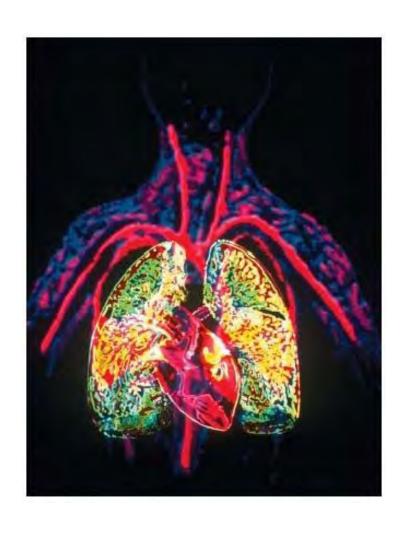
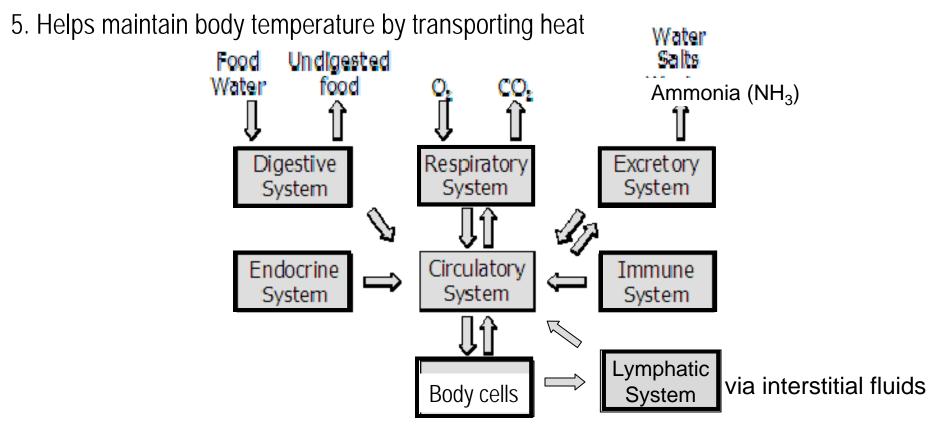
Circulation Ch 42



2

Functions of Circulatory System

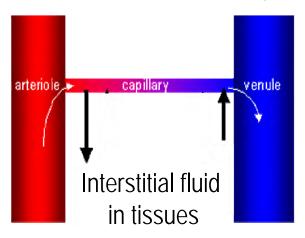
- 1. Allows all cells to exchange materials w/ the environment Gasses (respiratory syst), nutrients (digestive syst), NH₃ water, salt (excretory syst)
- 2. Allows communication btwn different organs/organ systems (hormones (Endocrine syst))
- 3. Transport of cells that fight infection, form clots & repair damaged tissue WBCs and platelets (Immune system, lymphatic system)
- 4. Helps stabilize body fluid pH & [ion] (kidneys, brain, pituitary (excretory syst))



Interstitial fluids (IF)

Fluid between cells

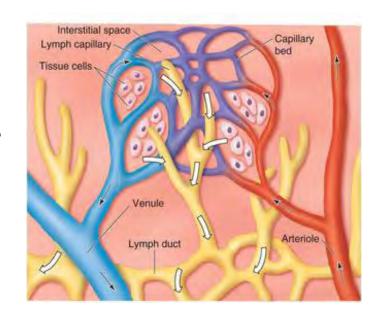
IF Derived from blood plasma



Moves substances btwn cells and circulatory system



- Most IF returns to circ syst directly into capillaries
- Some IF returns to circ syst indirectly via lymph vessels which connect to circ syst



Circulation in Complex Animals

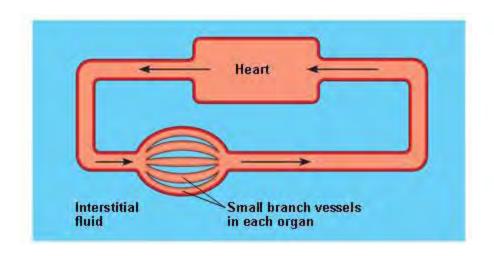
All complex animals have a circulatory system w/ 3 components:

