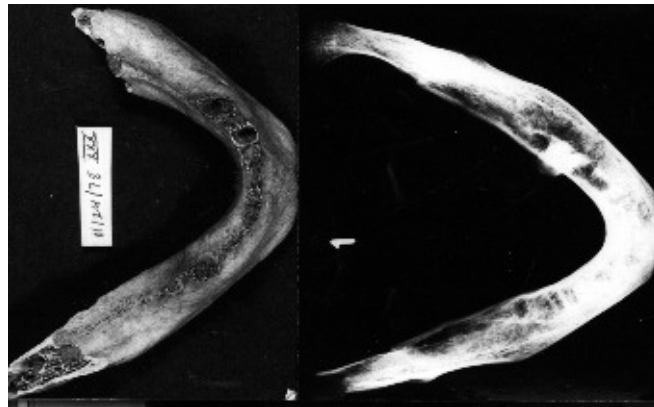
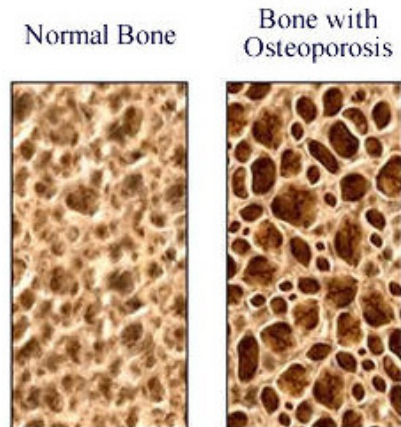


🔔 Metabolic & Endocrine disorders of bone:

∅ Osteoporosis:

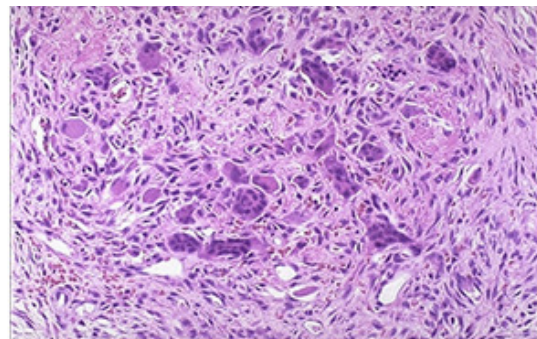
- Bone apposition < bone resorption
- **Risk factors:**
 - Postmenopausal women
 - Hyperthyroidism
 - Hyperparathyroidism
 - Cushing's syndrome
- **↓ bone quantity:** thin cortex & trabeculae & ↑ marrow
- **Dental aspects:** denture, surgery, sinus



∅ Hyperparathyroidism:

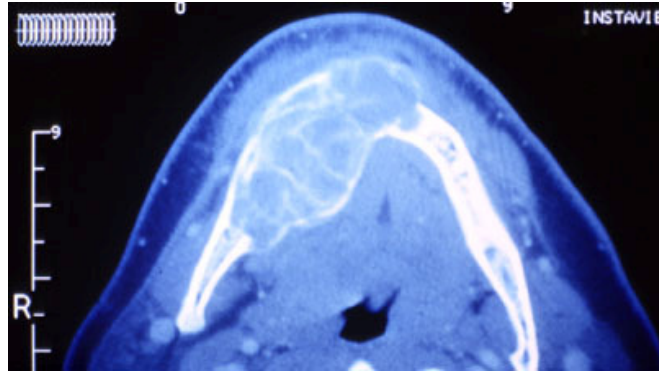
∇ Primary:

- Adenoma, Ca, Hyperplasia
- Postmenopausal women
- Hypercalcaemia → metastatic calcification
- **Bone disease:**
 - ↑ Osteoclastic activity
 - Brown tumours



- **Rx:**

- Osteoporosis
- Mottled areas of racy & thinning of cortical plates
- **Jaws:** loss of n trabecular pattern, LD, brown tumours



- **Biochemistry:**

- \uparrow Ca, \downarrow PO₄, \uparrow PTH, \uparrow ALKP, \uparrow urinary Ca & PO₄

∇ **Secondary:**

- RF → low Ca
- Rickets & Osteomalacia:

