The Cardiovascular System: Blood

The Functions of Blood

Blood Overview:

Provides a system for *rapid* transport within the body

Nutrients

Hormones

Waste products

Respiratory gases

Cells

Heat

The Functions of Blood

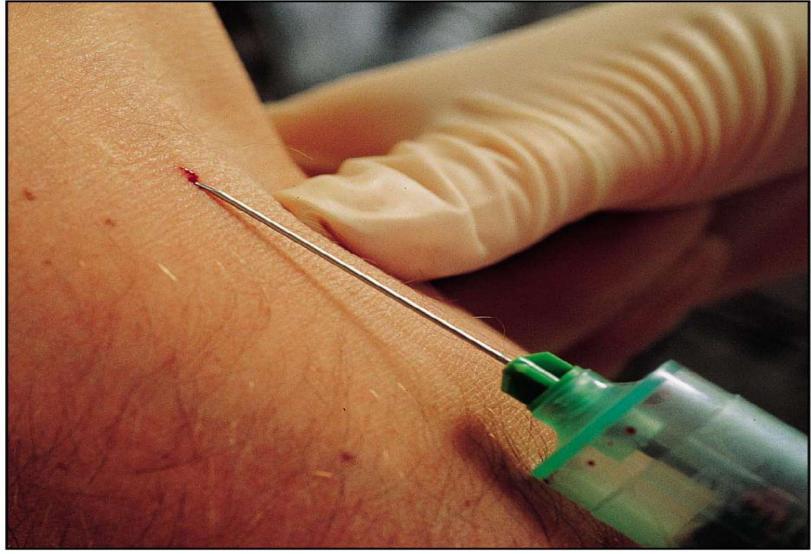
Blood Functions Include:

- Transport of cells and compounds Regulate pH and electrolytes of *interstitial fluids*
- Limit blood loss through damaged vessels
- Defend against pathogens, toxins Absorb, distribute heat as part of temperature regulation

The Composition of Blood

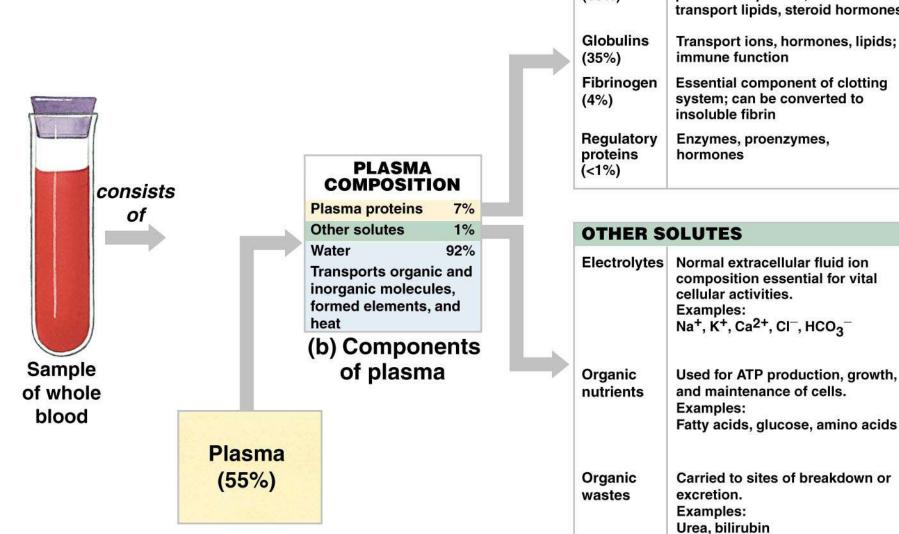
Blood Collection and Analysis Whole blood can be *fractionated* into: Plasma (liquid component) *Formed elements* (cellular components) Red blood cells (RBCs) White blood cells (WBCs) Platelets

Blood Sample



(a) Taking a blood sample

The Composition of Whole Blood



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PLASMA PROTEINS

Albumins

(60%)	pressure of plasma; transport lipids, steroid hormones
Globulins (35%)	Transport ions, hormones, lipids; immune function
Fibrinogen (4%)	Essential component of clotting system; can be converted to insoluble fibrin
Regulatory proteins (<1%)	Enzymes, proenzymes, hormones

Major contributors to osmotic

Plasma Basics

- Makes up about 55% of whole blood *Water* makes up about 92% of plasma
- Has more protein and oxygen than interstitial fluid
- Plasma proteins fall in three classes
 - Albumins
 - Globulins
 - Fibrinogen

Hemopoiesis—The cellular pathways by which the formed elements are produced.

