

Plasma Proteins

GIT Block

1 Lecture

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Overview:

- **Functions and characteristics of plasma proteins**
- **Measurement of plasma proteins and diagnosis of diseases**
- **Electrophoretic patterns of plasma proteins**
- **Acute phase proteins**

Plasma Proteins (pps)

- **Plasma contains >300 different proteins**
- **Many pathological conditions affect level of plasma proteins**
- **Mostly synthesized in the liver**
- **Some are produced in other sites**
- **A normal adult contains ~70 g/L of pps**

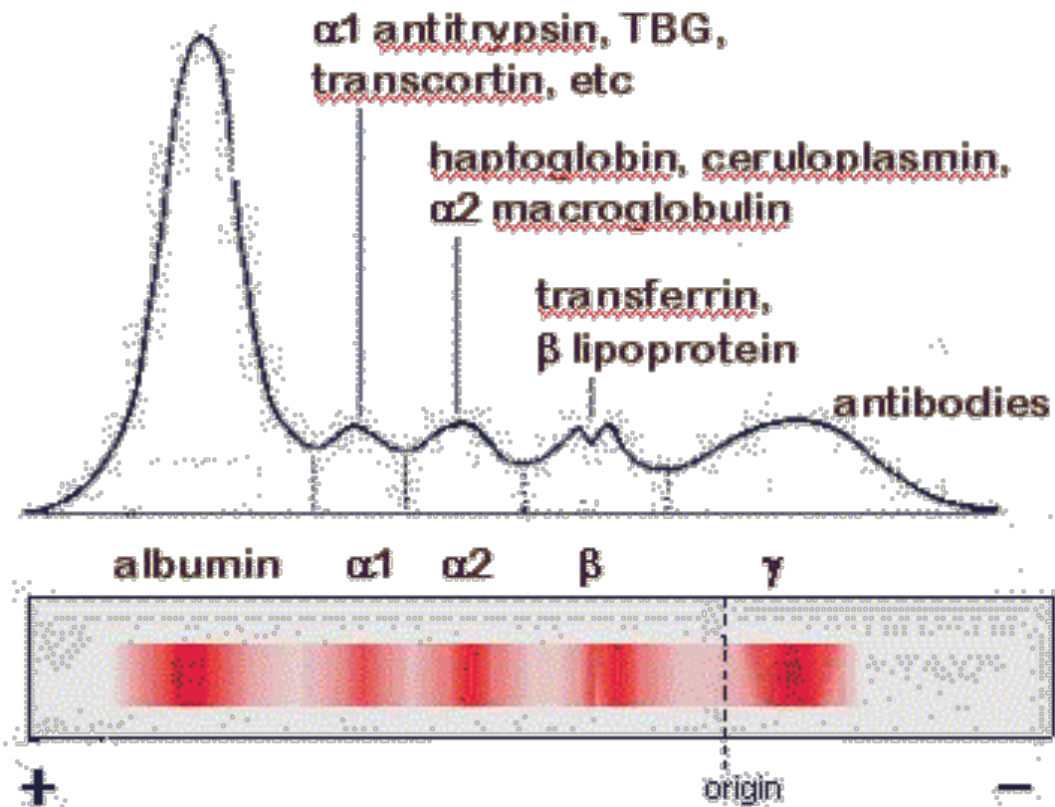
Functions of plasma proteins

- **Transport (Albumin, prealbumin, globulins)**
- **Maintain plasma oncotic pressure (Albumin)**
- **Defense (Immunoglobulins and complement)**
- **Clotting and fibrinolysis (Thrombin and plasmin)**

Measurement of Plasma Proteins

- A) Quantitative measurement of a specific protein:
Chemical or immunological reactions
- B) Semiquantitative measurement by electrophoresis:
- Proteins are separated by their electrical charge in electrophoresis
 - **Five separate bands of proteins** are observed
 - These bands change in disease

Normal Pattern of Plasma Protein Electrophoresis



Types of Plasma Proteins

- Prealbumin
- Albumin
- α_1 -Globulins:
 - α_1 -Antitrypsin, α -fetoprotein
- α_2 -Globulins:
 - Ceruloplasmin, haptoglobin
- β -Globulins:
 - CRP, transferrin, β_2 -microglobulin
- γ - Globulins

Prealbumin (Transthyretin)