∅ **Rickets & Osteomalacia:**

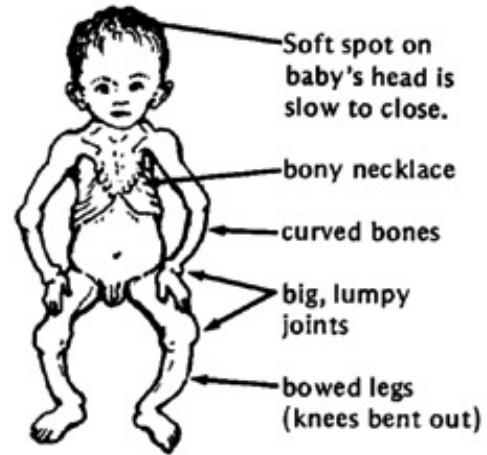
- \downarrow Vit D or resistance to its action
- RF, malabsorption of Ca
- Failure of mineralization of Osteoid & cartilage
- Weak bones, bending
- **Biochemistry:** n/ \downarrow ca, \downarrow PO₄, \uparrow ALKP



○ **Dental aspects:**

- E hypoplasia
- Dentine hypocalcification
- Delayed eruption
- Condyle

SIGNS OF RICKETS



Ø **Acromegaly:**

- Prognathism
- Macroglossia
- Lips & nose
- Hands & feet



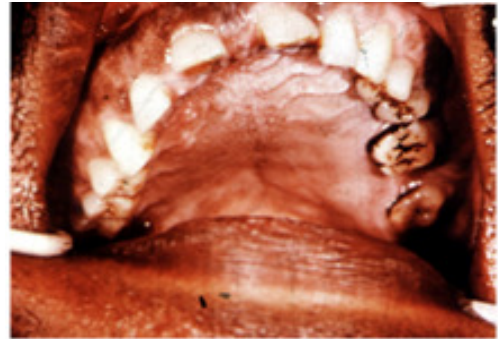
🔔 **Paget's disease:**

- Uncoordinated ↑ in Osteoclastic & osteoblastic activity of bone Cs
- Primary dysfunction of osteoclasts
- Viral inclusions
- **Abs:** measles & RCV
- **Aetiology:** slow viral infection + genetic factors

- **Three phases:** osteolytic, mixed & osteoblastic

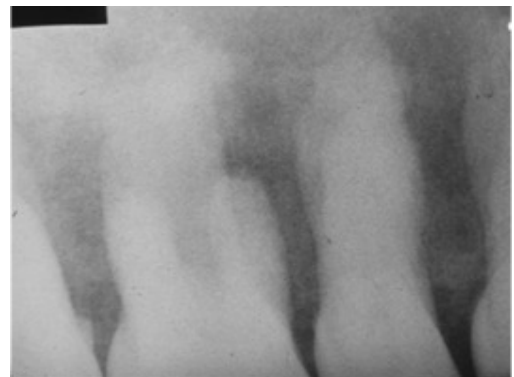
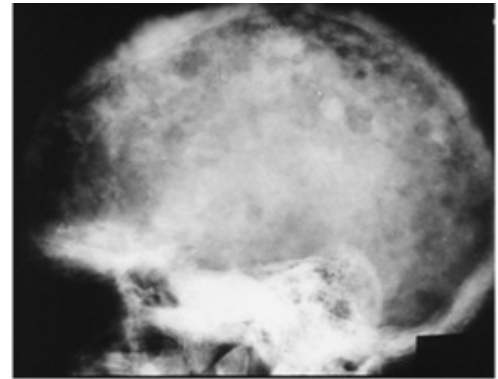
- **Clinically:**

- M>F, > 40y
- Geographical variation
- Monostotic or polyostotic
- Axial skeleton then skull & femur
- Max > Mand
- Deformity of spine & legs
- Fracture
- Enlargement of skull & facial bones
- Bone pain & joint disease
- Skull base; cranial nerves & spinal cord
- Symmetrical & gross enlargement of alveolar process, flat palate
- Spaces between teeth, malocclusion, incompetent lips
- Hypercementosis & ankylosis, root resorption
- Haemorrhage
- Infection