Stem cells (hemocytoblasts)—Cells that divide and mature to produce all three classes of formed elements.

Key Note:

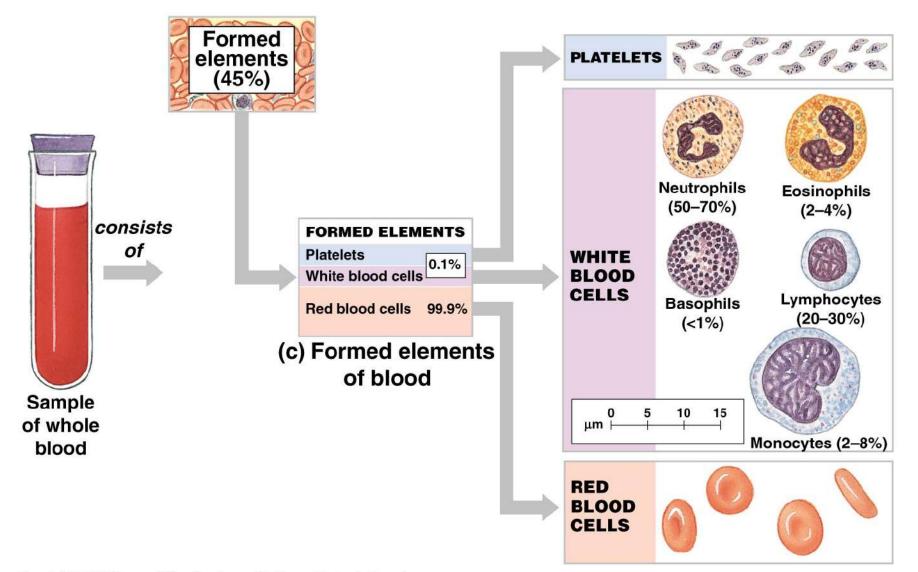
Approximately half the volume of whole blood consists of cells and cell products (the formed elements). Plasma resembles interstitial fluid but contains a unique mixture of proteins not found in other extracellular fluids

Red Blood Cells:

Also called, *erythrocytes* or *RBCs* Make up about 45% of whole blood volume

Make up 99.9% of the formed elements

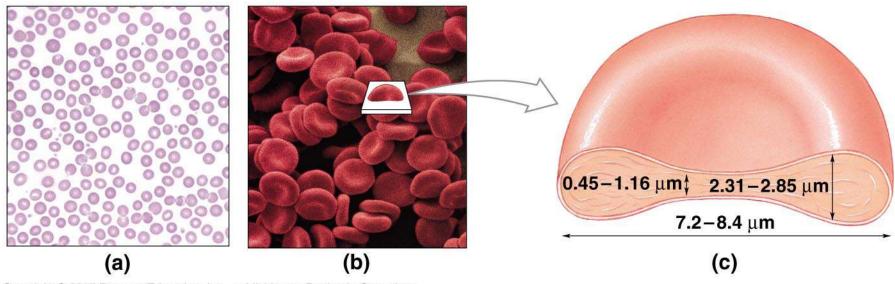
Hematocrit—Percentage of whole blood volume taken up by formed elements (mostly RBCs). In clinical shorthand, it's called, the "crit."



Properties of RBCs Transport oxygen and carbon dioxide in blood stream Have large surface to volume ratio Speeds up gas loading/unloading Lack most organelles

Makes more room for *hemoglobin* Degenerate after about 120 days

The Anatomy of Red Blood Cells



Red Blood Cell Composition:

Hemoglobin makes up 95% of RBC protein Globular protein composed of four *subunits*

Each subunit contains:

