

Chapter 9

Musculoskeletal System Disorders

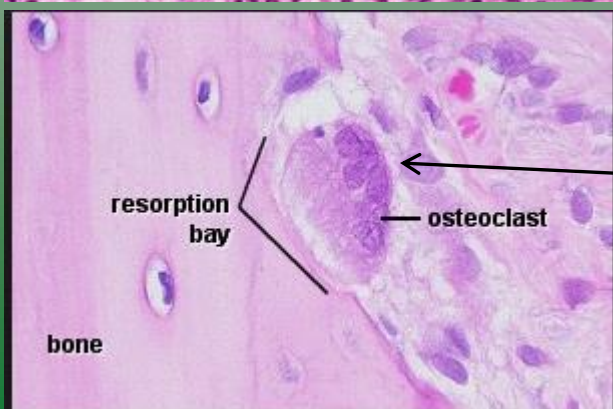
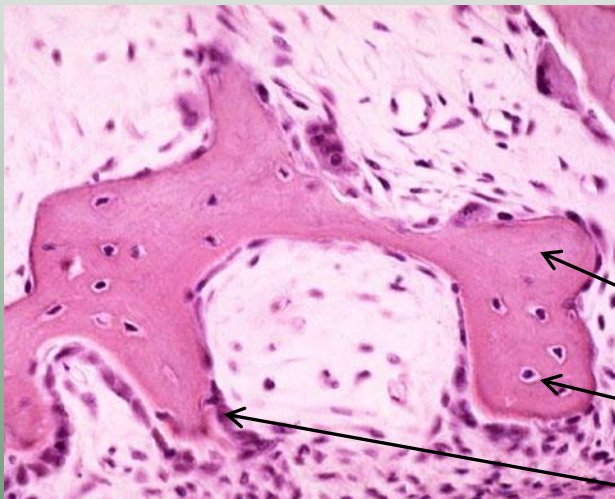
Review of Musculoskeletal System: Bones

- Classified by shape

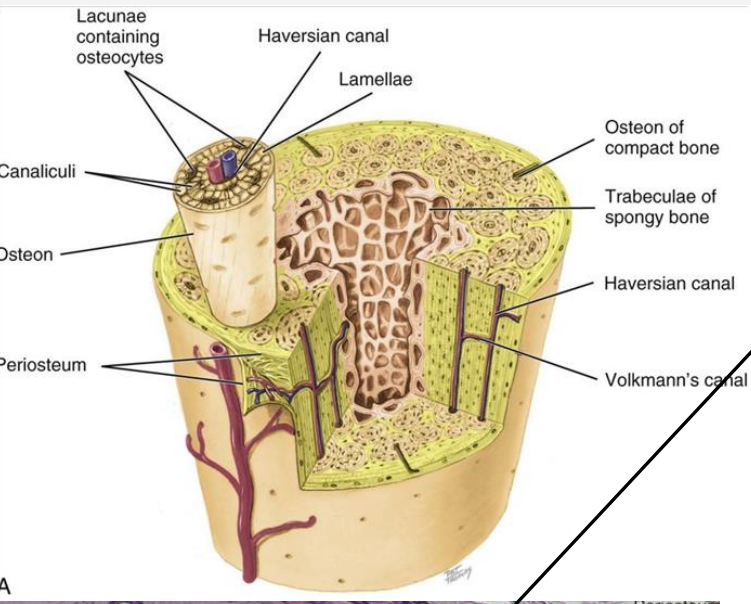
- Long bones
- Short bones
- Flat bones
- Irregular bones

- Bone tissue consists of:

- Matrix
- Mature bone cells (osteocytes)
- Bone-producing cells (osteoblasts)
- Bone-resorbing cells (osteoclasts)



Bone

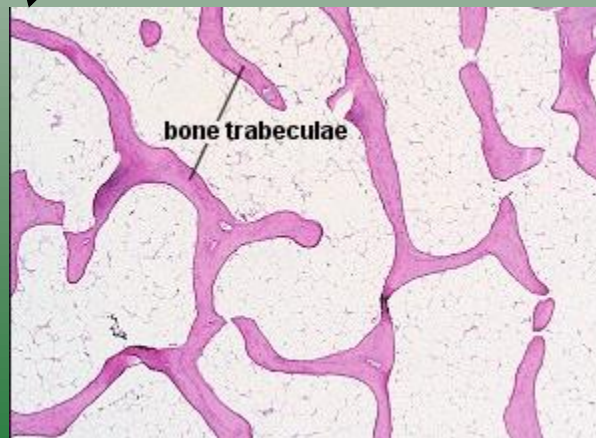
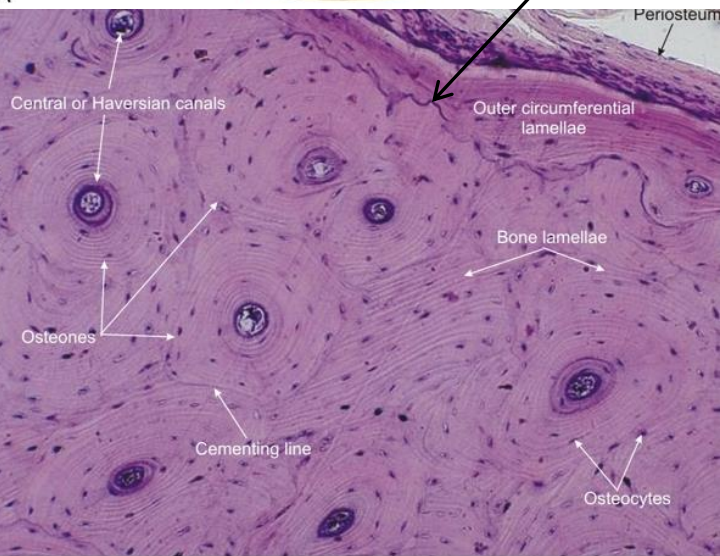


- Types of bone tissue

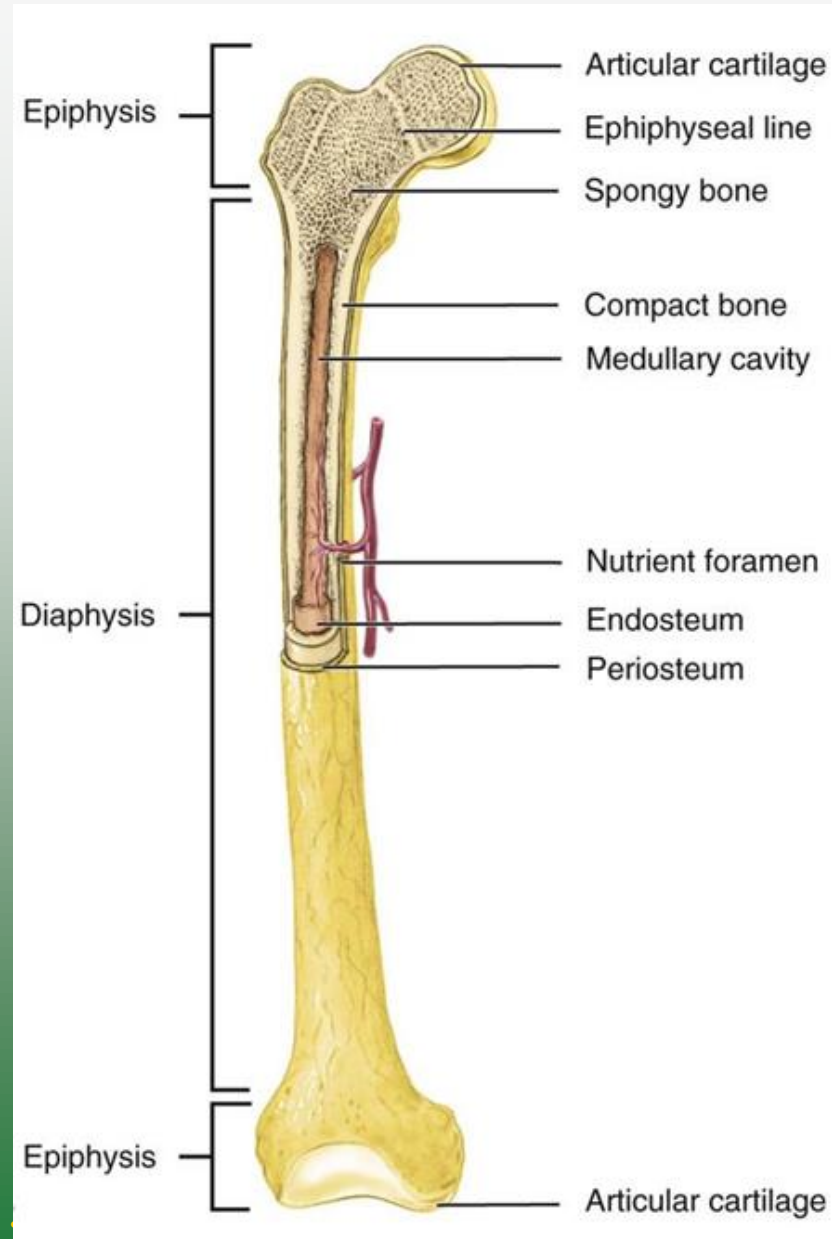
- Compact—outer covering of bone
- Cancellous (spongy)—interior of bone

- Other bone structures

- Periosteum—connective tissue covering over the bone
- Endosteum—osteoblast-rich lining of medullary cavity



Structure of a Long Bone

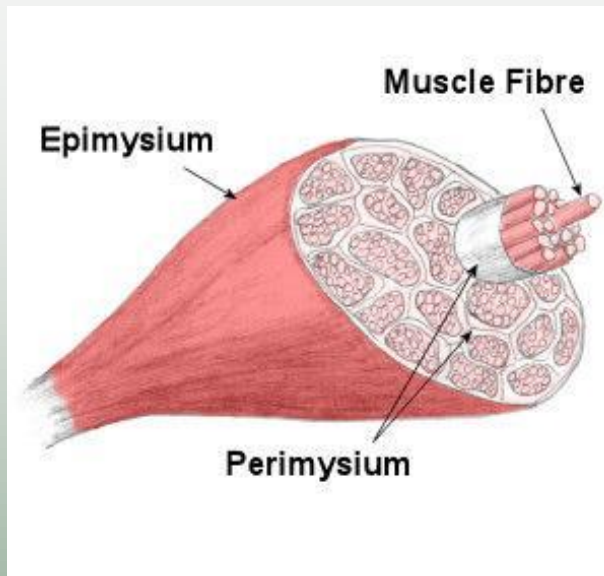


Skeletal Muscle

Functions of Skeletal Muscle

- Facilitate body movement.
- Maintain body position.
- Stabilize joints.
- Produce heat.

