

# Bone Tumors

Fifth Year – Tikrit Medical College

Orthopedic Surgery Lecture

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# Objectives

- ▶ Introduction & Classification Of Bone Tumors
- ▶ Defining Primary And Secondary Bone Tumor
- ▶ Clinical Presentation (Signs And Symptoms )
- ▶ X-ray Features Of Benign And Malignant Bone Tumor
- ▶ Other Imaging Modalities In Assessing Bone Lesion
- ▶ Lab Investigations And Biopsy Principles
- ▶ Staging ( Enneking's Staging )
- ▶ Goals Of Treatment In Benign And Malignant ( Both Primary And Secondary )
- ▶ Examples Of Common Benign Bone Tumors
- ▶ Examples Of Common Malignant Bone Tumor

# Bone Tumors

▶ A tumor is a lump or mass of tissue that divide uncontrollably.

▶ Classification:

1. primary bone tumor .. Arise originally from the bone

- ▶ benign or Malignant
- ▶ in first 3 decades of life.
- ▶ Benign tumors > malignant.
- ▶ commonest sites around the knee , distal femur and proximal tibia

2. secondary bone tumor ..

- ▶ metastasize to the bone form ( breast , prostate etc ..)
- ▶ malignant transformation of benign lesions.
- ▶ Most commonly noticed above the fifth decade of life

► **Histological classification based on dominant tissue:**

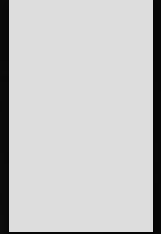
<b>Cell type</b>	<b>Benign</b>	<b>Malignant</b>
<b>Chondrogenic</b>	<b>Osteochondroma Enchondroma Chondroblastoma Chondromyxoid fibroma</b>	<b>Chondrosarcoma</b>
<b>Osteogenic</b>	<b>Osteoid osteoma Osteoblastoma</b>	<b>Osteosarcoma</b>
<b>Histocytic</b>	<b>Fibrous histiocyctoma</b>	<b>Malignantf fibrous histiocyctoma</b>
<b>Fibrogenic</b>	<b>fibrous cortical defect(non-ossifying fibroma), fibrous dysplasia, fibroma</b>	<b>Fibrosarcoma</b>
<b>Vascular</b>	<b>Hemangioma</b>	<b>Angiosarcoma</b>
<b>Others</b>	<b>Giant cell tumor, aneurismal bone cyst, simple bone cyst</b>	<b>Malignant Giant cell tumor</b>

# Clinical presentation:

## *History:*

1. **asymptomatic** accidentally discovered on x-ray, more likely with benign lesions.
2. **Pain:** it may be caused by:
  1. rapid expansion
  2. central hemorrhage.
  3. pathological fracture.
3. **Swelling or a lump.**
4. **Neurological symptoms .. Compression by mass**
5. **Pathological fracture**

# Examination:



- ▶ **Possible mass**
- ▶ **Joint effusion and \or limitation of movemet** in tumor around joint
- ▶ **muscle spasm and back stiffness, or painful scoliosis** in case of .Spinal lesions
- ▶ **Lymphadenopathy** should be checked
- ▶ **neurovascular check** for tumors in the limbs