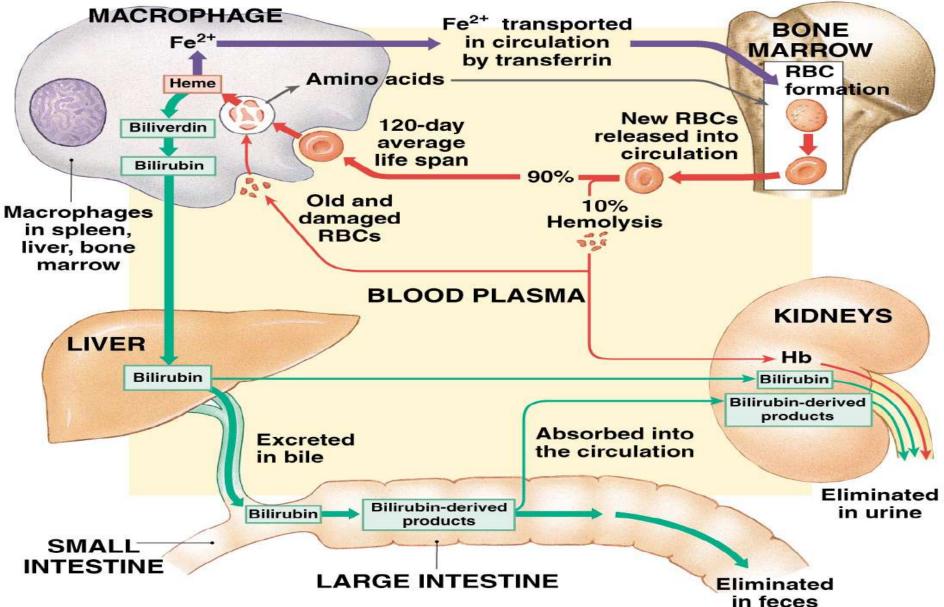
- A globin protein chain
- A molecule of heme

An atom of iron

A binding site for one oxygen molecule

Phagocytes recycle hemoglobin from damaged or dead RBCs

Hemoalobin Recyclina



Erythropoiesis—Process for formation of red blood cells

Occurs mainly in the bone marrow

Stimulated by *erythropoietin* (EPO)

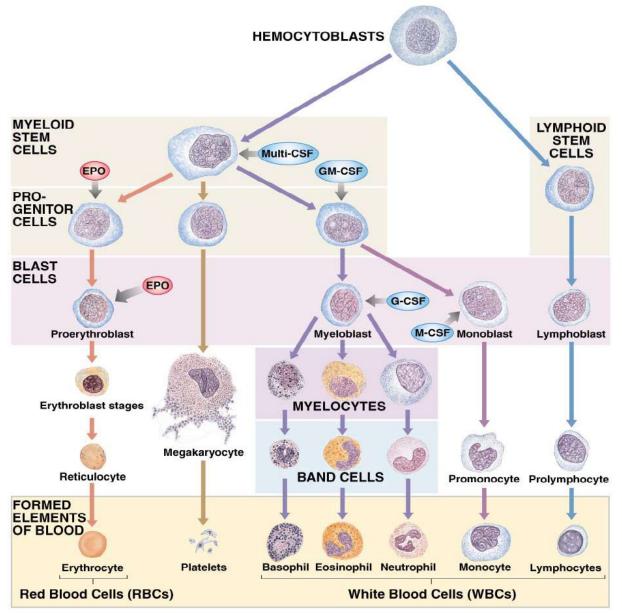
EPO increases when oxygen levels are low

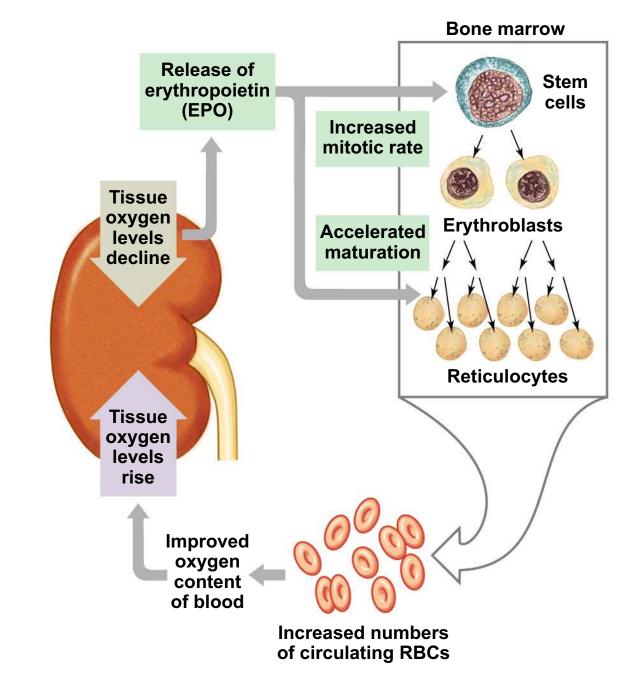
Development stages include:

Erythroblasts

Reticulocytes (after nucleus is expelled)

Differentiation of RBCs, Platelets, and WBCs





Key Note:

Red blood cells (RBCs) are the most numerous cells in the body. They circulate for about four months before being recycled; millions are produced each second. The hemoglobin inside transports oxygen from the lungs to peripheral tissues and carbon dioxide from the tissues to the lungs.