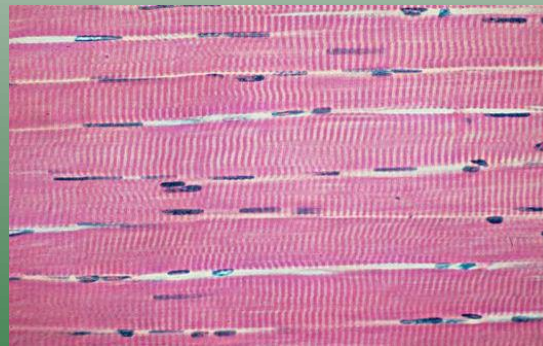
Connective Tissue Coverings of Skeletal Muscle



- **Epimysium**
 - Connective tissue surrounding the entire muscle
- **Perimysium**
 - Connective tissue surrounding muscle fascicles
- **Endomysium**
 - Connective tissue around individual muscle fibers

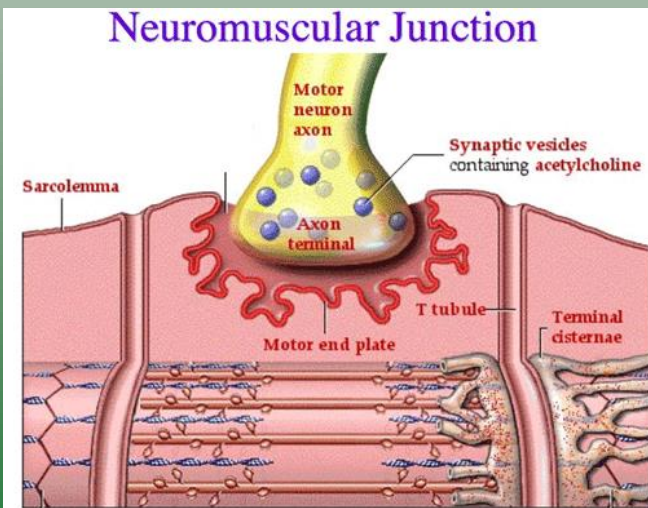
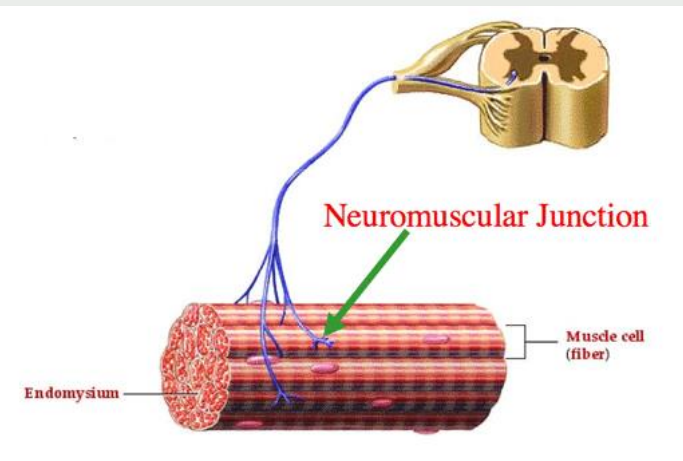


Cross sections



Long sections, shows striations

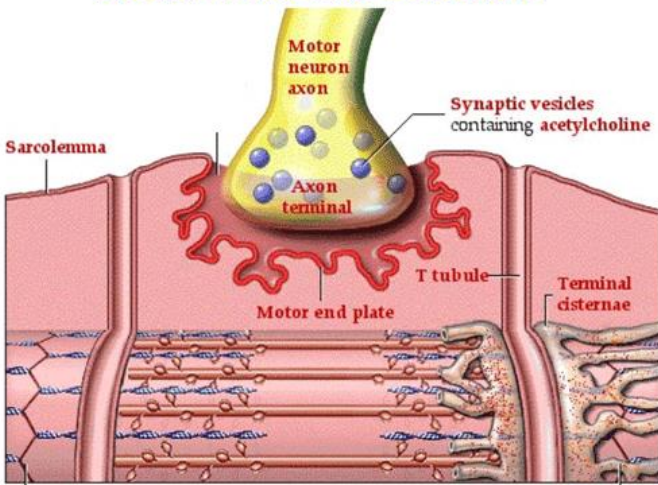
Skeletal Muscle (Cont.)



- Motor unit
 - Motor neuron in the spinal cord and all muscle fibers innervated by the neurons
- Neuromuscular junction
 - Synapse between the motor neuron nerve fiber and muscle fiber
 - Neurotransmitter—acetylcholine (ACh)
 - Breakdown of ACh by cholinesterase; anticholinesterases will interfere with the breakdown of ACh.

Neuromuscular Transmission and Muscle Contraction

Neuromuscular Junction

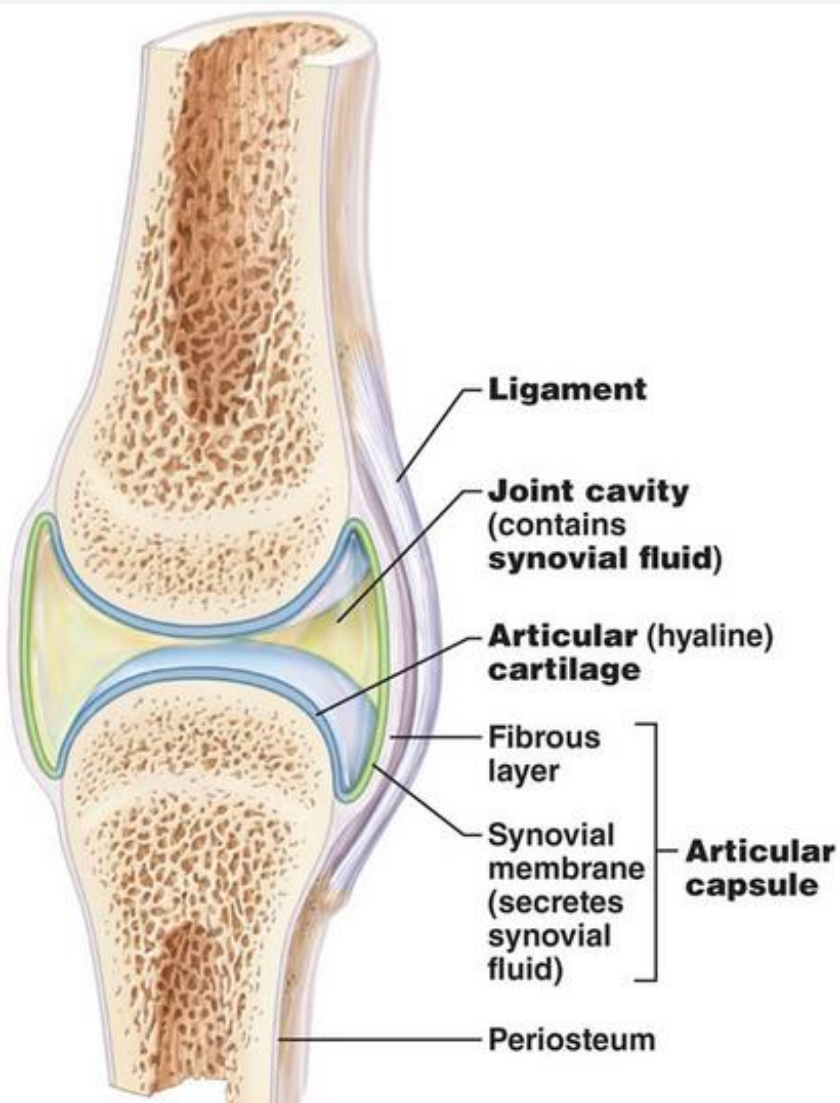


- Arrival of the action potential
- Depolarization of the presynaptic terminal
- Calcium influx
- Exocytosis of the neurotransmitter (Ach)
- Diffusion of neurotransmitter to postsynaptic receptors—attachment
- Generation of muscle action potential
- Release of calcium from sarcoplasmic reticulum
- Power stroke—contraction of muscle fiber