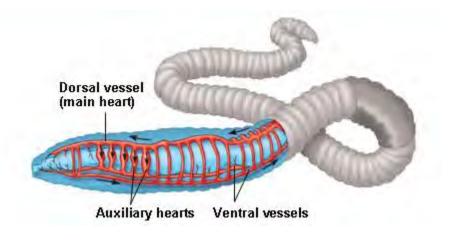
- 1. Fluid
- 2. Tubes
- 3. Pump

2 types of Circulation in Complex Animals

Closed Circulation

- •Blood never leaves tubes and moves in one direction:
- •O₂ rich blood, O₂ poor blood, & waste (NH₃) generally do not mix
- •Interstitial fluids move gasses (O₂/CO₂), nutrients, waste (NH₃), btwn blood & cells
- Occurs in some invertebrates & ALL vertebrates.

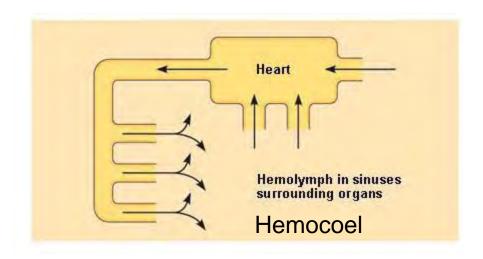


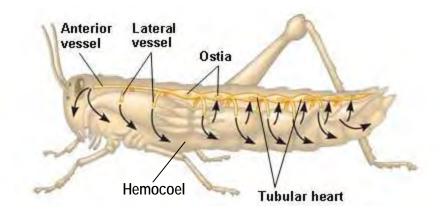


2 types of Circulation in Complex Animals

Open Circulation

- Tubes open-ended;
- Tissues/organs in hemocoel bathe in hemolymph
 - Hemolymph: analogous to blood + interstitial fluid
- •O₂ rich hemolymph + O₂ poor hemolymph + waste (NH₃) mix in hemocoel
- Occurs in arthropods & mollusks (except cephalopods)





Advantages / Disadvantages



Closed circulatory system

- -Higher pressure allows:
 - rapid delivery and removal
 - distant delivery (i.e. bigger bodies)