Blood Type

Determined by presence or absence of specific *antigens* (*agglutinogens*) on outside *surface* of RBC

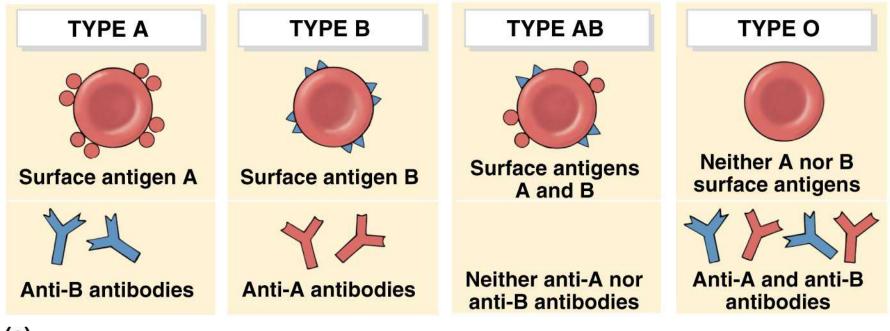
Antigens are called A, B, and Rh

Antibodies (agglutinins) in plasma react with foreign antigens on RBCs

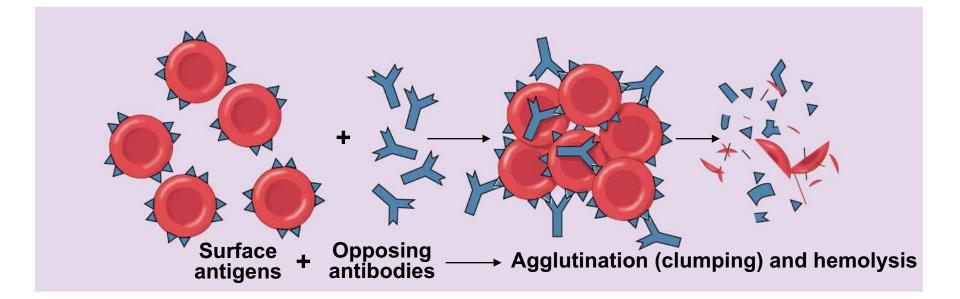
RBCs clump and break open

Anti-Rh antibody made after exposure to Rhpositive blood cells

Blood Types and Cross-Reactions



(a)



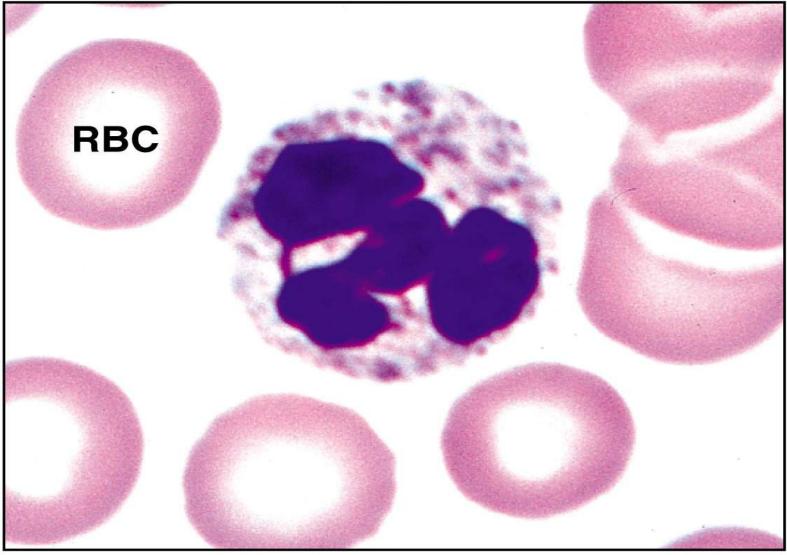
White Blood Cells (WBCs) Also called, *leukocytes* Defend the body against: Pathogens Toxins Abnormal cells Damaged cells