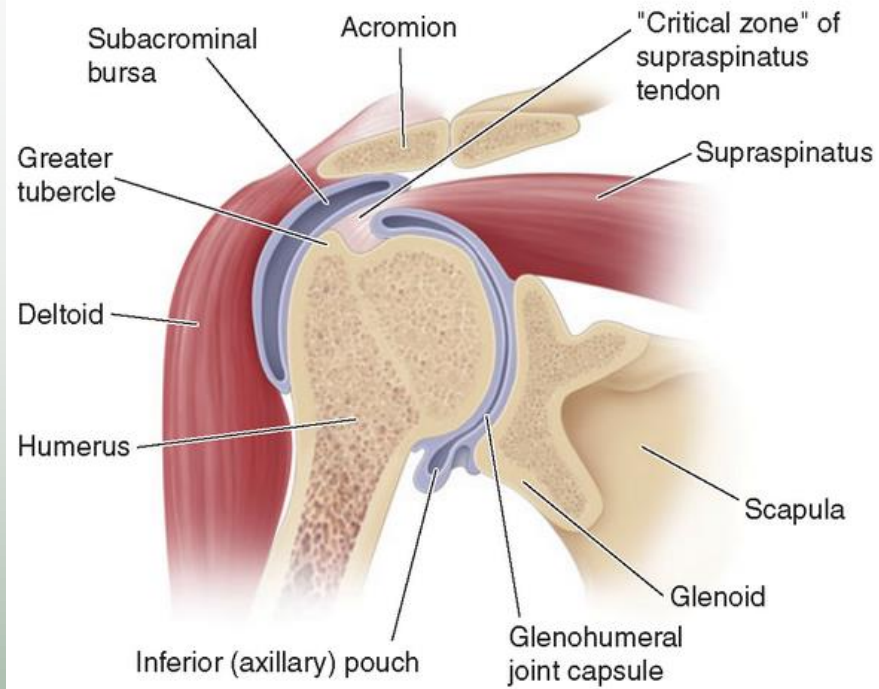
Joints

- Classified by degree of movement:
 - Synarthroses—immovable (e.g., cranial sutures)
 - Amphiarthroses—slightly movable (e.g., ribs to sternum)
 - Diarthroses (synovial)—freely movable (e.g., shoulder)

Structure of Synovial Joint



- Articular cartilage—covering over ends of bones
- Synovial membrane—produces synovial fluid that fills space between ends of bones
- Articular capsule—consists of synovial membrane, a fibrous capsule
- Ligaments—reinforces capsule, links bones, supports joint
- Menisci—lateral pads in some joints to stabilize
- Bursae—fluid-filled sacs to add extra cushion



Musculoskeletal Diagnostic Tests

- For bone disorders: radiography, bone scanning
- For muscular disorders: electromyography (EMG), biopsy
- For joint disorders: radiography, arthroscopy, magnetic resonance imaging (MRI), examination of synovial fluid

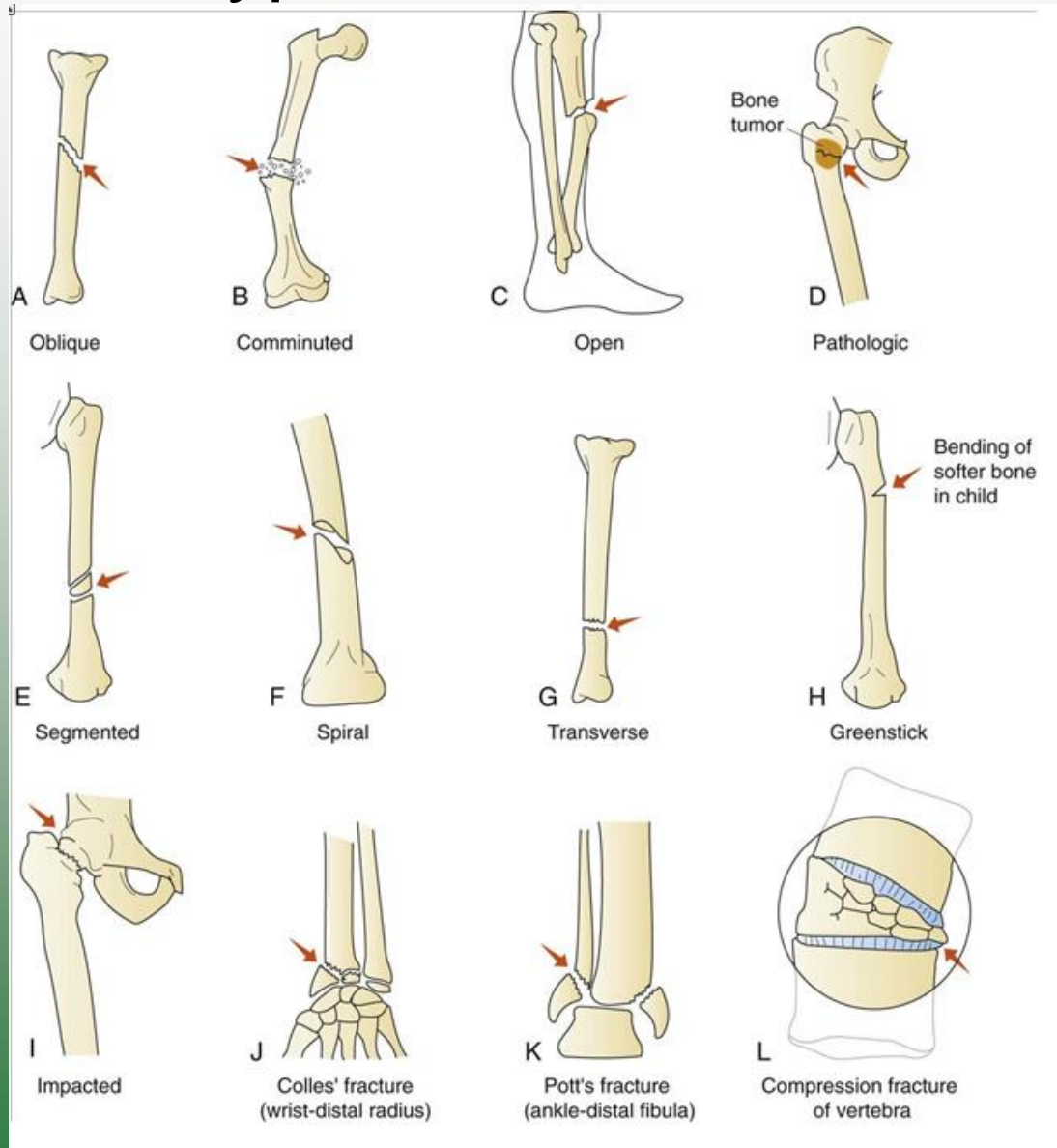
Fracture

- A fracture is a break in the integrity of a bone.
- Fractures occur because of trauma, neoplasms, or increased stress on bones.



Courtesy of Dr. Mercer Rang, The Hospital for Sick Children, Toronto, Ontario, Canada.

Types of Fractures



Classification of Fractures

- Complete: bone broken, forming separate pieces
- Incomplete: bone only partially broken
- Open (compound): skin broken
- Closed: skin not broken
- Simple: single break, maintaining alignment and position
- Comminuted: multiple fractures and bone fragments
- Compression: bone crushed or collapsed into small pieces

Trauma—Fractures

- Other types:
 - Impacted—one end forced into adjacent bone
 - Pathologic—results from weakness; occurs with little stress
 - Stress—fatigue fractures
 - Depressed—skull fractured and forced into brain