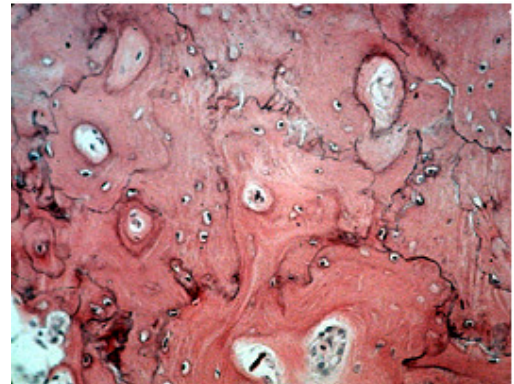
- **Rx:**

- Diffuse radcy
- Patchy Osteosclerosis “**cotton wool**” appearance
- Thick cortical plates of skull
- Enlarged maxilla & mandible
- Loss of LD, hypercementosis & ankylosis



- **Hist:**

- Rapid bone resorption & replacement
- **Initially:** ↑ Osteoclastic activity → vascular fibrous marrow
- **Then:** Osteoclastic & osteoblastic activity, mosaic pattern
- **Finally:** Osteoblasts predominate → dense lamellar bone
- **Jaws:** masses of cementum-like tissue, cementum (mosaic)



- **Biochemistry:**

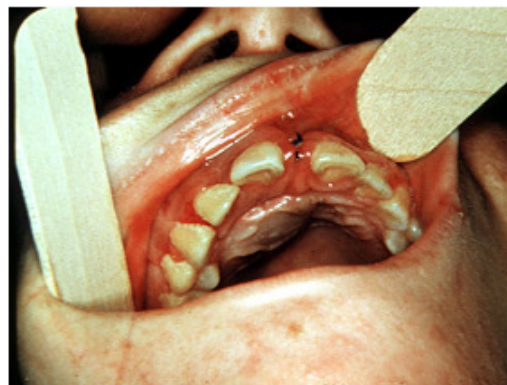
- ↑ ALKP:
- Normal Ca & PO₄

- **Complications:**

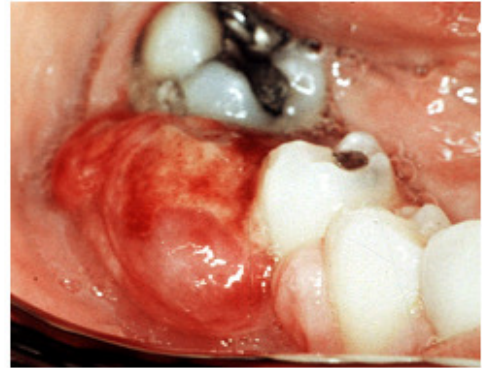
- ↑ A-V shunt → ↑-output HF
- Osteosarcoma

🔔 Central Giant cell granuloma:

- Giant cell lesions
- Less aggressive & destructive than in other bones
- **Clinically:**
 - Less common than PGCG
 - Majority: 10-30 ys, F>M

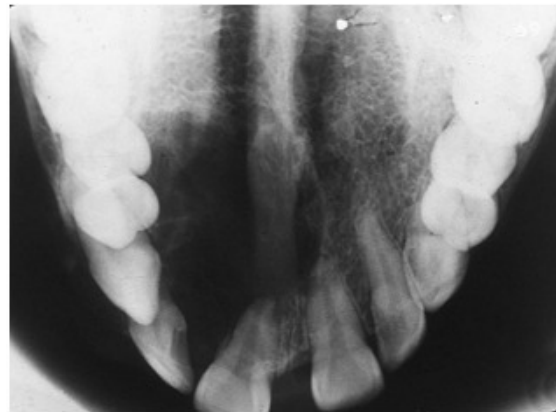


- Mand > Max, Anterior
- 75% in Mand & crossing midline
- Expansion
- ± Perforation & root resorption



