

IRON DEFICIENCY ANEMIA/ ANEMIA OF CHRONIC DISEASE

ANEMIA *Definition*

- Decrease in the number of circulating red blood cells
- Most common hematologic disorder by far

ANEMIA

Causes

- Blood loss
- Decreased production of red blood cells (Marrow failure)
- Increased destruction of red blood cells
 - Hemolysis
- Distinguished by reticulocyte count
 - Decreased in states of decreased production
 - Increased in destruction of red blood cells

ANEMIA

Causes - Decreased Production

- Cytoplasmic production of protein
 - Usually normocytic (MCV 80-100 fl) or microcytic (MCV < 80 fl)
- Nuclear division/maturation
 - Usually macrocytic (MCV > 100 fl)

ANEMIA

Causes - Cytoplasmic Protein Production

- Decreased hemoglobin synthesis
 - Disorders of globin synthesis
 - Disorders of heme synthesis
- Heme synthesis
 - Decreased Iron
 - Iron not in utilizable form
 - Decreased heme synthesis

IRON DEFICIENCY ANEMIA

Prevalence

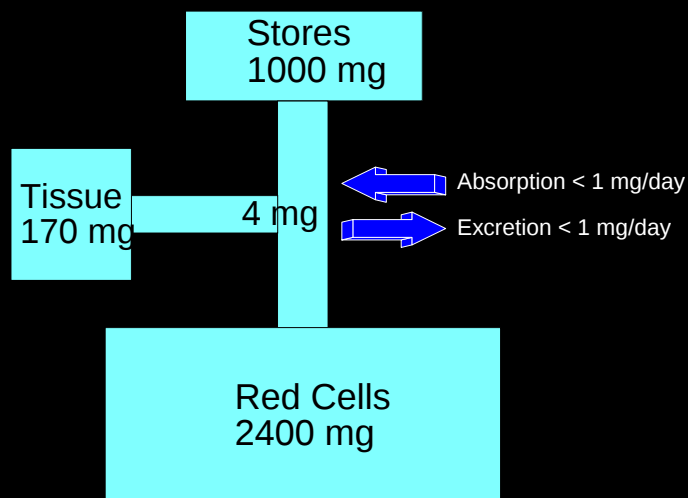
Country	Men (%)	Women (%)	Pregnant Women (%)
S. India	6	35	56
N. India		64	80
Latin America	4	17	38
Israel	14	29	47
Poland			22
Sweden		7	
USA	1	13	