Pathophysiology of Bone Fracture

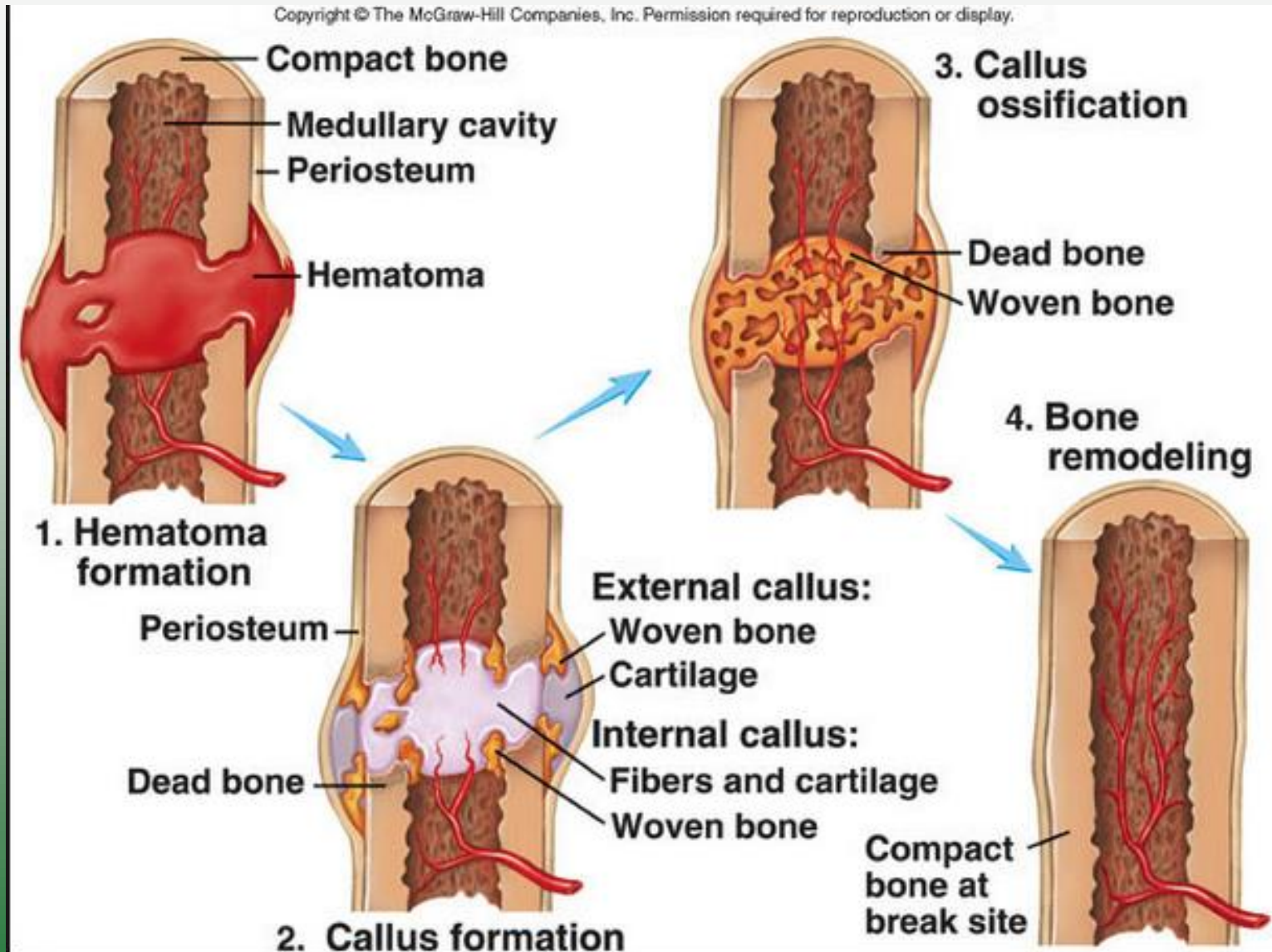
- Events at a fracture site.
 - **Bleeding**
 - Edema causes stretching of periosteum (if it is intact) and swelling of soft tissues → severe pain
 - Release of bradykinin and other chemical mediators also contributes to pain
 - **Clot forms at fracture site**
 - Systemic signs of inflammation may occur.

Healing of Bone Fracture

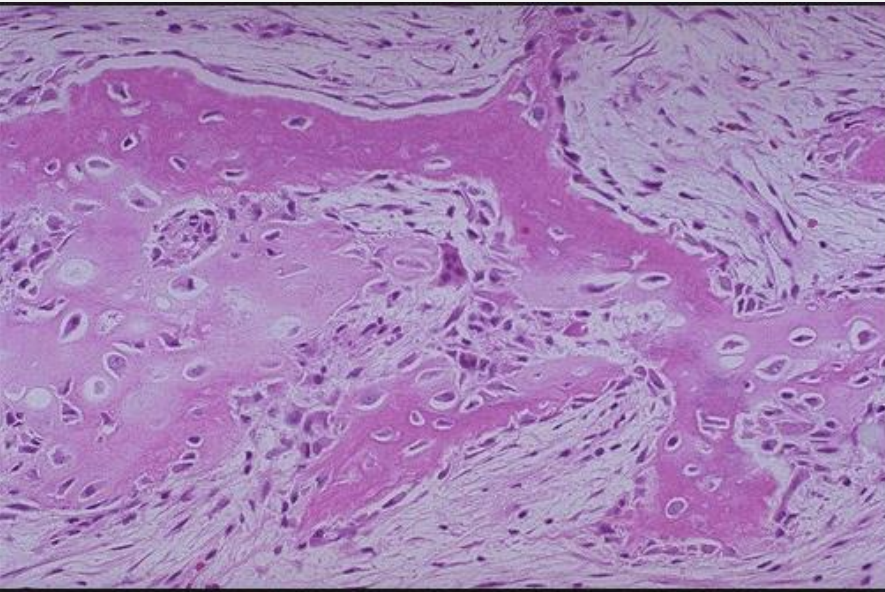
- **Hematoma**—fibrin network is formed.
- Phagocytic cells remove debris, **granulation tissue** forms.
- Fibroblasts lay down new collagen fibers.
- Chondroblasts form new cartilage.
- Formation of procallus or **soft callus of fibrous tissue and cartilage**.
- Osteoblasts generate new bone.
- Procallus is replaced by **bony or hard callus**.
- **Remodeling** of bone, with return to use

Healing of Fracture

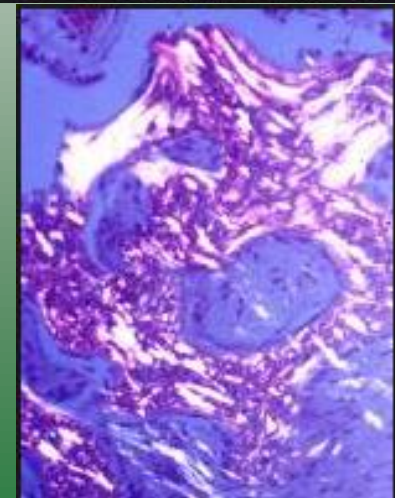
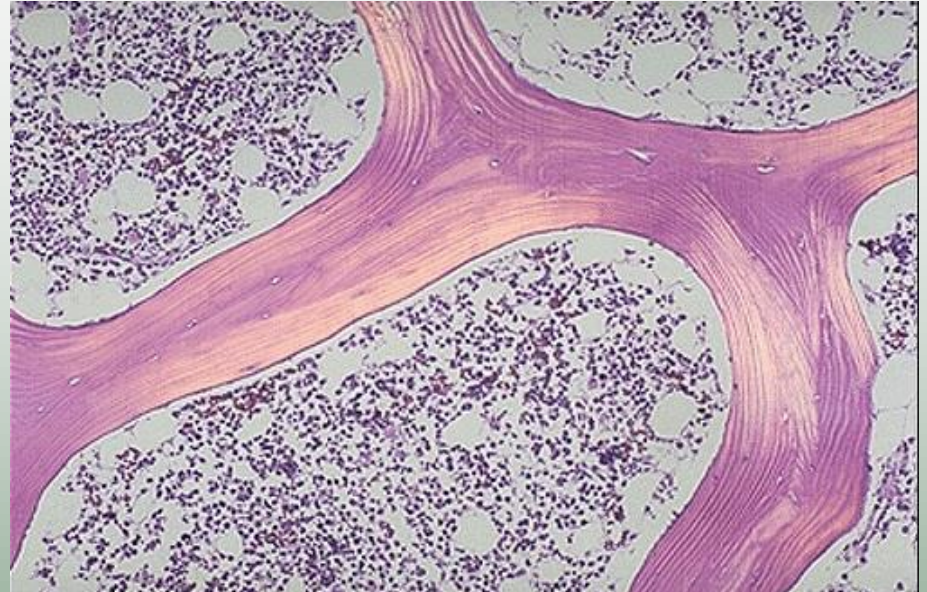
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Woven Bone



Lamellar Bone



Factors Affecting Bone Healing

- Amount of local damage
- Proximity of bone ends
- Presence of foreign material or infection
- Blood supply to fracture site
- Systemic factors, such as age, nutrition, anemia

Complications

- Muscle spasms
- Infections
- Ischemia
- Fat emboli
- Nerve damage
- Failure to heal or development of deformity during healing
- Residual effects such as osteoarthritis

Treatments of Fractures

- Closed reduction—pressure to restore bone position (splints and casts).
- Open reduction—surgery to align and/or Insert pins, screw rods, or plates to align



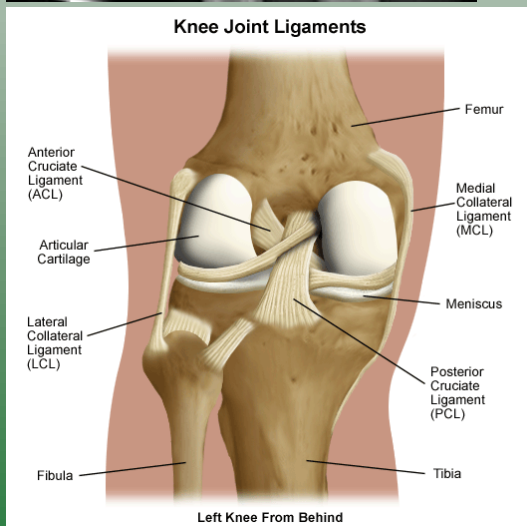
Compartment Syndrome

- Edema within one area of compartment of the limb that is between layers of dense connective tissue (fascia)
- Ischemia and infarction of tissue may occur because of compression of arterial blood supply.
- Dead tissue may become gangrenous, requiring amputation.
- ***A tight cast can cause compartment syndrome!***

Dislocations



- Separation of two bones at a joint, with loss of contact between articulating surfaces
- Usually accompanied by significant soft tissue damage to ligaments and tendons
- Distortion of joint usually evident
- May recur repeatedly, requiring surgery