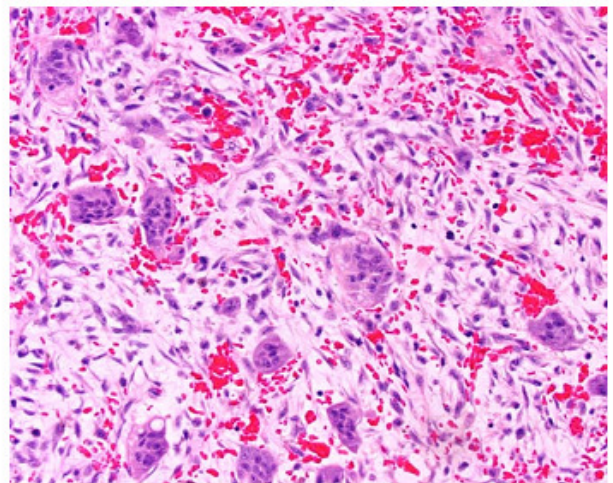
○ **Rx:**

- Cyst-like radcy w expansion
- “Soap-bubble” appearance
- Displacement & occasional resorption of roots



○ **Hist:**

- Giant Cs, mononuclear Cs, FV stroma
- Hemosiderin, extravasated RBCs
- Strands of collagen w Osteoid or bone
- Aggressive lesions



○ **Aetiology?**

🔔 **Torus/exostosis:**

- Nodular, non-neoplastic growth of bone
- Developmental, some AD
- **Torus palatinus:**
 - Develops after puberty in susceptible pts
 - Grows slowly over enter life



- Rounded, symmetrical, can become large & pedunculated
- Thin mucosa
- **Denture, speech, O.H**



- **Torus mandibularis:**

- Less common
- Lingual to premolar
- Frequently bilateral
- Usually multiple-lobed
- Tongue movement, denture, O.H



- **Hist:** cortical bone ± cancellous bone
- Buccal alveolus of Max in molar region



☒ **Bone tumours:**

- **Classification:**

- **Bone-forming tumours:**

- **Benign:** osteoma & osteoblastoma
- **Malignant:** osteosarcoma

- **Cartilage forming tumours:**

- **Benign:** chondroma
- **Malignant:** chondrosarcoma

- **Marrow tumours:** myeloma

- **Fibrous tumours:**

- **Benign:** cemento-ossifying fibroma

▪ **Tumour-like lesions in bone:** Langerhans cell histiocytosis, haemangioma

▪ **Metastatic tumours**

☒ **Osteoma:**

- Benign & slow growing
- Mand > Max
- Single or multiple
- Superficial or intraosseous



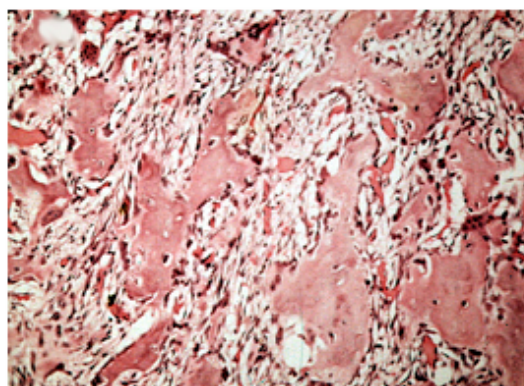
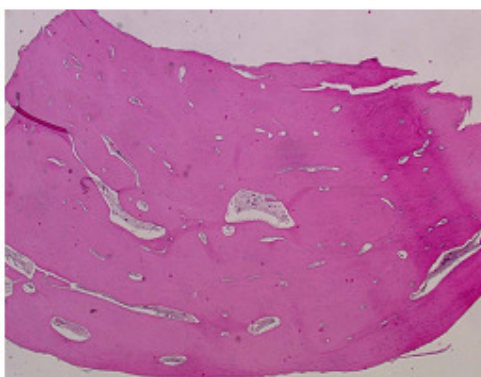
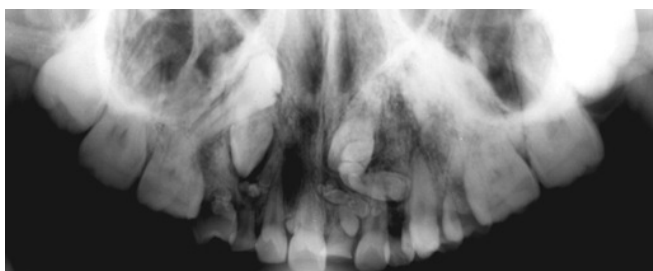
○ **Gardner Syndrome:**

