## **BLOOD PHYSIOLOGY**

# TEXTBOOK OF MEDICAL PHYSIOLOGYGUYTON & HALL 12<sup>TH</sup> EDITIONUNIT VI CHAPTERS 32-36

## L2Topic: Hemoglobin, Iron,)

- **1. Essential elements for RBC formation**
- 2. Vitamin B12
- **3. Structure & functions of Hb**
- 4. Iron Metabolism.
- 5. Anemia
- 6. Polycythemia

## Essential elements for RBCs formation and Maturation

#### **Certain elements are essential for RBC formation and maturation:**

- 1. Amino acid: formation of globin in Hb, sever protein deficiency leads to anaemia
- 2. Iron: formation of Hb, iron deficiency results in small cells (microcytic) anaemia

### Essential elements for RBCs formation and Maturation cont

- 3. Vitamins
  - Vit B12 and Folic acid
    - Synthesis of nucleoprotein
    - Deficiency of both causes anemia
  - Vit B6, Riboflavin, nicotinic acid, biotin,
    Vit C, Vit E
- 4. Essential elements
  - Copper, Cobalt, zinc, manganese
- 5. Hormones
  - Androgens, Thyroid, cortisol & growth hormones
  - Deficiencies of any one results in anaemia

## Vitamin B12 & Folic acid

- Important for DNA synthesis and final maturation of RBC
- Dietary source: meat, milk, liver, fat, green vegetables
- Deficiency of VIT B12 & folic acid leads to:
  - Failure of nuclear maturation & division
  - Abnormally large & oval shape RBC
  - Short life span
  - reduced RBC count & Hb content
  - Macrocytic (megaloblastic) anemia

#### Malabsorption of Vit. B12 Pernicious Anemia

- VB12 absorption needs intrinsic factor secreted by parietal cells of stomach
- VB12 + intrinsic factor is absorbed in the terminal lleum
- Causes of deficiencies
  - Inadequate intake
  - Poor absorption due to Intestinal disease
- Give rise to megaloblastic anaemia

## HAEMOGLOBIN

- Hb molecules consist 4 chains each formed of heme & polypeptide chain (globin)
- Heme consist of protoporphyrin ring + iron
- Abnormality in the polypeptide chain abnormal Hb (hemoglobinopathies) e.g thalassemias, sickle cell





# **Functions of Hemoglobin**

- Carriage of O2
  - Hb reversibly bind O<sub>2</sub> to form oxyhemoglobin, affect by pH, temperatre, H<sup>+</sup>
- Carriage of CO2
  - Hb bind CO<sub>2</sub> = carboxyhemaglobin
- Buffer

## **Iron metabolism**

Iron is needed for the synthesis of Hb, myoglobin cytochrome oxsidase, peroxidase & catalase

- Total Iron in the body = 4-5g
  - 65% Haemoglobin
  - 5% other hems
  - 1% bound to transferrin (betaglobulin) in blood
  - 15-30% stored iron in the form of ferritin in the liver, spleen and bone marrow.

# **Iron absorption**

- Iron in food mostly in oxidized form (Ferric)
- Better absorbed in reduced form (Ferrious)
- Iron in stomach is reduced by gastric acid, Vit.
  C.
- Rate of iron absorption depend on the amount of iron stored

## Transport and storage of iron

- Iron is transport in plasma in the form of Transferrin (apotransferrin+iron)
- Iron is stored in two forms
  - Ferritin (apoferritin+iron)
  - Haemosiderin (insoluble complex molecule)
- Daily loss of iron is 0.6 mgm in male & 1.3mgm/day in females

## **Destruction of RBC**

- **RBC** life span in circulation = 120 days
- Metabolic active cells
- Old cell has a fragile cell membrane, cell will rupture as it pass in narrow capillaries (spleen)
- Released Hb is taken up by macrophages in liver, spleen & bone marrow
- Hb is broken into its component:
  - Polypeptide broken to aminoacids to storage
  - Iron degraded to ferrtin and stored
  - Porphyrin ring transfer to bilirubin, secreted by the liver into bile

### ANAEMIAS

#### – Definiation

- Decrease number of RBC
- Decrease Hb

# Symptoms: Tired, Fatigue, short of breath, heart failure

## **Causes of anaemia**

- 1. Blood Loss
  - acute→ accident (RBC return to normal 3-6w)
  - Chronic  $\rightarrow$  microcytic hypochromic anaemi (ulcer, worms)
- 2. Decrease RBC production
  - Nutritional causes
    - Iron  $\rightarrow$  microcytic anaemia
    - VB12 & Folic acid  $\rightarrow$  megaloblastic anaemia
  - Bone marrow destruction by cancer, radiation, drugs  $\rightarrow$  Aplastic anaemia.

## **Causes of anaemia**

- **3.** Haemolytic  $\rightarrow$  excessive destrction
  - Abnormal cells or Hb
    - Spherocytosis
    - sickle cells
  - Incompatible blood transfusion
  - Erythroblastosis fetalis

## **Types of Anemia's**





#### Microcytic anemia Macrocytic anemia

# Polycythemia

#### **Increased number of RBC**

#### Types:

- 1. Primary (polycythemia rubra vera): uncontrolled RBC production
- 2. Secondary to hypoxia: high altitude (physiological), chronic respiratory or cardiac disease

# Objectives

# At the end of this lecture student should be able to:

- 1. Describe essential elements needed for RBC formation.
- 2. Describe the process of VB12 absorption and its malabsorption
- 3. Recognoise haemaglobin structure and its functions.
- 4. Discuss iron metabolism (absorption, storage and transport)

## **Objectives**

- At the end of this lecture student should be able to:
- 5. Describe the fate of old RBC .
- 6. Describe anemia and its causes .
- 7. Recognise causes of polycythemia.