



# Chapter 13

---

## Heart and Circulation

# Functions of the Circulatory System

---

- Transportation:
  - Respiratory:
    - Transport  $O_2$  and  $CO_2$ .
  - Nutritive:
    - Carry absorbed digestion products to liver and to tissues.
  - Excretory:
    - Carry metabolic wastes to kidneys to be excreted.



# Functions of the Circulatory System (continued)

---

- Regulation:
  - Hormonal:
    - Carry hormones to target tissues to produce their effects.
  - Temperature:
    - Divert blood to cool or warm the body.
  - Protection:
    - Blood clotting.
  - Immune:
    - Leukocytes, cytokines and complement act against pathogens.



# Components of Circulatory System

---

- Cardiovascular System (CV):

- Heart:

- Pumping action creates pressure needed to push blood through vessels.

- Blood vessels:

- Permits blood flow from heart to cells and back to the heart.
  - Arteries, arterioles, capillaries, venules, veins.

- Lymphatic System:

- Lymphatic vessels transport interstitial fluid.

- Lymph nodes cleanse lymph prior to return in venous blood.



# Composition of Blood

---

- Plasma:
  - Straw-colored liquid.
    - Consists of H<sub>2</sub>O and dissolved solutes.
      - Ions, metabolites, hormones, antibodies.
        - Na<sup>+</sup> is the major solute of the plasma.
- Plasma proteins:
  - Constitute 7-9% of plasma.
    - Albumin:
      - Accounts for 60-80% of plasma proteins.
      - Provides the colloid osmotic pressure needed to draw H<sub>2</sub>O from interstitial fluid to capillaries.
        - Maintains blood pressure.



# Composition of the Blood (continued)

---

- Plasma proteins (continued):
  - Globulins:
    - $\alpha$  globulin:
      - Transport lipids and fat soluble vitamins.
    - $\beta$  globulin:
      - Transport lipids and fat soluble vitamins.
    - $\gamma$  globulin:
      - Antibodies that function in immunity.
  - Fibrinogen:
    - Constitutes 4% of plasma proteins.
    - Important clotting factor.
      - Converted into fibrin during the clotting process.



# Composition of the Blood (continued)

---

- Serum:
  - Fluid from clotted blood.
    - Does not contain fibrinogen.
- Plasma volume:
  - Number of regulatory mechanisms in the body maintain homeostasis of plasma volume.
    - Osmoreceptors.
    - ADH.
    - Renin-angiotensin-aldosterone system.



# Erythrocytes

---

- Flattened biconcave discs.
- Provide increased surface area through which gas can diffuse.
- Lack nuclei and mitochondria.
  - Half-life ~ 120 days.
- Each RBC contains 280 million hemoglobin with 4 heme chains (contain iron).
- Removed from circulation by phagocytic cells in liver, spleen, and bone marrow.





# Leukocytes

---

- Contain nuclei and mitochondria.
- Move in amoeboid fashion.
  - Can squeeze through capillary walls (diapedesis).
- Almost invisible, so named after their staining properties.
  - Granular leukocytes:
    - Help detoxify foreign substances.
      - Release heparin.
  - Agranular leukocytes:
    - Phagocytic.
      - Produce antibodies.



# Platelets (thrombocytes)

---

- Smallest of formed elements.
  - Are fragments of megakaryocytes.
  - Lack nuclei.
- Capable of amoeboid movement.
- Important in blood clotting:
  - Constitute most of the mass of the clot.
  - Release serotonin to vasoconstrict and reduce blood flow to area.
- Secrete growth factors:
  - Maintain the integrity of blood vessel wall.
- Survive 5-9 days.

# Blood Cells and Platelets

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.



**Neutrophils**



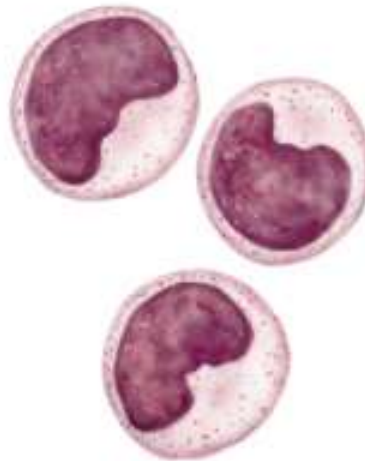
**Eosinophils**



**Basophils**



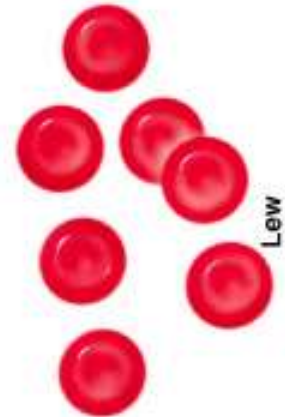
**Lymphocytes**



**Monocytes**



**Platelets**



**Erythrocytes**

Lew



# Hematopoiesis

---

- Undifferentiated cells gradually differentiate to become stem cells, that form blood cells.
- Occurs in myeloid tissue (bone marrow of long bones) and lymphoid tissue.
- 2 types of hematopoiesis:
  - Erythropoiesis:
    - Formation of RBCs.
  - Leukopoiesis:
    - Formation of WBCs.



# Erythropoiesis

---

- Active process.
  - 2.5 million RBCs are produced every second.
- Primary regulator is erythropoietin.
  - Binds to membrane receptors of cells that will become erythroblasts.
  - Erythroblasts transform into normoblasts.
  - Normoblasts lose their nuclei to become reticulocytes.
  - Reticulocytes change into mature RBCs.
    - Stimulates cell division.
- Old RBCs are destroyed in spleen and liver.
  - Iron recycled back to myeloid tissue to be reused in hemoglobin production.
- Need iron, vitamin B<sub>12</sub> and folic acid for synthesis.



# Leukopoiesis

---

- Cytokines stimulate different types and stages of WBC production.
- Multipotent growth factor-1, interleukin-1, and interleukin-3:
  - Stimulate development of different types of WBC cells.
- Granulocyte-colony stimulating factor (G-CSF):
  - Stimulates development of neutrophils.
- Granulocyte-monocyte colony stimulating factor (GM-CSF):
  - Stimulates development of monocytes and eosinophils.



# RBC Antigens and Blood Typing

- Each person's blood type determines which antigens are present on their RBC surface.
- Major group of antigens of RBCs is the ABO system:
  - Type A:
    - Only A antigens present.
  - Type B:
    - Only B antigens present.
  - Type AB:
    - Both A and B antigens present.
  - Type O:
    - Neither A or B antigens present.



# RBC Antigens and Blood Typing

(continued)

---

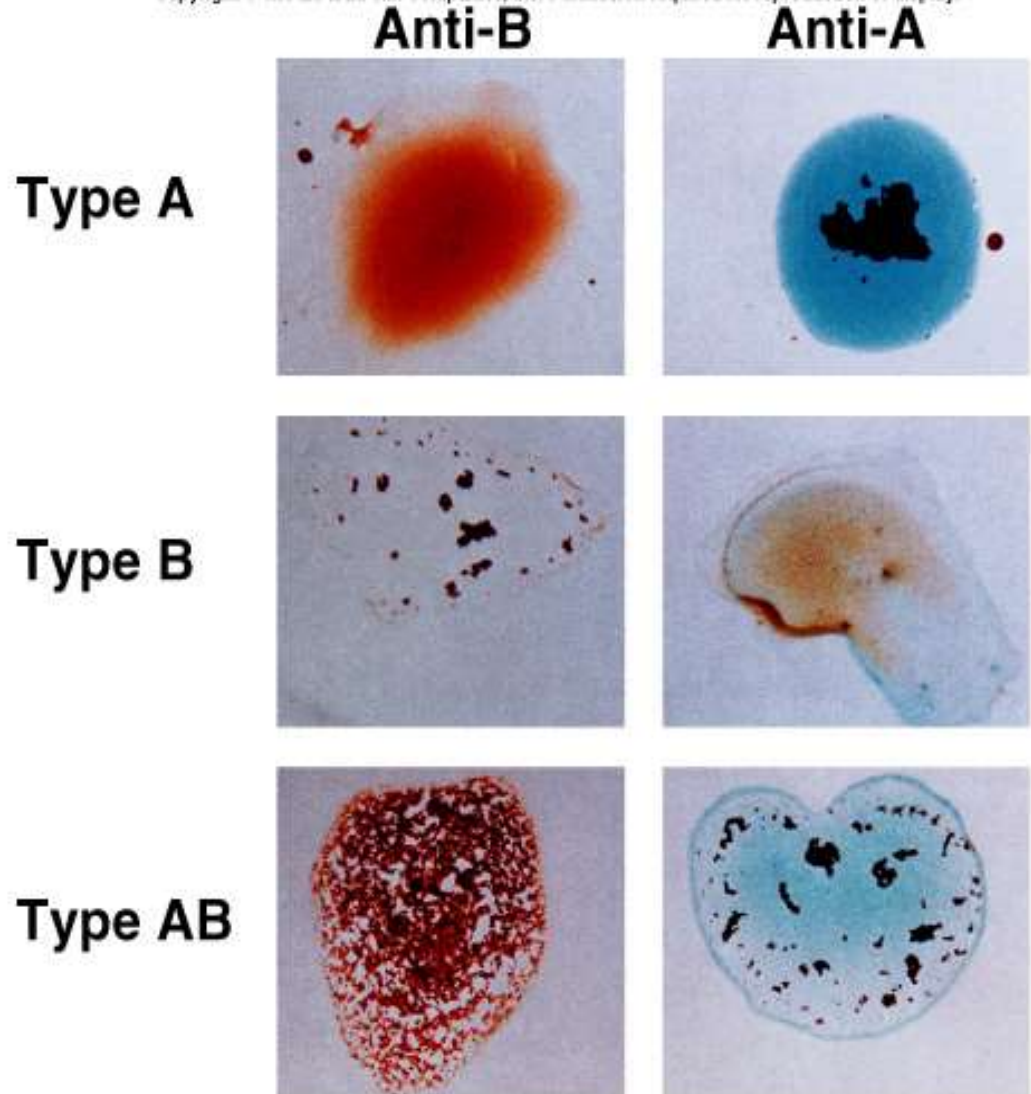
- Each person inherits 2 genes that control the production of ABO groups.
- Type A:
  - May have inherited A gene from each parent.
  - May have inherited A gene from one parent and O gene from the other.
- Type B:
  - May have inherited B gene from each parent.
  - May have inherited B gene from one parent and O gene from the other parent.
- Type AB:
  - Inherited the A gene from one parent and the B gene from the other parent.
- Type O:
  - Inherited O gene from each parent.