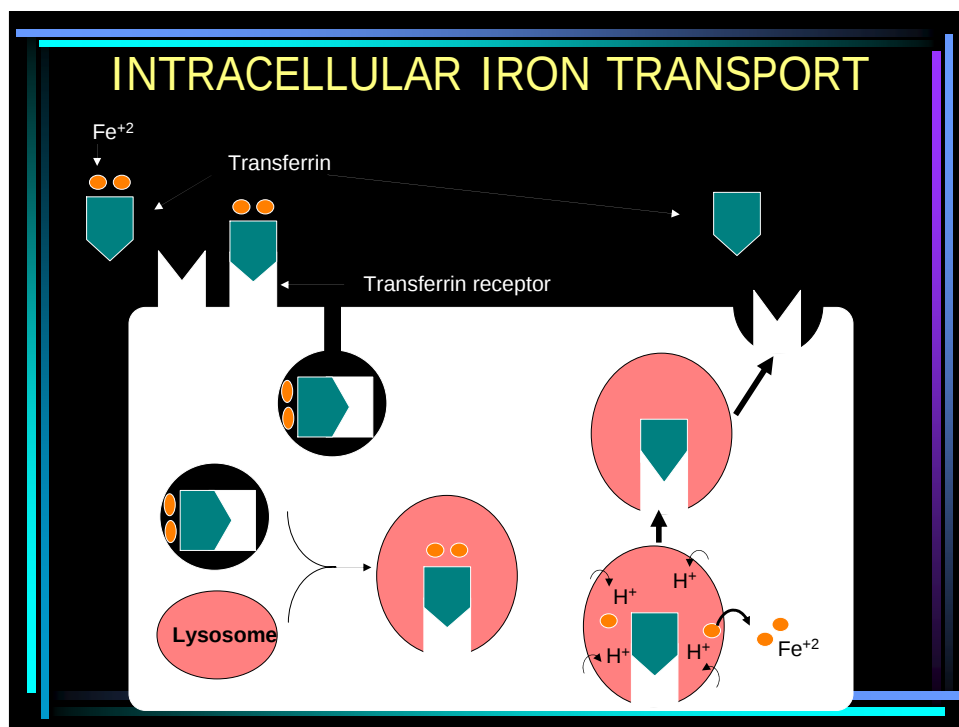
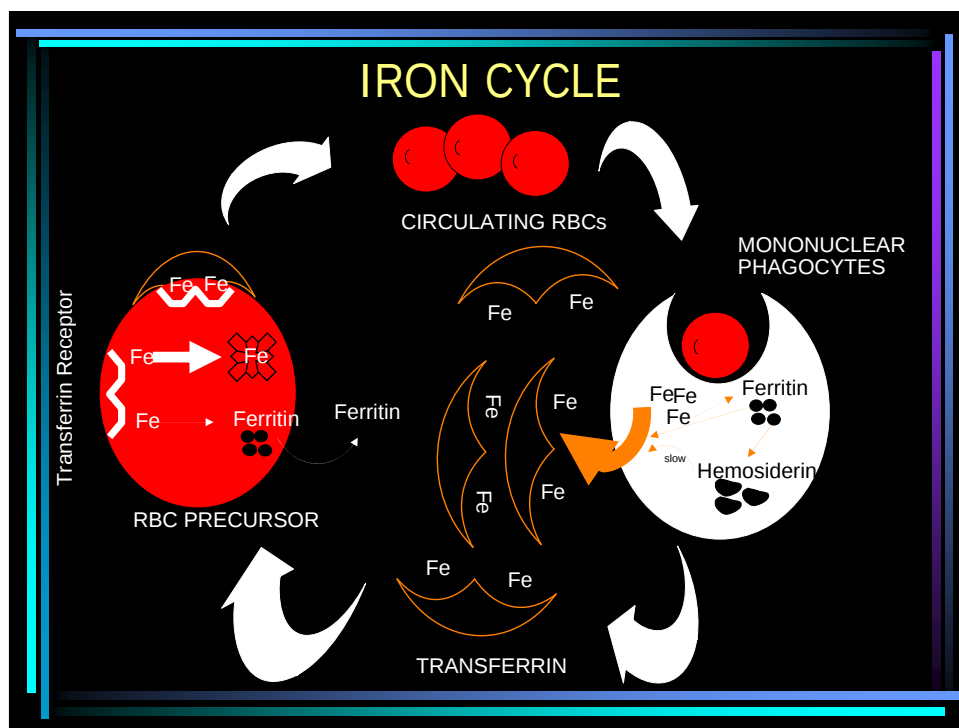
IRON

- Functions as electron transporter; vital for life
- Must be in ferrous (Fe^{+2}) state for activity
- In anaerobic conditions, easy to maintain ferrous state
- Iron readily donates electrons to oxygen, $\rightarrow$ superoxide radicals, H_2O_2 , $\text{OH}\cdot$ radicals
- Ferric (Fe^{+3}) ions cannot transport electrons or O_2
- Organisms able to limit exposure to iron had major survival advantage