# Imaging:



## ► X-ray : check for

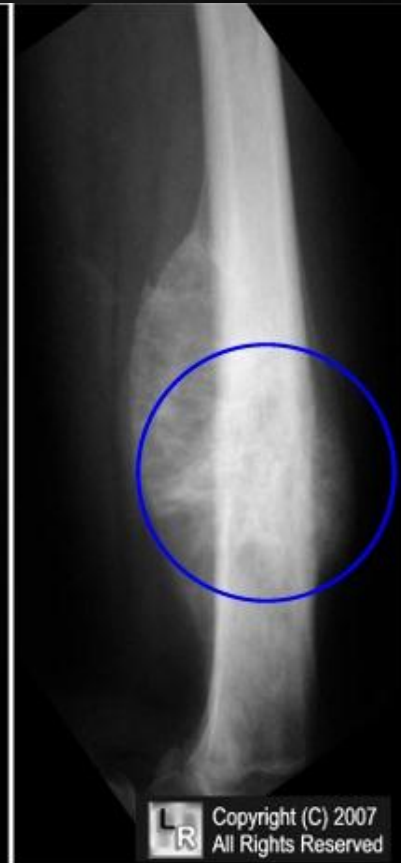
1. Which bone is involved
2. Where is the lesion in the bone? (epiphysis , metaphysis or diaphysis.)
3. lesion solitary or multiple?
4. centric or eccentric.
5. osteolytic or osteoblastic / is center calcified?
6. margins of the lesion well- or ill-defined?
7. Any cortical destruction?
8. Any periosteal reaction?
9. Any soft tissue extension