# Transfusion Reactions

- If blood types do not match, the recipient's antibodies attach to donor's RBCs and agglutinate.
- Type O:
  - Universal donor:
    - Lack A and B antigens.
    - Recipient's antibodies cannot agglutinate the donor's RBCs.
- Type AB:
  - Universal recipient:
    - Lack the anti-A and anti-B antibodies.
  - Cannot agglutinate donor's RBCs.

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.





# Rh Factor

---

- Another group of antigens found on RBCs.
- Rh positive:
  - Has Rho(D) antigens.
- Rh negative:
  - Does not have Rho(D) antigens.
- Significant when Rh- mother gives birth to Rh+ baby.
  - At birth, mother may become exposed to Rh+ blood of fetus.
    - Mother at subsequent pregnancies may produce antibodies against the Rh factor.
- Erythroblastosis fetalis:
  - Rh- mother produces antibodies, which cross placenta.
    - Hemolysis of Rh+ RBCs in the fetus.



# Blood Clotting

---

- Function of platelets:
  - Platelets normally repelled away from endothelial lining by prostacyclin (prostaglandin).
    - Do not want to clot normal vessels.
- Damage to the endothelium wall:
  - Exposes subendothelial tissue to the blood.



# Blood Clotting (continued)

---

- Platelet release reaction:
  - Endothelial cells secrete von Willebrand factor to cause platelets to adhere to collagen.
  - When platelets stick to collagen, they degranulate as platelet secretory granules:
    - Release ADP, serotonin and thromboxane  $A_2$ .
      - Serotonin and thromboxane  $A_2$  stimulate vasoconstriction.
      - ADP and thromboxane  $A_2$  make other platelets “sticky.”
        - Platelets adhere to collagen.
        - Stimulates the platelet release reaction.
    - Produce platelet plug.
      - Strengthened by activation of plasma clotting factors.



# Blood Clotting (continued)

---

- Platelet plug strengthened by fibrin.
- Clot reaction:
  - Contraction of the platelet mass forms a more compact plug.
  - Conversion of fibrinogen to fibrin occurs.
- Conversion of fibrinogen to fibrin:
  - Intrinsic Pathway:
    - Initiated by exposure of blood to a negatively charged surface (collagen).
      - This activates factor XII (protease), which activates other clotting factors.
    - $\text{Ca}^{2+}$  and phospholipids convert prothrombin to thrombin.
      - Thrombin converts fibrinogen to fibrin.
        - Produces meshwork of insoluble fibrin polymers.



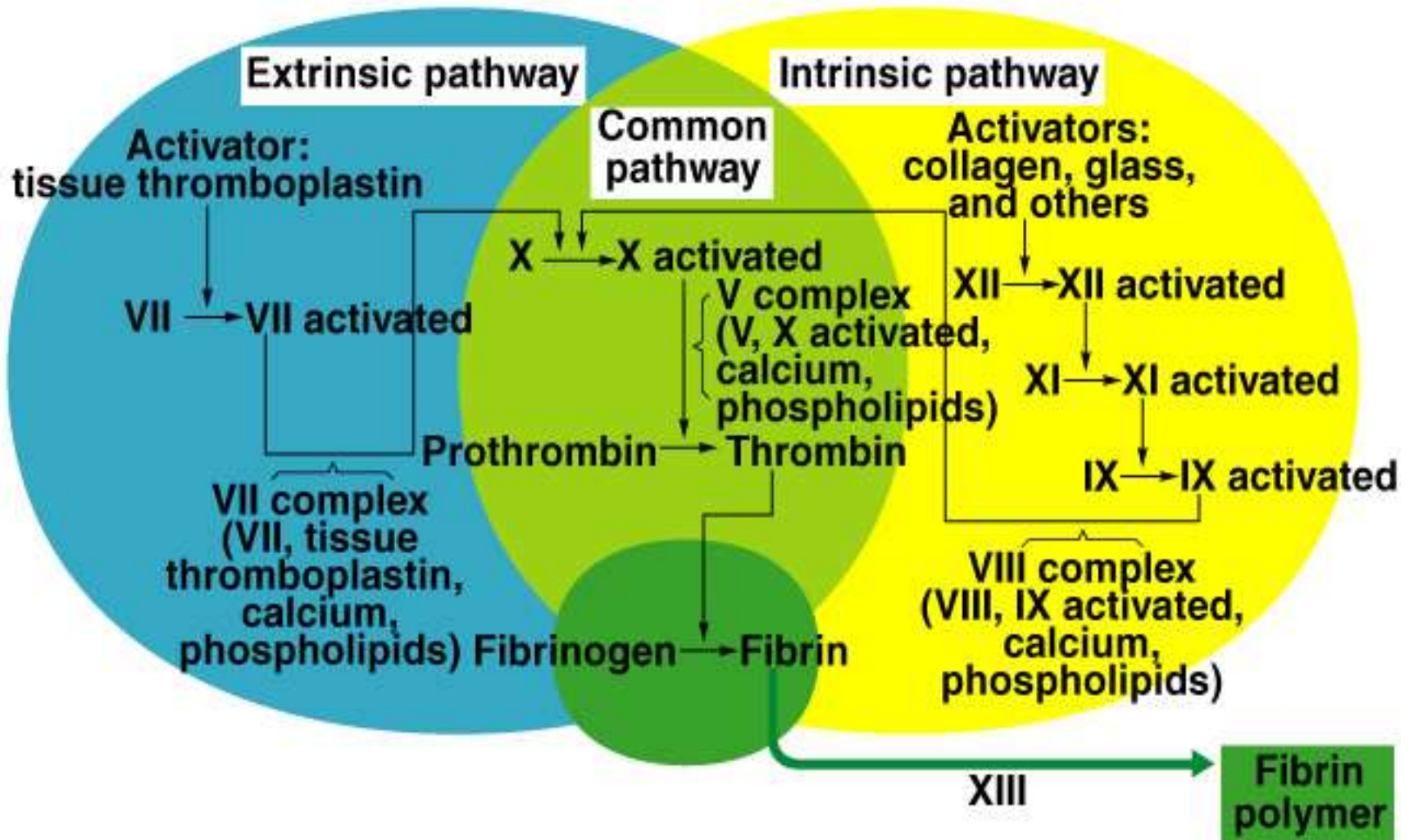
# Blood Clotting (continued)

---

- Extrinsic pathway:
  - Thromboplastin is not a part of the blood, so called extrinsic pathway.
  - Damaged tissue releases thromboplastin.
    - Thromboplastin initiates a short cut to formation of fibrin.

# Blood Clotting (continued)

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.





# Dissolution of Clots

---

- Activated factor XII converts an inactive molecule into the active form (kallikrein).
  - Kallikrein converts plasminogen to plasmin.
- Plasmin is an enzyme that digests the fibrin.
  - Clot dissolution occurs.
- Anticoagulants:
  - Heparin:
    - Activates antithrombin III.
  - Coumarin:
    - Inhibits cellular activation of vitamin K.





# Acid-Base Balance in the Blood

---

- Blood pH is maintained within a narrow range by lungs and kidneys.
- Normal pH of blood is 7.35 to 7.45.
- Some H<sup>+</sup> is derived from carbonic acid.
- **H<sub>2</sub>O + CO<sub>2</sub> ⇌ H<sub>2</sub>CO<sub>3</sub> ⇌ H<sup>+</sup> + HCO<sub>3</sub><sup>-</sup>**

# Acid-Base Balance in the Blood

(continued)

- Types of acids in the body:
  - Volatile acids:
    - Can leave solution and enter the atmosphere as a gas.
      - Carbonic acid.



- Nonvolatile acids:
  - Acids that do not leave solution.
    - Byproducts of aerobic metabolism, during anaerobic metabolism and during starvation.
    - Sulfuric and phosphoric acid.



# Buffer Systems

---

- Provide or remove  $H^+$  and stabilize the pH.
- Include weak acids that can donate  $H^+$  and weak bases that can absorb  $H^+$ .
- $HCO_3^-$  is the major buffer in the plasma.
- **$H^+ + HCO_3^- \longrightarrow H_2CO_3$**
- Under normal conditions excessive  $H^+$  is eliminated in the urine.



# Acid Base Disorders

---

- Respiratory acidosis:
  - Hypoventilation.
    - Accumulation of  $\text{CO}_2$ .
      - pH decreases.
- Respiratory alkalosis:
  - Hyperventilation.
    - Excessive loss of  $\text{CO}_2$ .
      - pH increases.
- Metabolic acidosis:
  - Gain of fixed acid or loss of  $\text{HCO}_3^-$ .
    - Plasma  $\text{HCO}_3^-$  decreases.
      - pH decreases.
- Metabolic alkalosis:
  - Loss of fixed acid or gain of  $\text{HCO}_3^-$ .
    - Plasma  $\text{HCO}_3^-$  increases.
      - pH increases.