- AD
- Intestinal polyps
- Unerupted normal & supernumerary teeth
- Fibromas of skin
- Epidermal/sebaceous cysts



○ **Hist:**

- Compact vs. cancellous



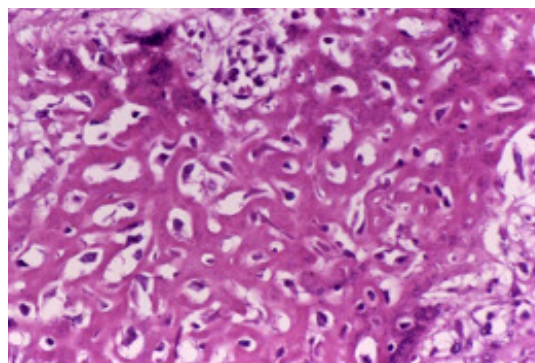
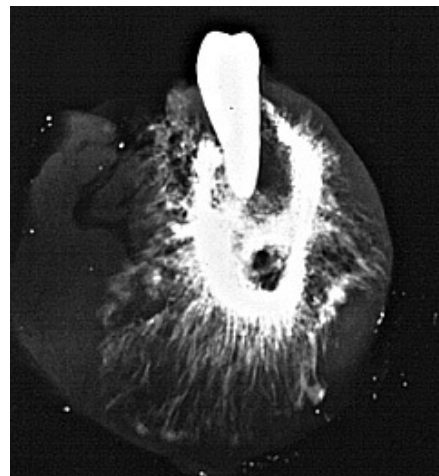
☒ Osteoblastoma:

- Rare
- Swelling & pain
- Rx: rounded, w-d w central radcy or speckling
- **Hist:** osteoblasts, MNGC, Osteoid & FV stroma



☒ Osteosarcoma:

- Most common bone sarcoma
- Long bones, 7% H & N
- Mand > Max, 10 ys later
 - Bony-hard swelling
 - W or without pain
 - Loosening of teeth, paraesthesia
 - Trismus
 - Nasal obstruction & eye symptoms
- **Rx:**
 - Variable
 - **Osteolytic tumours:** irregular radcy
 - **Sclerotic type:** irregular radio-opacity
 - “**Sun-ray**” appearance
 - PL
- **Hist:**
 - Osteoid
 - Fibroblastic type, chondroblastic type
- **Prognosis:** osteoblastic, less metastasis in jaws



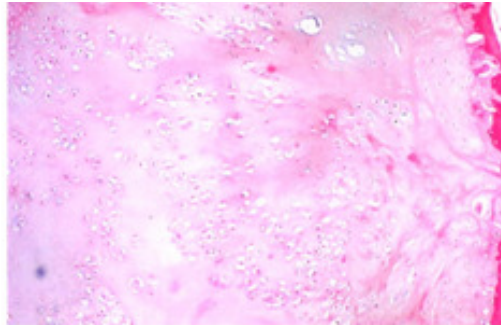
☒ Chondroma & Chondrosarcoma:

○ Ant Max & post Mand, symphysis, condyle, coronoid

○ Hist:

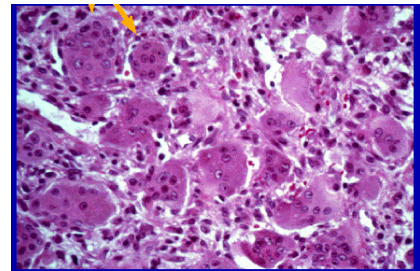
- Chondroma: mature cartilages
- Chondrosarcoma:
- Calcifications