-Allows for a design in which oxygenated and deoxygenated blood is not mixed

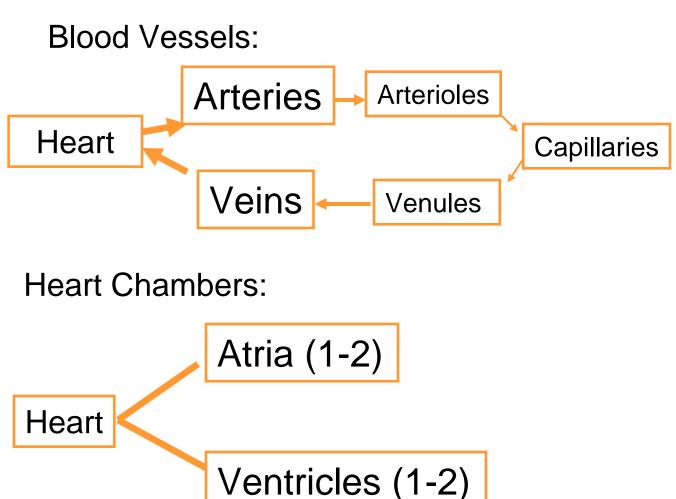


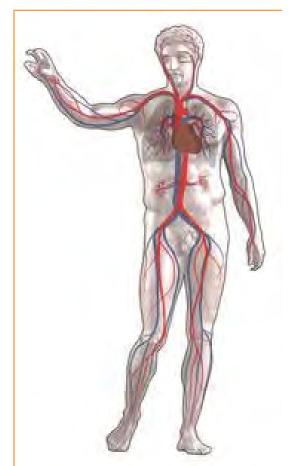
Open circulatory system

-Lower pressure so requires less energy

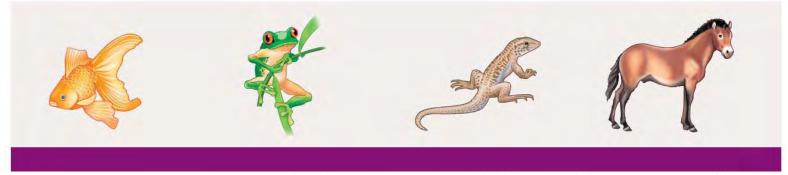
Closed Circulatory System

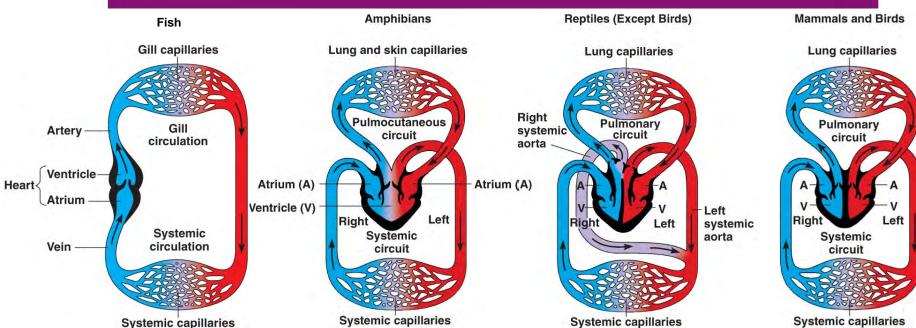
Consists of **blood vessels** and a 2 to 4 chambered **heart**





Vertebrate Circulatory Systems

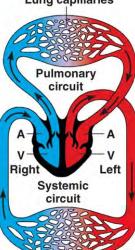




- 2-chambers (1A, 1V)
- Single circulation

- 3-chambers (2A, 1V)
- Double circulation
- Some O₂ rich/poor blood mixes but ridge keeps most separate
- When diving, O₂ poor blood diverted from lungs but not skin (i.e. cutaneous respiration)

- TSL: 3-chambers (2A, 1V)
- Double circulation
- Partial septum in ventricle keeps almost all O₂ rich/poor blood separate
- Crocs: 4-chambers (2A, 2V)
- •TWO systemic aortas in TSL & C
- •When diving, O₂ poor blood diverted from lungs in some reptiles



- 4-chambers (2A, 2V)
- Double circulation

Mammalian Circulation Pattern

- R atrium to R ventricle
- 2. to pulmonary arteries (O₂ poor)
- 3. to capillaries of lungs
- 4. to pulmonary vein (O₂ rich)
- 5. to L atrium to L venticle
- 6. to aorta
- 7. to arteries/ capillaries of the head, limbs and organs& coronary arteries of heart
- 8. to anterior and posterior vena cava
- 9. to R atrium

Know <u>order</u> of vessels and heart chambers

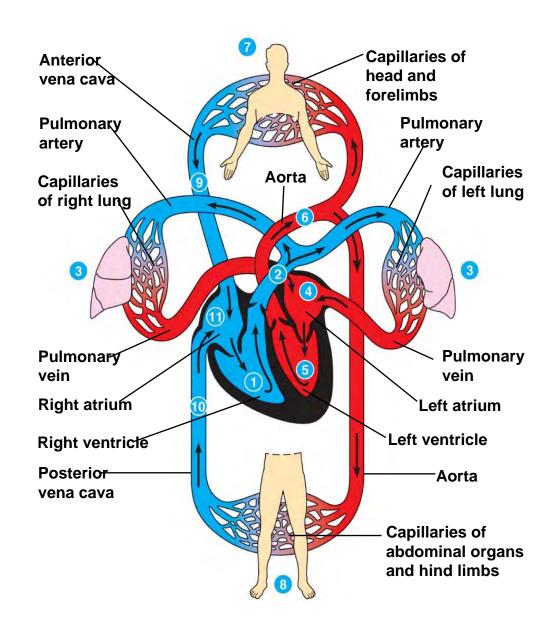
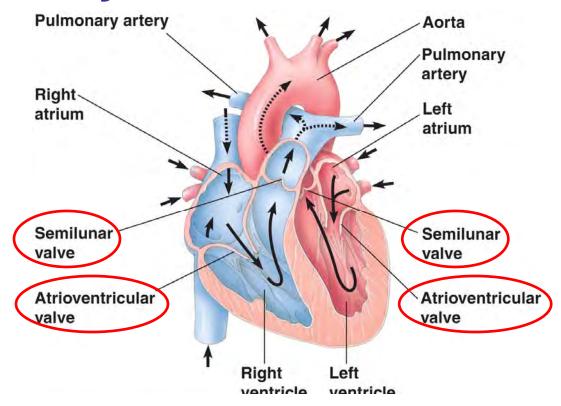