IRON

Body Compartments - 75 kg man



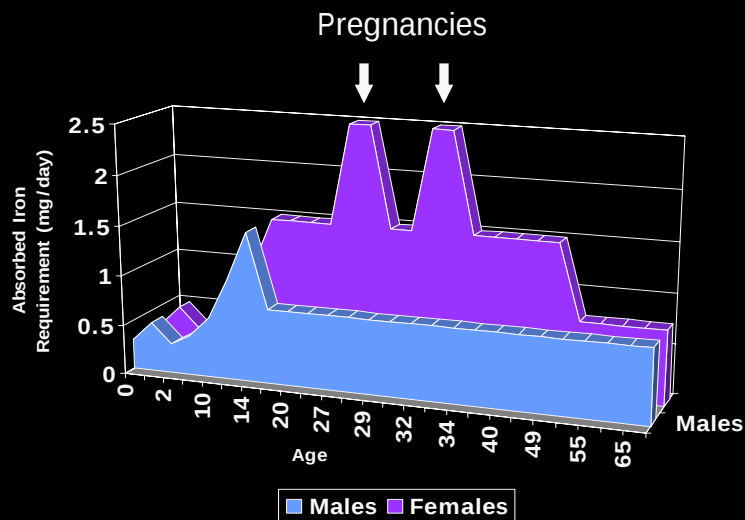


IRON

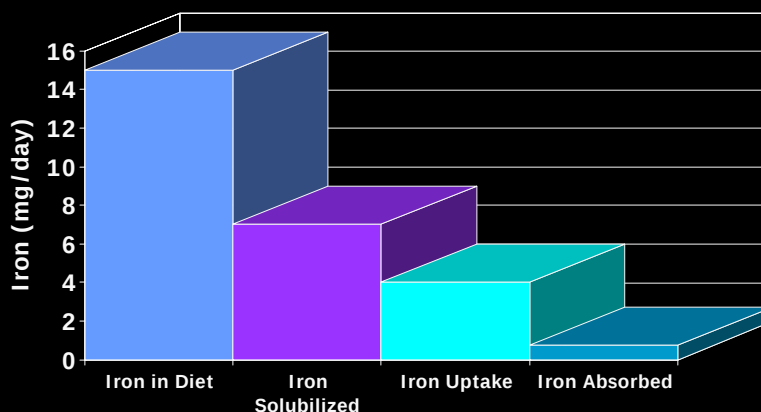
Causes of Iron Deficiency

- Blood Loss
 - Gastrointestinal Tract
 - Menstrual Blood Loss
 - Urinary Blood Loss (Rare)
 - Blood in Sputum (Rarer)
- Increased Iron Utilization
 - Pregnancy
 - Infancy
 - Adolescence
 - Polycythemia Vera
- Malabsorption
 - Tropical Sprue
 - Gastrectomy
 - Chronic atrophic gastritis
- Dietary inadequacy (almost never sole cause)
- Combinations of above