# pH

---

- Normal pH is obtained when the ratio of  $\text{HCO}_3^-$  to  $\text{CO}_2$  is 20:1.
- Henderson-Hasselbalch equation:
- $$\text{pH} = 6.1 + \log \frac{[\text{HCO}_3^-]}{[0.03P_{\text{CO}_2}]}$$

# Pulmonary and Systemic Circulations

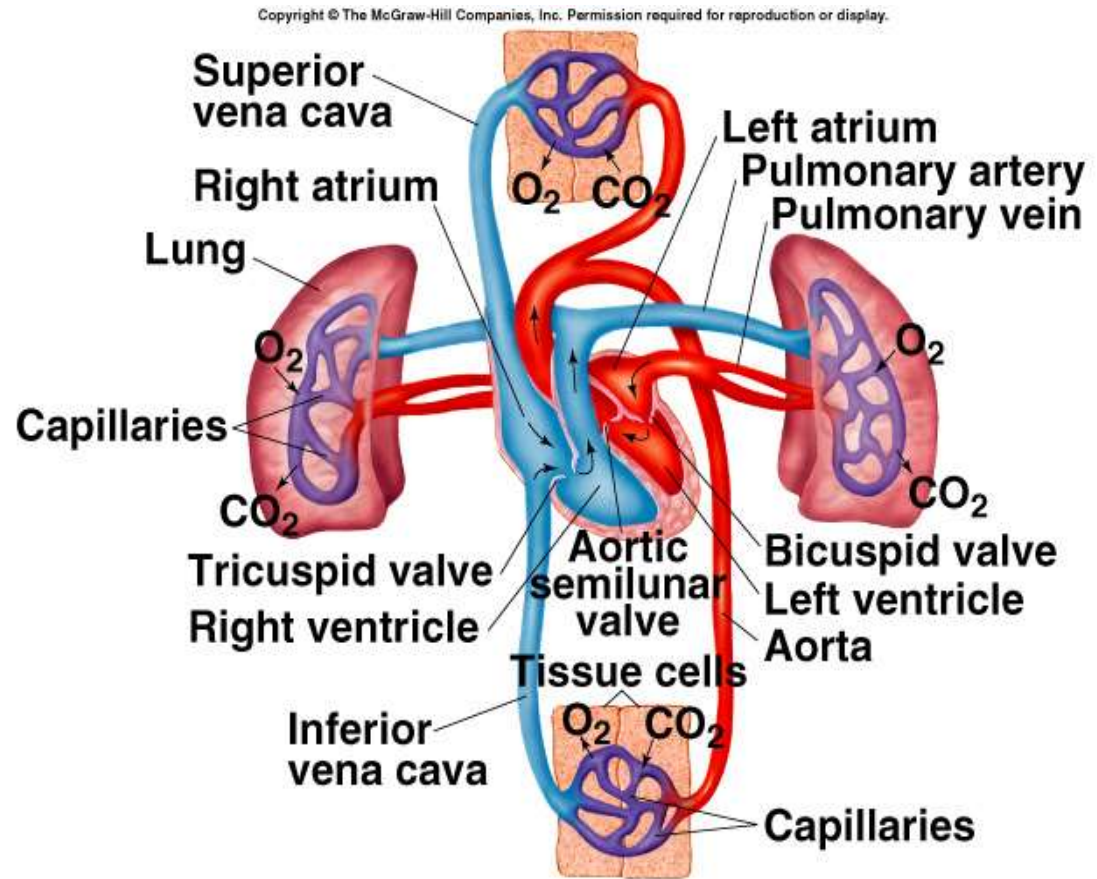
## Pulmonary circulation:

- Path of blood from right ventricle through the lungs and back to the heart.

## Systemic circulation:

- Oxygen-rich blood pumped to all organ systems to supply nutrients.

■ Rate of blood flow through systemic circulation = flow rate through pulmonary circulation.





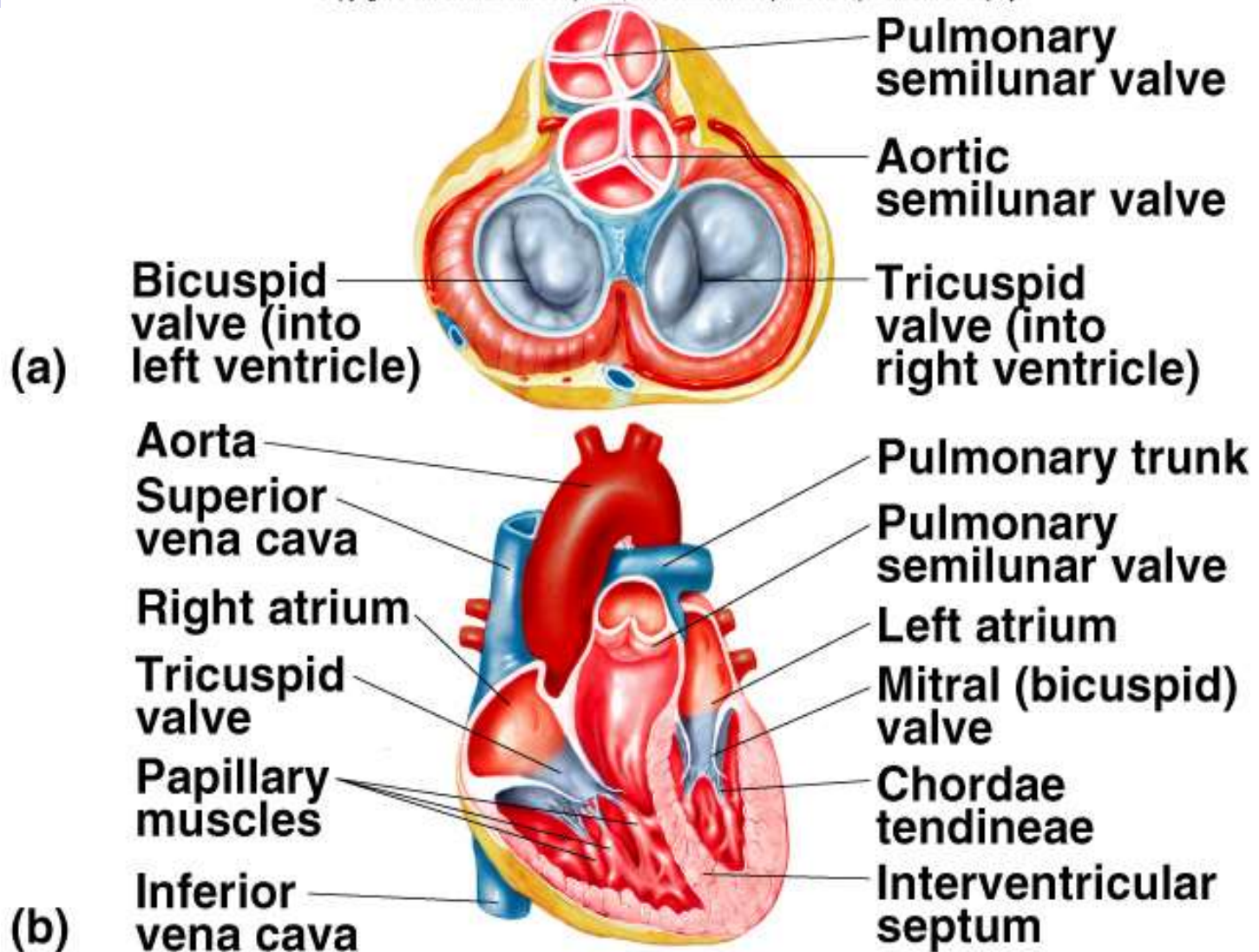
# Atrioventricular and Semilunar Valves

---

- Atria and ventricles are separated into 2 functional units by a sheet of connective tissue by AV (atrioventricular) valves.
  - One way valves.
  - Allow blood to flow from atria into the ventricles.
- At the origin of the pulmonary artery and aorta are semilunar valves.
  - One way valves.
  - Open during ventricular contraction.
- Opening and closing of valves occur as a result of pressure differences.

# Atrioventricular and Semilunar Valves

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.







# Cardiac Cycle

---

- Refers to the repeating pattern of contraction and relaxation of the heart.
  - Systole:
    - Phase of contraction.
  - Diastole:
    - Phase of relaxation.
  - End-diastolic volume (EDV):
    - Total volume of blood in the ventricles at the end of diastole.
  - Stroke volume (SV):
    - Amount of blood ejected from ventricles during systole.
  - End-systolic volume (ESV):
    - Amount of blood left in the ventricles at the end of systole.



# Cardiac Cycle (continued)

---

- Step 1: Isovolumetric contraction:
  - QRS just occurred.
  - Contraction of the ventricle causes ventricular pressure to rise above atrial pressure.
    - AV valves close.
  - Ventricular pressure is less than aortic pressure.
    - Semilunar valves are closed.
      - Volume of blood in ventricle is EDV.
- Step 2: Ejection:
  - Contraction of the ventricle causes ventricular pressure to rise above aortic pressure.
    - Semilunar valves open.
  - Ventricular pressure is greater than atrial pressure.
    - AV valves are closed.
      - Volume of blood ejected: SV.