Figure 42.6 (Campbell 9th ed)

Anatomy of the Mammalian Heart



Walls:

- a. Ventricles need greater pressure thus have thicker walls
 - Left ventricle thicker than right

Valves prevent backflow:

- a. Atrioventricular from ventricles back into atria
- b. Semilunar from aorta/pulmonary arteries back into ventricle

Heart Pumping

Cardiac cycle - The sequence of contraction and relaxation that makes up the heartbeat.

- A. Diastole relaxation
- B. Systole contraction

Cardiac output

- Blood volume pumped/ minuteDepends on:
 - A. Heart rate beats/ minute
 - B. Stroke volume
 - Amount of blood pumped from ventricle per contraction

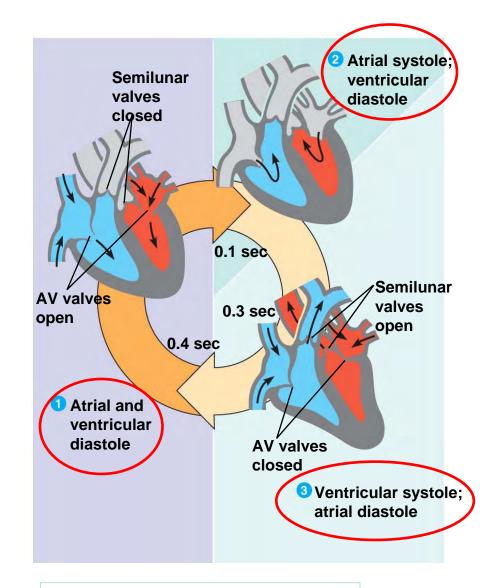
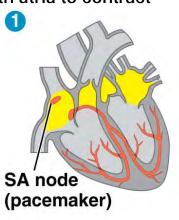
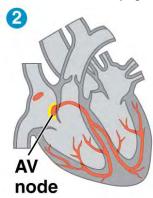


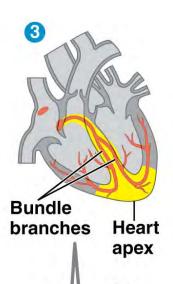
Figure 42.8 (Campbell 9th ed)

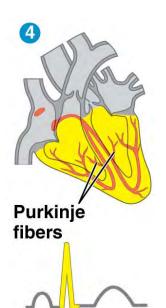
Control of Heart Rhythm

- 1. Sinoatrial node generates signal causing both atria to contract
- 2. Signal delayed at atrioventricular node to let atria empty
- 3. AV node passes signal to heart apex
- 4. Signal spreads throughout ventricle walls

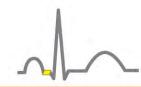










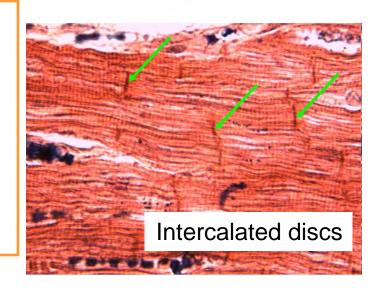


Sinoatrial node (SA):

- = Group of autorhythmic cells in wall of RA
 - a. Sets contraction rate and timing
 - b. Regulated by signals from nervous & endocrine (hormones) systems

Atrioventricular Node (AV):