Injuries

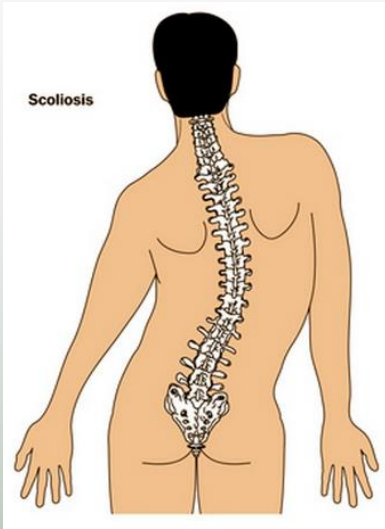
- Sprain—a tear in a ligament
- Strain—a tear in a tendon
- Avulsion—ligaments or tendons completely separated from bony attachments
- Immobilization often used to prevent tissue damage and promote healing
- Overuse injuries
 - Muscle tears

Bone Disorders

Bone Disorders—Osteoporosis

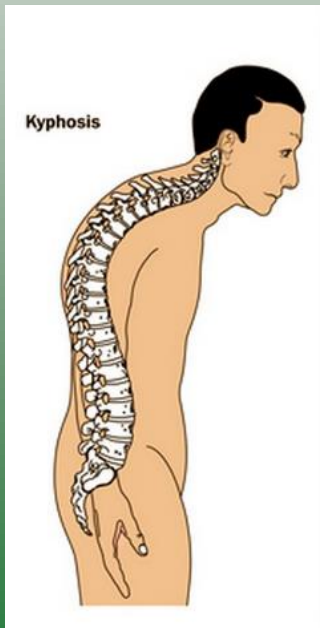
- Decrease in bone mass and density
- Occurs in two forms:
 - Primary
 - Idiopathic
 - Age 50+ years
 - Decreased sex hormones
 - Decreased calcium intake
 - Secondary
 - As a complication of another disorder

Osteoporosis



- Pathophysiology

- Bone resorption exceeds formation.
- Results in loss of compact bone
- Diagnosed with bone density scans
- Can cause compression fractures of vertebrae, wrist, or hip
- Can lead to kyphosis and scoliosis



Osteoporosis (Cont.)

- Predisposing factors
 - Age 50+ years
 - Decreased mobility or sedentary lifestyle
 - Hormonal factors
 - Excess corticosteroids or parathyroid hormone (PTH)
 - Deficit of estrogen or testosterone
 - Deficits of calcium, vitamin D, or protein
 - Cigarette smoking
 - Lower BMI
 - Asian or European ancestry
 - Excessive caffeine intake

Osteoporosis (Cont.)

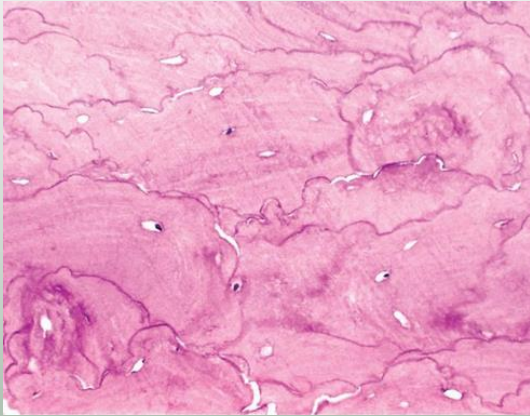
- Treatment

- Dietary supplements
- Weight-bearing exercise
- Physiotherapy to reduce pain and maintain function
- Bisphosphonates, inhibits osteoclasts.
- Calcitonin

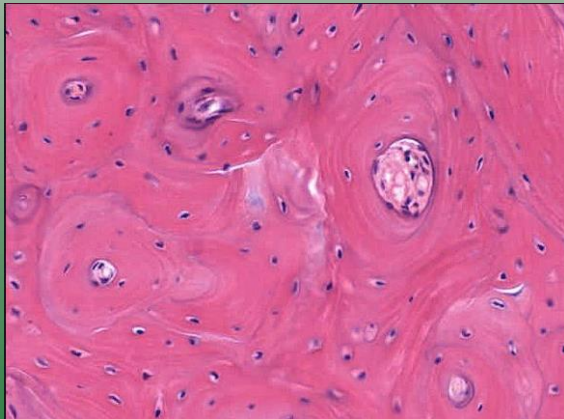
Rickets and Osteomalacia

- Result from deficit of vitamin D and phosphates
- Causes—dietary deficits, malabsorption, intake of phenobarbital, lack of sun exposure
- In children, leads to weak bones and other skeletal deformities
- In adults, may lead to soft bones, resulting in compression fractures

Paget's Disease



Paget's (above): Odd pattern of cement lines, normal below.

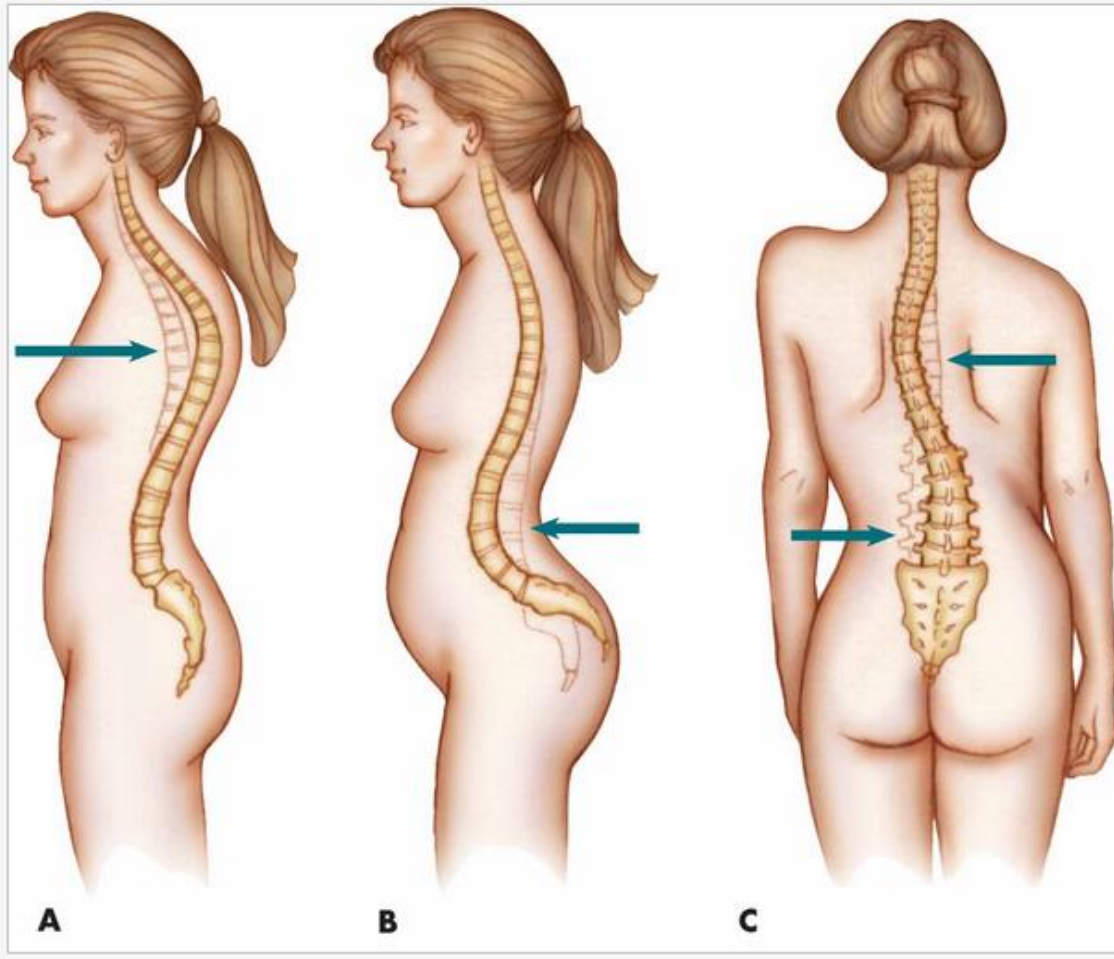


- Occurs in adults older than 40 years
- Exact cause has not yet been established.
- Bone destruction replaced by fibrous tissue
- Pathologic fractures are common.
- In vertebrae—can result in compression fractures and kyphosis
- In skull—increased pressure resulting in headaches and compression of cranial nerves → severe pain

Osteomyelitis

- Bone infection caused by bacteria or fungi
- Signs and symptoms
 - Local inflammation and bone pain
 - Fever and excessive sweating
 - Chills
 - General malaise
- Treatment
 - Antibiotics
 - Surgery may be required.

