# Upper limb tumours

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#### > MCQs

- > Tumour Basics
- Common tumours
  - Bone
  - Soft tissue
- Considerations specific to Upper Limb tumours
- Proximal humeral resection/reconstruction

#### Cases

MCQ answers





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#### **Definition:**

Mass of tissue formed as a result of abnormal excessive and inappropriate proliferation of cells, growth of which occurs indefinitely regardless of the mechanisms that control normal cellular proliferation

History/Examination

> Pain

- Persistent, night, analgesia
- > Swelling/mass
- Rate of progression
- > Age
  - Young Benign Vs Ewings / osteosarcoma
  - 40-60 Chondrosarcoma / haemopoetic tumours
  - 70s Metastasis / osteosarcoma / myeloma / lymphoma
- > Neurologic symptoms
- > Previous malignancy / radiotherapy / +ve FH

Imaging: Image whole bone affected

- What is the effect of the lesion on the bone?
  - Zone of transition / margin
  - Slow growing narrow / sclerotic
  - Rapid growing permeative / codmans traiangle / sunray spiculation
- > What is the effect of the bone on the lesion?
- Is the lesion solitary or multiple?
- Where in the bone is the lesion

Imaging: Other studies
Characterising and Staging tumour
CXR
USS
Bone Scan
CT/MRI
Others eg PET Scan

#### Bloods

- > Ca<sup>2+</sup>, ALP
- PSA, Electophoresis, Urine Bence-Jones

#### Biopsy

- Ideally surgeon who will perform resection
- > Performed through muscle
- Don't expose Neurovascular bundles
- Stay within compartment
- Longitudinal not transverse incision
- Don't lift skin/tissue flaps
- Send sample for culture
- Meticulous haemostasis
- Fresh Vs Fixed

Needle (Jamshedi/Trucut/Islam) Vs Open Vs Excisional

Staging

- > Why?
  - Prognostic / Guide treatment & adjuvant therapies
- Enneking
  - Grade (From Biopsy): Low (G1) Vs High (G2) grade
  - Site (From local imaging): Intracompartmental (T1) Vs Extracompartmental (T2)
  - Metastasis (From staging CT): No Mets (M0) Vs Mets (M1)

Stage	Grade	Site	Metastasis
IA	G1	T1	M0
IB	G1	T2	M0
IIA	G2	T1	M0
IIB	G2	T2	M0
]]]	Any	Any	M1

Plus graph sarcoma.org Ch 1

**Tumour Excision** 

Diagram sarcoma.org Ch 1

Osteosarcoma – Malignant spindle cell tumour

- > Bimodal age distribution
- Distal femur (50%) & Prox Humerus (25%)
- X-ray lytic/sclerotic, permeative margins, Codmans triangle, sunray spiculation
- $> \approx 10\%$  have lung mets at presentation
- Poor prognosis if develops in Pagetic bone

Chondrosarcoma – Malignant cartilage tumour

- > 4<sup>th</sup>/5<sup>th</sup> Decade
- > M>F
- X-ray patchy calcification: Popcorn appearance, endosteal scalloping
- Often slow growing with late metastasis
- > Not chemo/radiosensitive

Ewings – Malignant small round blue cell tumour

- Assoc with (11:22) chromosome translocation
- Occurs in kids (median age 13)
- Mainly femoral / tibial diaphysis
- Often have soft tissue invasion leading to Onion skin appearance on x-ray
- Neoadjuvant Chemo highly effective in ↓tumour bulk

Giant cell tumour – Benign but aggressive tumour

- 80% occur in the mature skeleton
- Varied behaviour
  - Latent vs active vs aggressive
- Pathology: Multinucleated giant cells
   & stomal cells
- Epiphyseal abutting subchondral bone
- Treatment: excision preserving joint / reconstruction

Enchondroma – Benign Cartilage tumour

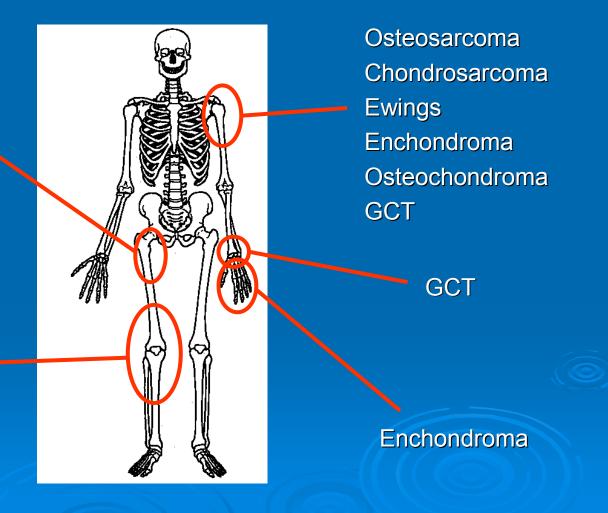
- Islands of persisent cartilage in metaphysis due to defective endochonral ossification
- Lesions in hand/feet benign
- Lesions in pelvis/long bones more concerning
- Single Vs Multiple (Olliers) Vs + Haemangiomas (Maffuccis)