= Autorhythmic cells in wall btwn RA and LA

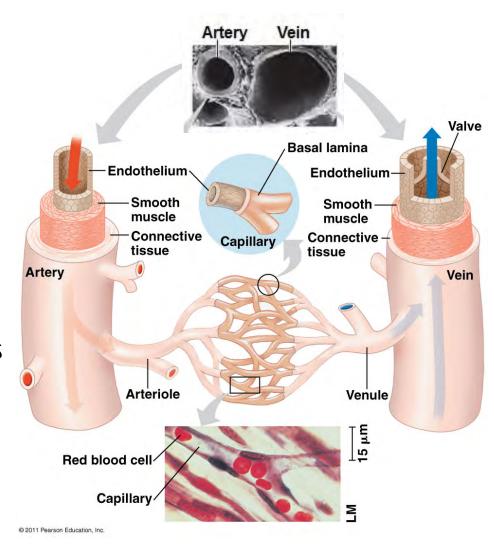




Vessel Architecture

A. Veins vs Arteries structure

- Both with <u>three</u> tissue layers
 - 1. Outer connective tissue
 - Thicker in arteries
 - 2. Middle smooth muscle
 - Thicker in arteries
 - 3. Inner epithelium (endothelium)
 - Similar in arteries and veins
- Lumen
 - Larger in veins
- Valves
 - •Only in veins (Why?)



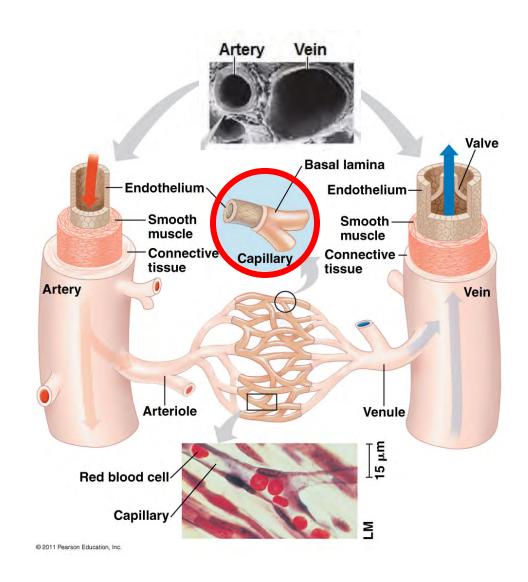
Vessel Architecture

A. Veins vs Arteries structure

B. Capillary structure

- One tissue layer
 - Epithelium (endothelium)
 - Includes outer basement membrane
- Lumen thickness of 1 RBC

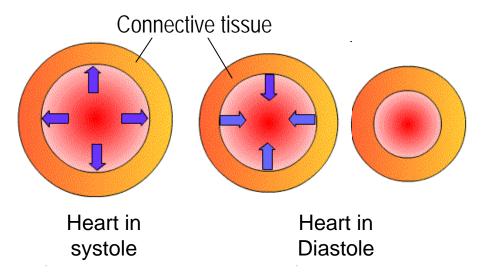
Thus capillaries are more permeable to substances than arteries/veins



What causes blood to flow?

Arteries

- A. Heart pumping
- B. Contraction of thick <u>smooth</u> muscle
- C. Recoil of thick elastic connective tissue



Veins

- A. Contraction of smooth muscle
- B. Contraction of abutting skeletal muscle
- C. Expansion due to pressure change during inhalation (veins near heart only)

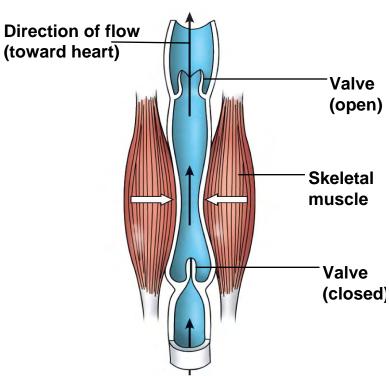
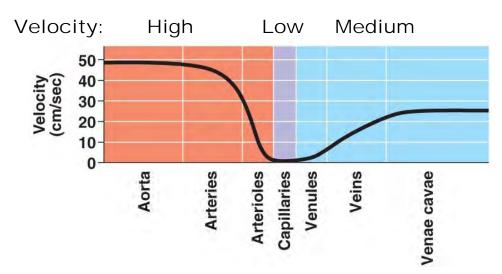


Figure 42.13 (Campbell 9th ed)

Blood velocity decreases as it goes from thick arteries into thin capillaries. Why?