WBC Properties

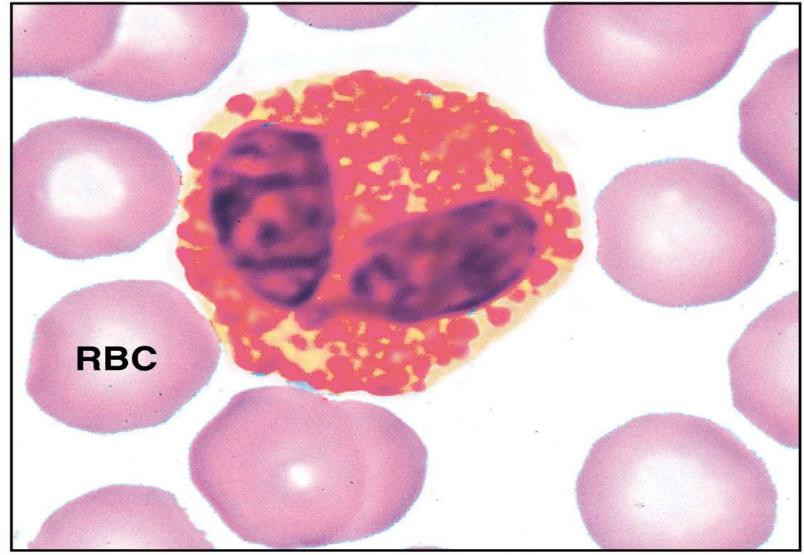
- Perform *diapedesis*—Push between cells to cross blood vessel walls and enter the tissues
- Exhibit *chemotaxis*—Move toward specific chemicals released by bacteria or injured cells
- Consist of two groups:
 - *Granulocytes* (cytoplasmic granules) *Agranulocytes* (no granules)

Three Types of *Granulocytes* Neutrophils 50–70% of circulating WBCs Phagocytic Eosinophils Less common Phagocytic Attracted to foreign proteins Basophils Release histamine Promote inflammation

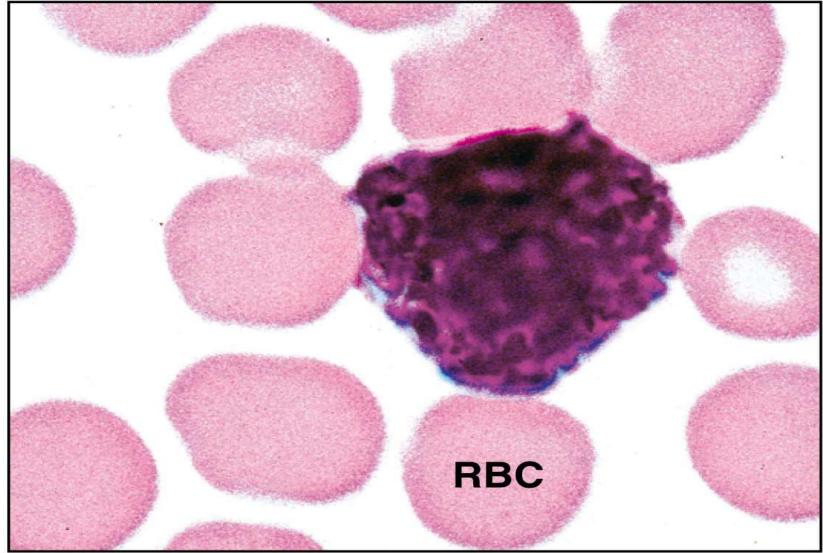
Two Types of Agranulocytes Lymphocytes Found mostly in *lymphatic system* Provide *specific* defenses Attack foreign cells Produce antibodies Destroy abnormal (cancer) cells Monocytes Migrate into tissues Become *macrophages* Live as phagocytic amoeba



