #2 and #5. No hemolysis should be seen in tube #3. Key: * = Serum is heat inactivated; 0 = no hemolysis; + = hemolysis.



Diagnosis of anemia

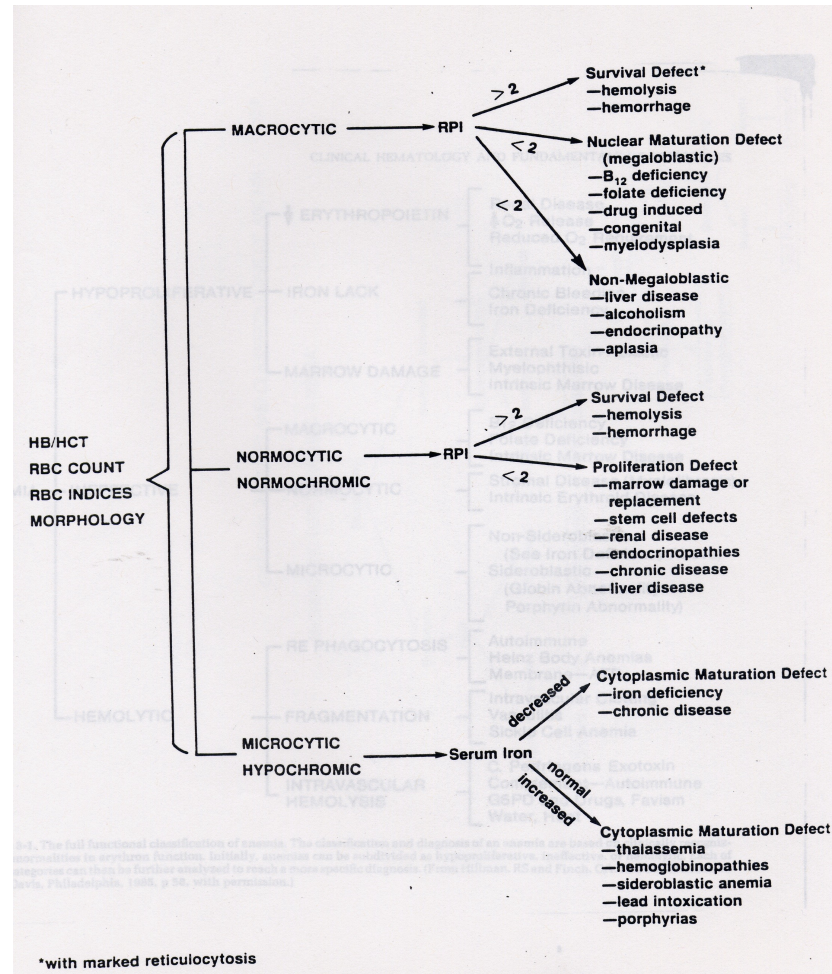
- Evaluation of RBC enzymes and metabolic pathways – enzyme deficiencies in carbohydrate metabolic pathways are usually associated with a hemolytic anemia.
- Evaluation of erythropoietin levels – is used to determine if a proper bone marrow response is occurring.
 - Low levels of RBCs could be due to a bone marrow problem or to a lack of erythropoietin production.
- Serum iron, iron binding capacity and % saturation – used to diagnose iron deficiency anemias (more on this later)
- Bone marrow cultures – used to determine the viability of stem cells.