# Benign bone lesion

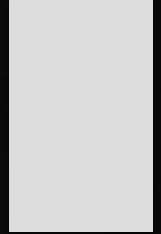




# malignant bone lesion

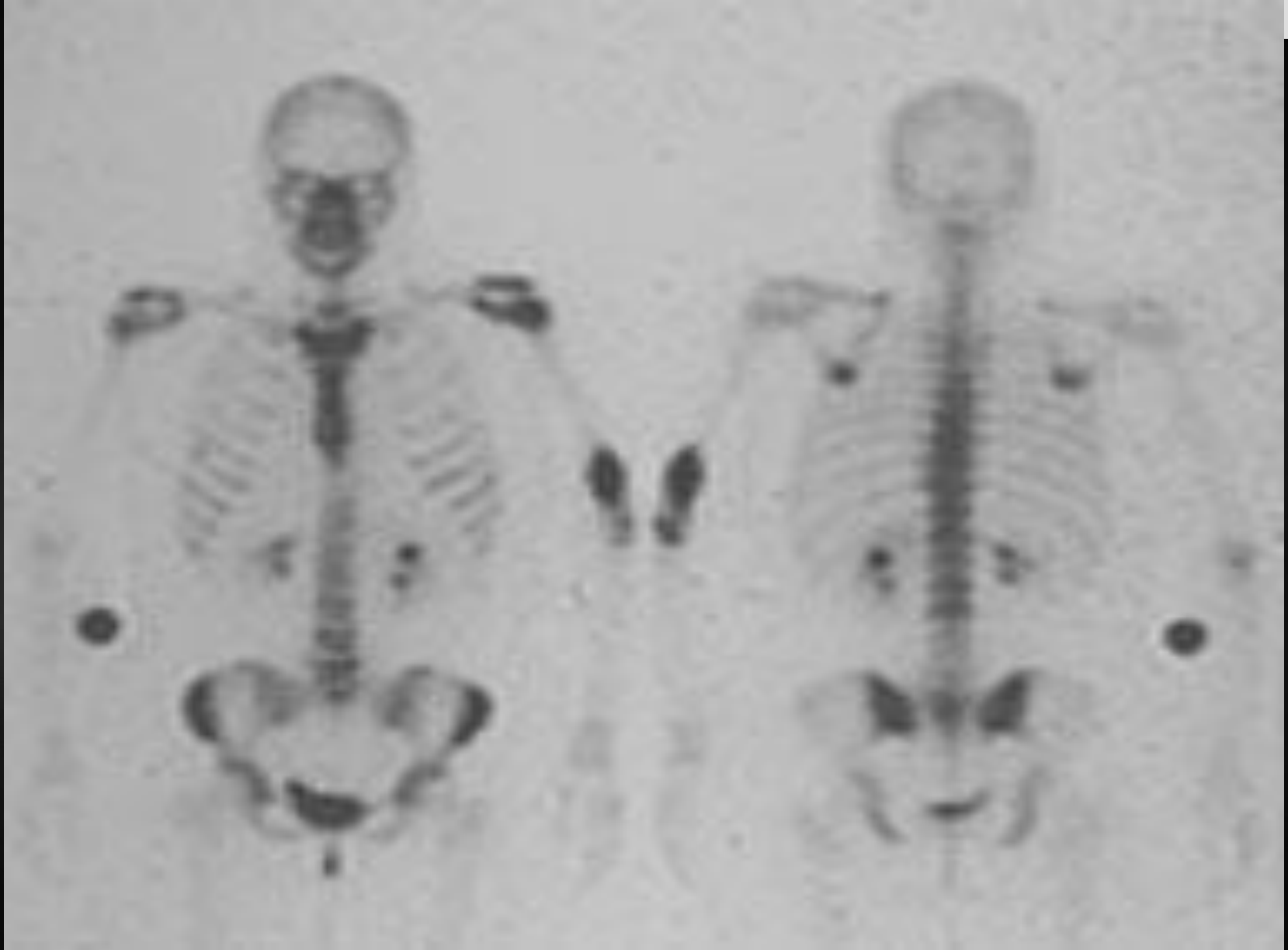


# Other imaging:

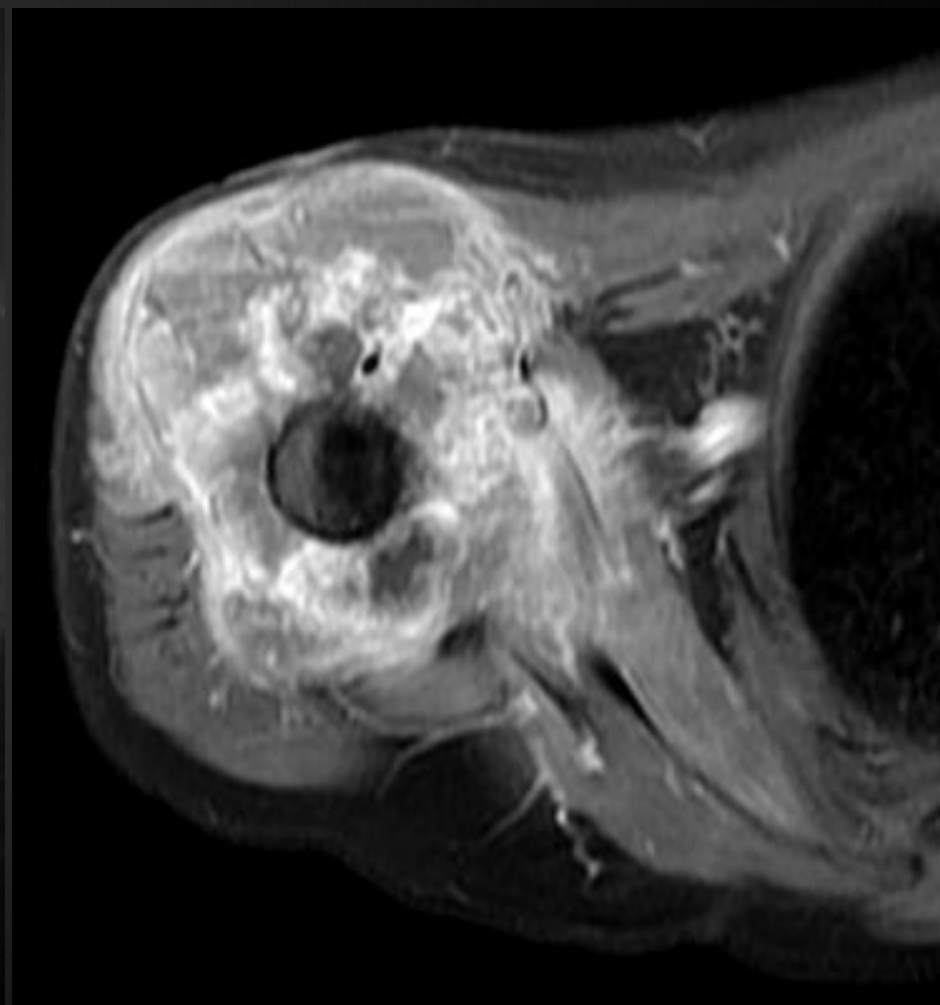
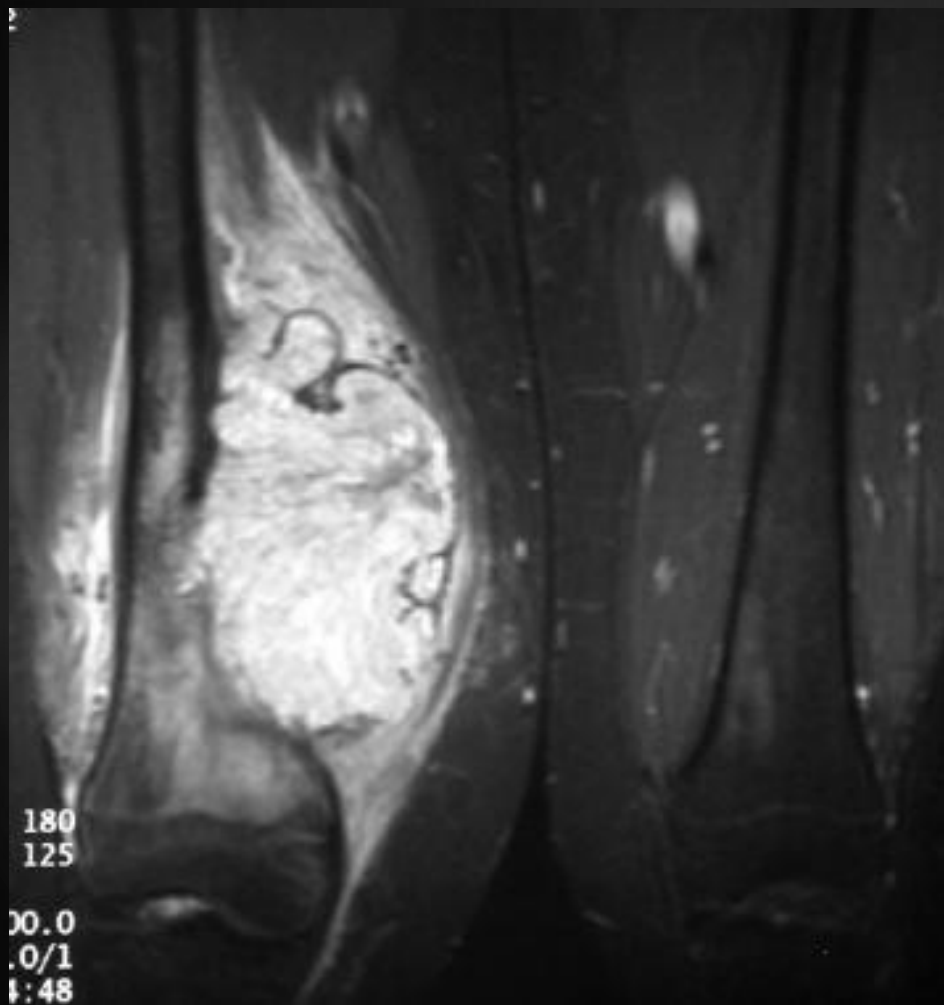


- ▶ **Radioisotope scanning** : Helpful in metastatic and skip lesions.
  
- ▶ **CT & MRI** : can determine:
  - ✓ The intra osseous and extra osseous extension of the tumor.
  - ✓ Skip lesion in the same bone
  - ✓ Lesions in inaccessible sites ,like the spine or pelvis.
  - ✓ Pulmonary metastasis.

# Bone Isotope Scan



# MRI



# Lab. investigations

- ▶ **Blood tests to exclude other conditions e.g.**
  - ▶ infection
  - ▶ metabolic bone disorders
  - ▶ “brown tumor” in hyperparathyroidism.
- ▶ **Serum and urine protein electrophoresis**
  - ▶ for abnormal globulin and Bence-Jones protein in myeloma.
- ▶ **serum acid phosphatase**
  - ▶ for prostatic carcinoma.