- **A transport protein for:**
 - **Thyroid hormones**
 - **Retinol (vitamin A)**
- **Migrates faster than albumin in electrophoresis**
- **Separated by immunoelectrophoresis**
- **Lower levels found in:**
 - **liver disease, nephrotic syndrome, acute phase inflammatory response, malnutrition**
- **Short half-life (2 days)**

Albumin

- **Most abundant plasma protein (~40 g/L) in normal adult**
- **Synthesized in the liver as preproalbumin and secreted as albumin**
- **Half-life in plasma: 20 days**
- **Decreases rapidly in injury, infection and surgery**

Functions

- Maintains oncotic pressure:
 - The osmotic pressure exerted by plasma proteins that pulls water into the circulatory system
 - Maintains fluid distribution in and outside cells and plasma volume
- 80% of plasma oncotic pressure is maintained by albumin

Functions

- **A non-specific carrier of**
 - hormones, calcium, free fatty acids, drugs, etc.
- **Tissue cells can take up albumin by pinocytosis where it is hydrolyzed to amino acids**
- **Useful in treatment of liver diseases, hemorrhage, shock and burns**

Hypoalbuminemia

- **Causes**

- Decreased albumin synthesis (liver cirrhosis, malnutrition)
- Increased losses of albumin
 - Increased catabolism in infections
 - Excessive excretion by the kidneys (nephrotic syndrome)
 - Excessive loss in bowel (bleeding)
 - Severe burns (plasma loss in the absence of skin barrier)

Hypoalbuminemia

Effects

- Edema due to low oncotic pressure
 - Albumin level drops in liver disease causing low oncotic pressure
 - Fluid moves into the interstitial spaces causing edema
- Reduced transport of drugs and other substances in plasma
- Reduced protein-bound calcium
 - Total plasma calcium level drops
 - Ionized calcium level may remain normal

Hyperalbuminemia

- No clinical conditions are known that cause the liver to produce large amounts of albumin
- The only cause of hyperalbuminemia is dehydration

α_1 -Antitrypsin

- **Synthesized by the liver and macrophages**
- **An acute-phase protein that inhibits proteases**
- **Proteases are produced endogenously and from leukocytes and bacteria**
 - Digestive enzymes (trypsin, chymotrypsin)
 - Other proteases (elastase, thrombin)
- **Infection leads to protease release from bacteria and from leukocytes**

Types of α_1 -Antitrypsin

- Over 30 types are known
- The most common is M type
- Genetic deficiency of α_1 -Antitrypsin
 - Synthesis of the defective α_1 -Antitrypsin occurs in the liver but it cannot secrete the protein
 - α_1 -Antitrypsin accumulates in hepatocytes and is deficient in plasma

Clinical Consequences of α_1 -Antitrypsin Deficiency

- Neonatal jaundice with evidence of cholestasis
- Childhood liver cirrhosis
- Pulmonary emphysema in young adults

Laboratory Diagnosis

- Lack of α_1 -globulin band in protein electrophoresis
- Quantitative measurement of α_1 -Antitrypsin by:
 - Radial immunodiffusion, isoelectric focusing or nephelometry

α -Fetoprotein (AFP)

- Synthesized in the developing embryo and fetus by the parenchymal cells of the liver
- AFP levels decrease gradually during intra-uterine life and reach adult levels at birth
- Function is unknown but it may protect fetus from immunologic attack by the mother
- No known physiological function in adults

α -Fetoprotein (AFP)

- **Elevated maternal AFP levels are associated with:**
 - **Neural tube defect, anencephaly**
- **Decreased maternal AFP levels are associated with:**
 - **Increased risk of Down's syndrome**
- **AFP is a tumor marker for:**
Hepatoma and testicular cancer

Ceruloplasmin

- **Synthesized by the liver**
- **Contains >90% of serum copper**
- **An oxidoreductase that inactivates ROS causing tissue damage in acute phase response**
- **Important for iron absorption from the intestine**
- **Wilson's disease:**
 - **Due to low plasma levels of ceruloplasmin**
 - **Copper is accumulated in the liver and brain**

Haptoglobin

- **Synthesized by the liver**
- **Binds to free hemoglobin to form complexes that are metabolized in the RES**
- **Limits iron losses by preventing Hb loss from kidneys**
- **Plasma level decreases during hemolysis**