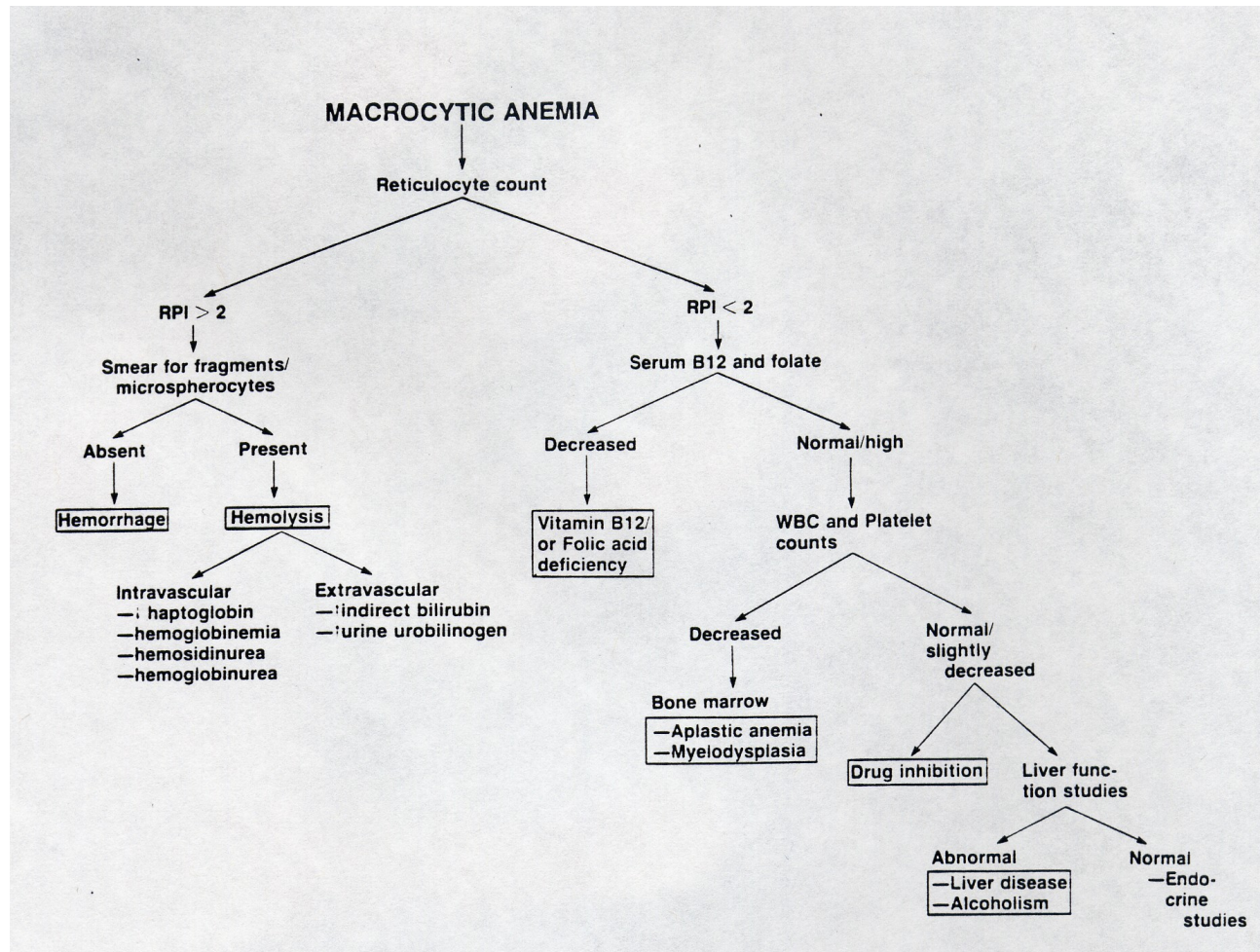
Classification of anemias

- Anemias may be classified morphologically based on the average size of the cells and the hemoglobin concentration into:
 - Macrocytic
 - Normochromic, normocytic
 - Hypochromic, microcytic