# Cardiac Cycle (continued)

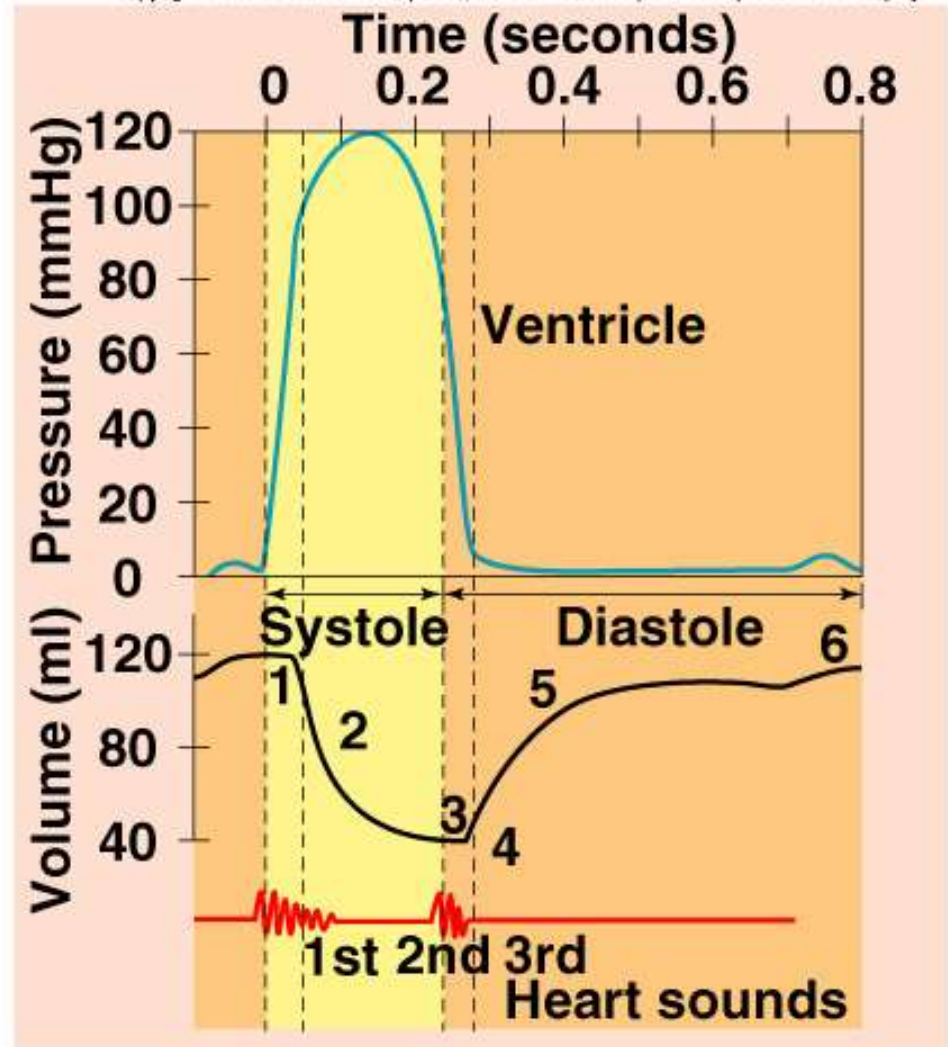
---

- Step 3: T wave occurs:
  - Ventricular pressure drops below aortic pressure.
- Step 4: Isovolumetric relaxation:
  - Back pressure causes semilunar valves to close.
    - AV valves are still closed.
      - Volume of blood in the ventricle: ESV.
- Step 5: Rapid filling of ventricles:
  - Ventricular pressure decreases below atrial pressure.
    - AV valves open.
      - Rapid ventricular filling occurs.

# Cardiac Cycle (continued)

- Step 6: Atrial systole:
  - P wave occurs.
  - Atrial contraction.
    - Push 10-30% more blood into the ventricle.

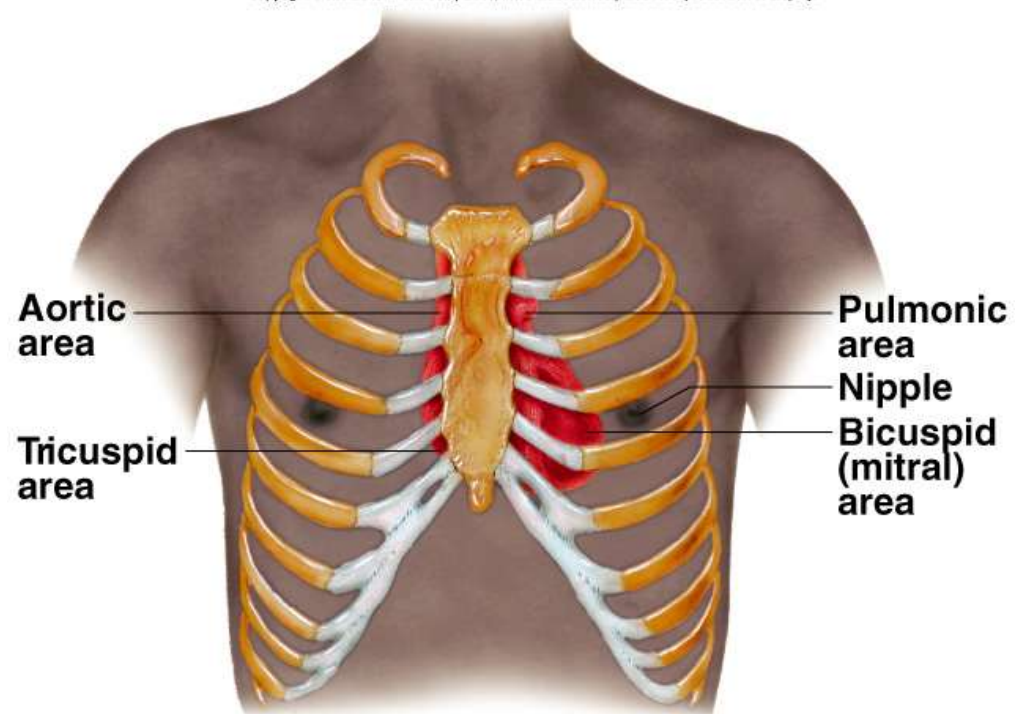
Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.



# Heart Sounds

- Closing of the AV and semilunar valves.
- Lub (first sound):
  - Produced by closing of the AV valves during isovolumetric contraction.
- Dub (second sound):
  - Produced by closing of the semilunar valves when pressure in the ventricles falls below pressure in the arteries.

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.





# Heart Murmurs

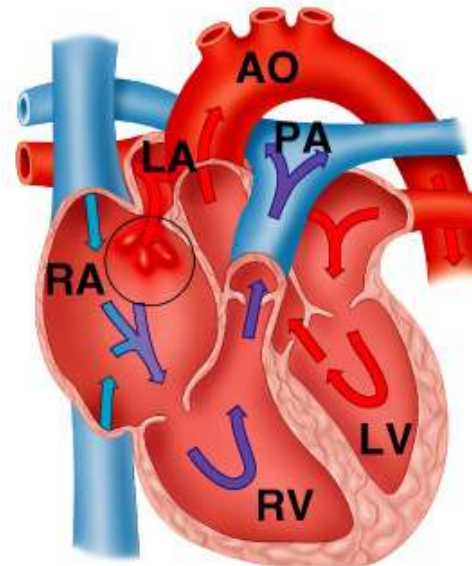
---

- Abnormal heart sounds produced by abnormal patterns of blood flow in the heart.
- Defective heart valves:
  - Valves become damaged by antibodies made in response to an infection, or congenital defects.
- Mitral stenosis:
  - Mitral valve becomes thickened and calcified.
    - Impairs blood flow from left atrium to left ventricle.
    - Accumulation of blood in left ventricle may cause pulmonary HTN.
- Incompetent valves:
  - Damage to papillary muscles.
    - Valves do not close properly.
      - Murmurs produced as blood regurgitates through valve flaps.

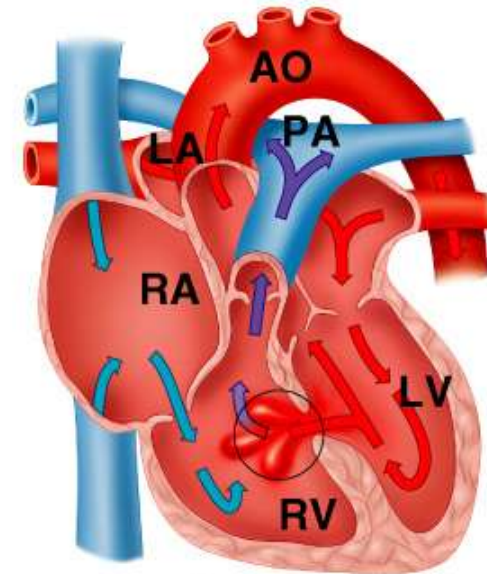
# Heart Murmurs

- Septal defects:
  - Usually congenital.
    - Holes in septum between the left and right sides of the heart.
    - May occur either in interatrial or interventricular septum.
  - Blood passes from left to right.

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.



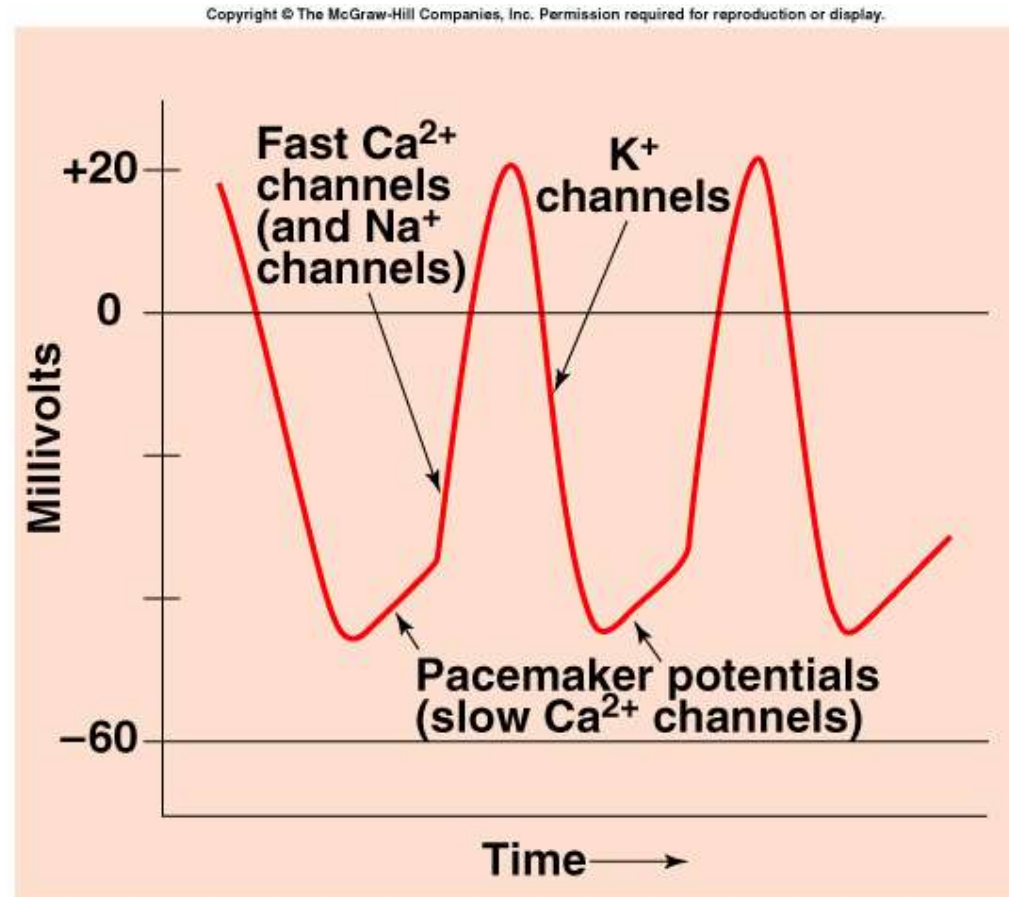
Septal defect  
in atria



Septal defect  
in ventricles

# Electrical Activity of the Heart

- SA node:
  - Demonstrates automaticity:
    - Functions as the pacemaker.
  - Spontaneous depolarization (pacemaker potential):
    - Spontaneous diffusion caused by diffusion of  $\text{Ca}^{2+}$  through slow  $\text{Ca}^{2+}$  channels.
      - Cells do not maintain a stable RMP.