DAILY IRON REQUIREMENTS



IRON ABSORPTION



GI ABSORPTION OF IRON

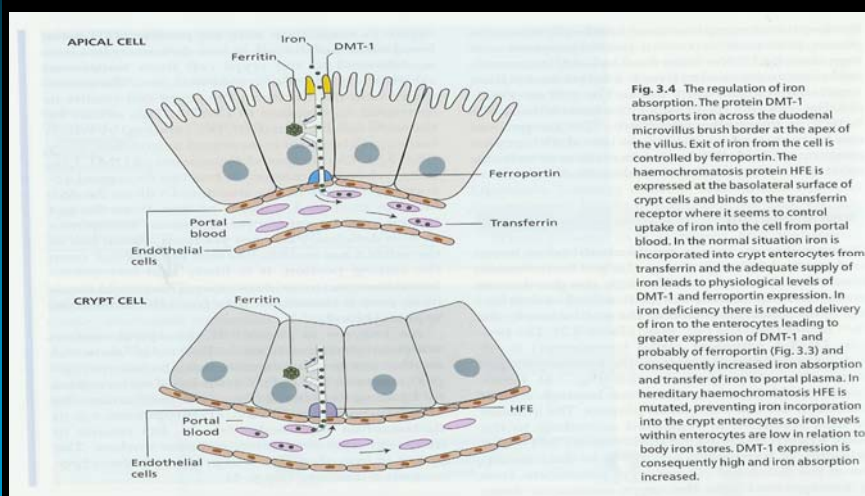


Fig. 3.4 The regulation of iron absorption. The protein DMT-1 transports iron across the duodenal microvillus brush border at the apex of the villus. Exit of iron from the cell is controlled by ferroportin. The haemochromatosis protein HFE is expressed at the basolateral surface of crypt cells and binds to the transferrin receptor where it seems to control uptake of iron into the cell from portal blood. In the normal situation iron is incorporated into crypt enterocytes from transferrin and the adequate supply of iron leads to physiological levels of DMT-1 and ferroportin expression. In iron deficiency there is reduced delivery of iron to the enterocytes leading to greater expression of DMT-1 and probably of ferroportin (Fig. 3.3) and consequently increased iron absorption and transfer of iron to portal plasma. In hereditary haemochromatosis HFE is mutated, preventing iron incorporation into the crypt enterocytes so iron levels within enterocytes are low in relation to body iron stores. DMT-1 expression is consequently high and iron absorption increased.

FERRITIN/TRANSFERRIN REGULATION

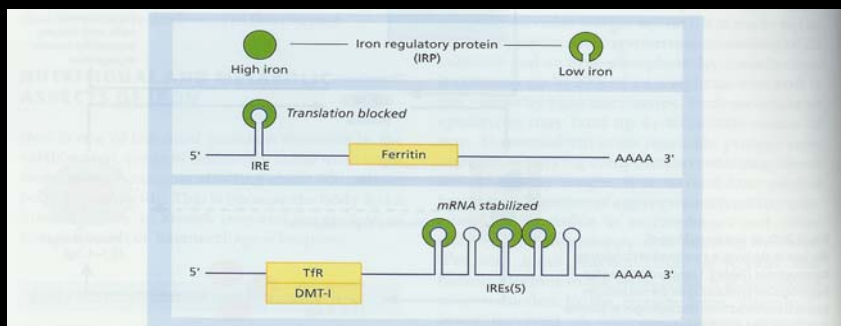
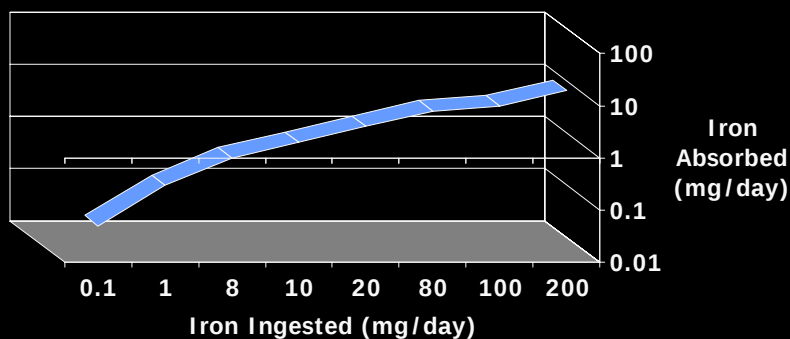


Fig. 3.3 Regulation of transferrin receptor (TfR), DMT-1 (divalent metal transporter), ferroportin and ferritin expression by iron regulatory protein (IRP) sensing of intracellular iron levels. IRPs (●) are able to bind to stem-loop structures called iron response elements (IREs) (●) in transferrin receptor or ferritin messenger RNAs (mRNAs). IRP binding to the IRE within the 3' untranslated region of the former leads to stabilization of mRNA levels and

increased protein synthesis whereas IRP binding to the IRE within the 5' untranslated region of ferritin mRNA reduces translation. IRPs can exist in two states — at times of high iron levels the IRP binds iron and exhibits a reduced affinity for the IREs whereas when iron levels are low the binding of IRPs to IREs is increased. In this way synthesis of TfR, DMT-1 and ferritin is coordinated to physiological requirements.