Transferrin

- **A major iron-transport protein in plasma**
 - **30% saturated with iron**
- **Plasma level drops in:**
 - **Malnutrition, liver disease, inflammation, malignancy**
- **Iron deficiency results in increased hepatic synthesis**
- **A negative acute phase protein**

β_2 -Microglobulin

- A component of human leukocyte antigen (HLA)
- Present on the surface of lymphocytes and most nucleated cells
- Filtered by the renal glomeruli due to its small size but most (>99%) is reabsorbed
- Elevated serum levels are found in
 - Overproduction in disease
- May be a tumor marker for:
 - Leukemia, lymphomas, multiple myeloma

C-Reactive Protein (CRP)

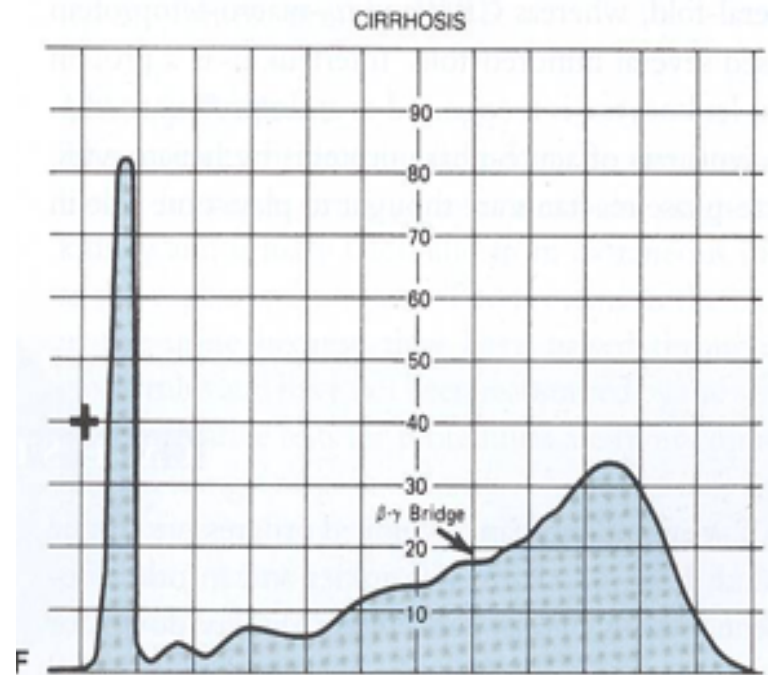
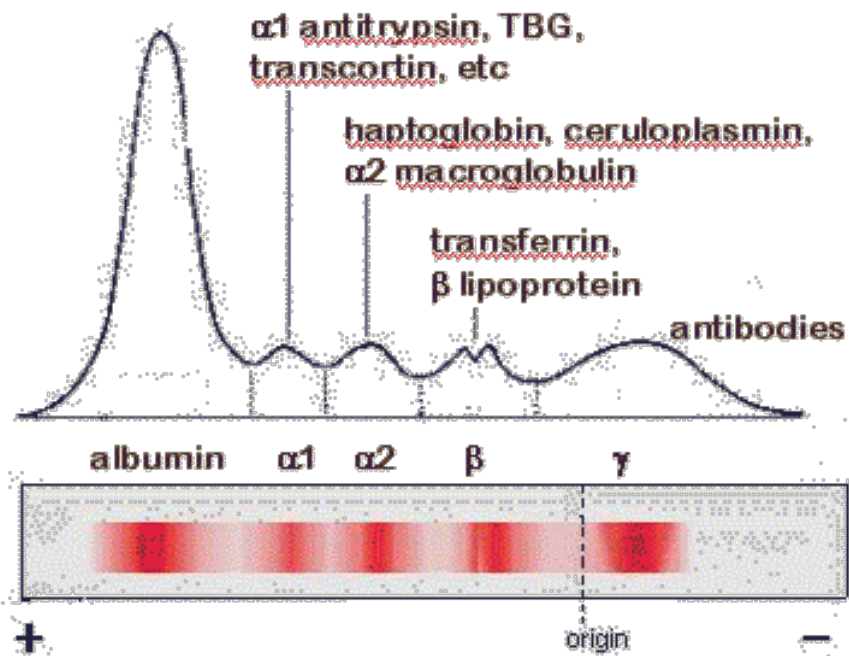
- **An acute-phase protein synthesized by the liver**
- **Important for phagocytosis**
- **High plasma levels are found in many inflammatory conditions such as rheumatoid arthritis**
- **A marker for ischemic heart disease**