Area:

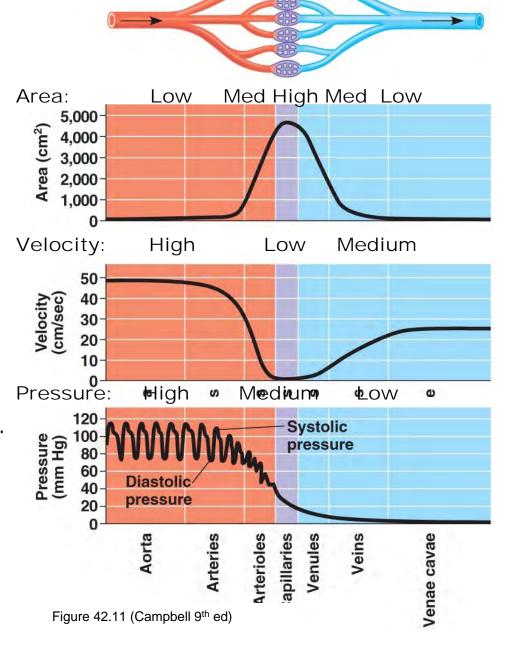


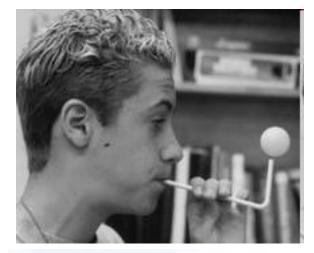


Blood velocity decreases as it goes from thick arteries into thin capillaries. Why?

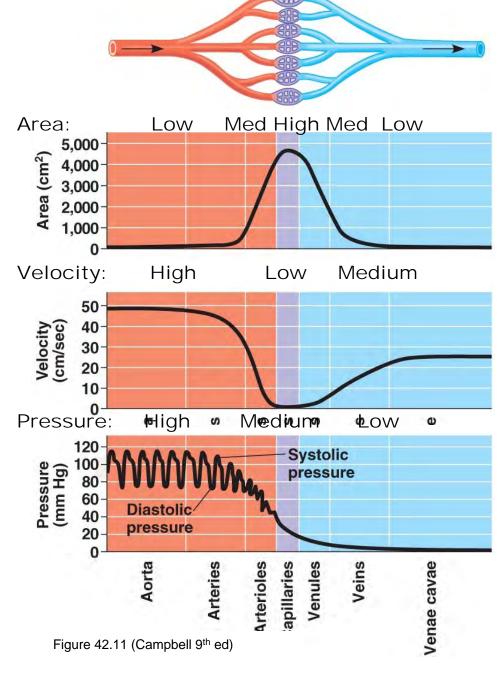
Though small, there are many capillaries so their total volume is high and far exceeds the arteries.

Low area in the arteries \rightarrow high pressure. High area in capillaries \rightarrow low pressure.









Pressure must be low in <u>capillaries</u> since they must be <u>thin</u> to transmit substances between the blood and interstitial fluids.

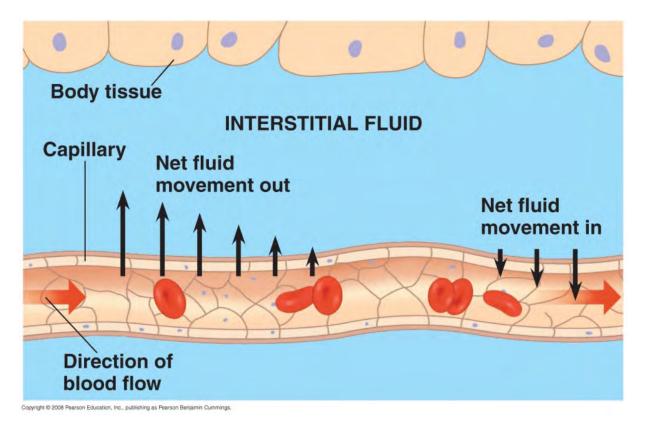


Figure 42.11 (Campbell 9th ed)

Capillary exchange

- Osmotic pressure (OP) and blood pressure (BP) affect fluid exchange btwn capillaries and interstitial fluid
- 2. Venous end of capillary bed:
 - a) BP low (far from heart).
 - b) OP high due to large proteins & RBCs in blood making [solute] higher than interstitial fluids.
 - c) Since OP>BP, interstitial fluid (H₂O) + waste (CO₂, NH₃) flow from tissues <u>into</u> capillary due to osmosis
- 3. Arteriole end of capillary bed:
 - a) BP>OP
 - b) Net flow of H₂O, nutrients, O₂
 out of capillary into tissue