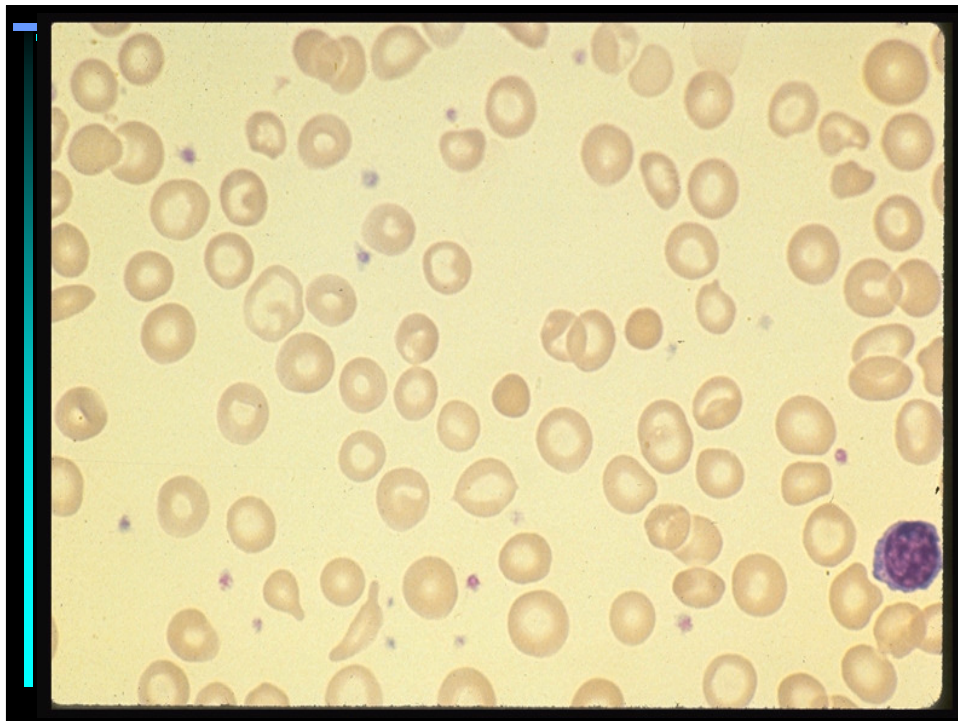
IRON ABSORPTION

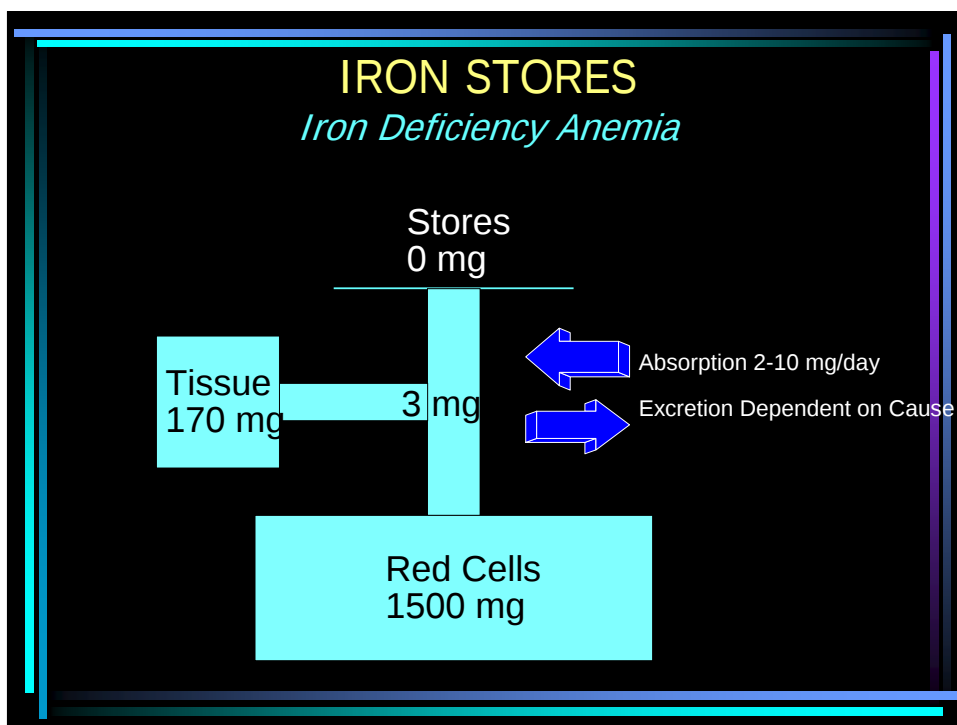
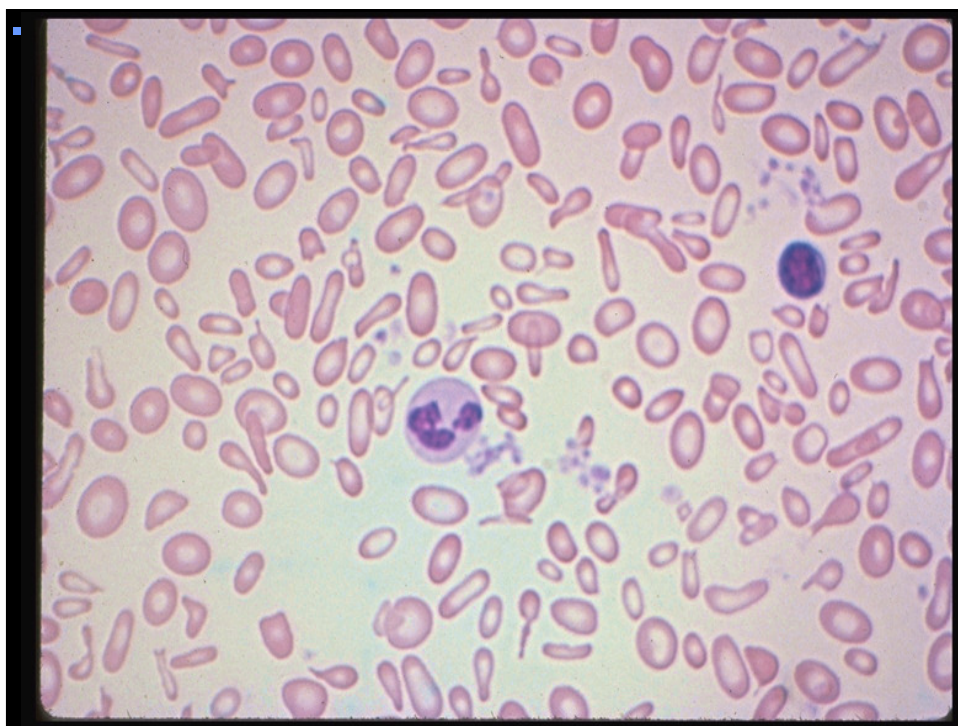


IRON DEFICIENCY ANEMIA

Progression of Findings

- Stainable Iron, Bone Marrow Aspirate
- Serum Ferritin - Low in Iron Deficiency
- Desaturation of transferrin
- Serum Iron drops
- Transferrin (Iron Binding Capacity) Increases
- Blood Smear - Microcytic, Hypochromic; Aniso- & Poikilocytosis
- Anemia





IRON DEFICIENCY

Symptoms

- Fatigue - Sometimes out of proportion to anemia
- Atrophic glossitis
- Pica
- Koilonychia (Nail spooning)
- Esophageal Web

IRON

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IRON REPLACEMENT THERAPY Response

- Usually oral; usually 300-900 mg/day
- Requires acid environment for absorption
- Poorly absorbed

IRON THERAPY Response

- Initial response takes 7-14 days
- Modest reticulocytosis (7-10%)
- Correction of anemia requires 2-3 months
- 6 months of therapy beyond correction of anemia needed to replete stores, assuming no further loss of blood/iron
- Parenteral iron possible, but problematic