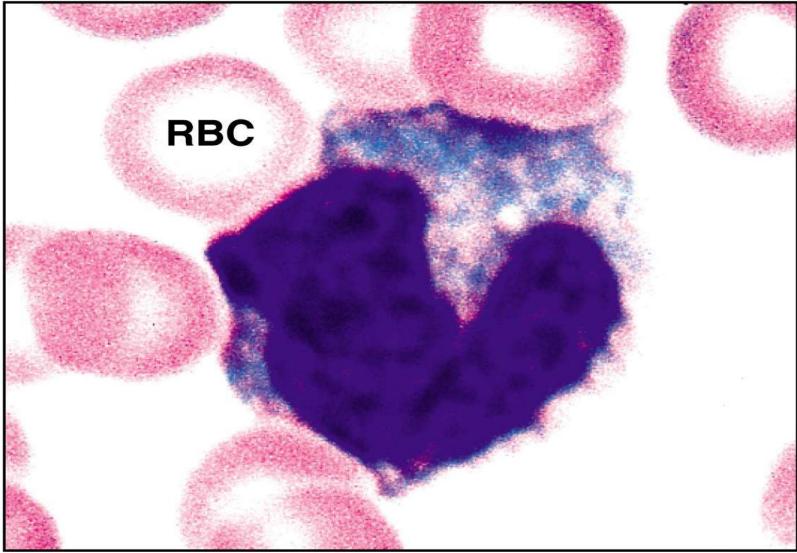
(a) Neutrophil



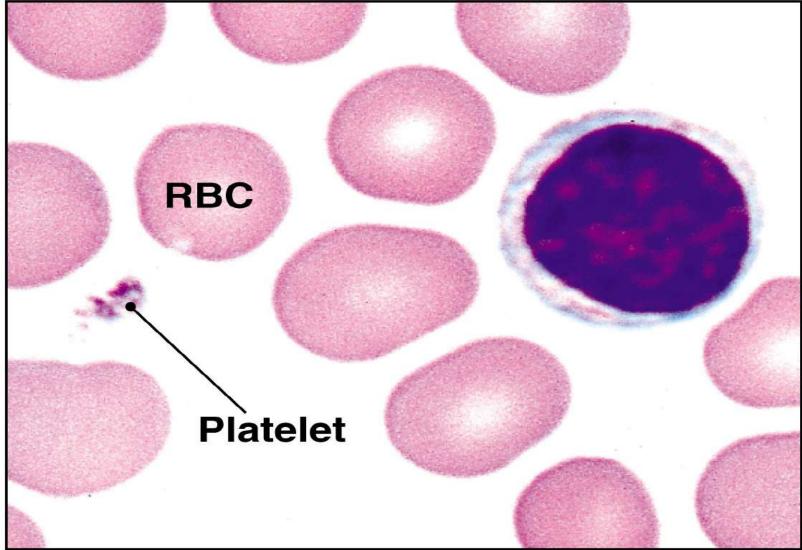
(b) Eosinophil



(c) Basophil



(d) Monocyte



(e) Lymphocyte

Production of WBCs in Bone Marrow *Myeloid stem cells* produce: Granulocytes (three types) Monocytes (future macrophages) Lymphoid stem cells produce lymphocytes Process called, *lymphopoiesis* Lymphocytes enter blood Migrate to lymphoid tissues

Regulation of WBC Maturation

- Colony-stimulating factors (CSFs)— Hormones which regulate certain WBC populations
 - Four CSFs are known
 - CSFs target stem cell lines
 - Several CSFs used with cancer patients with bone marrow suppression

Regulation of WBC Maturation

- Regulation of lymphocyte maturation is poorly understood
 - *Thymosins* (hormones in *thymus* gland) trigger *T cells* to develop

Key Note:

RBCs outnumber WBCs by a 1000 to 1. WBCs defend the body against infection, foreign cells, or toxins, and assist in the repair of damaged tissues. Most numerous are neutrophils, which engulf bacteria, and lymphocytes, which are responsible for the specific immune defenses.



Platelets:

- Produced in the bone marrow
- Released from *megakaryocytes* as cytoplasmic fragments into the blood
- Essential to clotting process

Hemostasis—Processes that stop the loss of blood from a damaged vessel. Largely dependent on platelets and soluble proteins (clotting factors).