# Biopsy



- ▶ for accurate diagnosis

## two basic methods of doing a biopsy:

### 1. Needle biopsy

- ▶ under local anesthesia or GA using an X-ray or CT guidance
- ▶ value is in sampling inaccessible tumors.

### 2. Open biopsy:

- ▶ done through a small incision under general anesthesia in an operating room.
- ▶ Several samples should be taken



# Needle biopsy





**Open biopsy**



# Differential Diagnosis tumor mimicker

1. myositis ossificans.
2. stress fracture.
3. bone infection.
4. brown tumor of hyperparathyroidism.

# Staging the lesion:

## ▶ **Enneking's staging of benign lesion:**

- ▶ *latent*
- ▶ *active*
- ▶ *aggressive.*

## ▶ **Enneking's staging of malignant tumor:**

### Stage 1: low grade sarcomas

- ▶ *1A: intracompartmental*
- ▶ *1B: extracompartmental*

### Stage 2: high grade lesions.

- ▶ *2A: intracompartmental*
- ▶ *2B: extracompartmental*

### Stage 3: sarcomas which have metastasized. e.g. to lung.

# Management of Primary Benign tumors:

- ▶ Observation only / might disappear over time (e.g. fibrous cortical defect, simple bone cyst)
- ▶ Excision to reduce the risk of pathological fracture
- ▶ Excision because its symptomatic / or have a risk of malignant potentials like Giant cell tumor

# Management of Primary Malignant tumors:

- ▶ If the lesion is suspected to be a malignant tumor ,the patient is admitted for
  - ▶ detailed examination
  - ▶ blood tests
  - ▶ CXR
  - ▶ pulmonary CT
  - ▶ biopsy.
- ▶ **Treatment goals include**
  - ▶ Removing the tumor
  - ▶ preserving the function of the body .

# Methods of treatment of malignant tumor

- ▶ Tumor excision with wide excision or radical excision.
  - ▶ Limb-sparing surgery: removes cancerous section of bone but keeps nearby muscles, tendons, nerves and blood vessels . The excised bone is replaced with a metallic implant (prosthesis) or bone transplant.
  - ▶ Amputation : removes all or part of an arm or leg when the tumor is large and/or nerves and blood vessels are involved.
- ▶ Radiotherapy: uses high-dose X-rays.
  - ▶ Shrinks the tumors
  - ▶ suitable for inaccessible sites
- ▶ Multi-agent chemotherapy : neoadjuvant for malignant bone tumors

# Benign Bone Tumors

# Osteochondroma

the most common benign bone tumor

- ▶ common locations include
  - ▶ knee (proximal tibia, distal femur)
  - ▶ proximal femur
  - ▶ proximal humerus
- ▶ Can be either
  1. solitary osteochondroma
  2. Multiple Hereditary Exostosis (MHE)
- ▶ Clinical presentation
  - ▶ Asymptomatic / painless mass
  - ▶ mechanical symptoms
  - ▶ symptoms of neurovascular compression