Abnormal Curvatures of the Spine



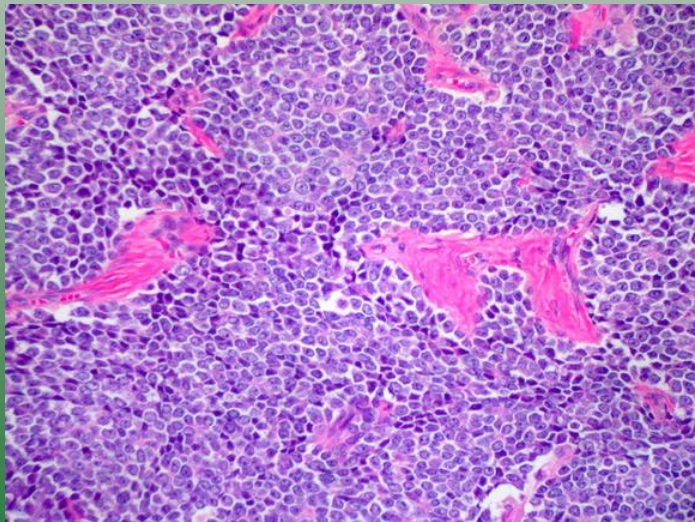
- **A. Kyphosis**
 - Hunchback or humpback—abnormally rounded upper back
- **B. Lordosis**
 - Swayback—curving inward at the lower back
- **C. Scoliosis**
 - S- or C-shaped—sideways curve to the spine

Tumors

- Common site of metastatic tumors from breast, lung, or prostate tumors
- Most primary tumors termed *sarcomas* are malignant.
- Osteosarcoma—most common primary neoplasm of bone
 - Occurs in the shaft of long bones of the leg
 - Common in children, adolescents, and young adults
 - Bone pain at rest is a warning sign for this cancer!

Tumors (Cont.)

- Chondrosarcomas—arise from cartilage cells; more common in adults
- Ewing's sarcoma is common in adolescents and usually occurs in the shaft (in marrow cavity) of long bones. Also known as PNET – Primitive Neuroectodermal Tumor.



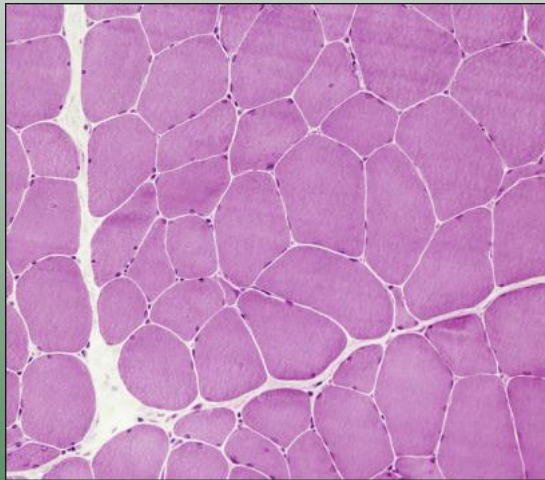
Small primitive cells with no bone or cartilage formation.

Bone Tumors

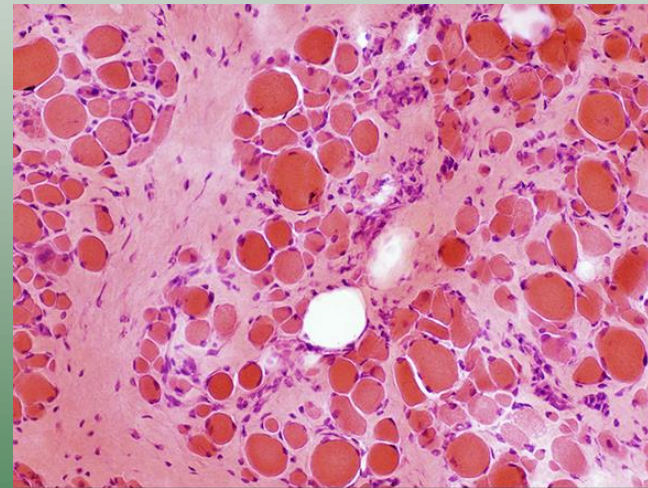
- Tumors metastasize to lungs early in the course of the disease.
- Treatment
 - Excision of tumor if possible
 - Surgical amputation if excision is not feasible
 - Chemotherapy to reduce metastasis

Muscular Dystrophy (MD)

- Group of autosomal recessive disorders
- Degeneration of skeletal muscle over time
- Duchenne's MD or pseudohypertrophic MD most common type; affects young boys



Normal Muscle: Uniform fiber size and tightly packed.

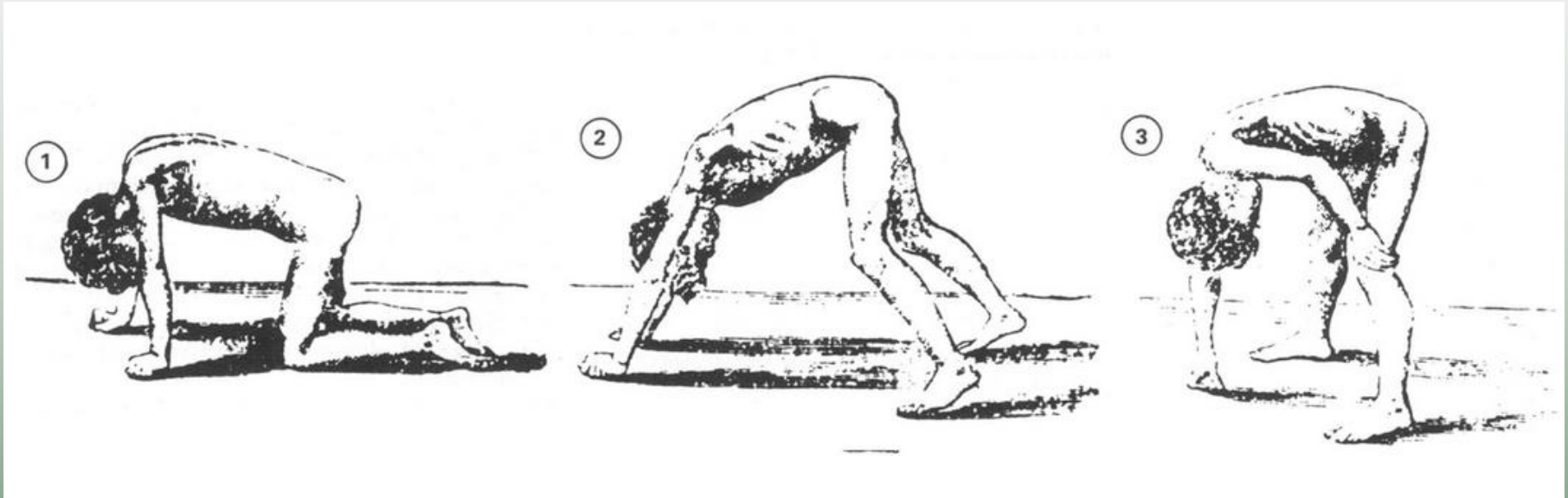


Muscular Dystrophy: Widely variable fiber size and fibers separated by fibrous tissue.

Muscular Dystrophy (MD) (Cont.)

- Signs and symptoms
 - With Duchenne's MD—early motor weakness
 - Weakness in pelvic girdle—waddling gait, difficulty climbing stairs
 - Gower maneuver—pushing up to erect position (using arms to stand up due to weak legs).
 - Tendon reflexes reduced
 - Deformities develop, such as kyphoscoliosis
 - Respiratory infections common
 - Cardiac myopathy occurs commonly.

Gower's Sign



Muscular Dystrophy (MD) (Cont.)

- Diagnostic tests
 - Identification of common genetic abnormalities
 - Elevated creatine kinase levels
 - Electromyography
 - Muscle biopsy
 - Genetic Tests

Muscular Dystrophy (MD) (Cont.)

- Treatment

- No curative treatment available
- Moderate exercise to maintain motor function
- Supportive appliances
- Physiotherapy and occupational therapy to maximize function and adaptation
- Massage—reduces pain and stiffness
- Ventilator—in case of respiratory failure
- Research being done on genetic therapies

Fibromyalgia

- Syndrome characterized by:
 - Pain in soft tissues
 - Stiffness affecting muscles, tendons, and soft tissue
 - No obvious inflammation or atrophy
 - Sleep disturbance and severe fatigue
 - Anxiety and/or depression may be present.
- Cause is not known but hypothesized to be imbalance in serotonin and other neurotransmitters or increased production of substance P

Fibromyalgia (Cont.)

- Predisposing and aggravating factors
 - Onset is higher in women age 20 to 50 years.
 - History of physical or psychological trauma or chronic pain
 - Sleep deprivation
 - Stress
 - Fatigue

Fibromyalgia (Cont.)