Hypergammaglobulinemia

- May result from stimulation of
 - B cells (Polyclonal hypergammaglobulinemia)
 - Monoclonal proliferation (Paraproteinemia)

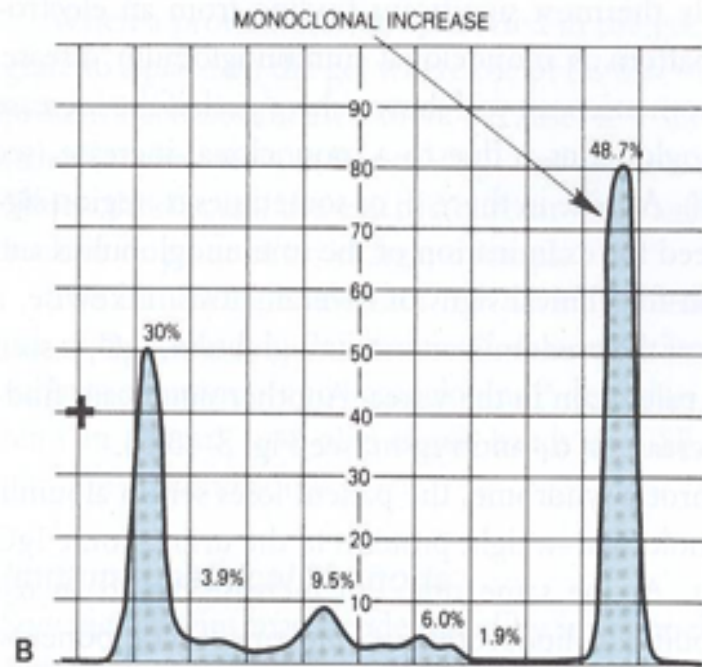
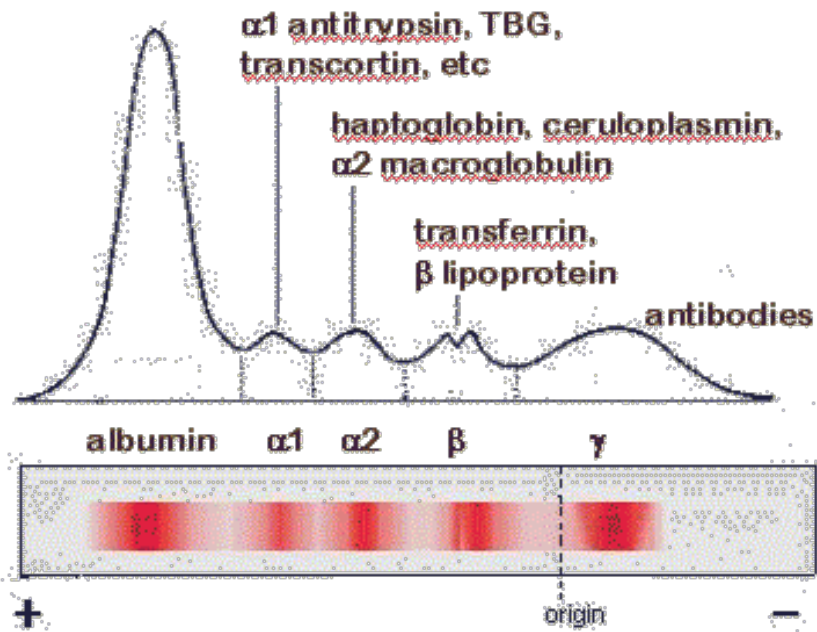
Polyclonal hypergammaglobulinemia:

- Stimulation of many clones of B cells produce a wide range of antibodies
- γ -globulin band appears large in electrophoresis
- Clinical conditions: acute and chronic infections, autoimmune diseases, chronic liver diseases



Monoclonal Hypergammaglobulinemia

- **Proliferation of a single B-cell clone produces a single type of Ig**
- **Appears as a separate dense band (paraprotein or M band) in electrophoresis**
- **Paraproteins are characteristic of malignant B-cell proliferation**
- **Clinical condition: multiple myeloma**