ANEMIA OF CHRONIC DISEASE

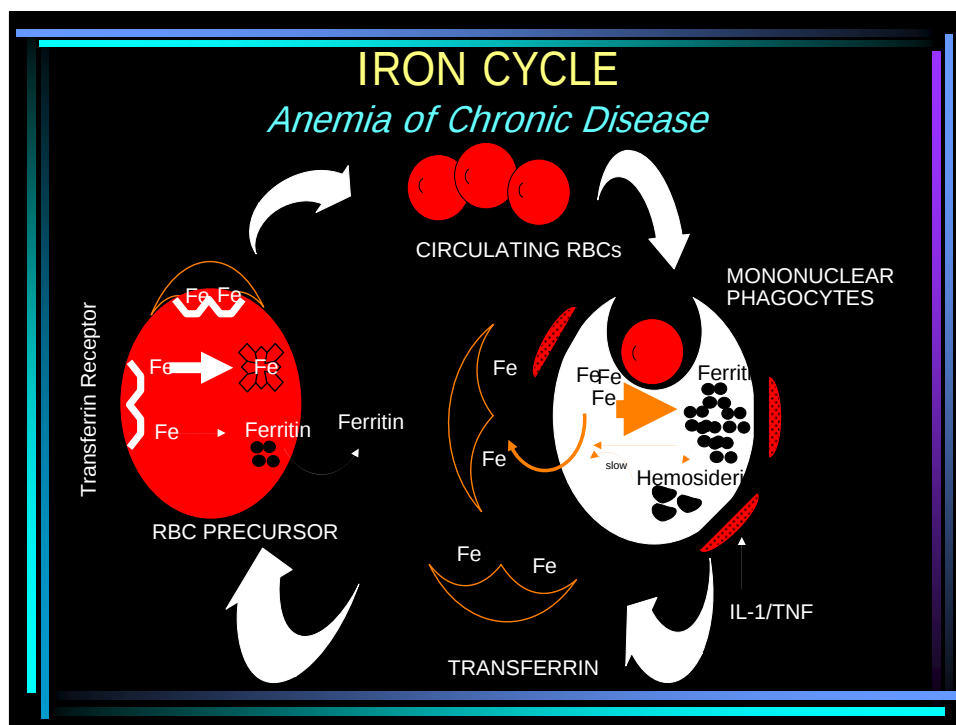
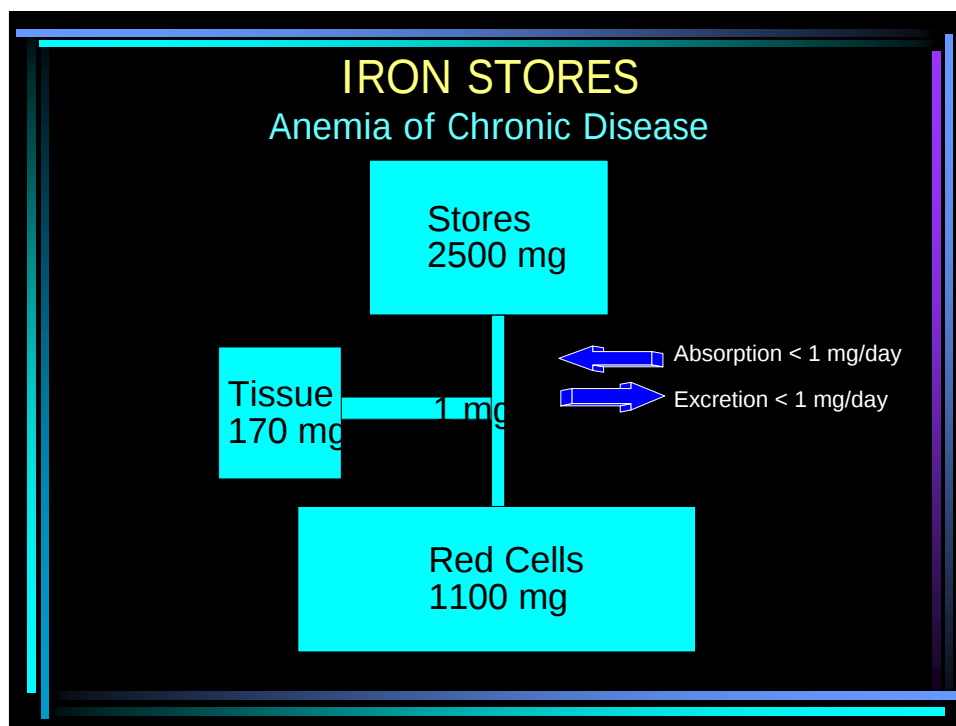
Findings