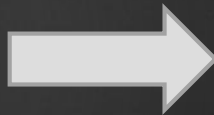
# Osteochondroma

## ► Radiograph

- sessile (broad base) or



- pedunculated



## ► Treatment

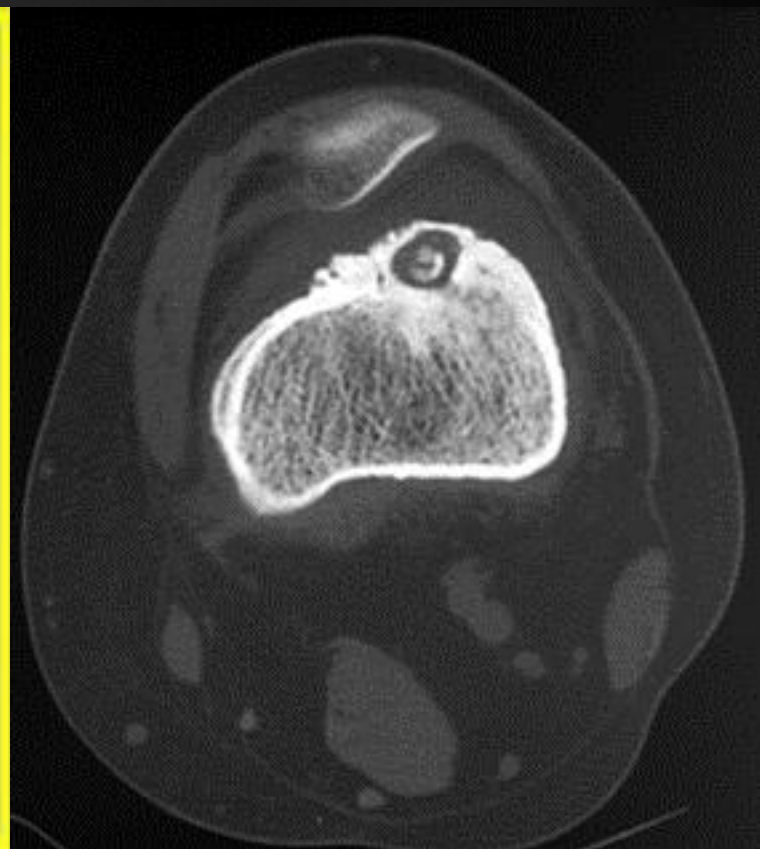
- Observation alone .. If Asymptomatic
- Operative .. If symptomatic or growing fast



# Osteoid osteoma

- ▶ A small, discrete, painful, benign bone lesion
- ▶ **Commonest** location
  - ▶ proximal femur > tibia diaphysis
    - ▶ usually within bone cortex
  - ▶ Spine .. Produce scoliosis
- ▶ Characterized by central nidus with surrounding sclerotic rim
- ▶ Pain is constant at night and relived with NSAIDS
- ▶ Radiographs
  - ▶ reactive bone around radiolucent nidus
- ▶ CT imaging is the study of choice

# Osteoid osteoma



# Osteoid osteoma

## Management

- ▶ **clinical observation and NSAID administration**
- ▶ **percutaneous radiofrequency ablation**
- ▶ **surgical resection/curettage**
  - ▶ complete marginal resection of nidus (sclerotic bone is normal and can be left behind)

# Non-ossifying Fibroma

- ▶ fibrogenic lesion /dysfunctional ossification
- ▶ Locations ... metaphysis of long bones

- ▶ **Symptoms**

- ▶ asymptomatic found incidentally
  - ▶ Or pathologic fracture

- ▶ **Radiographs is diagnostic**

- ▶ metaphyseal cortical eccentric "bubbly" lytic lesion surrounded by sclerotic rim

- ▶ **Treatment**

- ▶ **observation** .. most resolve spontaneously
  - ▶ **curettage and bone grafting** .. If symptomatic or at risk of fracture



# Unicameral bone cyst

## simple bone cyst

- ▶ A non-neoplastic, serous fluid-filled bone lesion / failure of bone formation
- ▶ usually found in the metaphysis of long bones in young patients <20 years
- ▶ found in the
  - ▶ proximal humerus
  - ▶ Proximal femur
  - ▶ Distal tibia and radius
- ▶ Symptoms
  - ▶ most asymptomatic unless fracture occurs (usually with minor trauma)
  - ▶ pathologic fracture in ~50%

# Unicameral bone cyst (simple bone cyst)

- ▶ Radiographs

- ▶ central, lytic, well-demarcated metaphyseal lesion
- ▶ thinning of cortices



- ▶ Treatment

- ▶ Observation if at low risk of fracture
- ▶ aspiration/methylprednisolone injection
- ▶ curettage and bone grafting +/- internal fixation based on tumor location