# Pacemaker AP

---

- Depolarization:
  - VG fast  $\text{Ca}^{2+}$  channels open.
    - $\text{Ca}^{2+}$  diffuses inward.
  - Opening of VG  $\text{Na}^{+}$  channels may also contribute to the upshoot phase of the AP.
- Repolarization:
  - VG  $\text{K}^{+}$  channels open.
    - $\text{K}^{+}$  diffuses outward.
- Ectopic pacemaker:
  - Pacemaker other than SA node:
    - If APs from SA node are prevented from reaching these areas, these cells will generate pacemaker potentials.



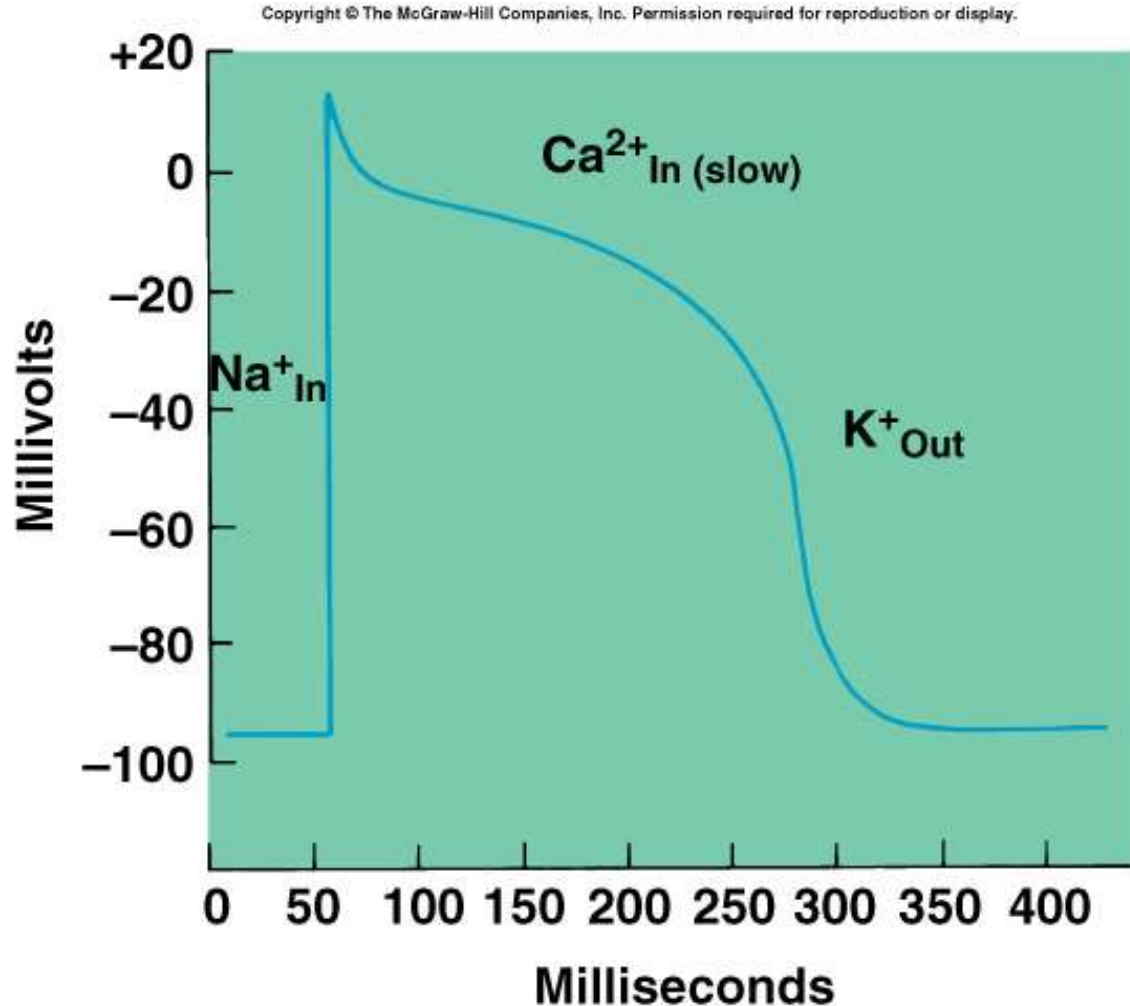
# Myocardial APs

---

- Majority of myocardial cells have a RMP of  $-90$  mV.
- SA node spreads APs to myocardial cells.
  - When myocardial cell reaches threshold, these cells depolarize.
- Rapid upshoot occurs:
  - VG  $\text{Na}^+$  channels open.
    - Inward diffusion of  $\text{Na}^+$ .
- Plateau phase:
  - Rapid reversal in membrane polarity to  $-15$  mV.
    - VG slow  $\text{Ca}^{2+}$  channels open.
      - Slow inward flow of  $\text{Ca}^{2+}$  balances outflow of  $\text{K}^+$ .

# Myocardial APs (continued)

- Rapid repolarization:
  - VG  $K^+$  channels open.
  - Rapid outward diffusion of  $K^+$ .





# Conducting Tissues of the Heart

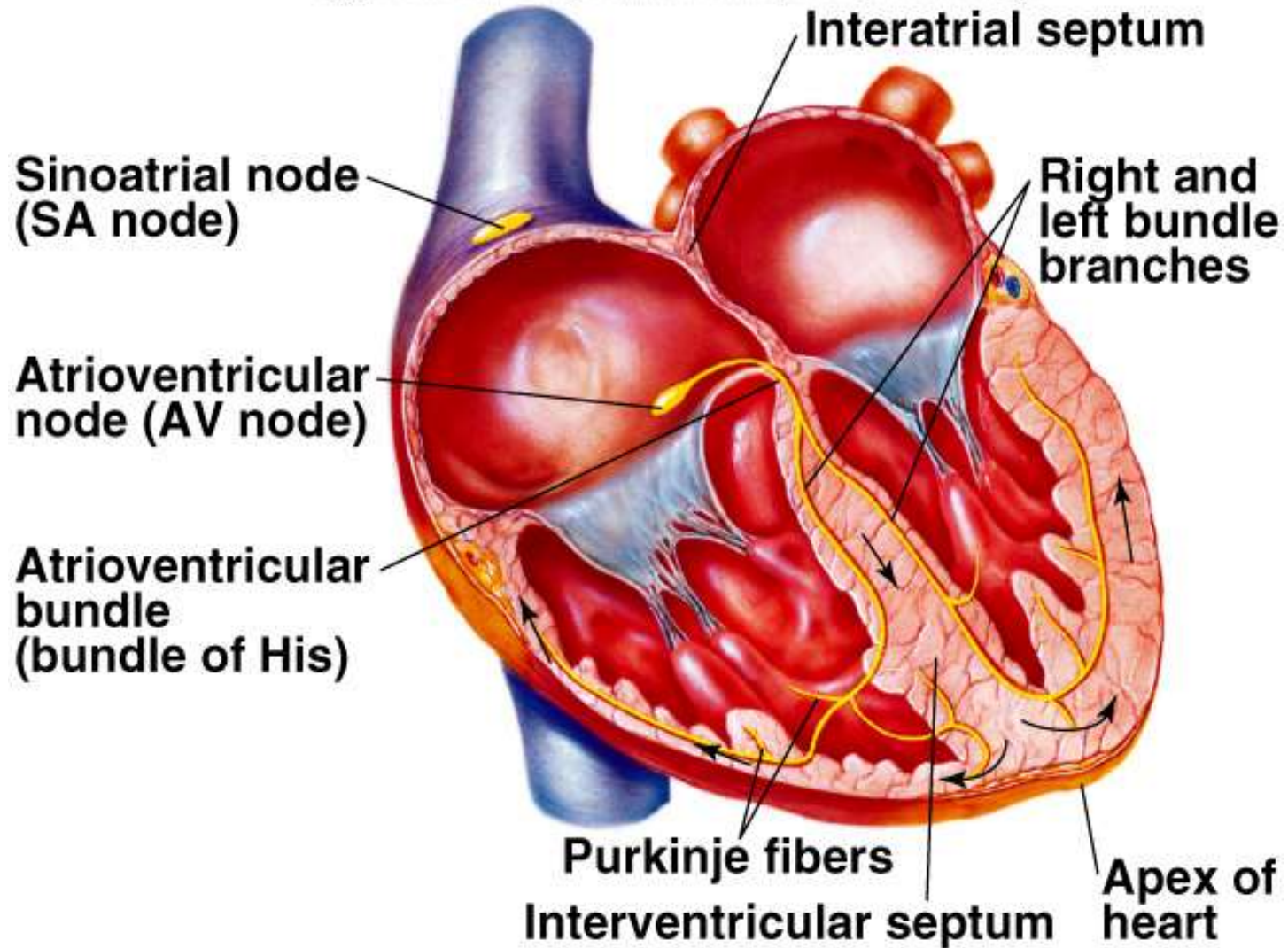
---

- APs spread through myocardial cells through gap junctions.
- Impulses cannot spread to ventricles directly because of fibrous tissue.
- Conduction pathway:
  - SA node.
  - AV node.
  - Bundle of His.
  - Purkinje fibers.
- Stimulation of Purkinje fibers cause both ventricles to contract simultaneously.