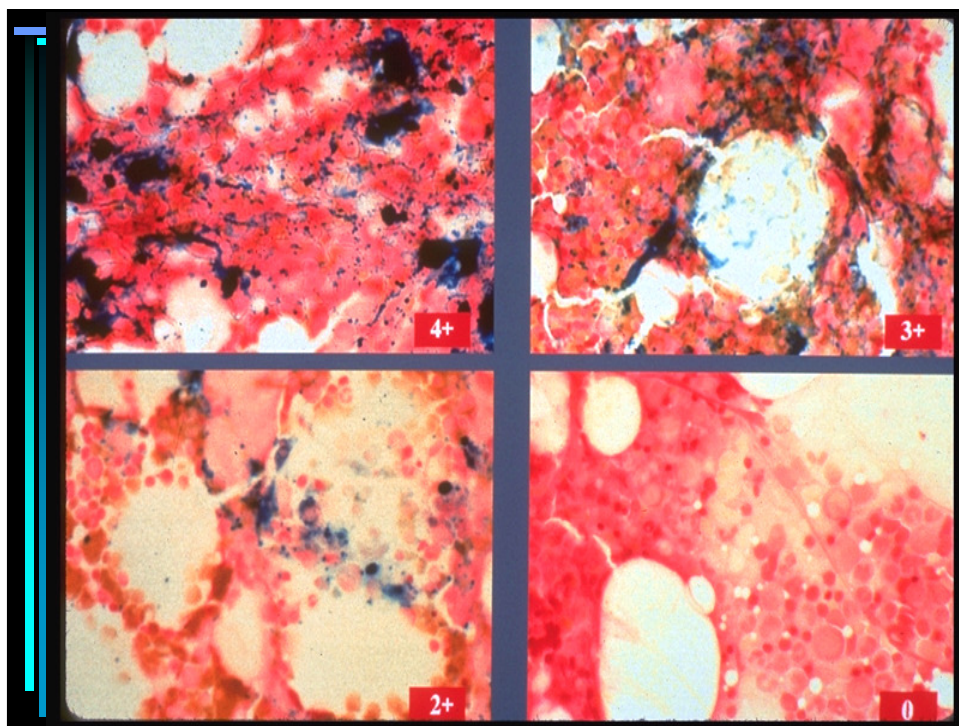
- Mild, non-progressive anemia (Hgb c. 10, Hct c. 30%)
- Other counts normal
- Normochromic/normocytic (30% hypochromic/microcytic)
- Mild aniso- & poikilocytosis
- Somewhat shortened RBC survival
- Normal reticulocyte count (Inappropriately low for degree of anemia)
- Normal bilirubin
- EPO levels increased but blunted for degree of anemia

ANEMIA OF CHRONIC DISEASE

Causes

- Thyroid disease
- Collagen Vascular Disease
 - Rheumatoid Arthritis
 - Systemic Lupus Erythematosus
 - Polymyositis
 - Polyarteritis Nodosa
- Inflammatory Bowel Disease
 - Ulcerative Colitis
 - Crohn's Disease
- Malignancy
- Chronic Infectious Diseases
 - Osteomyelitis
 - Tuberculosis
- Familial Mediterranean Fever





IRON DEFICIENCY <i>versus</i> ACD			
	Serum Iron	Transferrin	Ferritin
Iron Deficiency	↓	↑	↓
ACD	↓	↓	↑

SUMMARY

Iron-Related Anemias

- Most common anemia
- Symptom of pathologic process
- Primary manifestation is hematologic
- Treatment requires:
 - Replacement therapy
 - Correction of underlying cause (if possible)