- Treatment

- Stress avoidance or reduction
- Regular exercise in the morning
- Pace activity and rest as needed.
- Applications of heat or massage
- Analgesic drugs
- Low doses of antidepressants
- NSAIDs
- New drugs—pregabalin (Lyrica)

Joint Disorders

Osteoarthritis

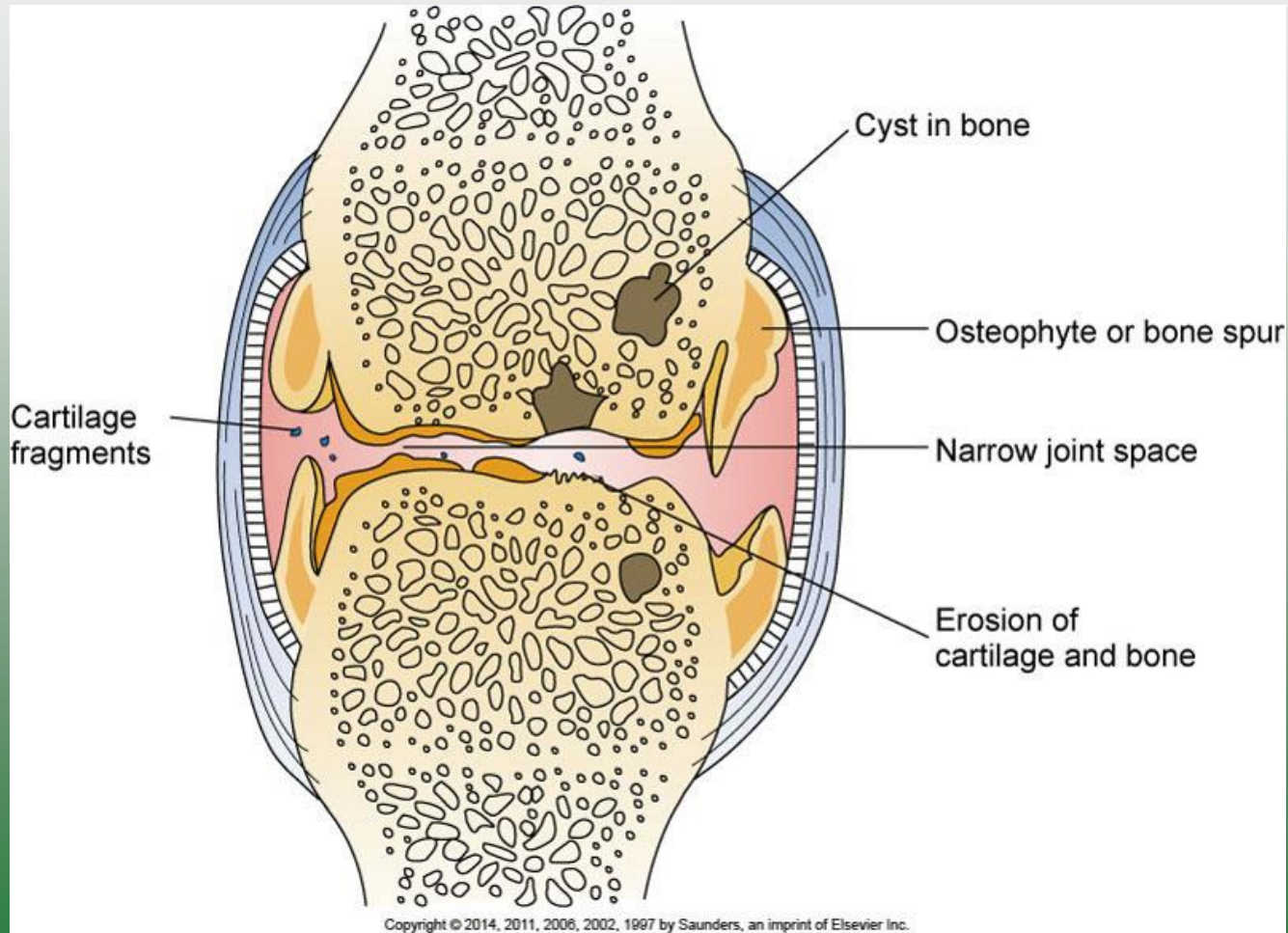
- Degenerative—wear and tear joint disease
- May be the result of increased weight-bearing or lifting
- Incidence increasing
- Genetic component identified in research with mice

Osteoarthritis (Cont.)

- Pathophysiology

- Articular cartilage is damaged.
- Surface of cartilage becomes rough and worn
- Tissue damage causes release of enzymes, accelerating disintegration of cartilage.
- Subchondral bone may be exposed.
- Osteophytes and cartilage break off.
- Joint space becomes narrower.
- Secondary inflammation of surrounding tissue
- Loss of normal range of joint motion
- Pain with weight-bearing and use

Pathologic Changes with Osteoarthritis



Osteoarthritis (Cont.)

- Causes

- Primary form—weight-bearing, obesity, aging
- Secondary form—follows trauma or repetitive use
- Genetic factors thought to play a role
- Weight-bearing joints most frequently affected but finger joints also involved

Osteoarthritis (Cont.)

- Signs and symptoms
 - Aching **pain with weight-bearing** and movement
 - **Pain gets worse as day goes on.**
 - Joint movement is limited.
 - Recreational and social activities become limited because of pain.
 - Walking is difficult.
 - Predisposition to falls
 - In temporomandibular joint (TMJ) syndrome, mastication and speaking are difficult.
 - Bony enlargement of distal interphalangeal joints

Osteoarthritis (Cont.)

- Treatment

- Minimize stress on joint by use of adaptive devices such as a cane
- Pacing activity
- Mild exercise program to maintain fitness and joint function
- Supports such as hand brace to facilitate movement
- Orthotic Inserts in shoes

Osteoarthritis (Cont.)

- Treatment (Cont.)
 - Massage therapy
 - Physiotherapy
 - Acupuncture
 - Occupational therapy
 - **Glucosamine chondroitin supplements**
 - Injection of synthetic synovial fluid (hyaluronic acid)
 - **NSAIDs**
 - **Analgesics**
 - Arthrotomy to stabilize joint
 - **Surgical joint replacement**

Rheumatoid Arthritis

- Considered an **autoimmune disorder**
- Causes chronic **systemic** inflammatory disease
- Higher incidence in women than in men
- Affects **all age groups**
- **Morning stiffness, gets better as day goes on.**

Rheumatoid Arthritis (Cont.)

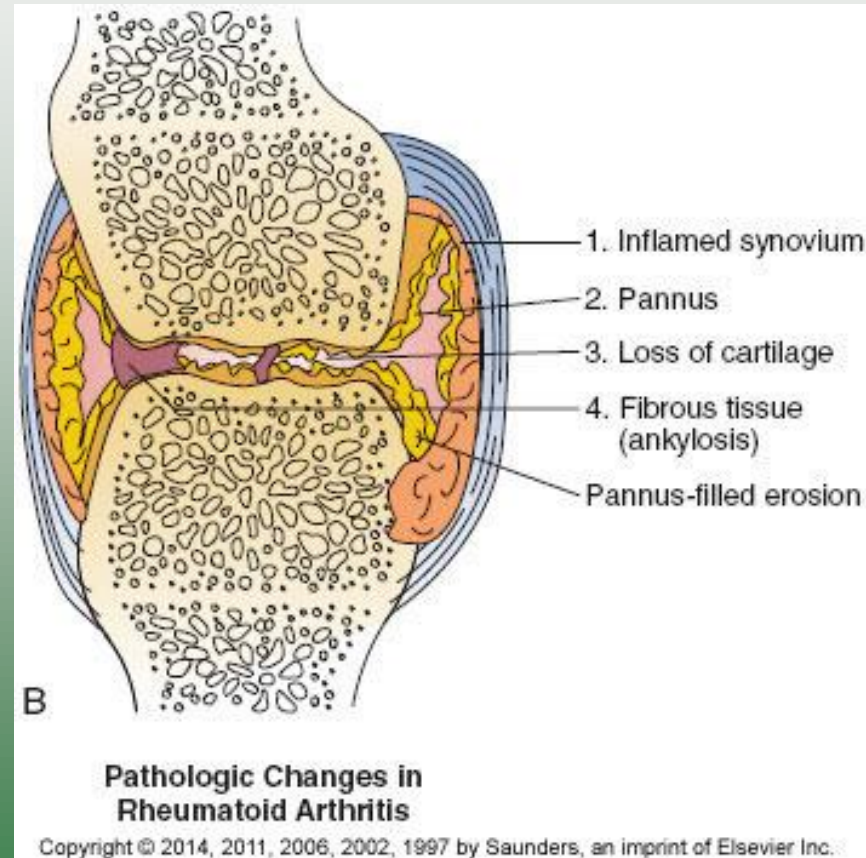
- Pathophysiology

- **Synovitis**—marked inflammation, cell proliferation
- Pannus formation—granulation tissue spreads.
- **Cartilage erosion**—creates unstable joint
- **Fibrosis**—calcifies and obliterates joint space
- Ankylosis—joint fixation and deformity develop if untreated.