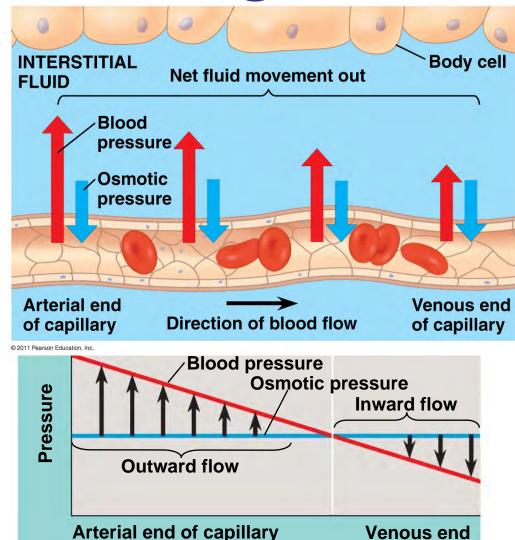
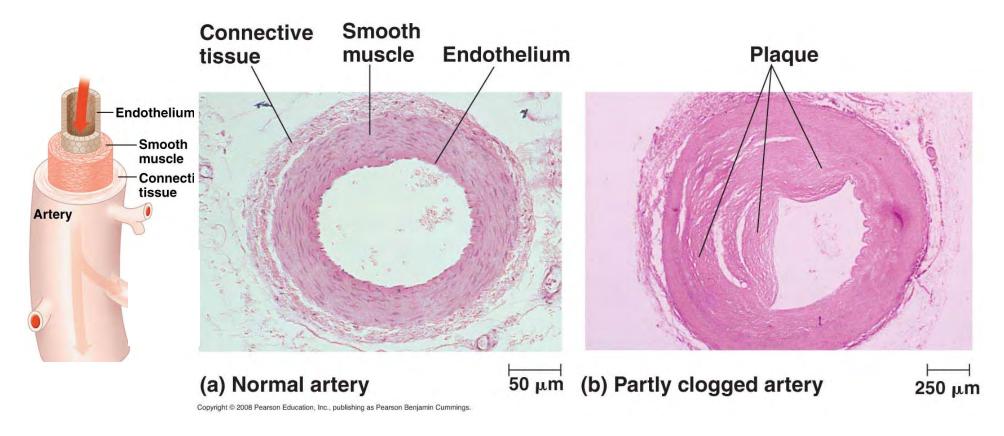


Figure 42.15 (Campbell 8th ed)

4. H₂0 etc also returned to blood via lymphatic system



Atherosclerosis aka 'Hardening of the arteries'



- 1. Vitamin C needed to make connective tissue collagen fibers.
- Deficiency damages connective tissue and leukocytes move in to repair damage.
- 3. Leukocytes take up lipids, primarily cholesterol, which joins with fibers.
- 4. Altered fibers make CT stiff causing higher blood pressure.