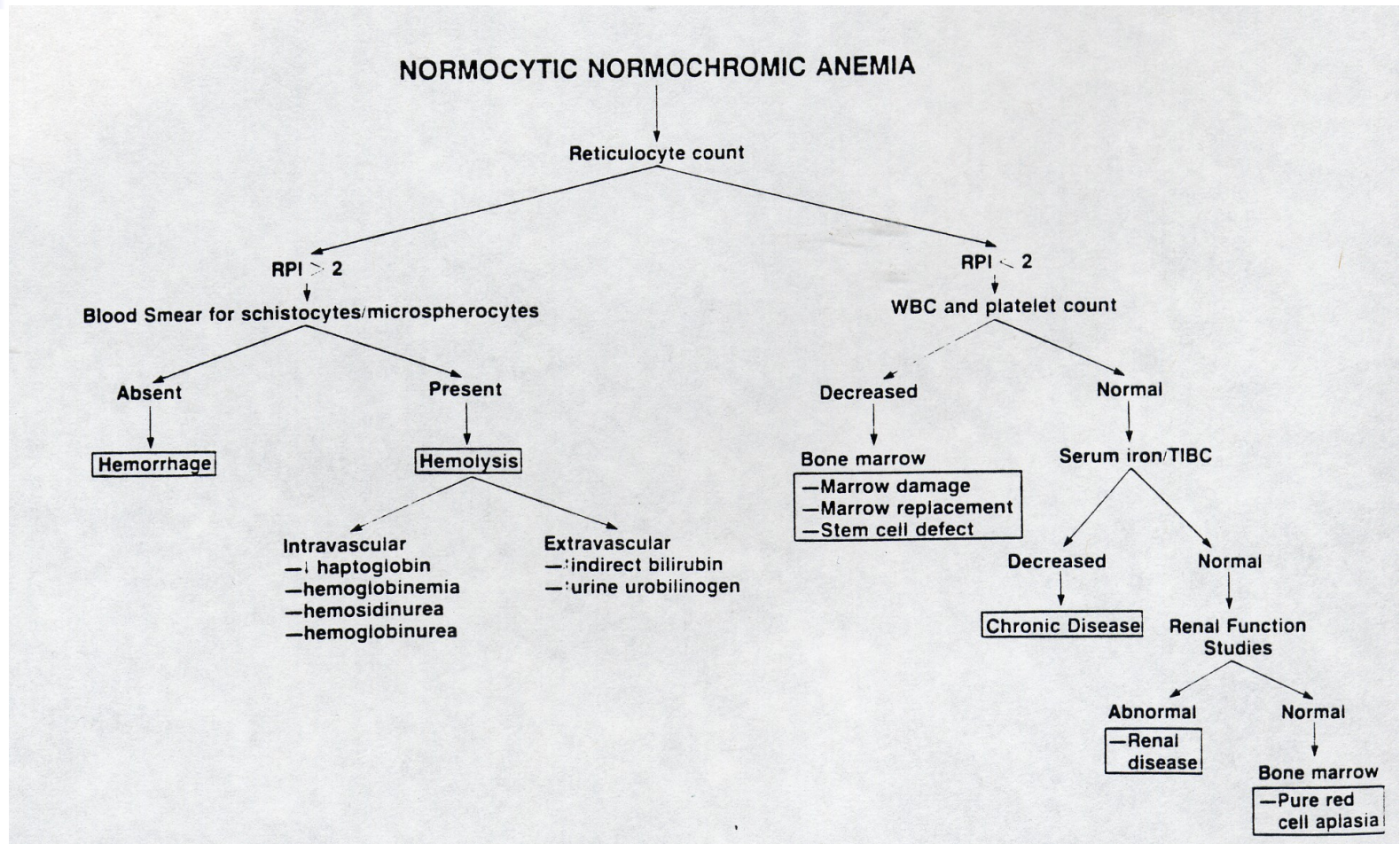
Morphological classification of anemias



Macrocytic anemias



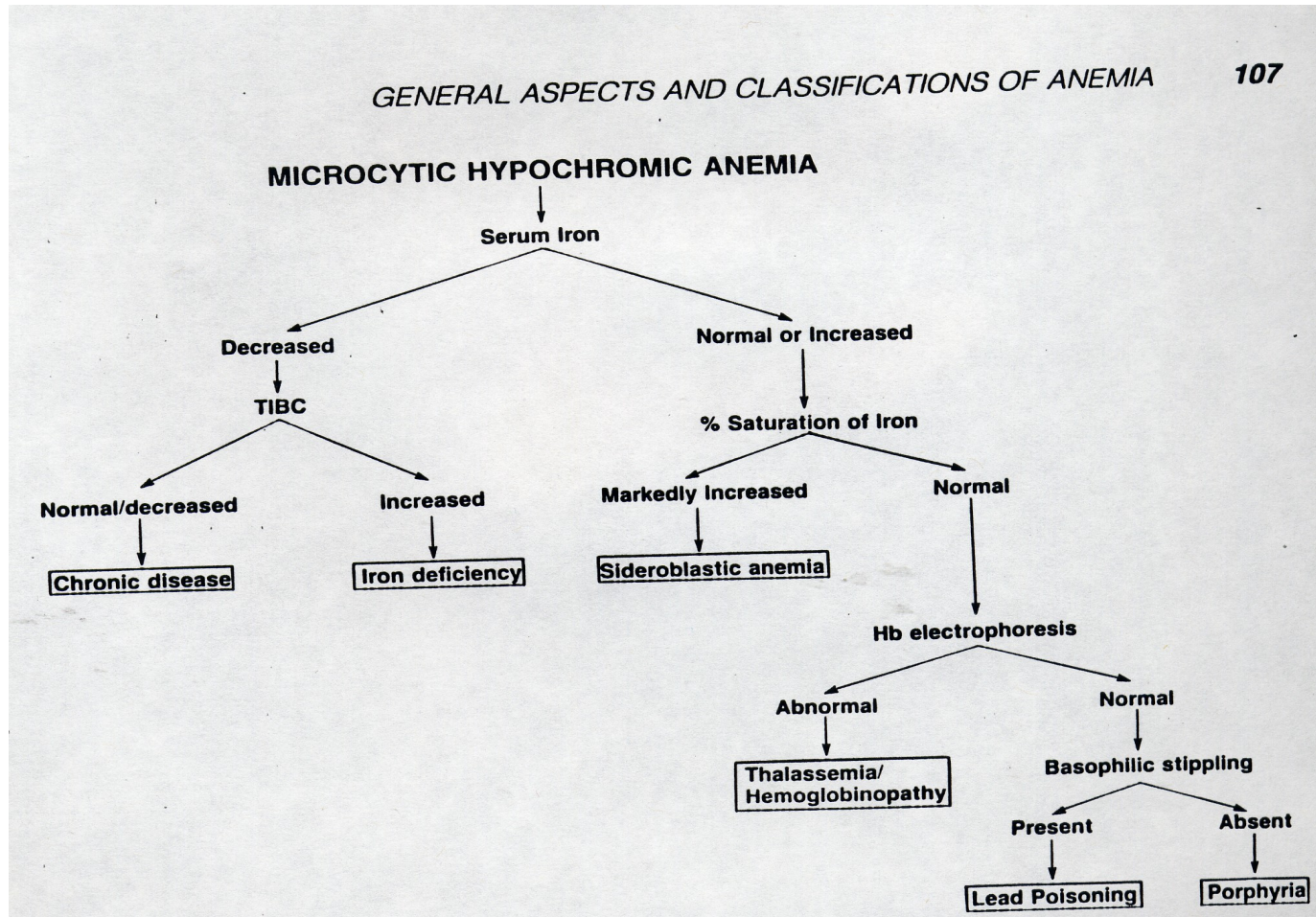
Normochromic, normocytic anemias



Hypochromic, microcytic anemias

GENERAL ASPECTS AND CLASSIFICATIONS OF ANEMIA

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Classification of anemias

- Anemias may also be classified functionally into:
 - Hypoproliferative (when there is a proliferation defect)
 - Ineffective (when there is a maturation defect)
 - Hemolytic (when there is a survival defect)

Functional classification of anemias

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CLINICAL HEMATOLOGY AND FUNDAMENTALS OF HEMOSTASIS

