

# Assessment and Management of Patients with hematologic Disorders

Dr. Wadad K. Mohammad

# Blood Study

## Methods of obtaining blood:

Venipuncture

Finger puncture

Bone marrow aspiration

# Anemia

A low red cell count and a below normal hemoglobin or hematocrit level.

# Pathophysiology of Anemia

## *Bone marrow failure*

**Excessive red cell loss or both.**

**(Erythropoiesis) may occur as a result of a nutritional deficiency, toxic exposure, tumor invasion, or unknown cause.**

Red blood cell may be lost through **hemorrhage** or **hyperhemolysis**. Increase destruction.

# Etiology of Anemia

## **Production defects:**

Nutritional deficiencies - Vitamin B12, folate or iron deficiency.

Inflammation/chronic disease.

Primary marrow disorders-

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**Blood loss.**

**Blood destruction.**

# Etiology of Anemia

## **Blood Loss:**

bleeding, such as from repeated venipuncture in patients undergoing a medical evaluation, blood losses associated with repeated hemodialysis procedures, or excessive blood donations

Bleeding during or after surgical procedures

# Clinical Manifestation

Hb level between 9g/100ml – 11g/100ml

No symptoms other than **tachycardia** on **exertion**

**Exertional dyspnea:**

Is likely to occur below, but not above 7.5mg/100ml

**Weakness** , only below 6g/100ml , **dyspnea** at rest  
,below 3gm/100ml. And cardiac failure.

# Classification of Anemias

Hypoproliferation anemia :The bone marrow is unable to produce adequate numbers of cells, the reticulocyte count is depressed.



# Hypoproliferation anemia

1. Aplastic Anemia
2. Iron Deficiency Anemia
3. Megaloblastic Anemia
4. Vitamin B<sub>12</sub> Deficiency
5. Folic Acid Deficiency

# classification of Anemia<sub>s</sub>

## Hemolytic Anemia

Abnormalities is usually within the red cell itself:

Sickle cell anemia

G-6-pD

# Hemolytic Anemia

1. Inherited Hemolytic Anemia : Spherocytosis , sickle cell anemia, other hemoglobinopathies( Thalassemia)
2. Acquired Hemolytic Anemia