Three phases in Hemostasis:

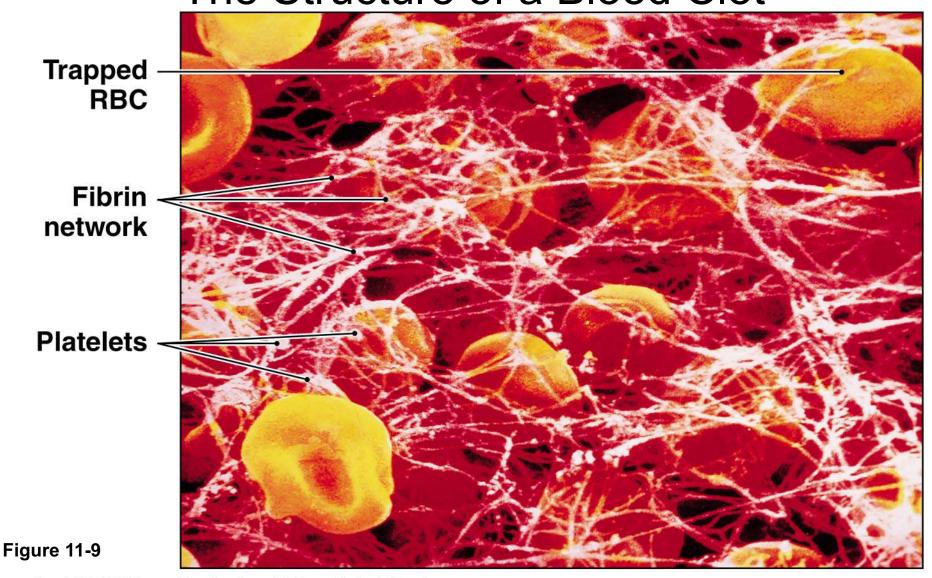
- Vascular phase
 - Local contraction of injured vessel
- Platelet phase
 - Platelets stick to damaged vessel wall
- Coagulation phase
 - Clotting factors in plasma form blood clot

Hemostasis

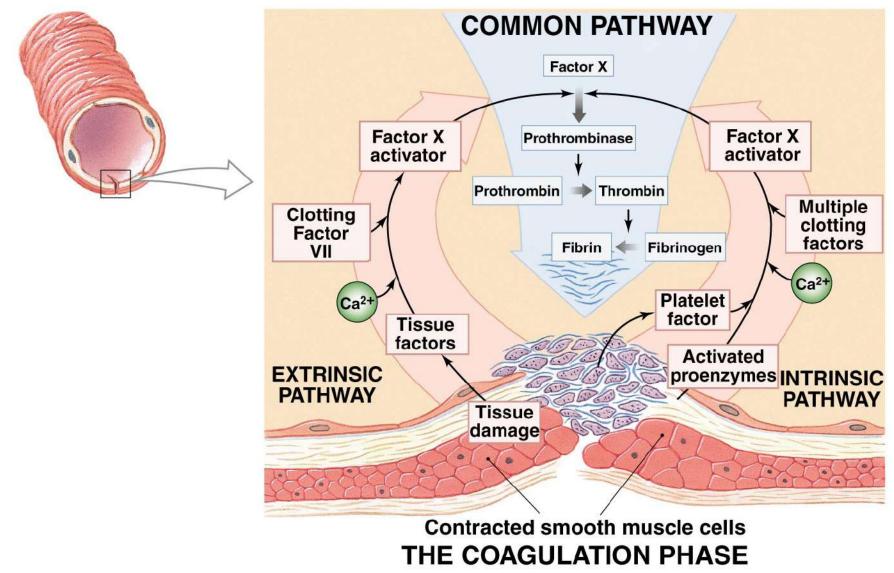
The Clotting Process

- Coagulation pathways require an external trigger
 - Extrinsic pathway
 - Triggered by factors released by injured endothelial cells or peripheral tissues
 - Intrinsic pathway
 - Triggered by factors released by platelets stuck to vessel wall
 - Both pathways lead to *common pathway*
 - *Thrombin* converts soluble *fibrinogen* subunits to an insoluble polymer, *fibrin*

Hemostasis The Structure of a Blood Clot



Coagulation Phase of Hemostasis



Clot Retraction and Removal Clot *retracts* because platelets *contract* Pulls broken vessel closed Clot gradually dissolves Called, *fibrinolysis* Plasmin, an enzyme derived from plasminogen in the plasma, cuts fibrin apart like a molecular scissors

Hemostasis

Key Note:

Platelets coordinate hemostasis (blood clotting). If they are activated by abnormal changes in their surroundings, platelets release clotting factors and other chemicals. Hemostasis is a complex cascade that produces a fibrous patch that is subsequently remodeled and then removed as repair proceeds.