# Conducting Tissues of the Heart

(continued)

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.





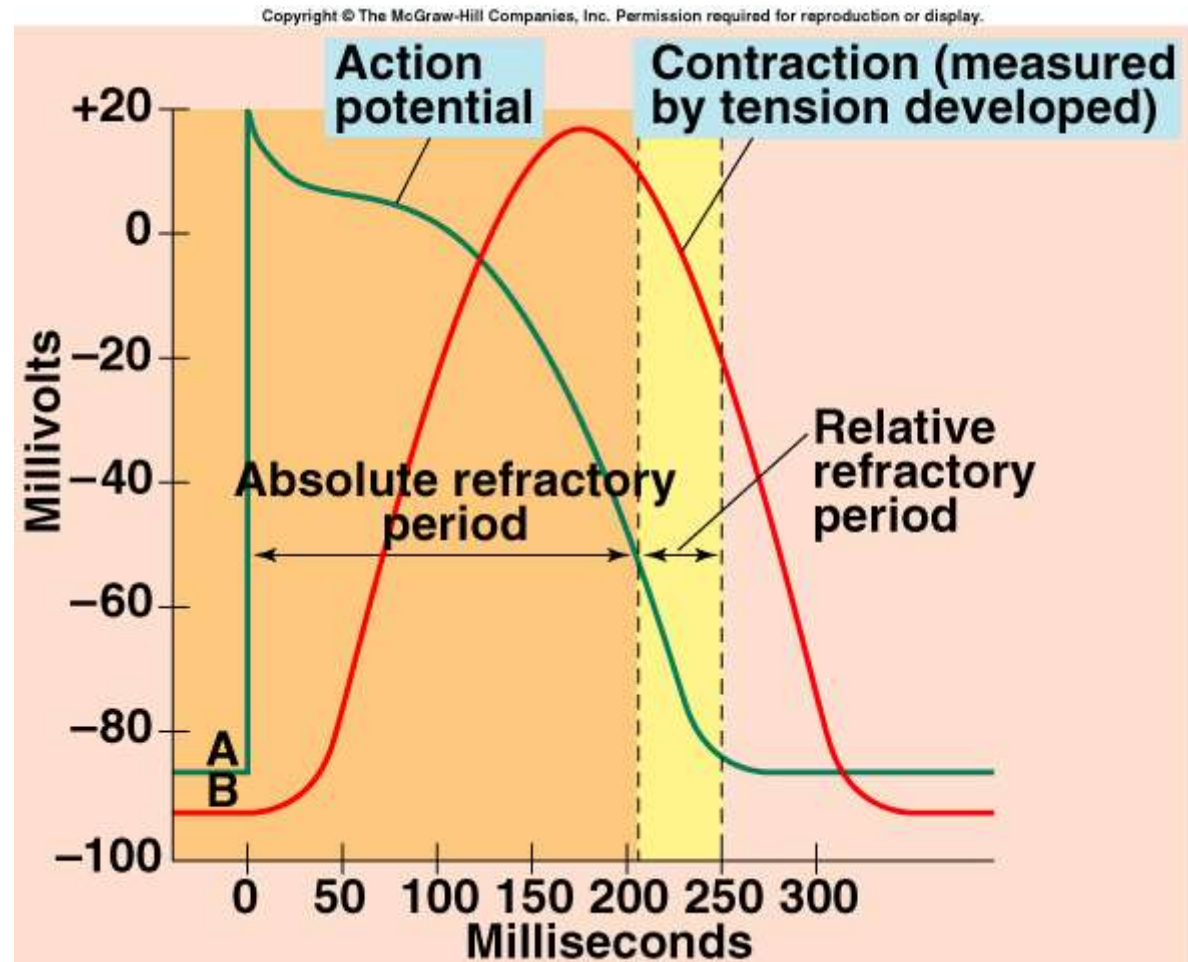
# Conduction of Impulse

---

- APs from SA node spread quickly at rate of 0.8 - 1.0 m/sec.
- Time delay occurs as impulses pass through AV node.
  - Slow conduction of 0.03 – 0.05 m/sec.
- Impulse conduction increases as spread to Purkinje fibers at a velocity of 5.0 m/sec.
  - Ventricular contraction begins 0.1–0.2 sec. after contraction of the atria.

# Refractory Periods

- Heart contracts as syncytium.
- Contraction lasts almost 300 msec.
- Refractory periods last almost as long as contraction.
- Myocardial muscle cannot be stimulated to contract again until it has relaxed.
  - Summation cannot occur.



# Excitation-Contraction Coupling in Heart Muscle



- Depolarization of myocardial cell stimulates opening of VG  $\text{Ca}^{2+}$  channels in sarcolemma.
  - $\text{Ca}^{2+}$  diffuses down gradient into cell.
    - Stimulates opening of  $\text{Ca}^{2+}$ -release channels in SR.
  - $\text{Ca}^{2+}$  binds to troponin and stimulates contraction (same mechanisms as in skeletal muscle).
- During repolarization  $\text{Ca}^{2+}$  actively transported out of the cell via a  $\text{Na}^{+}$ - $\text{Ca}^{2+}$ -exchanger.





# Electrocardiogram (ECG/EKG)

---

- The body is a good conductor of electricity.
  - Tissue fluids have a high [ions] that move in response to potential differences.
- Electrocardiogram:
  - Measure of the **electrical activity** of the heart per unit time.
    - Potential differences generated by heart are conducted to body surface where they can be recorded on electrodes on the skin.
- Does **NOT** measure the flow of blood through the heart.

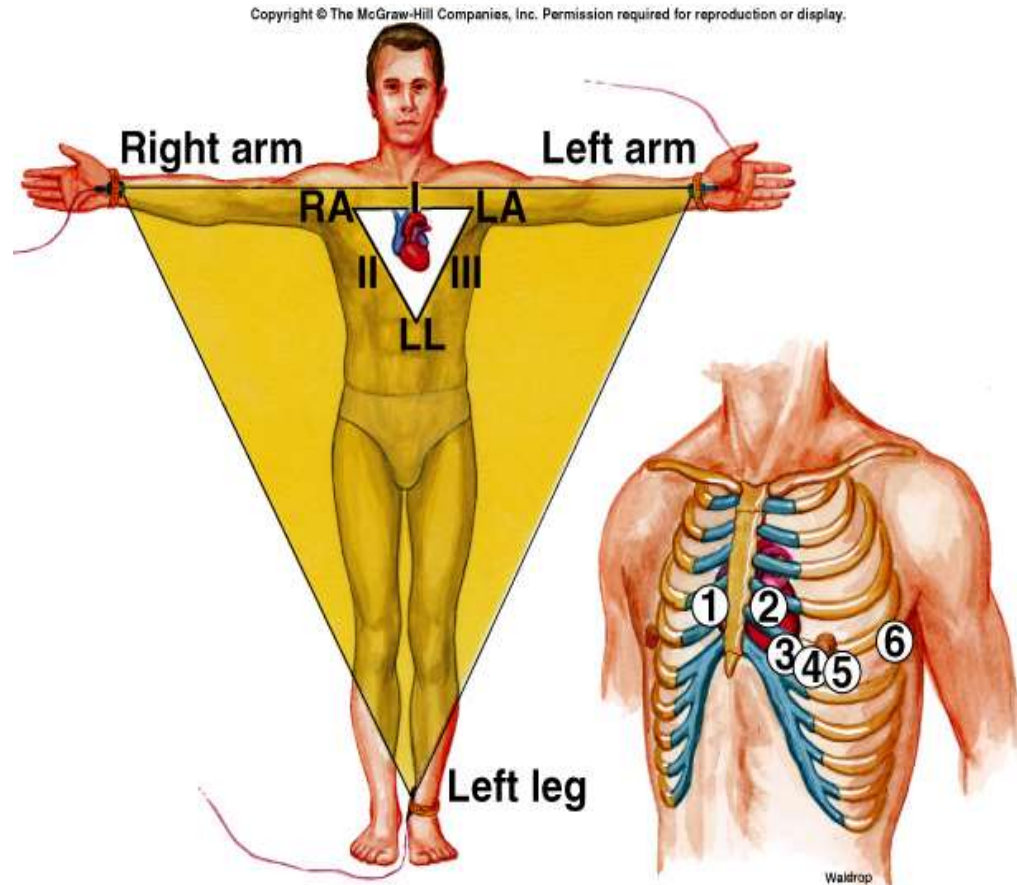
# ECG Leads

## ■ Bipolar leads:

- Record voltage between electrodes placed on wrists and legs.
- Right leg is ground.

## ■ Unipolar leads:

- Voltage is recorded between a single "exploratory electrode" placed on body and an electrode built into the electrocardiograph.
- Placed on right arm, left arm, left leg, and chest.
  - Allow to view the changing pattern of electrical activity from different perspectives.



# ECG

## ■ P wave:

- Atrial depolarization.