# Giant cell tumor

- ▶ A **benign** aggressive tumor found in the metaphysis of long bones in **mature** adults
- ▶ distal femur > proximal tibia > distal radius
- ▶ **Clinical features**
  - ▶ pain in the involved joint
  - ▶ palpable mass
- ▶ **Radiograph**

eccentric lytic epiphyseal/metaphyseal lesion extends subchondral bone

  - ▶ Chest radiograph or chest CT .. 5% pulmonary metastasis
  - ▶ Bone scan is very hot
  - ▶ MRI ,... signal change



# Giant cell tumor





# Giant cell tumor

## Treatment

- ▶ **medical management ??? New modality ..**
  - ▶ bisphosphonates
  - ▶ denosumab
- ▶ **Operative**
  - ▶ **extensive curettage and reconstruction (with adjuvant treatment)**
  - ▶ 10-30% recurrence with curettage alone verses 3% with adjuvant treatment (phenol , hydrogen peroxide , argon laser ..)



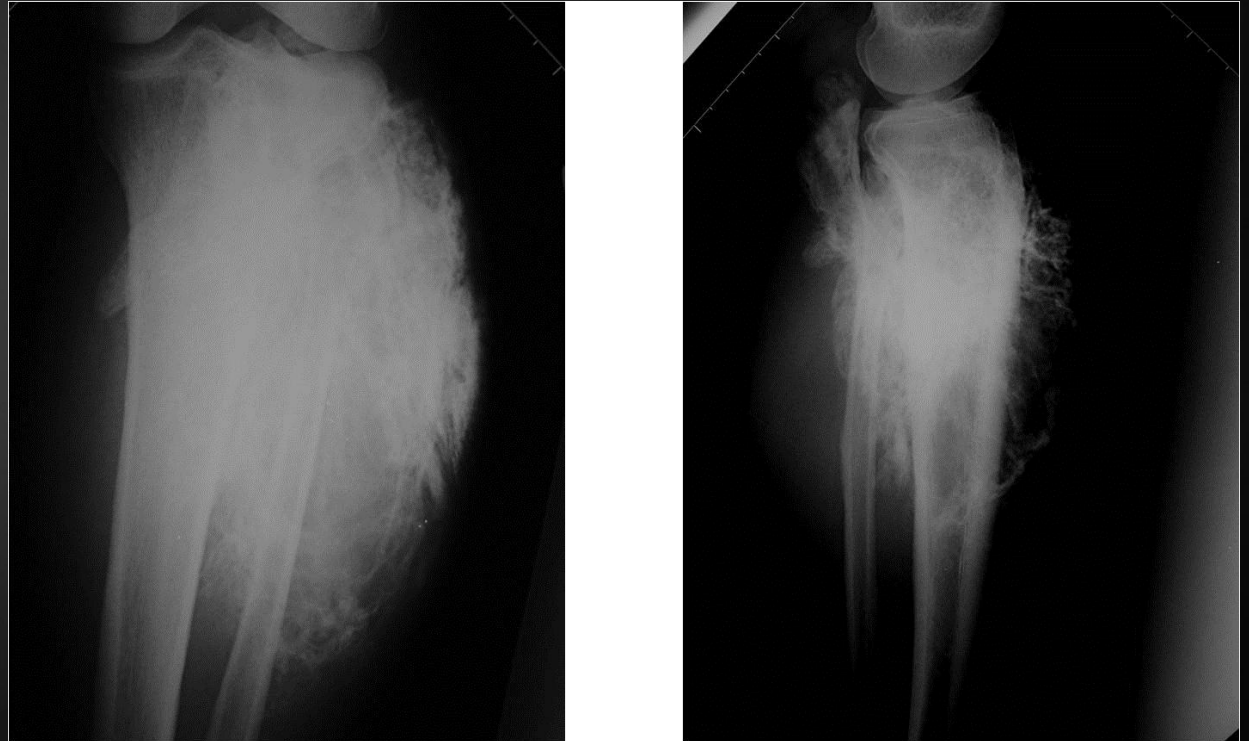
# **Malignant Bone Tumors**

# Osteosarcoma

- ▶ the most common primary sarcoma of bone
- ▶ in children and young adults <25 years
- ▶ common site / distal femur & proximal tibia
- ▶ commonly diagnosed at Stage IIB (high grade, extra-compartmental, no metastases)
- ▶ 10-20% of patients has pulmonary metastases
- ▶ **Presentation**
  - ▶ rapidly progressive pain, fever, and swelling
  - ▶ may feel mass on examination
- ▶ Radiographs
  - ▶ mixed blastic and destructive lesion
  - ▶ sun-burst or hair on end pattern
  - ▶ periosteal reaction (Codman's triangle)

# Osteosarcoma

- ▶ MRI must include entire involved bone to determine
  - ▶ soft tissue
  - ▶ neurovascular involvement
  - ▶ skip metastases in same bone
- ▶ Bone scan
- ▶ chest Ct for metastasis



# Osteosarcoma

## Treatment

- ▶ **multi-agent chemotherapy and limb salvage resection**
  - ▶ preoperative chemotherapy given for 8-12 weeks followed by ...
  - ▶ resection then ...
  - ▶ maintenance chemotherapy for 6-12 months after surgical resection
- ▶ Prognosis 76% long-term survival with modern treatment.

# Ewing's Sarcoma

- ▶ typically from 5-25 years of age
- ▶ second common malignant bone tumor in children
- ▶ ~50% are found in the diaphysis of long bones
- ▶ Genetics t(11:22) translocation in all cases