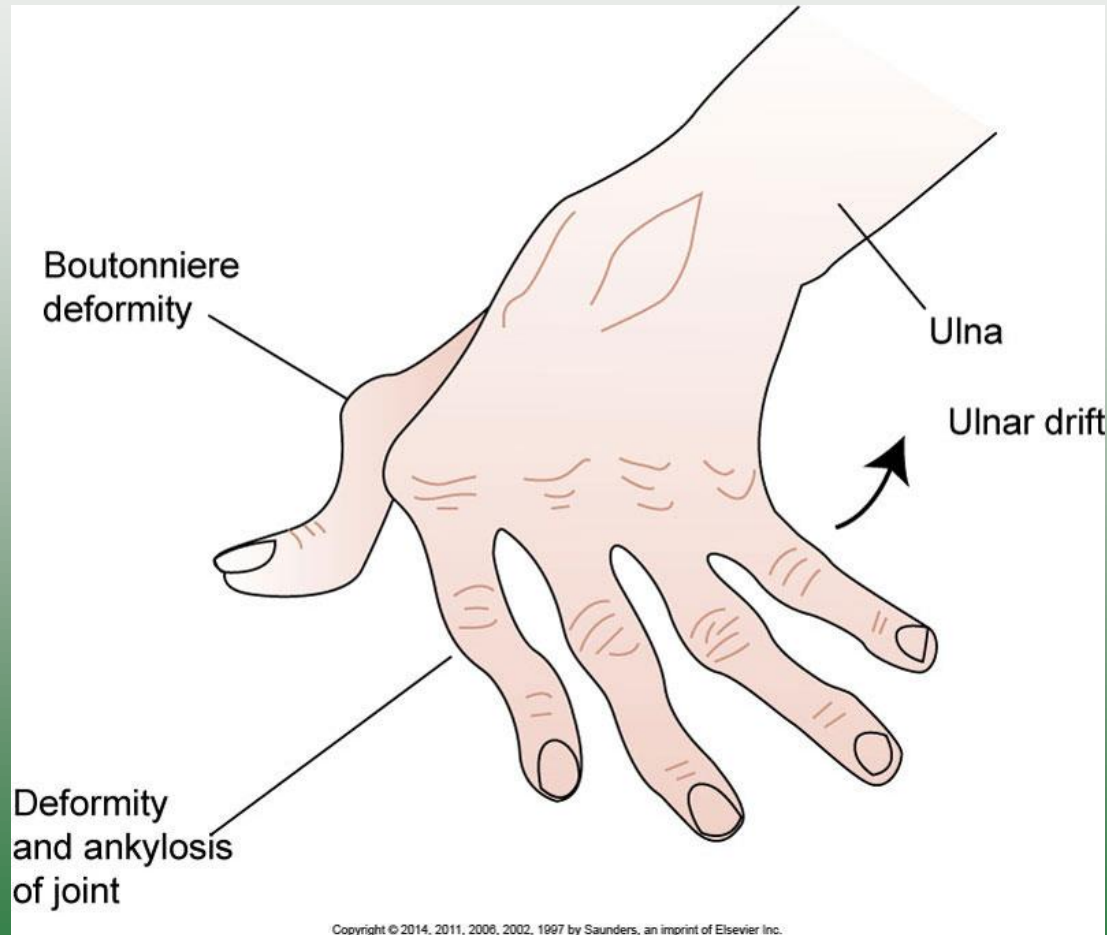
Rheumatoid Arthritis (Cont.)

- Other changes frequently occur around the joint.
 - Atrophy of muscles
 - Bone alignment shifts
 - Muscle spasms caused by inflammation and pain
 - Contractures and deformity develop.

Rheumatoid Arthritis (Cont.)



Typical Deformity in a Hand with Rheumatoid Arthritis



Joint Disorders: Rheumatoid Arthritis (Cont.)

- Systemic effects
 - Marked fatigue
 - Depression
 - Malaise
 - Anorexia
 - Low-grade fever
 - Iron deficiency anemia that is resistant to iron therapy

Rheumatoid Arthritis (Cont.)

- Cause

- Exact cause not known
- Genetic factor is present.
- Familial predisposition
- Some links to viral infections

Rheumatoid Arthritis (Cont.)

- Signs and symptoms
 - Affected joints are extremely painful.
 - Stiffness of joints
 - Redness and swelling of joints
 - Joint involvement includes small joints and is often bilateral.
 - Joint movement impaired
 - Eventually, the joint becomes fixed and deformed.

Rheumatoid Arthritis (Cont.)

- Systemic signs
 - Fatigue
 - Anorexia
 - Mild fever
 - Generalized lymphadenopathy
 - Generalized aching

Rheumatoid Arthritis (Cont.)

- Treatment

- Balance between rest and moderate activity
- Heat and cold applications
- Physical and occupational therapy
- NSAIDs
- Glucocorticoids for severe inflammation
- Analgesia for pain
- Disease-modifying antirheumatic drugs, such as gold salts, methotrexate, hydroxychloroquine
- Biological response–modifying agents, such as infliximab, rituximab, anakinra

Juvenile Rheumatoid Arthritis (JRA)

- Several different types
- Onset more acute than adult form
- Large joints frequently affected

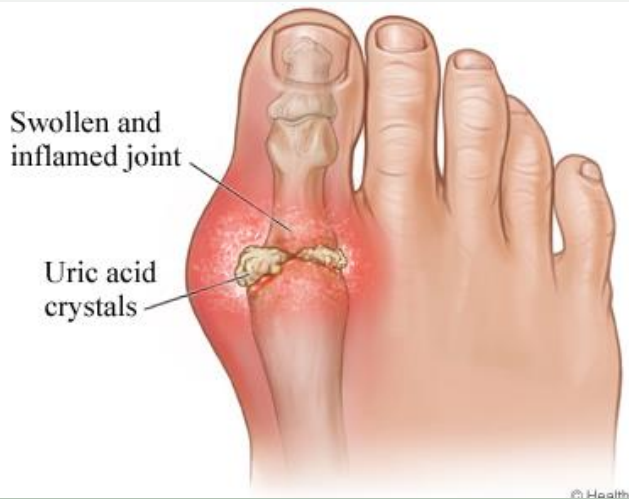
Infectious Arthritis

- Also known as septic arthritis
- Develops in single joint
- Joint is red, swollen, painful, with decreased movement
- Causes
 - Direct introduction of bacteria into joint, such as trauma, nonsterile injection, surgery
 - Secondary infection because of bacteremia
- Treatment with antimicrobials over sustained period; often requires IV administration

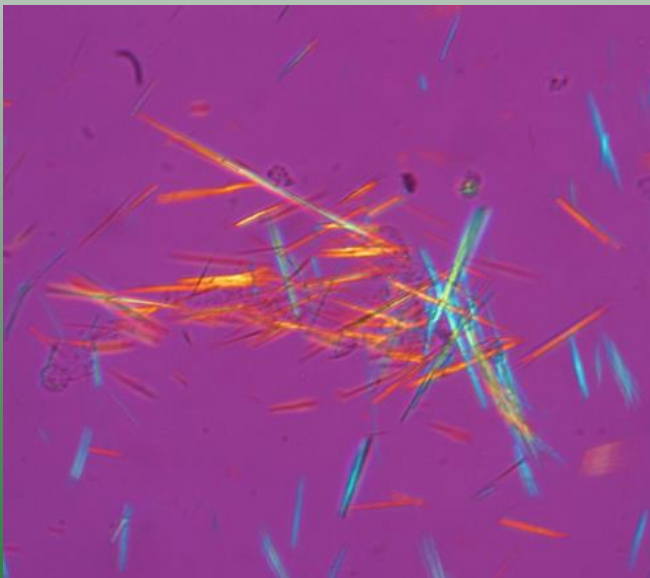
Gout

- Also known as gouty arthritis
- Results from deposits of uric acid and crystals in the joint, causing inflammation
- Formation of tophus—large hard nodule of urate crystals
- Tophi cause local inflammation and occur after the first attack of gout.

Gout (Cont.)



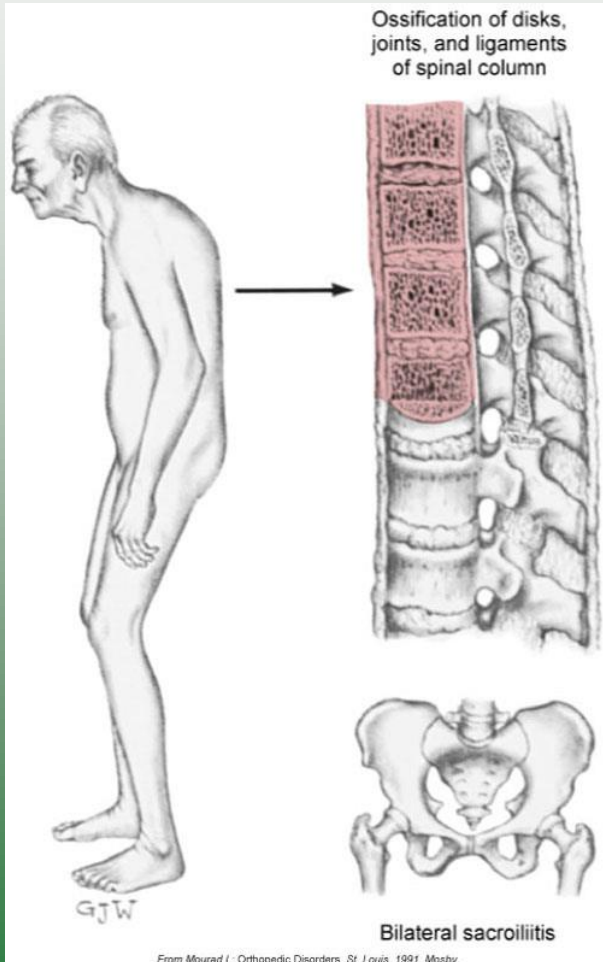
- Uric acid and crystals form because of inadequate renal excretion, chemotherapy, metabolic abnormality, and/or genetic factors.
- Inflammation causes redness, swelling, and pain. **Classic is red inflamed big toe.**
- Treated by reducing uric acid levels with drugs and dietary changes
- Diagnosed by examination of synovial fluid and blood tests
 - **NOTE:** Use of NSAIDs prior to blood tests will cause a false-negative result.



Ankylosing Spondylitis

- Chronic, progressive, inflammatory condition
- Affects sacroiliac joints, intervertebral spaces, costovertebral joints
- More common in men age 20 to 40 years
- Cause has not yet been determined—thought to be an autoimmune disorder with a genetic basis

Characteristic Posture and Sites of Ankylosing Spondylitis



- Progression of condition
 - Vertebral joints inflamed
 - Fusion of joint—loss of mobility
 - Inflammation in lower back, then up spine
 - Kyphosis develops
 - Osteoporosis common
 - Lung expansion limited