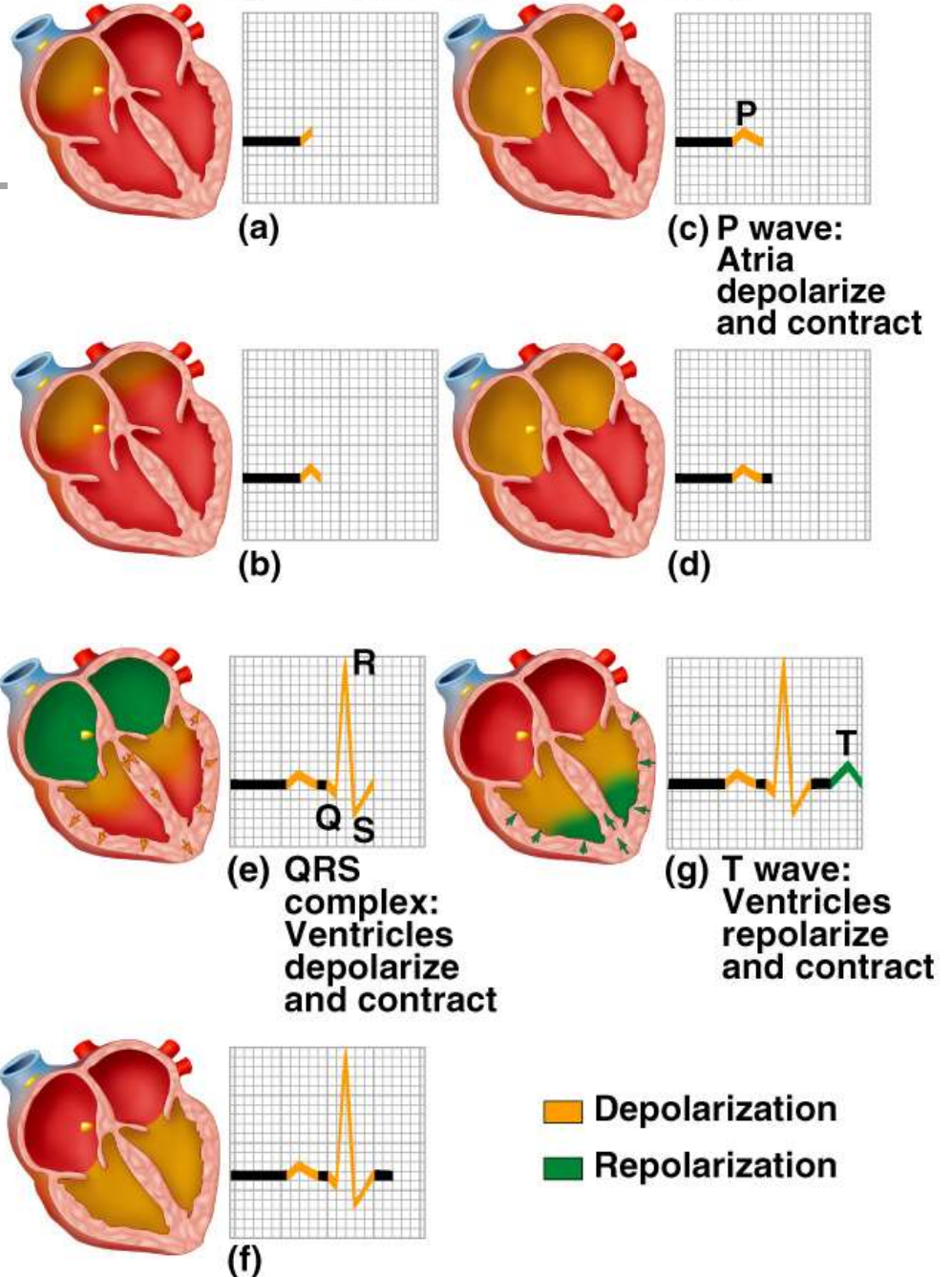
## ■ QRS complex:

- Ventricular depolarization.
- Atrial repolarization.

## ■ T wave:

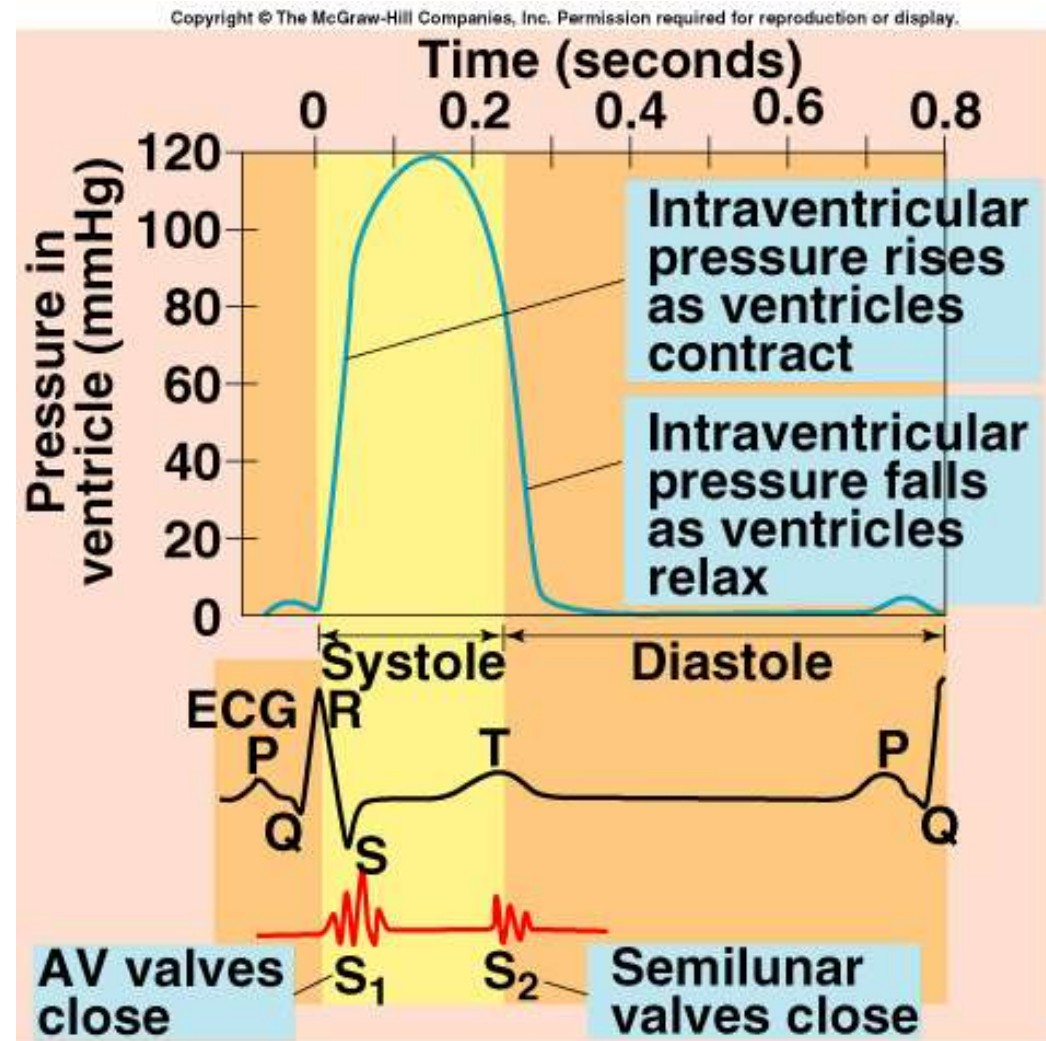
- Ventricular repolarization.

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.



# Correlation of ECG with Heart Sounds

- First heart sound:
  - Produced immediately after QRS wave.
  - Rise of intraventricular pressure causes AV valves to close.
- Second heart sound:
  - Produced after T wave begins.
  - Fall in intraventricular pressure causes semilunar valves to close.





# Systemic Circulation

---

- Arteries.
  - Arterioles.
  - Capillaries.
  - Venules.
  - Veins.
- Role is to direct the flow of blood from the heart to the capillaries, and back to the heart.



# Blood Vessels

---

- Walls composed of 3 “tunics:”
  - Tunica externa:
    - Outer layer comprised of connective tissue.
  - Tunica media:
    - Middle layer composed of smooth muscle.
  - Tunica interna:
    - Innermost simple squamous endothelium.
    - Basement membrane.
    - Layer of elastin.



# Blood Vessels (continued)

---

- Elastic arteries:
  - Numerous layers of elastin fibers between smooth muscle.
    - Expand when the pressure of the blood rises.
      - Act as recoil system when ventricles relax.
- Muscular arteries:
  - Are less elastic and have a thicker layer of smooth muscle.
  - Diameter changes slightly as BP raises and falls.
- Arterioles:
  - Contain highest % smooth muscle.
    - Greatest pressure drop.
      - Greatest resistance to flow.



# Blood Vessels (continued)

---

- Most of the blood volume is contained in the venous system.
  - Venules:
    - Formed when capillaries unite.
      - Very porous.
  - Veins:
    - Contain little smooth muscle or elastin.
      - Capacitance vessels (blood reservoirs).
    - Contain 1-way valves that ensure blood flow to the heart.
- Skeletal muscle pump and contraction of diaphragm:
  - Aid in venous blood return of blood to the heart.





# Types of Capillaries

---

- Capillaries:
  - Smallest blood vessels.
    - 1 endothelial cell thick.
      - Provide direct access to cells.
        - Permits exchange of nutrients and wastes.
  - Continuous:
    - Adjacent endothelial cells tightly joined together.
      - Intercellular channels that permit passage of molecules (other than proteins) between capillary blood and tissue fluid.
        - Muscle, lungs, and adipose tissue.
  - Fenestrated:
    - Wide intercellular pores.
      - Provides greater permeability.
        - Kidneys, endocrine glands, and intestines.
  - Discontinuous (sinusoidal):
    - Have large, leaky capillaries.
      - Liver, spleen, and bone marrow.