Positive Acute Phase Proteins

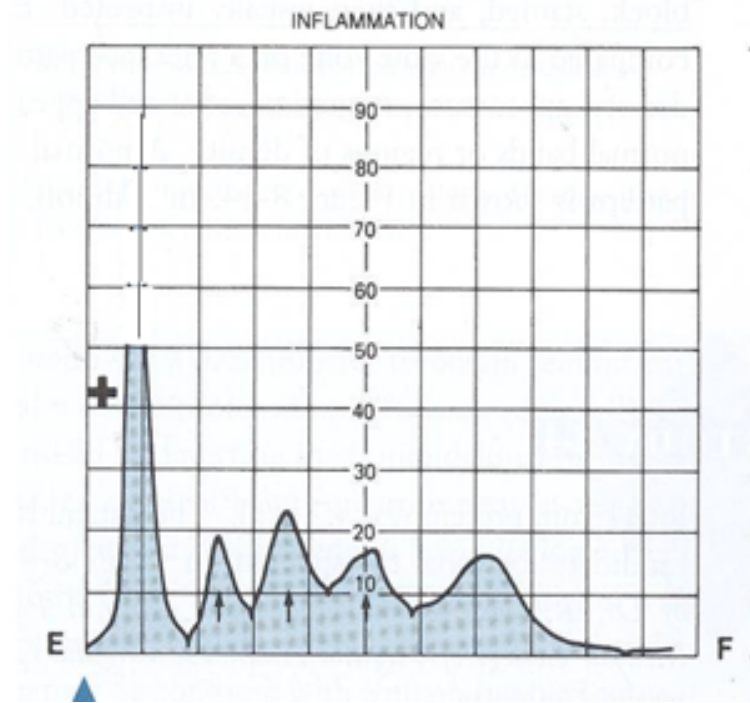
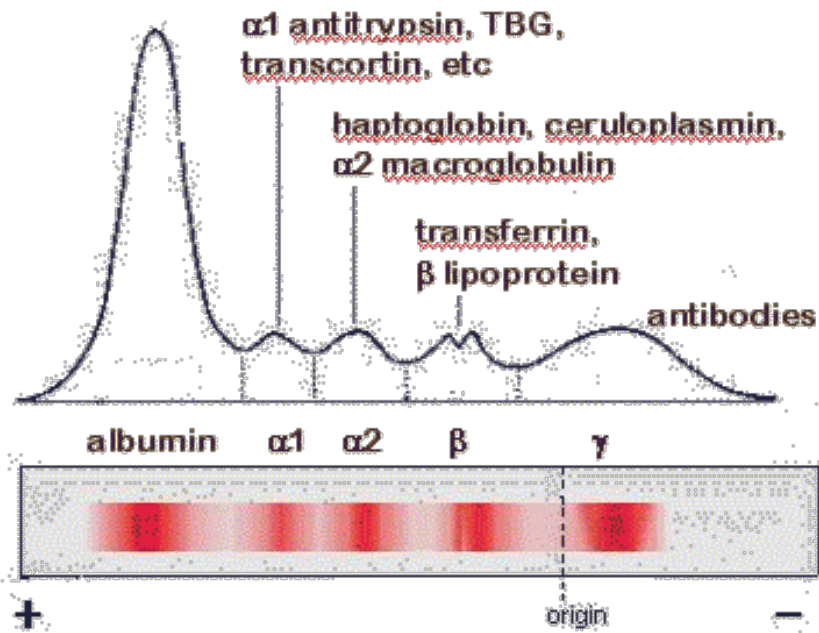
- Plasma protein levels **increase** in:
 - Infection, inflammation , malignancy, trauma, surgery
- These proteins are called acute phase reactants
- Synthesized due to body's response to injury
- Examples: α_1 -Antitypsin, haptoglobin, ceruloplasmin, fibrinogen, c-reactive protein

Positive Acute Phase Proteins

- Mediators cause these proteins to increase after injury
- Mediators: Cytokines (IL-1, IL-6), tumor necrosis factors α and β , interferons, platelet activating factor

Functions:

1. Bind to polysaccharides in bacterial walls
2. Activate complement system
3. Stimulate phagocytosis



Negative Acute Phase Proteins

- **These proteins decrease in inflammation**
 - **Albumin, prealbumin, transferrin**
- **Mediated by inflammatory response via cytokines and hormones**
- **Synthesis of these proteins decrease to save amino acids for positive acute phase proteins**

References

- Lecture Notes in Clinical Biochemistry 7th edition, pages 92-98
- Clinical Biochemistry An Illustrated Colour Text pages 48-51