○ **Clinically:** malignant → pain, loosening of teeth



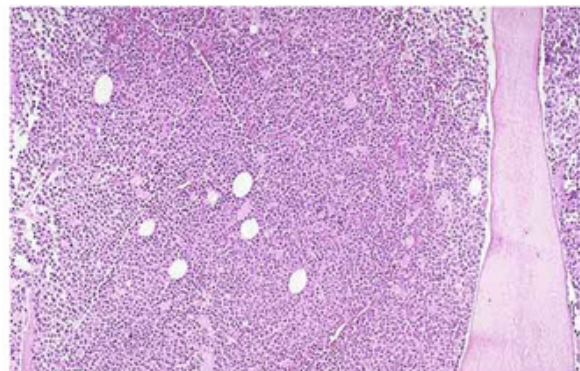
☒ Giant-cell tumour:

- Ends of long bones
- Aggressive & locally invasive W ↑ LRR ± metastasis
- **Hist** = CGCG
 - Uniform distribution of giant Cs
 - No Osteoid or bone
- Older age group



☒ Myeloma:

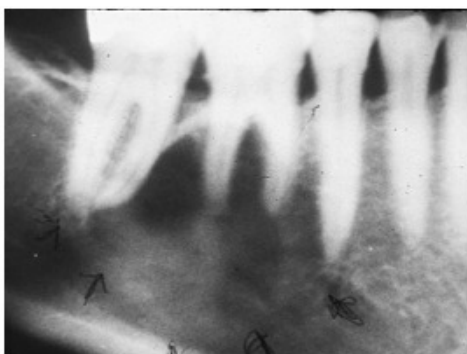
- Differentiated B lymphocytes or plasma Cs win bone marrow
- **Solitary plasmacytoma**
- Jaw or oral soft tissues
- 50-70 yrs
- Large amounts of a single homogenous type of Igs
- **Bence-Jones proteins**
- Multiple foci of bone destruction, bone pain, anaemia, thrombocytopenia
- Infection
- Hypercalcaemia
- Proteinuria
- Bones w red marrow
- **Jaw lesions:** Mand > Max, S&S



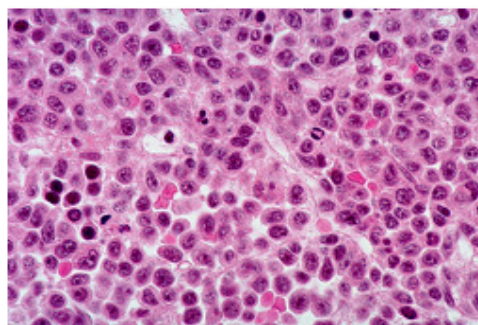
- Macroglossia



- **Rx:** w-d, round/oval punched-out radices



- **Hist:** sheets of plasma Cs, Immunohistochemistry



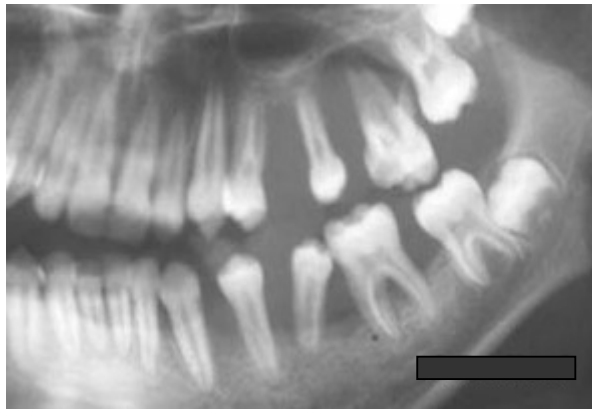
⊗ Langerhans cell histiocytosis:

- Langerhans cells

- **3 forms:**

1. Unifocal eosinophilic granuloma (Chronic focal LCH):

- < 20 yrs, M > F
- Cranium, jaws, ribs & long bones, Mand>Max
- **Jaws:** localized bone destruction w swelling & often pain
- **Rx: “floating in air”**
- Spontaneous regression, curettage, excision or Rxd
- → Multifocal lesion