- ▶ **Presentation**
  - ▶ pain with fever
  - ▶ mimics an infection !!!!!
  - ▶ swelling and local tenderness
- ▶ **Radiographs**
  - ▶ destructive lesion in the diaphysis or metaphysis with a moth-eaten appearance
  - ▶ periosteal reaction give "onion skin" or "sunburst" appearance
- ▶ **MRI** .. soft-tissue extension and marrow involvement
- ▶ CT chest and bone scan for metastasis

# Ewing's Sarcoma



# Ewing's Sarcoma

## Treatment

- ▶ **Neoadjuvant chemotherapy with limb salvage resection followed by postoperative chemotherapy**
- ▶ **the standard of therapy in most patients**
  - ▶ Neoadjuvant chemotherapy given for 8-12 weeks followed by surgical resection then maintenance chemotherapy for 6-12 months
- ▶ Prognosis
  - ▶ 60-70% long term survival with **isolated** extremity disease
  - ▶ 15% long term survival if patient presents with **metastatic** disease



# Bone Metastasis / Secondary Bone Tumor

- ▶ most common malignancy of bone is metastatic disease
- ▶ metastatic lesions are usually found in older patients (> 40 years)
- ▶ carcinomas commonly spread to bone include (Breast, lung, thyroid, renal, prostate)
- ▶ common sites of metastatic lesions include spine> proximal femur> humerus
- ▶ Symptoms
  - ▶ pain
  - ▶ pathologic fracture
  - ▶ metastatic hypercalcemia

# Evaluation of bone metastasis

Workup for older patient with bone lesion and unknown primary includes

▶ **Imaging:**

- ▶ plain radiographs in two planes of affected limb
- ▶ CT of chest / abdomen / pelvis
- ▶ bone scan to detect extent of disease

▶ **Labs**

- ▶ CBC , ESR
- ▶ LFTs, Ca, Phos, alkaline phosphatase
- ▶ serum and urine immuno-electrophoresis

- ▶ **biopsy** .. where a primary carcinoma is not identified, obtaining a biopsy is necessary to rule out a primary bone lesion.

# Treatment of metastatic bone disease

- ▶ **Nonoperative ...**
  - ▶ bisphosphonate therapy
  - ▶ chemotherapy, radiotherapy, and hormone therapy
- ▶ **Operative .. aim is not cure but to improve the quality of life !!!**
  - ▶ stabilization of complete fracture with postoperative radiotherapy
  - ▶ prophylactic stabilization of impending fracture, postoperative radiation