Blood Composition

Blood = cells suspended in liquid called plasma

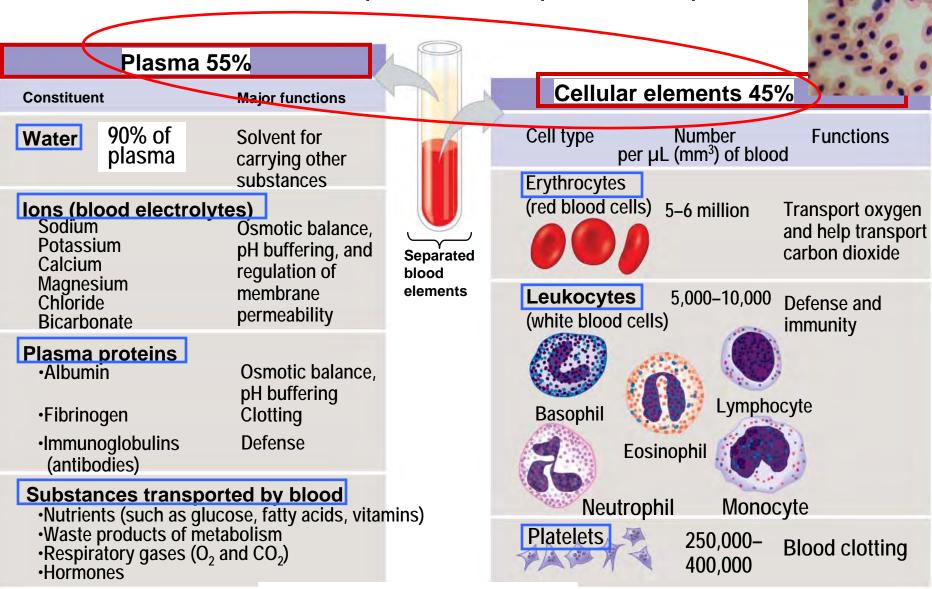


Figure 42.17 (Campbell 9th ed)

Regulation of Red Blood Cell Production

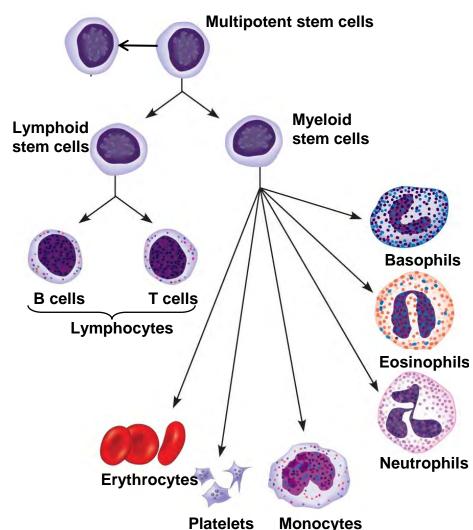
- Erythrocyte production controlled via <u>negative</u> feedback
- •Low O₂ signal <u>in kidney</u> causes release of hormone <u>erythropoietin</u> which acts on stem cells to increase RBCs
- •High O₂ stops release of erythropoietin

Erythropoletin Cycle

Epo stimulates stem cells in bone marrow in bone marrow increased O2 carrying capacity series of steps

(-) feedback on Epo RBC

 Stem cells found in red bone marrow (ribs, vertebrae, sternum, pelvis)



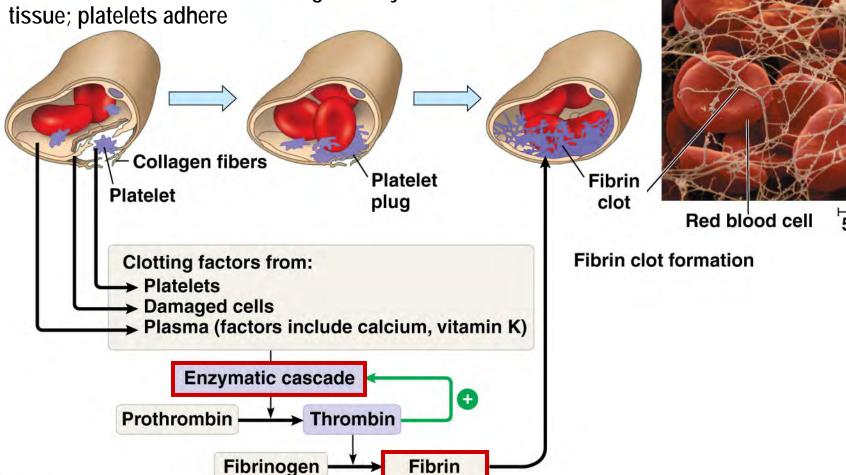
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Blood Clotting

Endothelium of vessel is damaged, exposing connective tissue; platelets adhe

Platelets form a plug
 & release clotting factor
 that begins enzyme cascade

Fibrin, formed from fibrinogen in blood, forms clot to reinforce plug



Readings on which you will NOT be tested

- Figure 42.12 (907)
- Heart attacks (914)
- Risk factors and Treatment (914-915)
- Figure 42.21 (914)
- Sections 42.5 to 42.7 (915-926)

In general:

- You are NOT responsible for definitions of terms or sections included in the text but which were not discussed in lecture
- You are not responsible for the details of examples used in the text but not discussed in lecture. HOWEVER, these additional examples will help your understanding of concepts discussed and may be used on exams to test if you understand the general concepts.
- You ARE responsible for material covered in lecture but not included in the readings

Next Lecture

• Immune System – Chapter 43