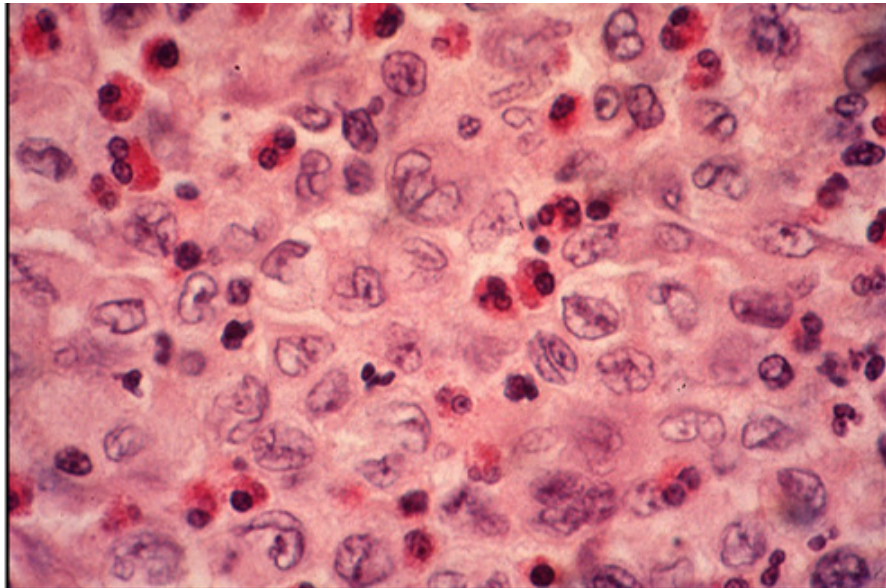
2. Multifocal eosinophilic granuloma (chronic disseminated LCH):

- Several bones & often other organs
- Skull
- Jaws
- ± Liver, spleen, LNs
- **Hand-Schuller-Christian syndrome**

3. Progressive (acute) disseminated histiocytosis:

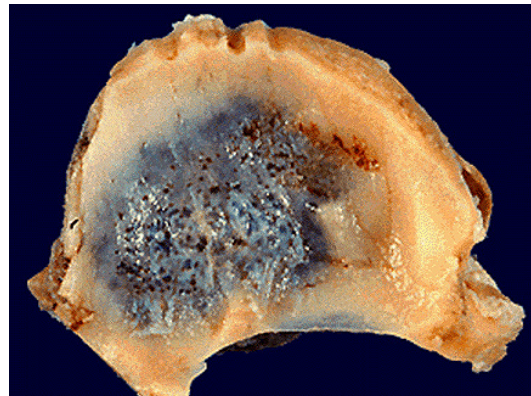
- **Letterer-Siwe disease**
- Aggressive

- Infants & young children
- Disseminated→ skin, viscera, bones
- **Clinically:** fever, malaise, ↑LNs, ↑ liver & spleen, pancytopenia
- **Rx:**
 - Osteolytic lesions
 - **Jaws:** diffuse bone destruction, floating, loosening of teeth
- **Hist:**
 - Histiocytes w pale, lobulated nuclei & eosinophilic cytoplasm
 - Eosinophils, neutrophils, lymphocytes
 - He, necrosis, fibrosis, giant Cs

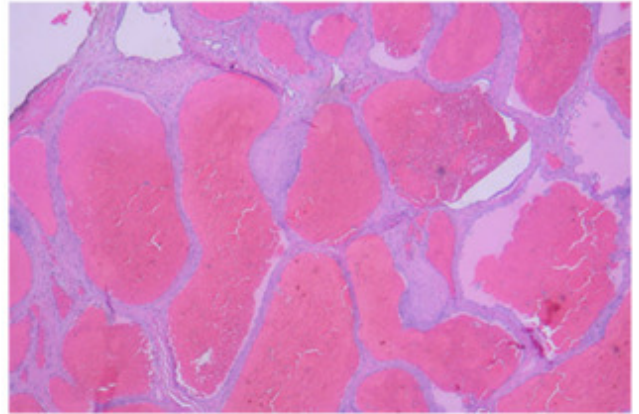


☒ Haemangioma of bone:

- **Mand>Max**
- **Clinically:**
 - Progressive painless swelling
 - Pulsatile
 - Loosening, bleeding
 - Hge



- **Rx:** osteolytic defect, multilocular
- **Hist:** usually cavernous



☒ **Metastatic tumours:**

- Breast, bronchus & kidney
- Bone & soft tissue
- Mand>Max
- S & S
- Osteoblastic tumours