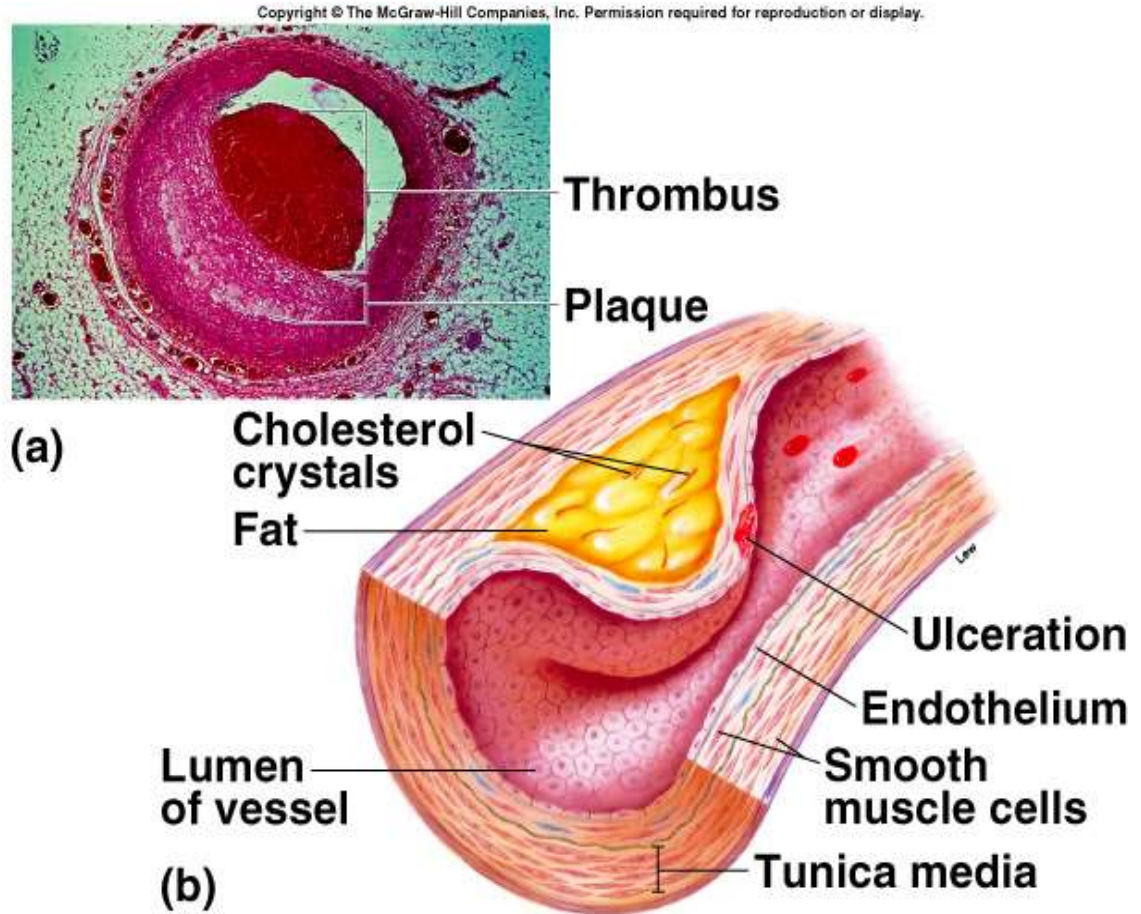
# Atherosclerosis

---

- Most common form of arteriosclerosis (hardening of the arteries).
- Mechanism of plaque production:
  - Begins as a result of damage to endothelial cell wall.
    - HTN, smoking, high cholesterol, and diabetes.
  - Cytokines are secreted by endothelium; platelets, macrophages, and lymphocytes.
    - Attract more monocytes and lymphocytes.

# Atherosclerosis (continued)

- Monocytes become macrophages.
  - Engulf lipids and transform into foam cells.
- Smooth muscle cells synthesize connective tissue proteins.
  - Smooth muscle cells migrate to tunica interna, and proliferate forming fibrous plaques.





# Cholesterol and Plasma Lipoproteins

---

- High blood cholesterol associated with risk of atherosclerosis.
- Lipids are carried in the blood attached to protein carriers.
- Cholesterol is carried to the arteries by LDLs (low-density lipoproteins).
  - LDLs are produced in the liver.
    - LDLs are small protein-coated droplets of cholesterol, neutral fat, free fatty acids, and phospholipids.

# Cholesterol and Plasma Lipoproteins (continued)

- Cells in various organs contain receptors for proteins in LDL.
  - LDL protein attaches to receptors.
    - The cell engulfs the LDL and utilizes cholesterol for different purposes.
    - LDL is oxidized and contributes to:
      - Endothelial cell injury.
      - Migration of monocytes and lymphocytes to tunica interna.
      - Conversion of monocytes to macrophages.
  - Excessive cholesterol is released from the cells.
    - Travel in the blood as HDLs (high-density lipoproteins), and removed by the liver.
      - Artery walls do not have receptors for HDL.

# Ischemic Heart Disease

## ■ Ischemia:

- Oxygen supply to tissue is deficient.
  - Most common cause is atherosclerosis of coronary arteries.
- Increased [lactic acid] produced by anaerobic respiration.

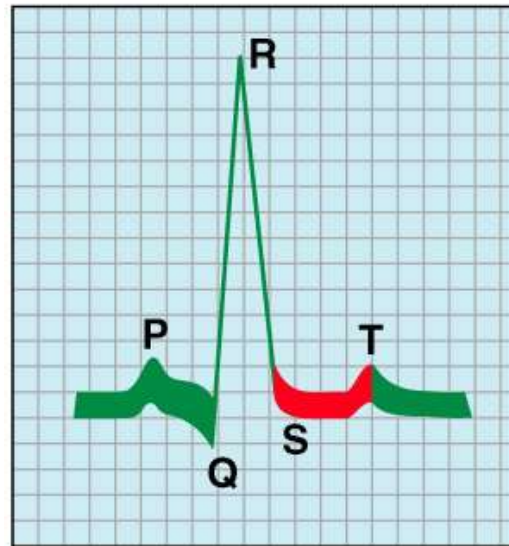
## ■ Angina pectoris:

- Substernal pain.

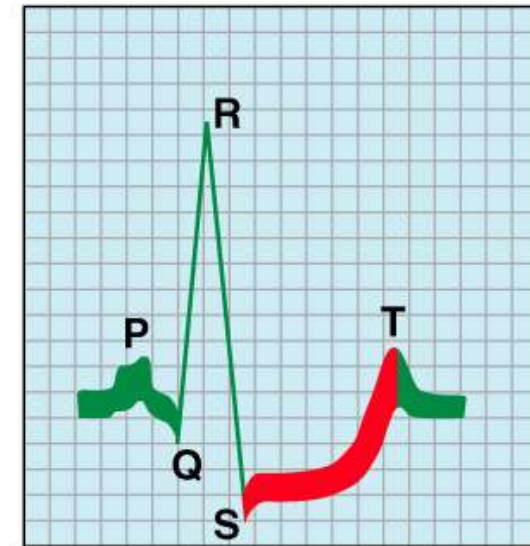
## ■ Myocardial infarction (MI):

- Changes in T segment of ECG.
- Increased CPK and LDH.

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.



Normal



Ischemia

# Arrhythmias Detected on ECG

- Arrhythmias:
  - Abnormal heart rhythms.
- Flutter:
  - Extremely rapid rates of excitation and contraction of atria or ventricles.
    - Atrial flutter degenerates into atrial fibrillation.
- Fibrillation:
  - Contractions of different groups of myocardial cells at different times.
    - Coordination of pumping impossible.
      - Ventricular fibrillation is life-threatening.

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.



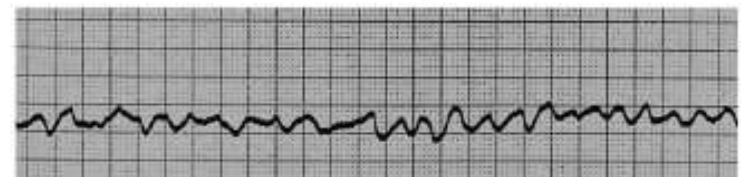
**Sinus bradycardia**



**(a) Sinus tachycardia**



**Ventricular tachycardia**



**(b) Ventricular fibrillation**



# Arrhythmias Detected on ECG

(continued)

---

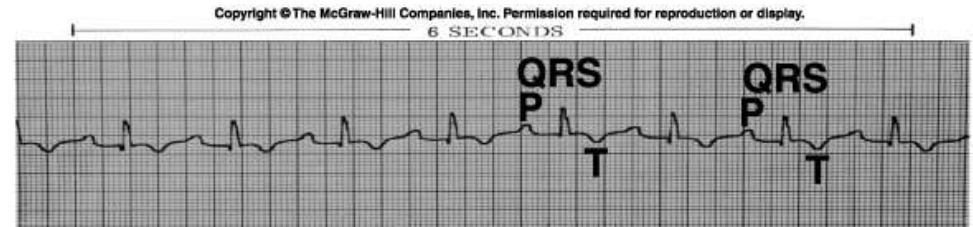
- Bradycardia:
  - HR slower < 60 beats/min.
- Tachycardia:
  - HR > 100 beats/min.
- First-degree AV nodal block:
  - Rate of impulse conduction through AV node exceeds 0.2 sec.
    - P-R interval.
- Second-degree AV nodal block:
  - AV node is damaged so that only 1 out of 2-4 atrial APs can pass to the ventricles.
    - P wave without QRS.



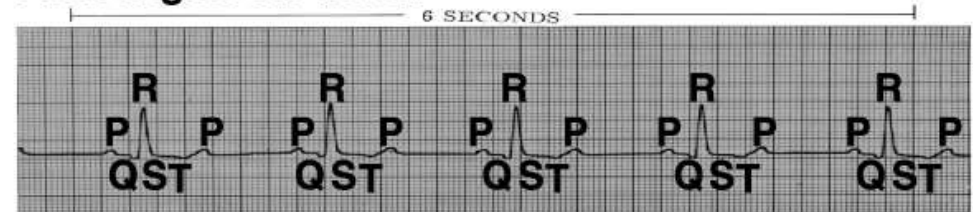
# Arrhythmias Detected on ECG

(continued)

- Third-degree (complete) AV nodal block:
  - None of the atrial waves can pass through the AV node.
  - Ventricles paced by ectopic pacemaker.



First-degree AV block



Second-degree AV block



Third-degree AV block



# Lymphatic System

---

- 3 basic functions:
  - Transports interstitial (tissue) fluid back to the blood.
  - Transports absorbed fat from small intestine to the blood.
  - Helps provide immunological defenses against pathogens.

# Lymphatic System (continued)

## Lymphatic capillaries:

- Closed-end tubules that form vast networks in intercellular spaces.

## Lymph:

- Fluid that enters the lymphatic capillaries.
  - Lymph carried from lymph capillaries, to lymph ducts, and then to lymph nodes.
- Lymph nodes filter the lymph before returning it to the veins.