Osteochondroma (Exostosis) – Benign bone surface tumour

- Commonest benign bone tumour
- Solitary Vs Multiple (Diaphyseal acalasis)
- Bone stalk with cartilage cap
- Should stop growing when parent bone stops growing
- Low risk of malignant change
- Concern if *fsize* or cap >1cm

Osteosarcoma Chondrosarcoma Ewings Enchondroma Osteochondroma

Osteosarcoma Chondrosarcoma Ewings Enchondroma Osteochondroma GCT



#### Metastasis

- Lung, Breast, Prostate, Kidney, Thyroid Principles:
- Control pain
- Control mass of deposits
- Treat fractures
- > Treat ↑Ca<sup>2+#</sup>
- Prophylactic stabilisation

### Soft Tissue Tumours

#### > Soft Tissue Sarcomas in upper limb

	UL (%)	LL (%)
Malignant fibrous histiocytoma	40	31
Liposarcoma	15	25
Synovial Sarcoma	10	5
Malignant PNST	7	7
Lieomyosarcoma	7	8
Fibrosarcoma	7	3
Epithelioid sarcoma	3	1
Other	11	20

Gerrand et al: The influence of Anatomic location in patients with STS of the extremity. CANCER 2003 97(2): 485

#### Soft Tissue Tumours

#### Benign "lumps and bumps"

- Synovium : Ganglia, GCT tendon sheath, PVNS
- Fat : Lipoma
- Vascular : AV malformations, Haemangiomas, Glomus tumour
- Fibrous tissue : Fibroma, Fibromatosis
- Neural : Schwanoma, Neurofibroma
- Others : Post traumatic conditions, epidermal cysts, CMC Boss

#### Considerations specific to UL STS

#### > UL Vs LL

- Smaller lesions at presentation
- Less likely to be deep to or involving the investing fascia
- Higher rate of unplanned excision before referral
- Tumours of different histological types
- Higher rate of local recurrence
  - ?Related to:
  - Histological type
  - Unplanned excisions
  - Anatomy
  - Use of adjuvant modalities

#### Considerations specific to UL STS

Preservation of function is key consideration

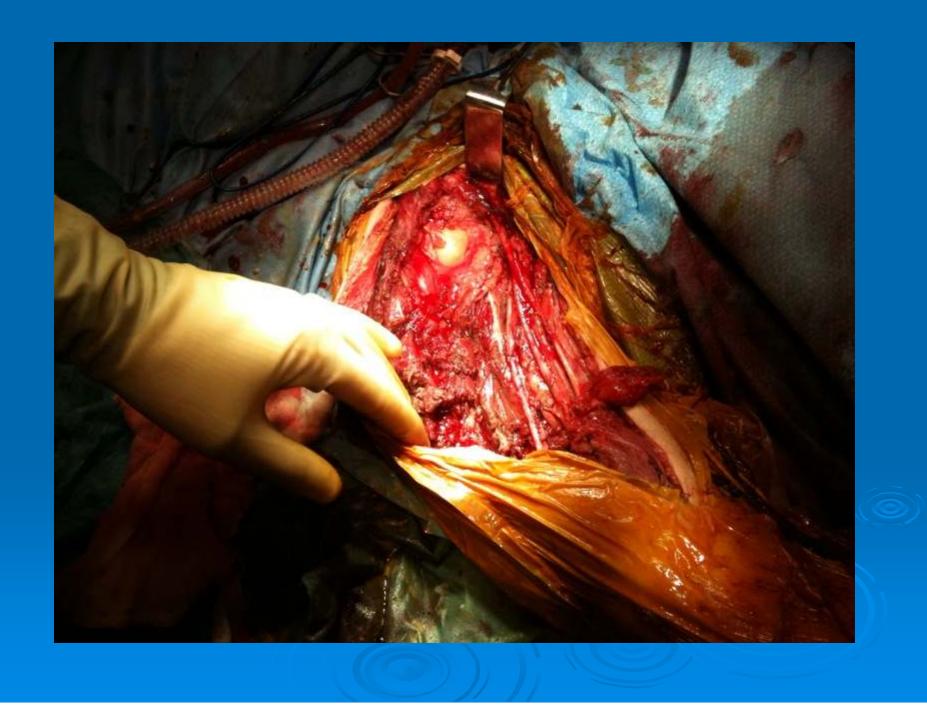
- Less likely to amputate
- Preference for WLE and reconstruction
- Treatment of 2ndry boney metastasis
  - UL not weight bearing
  - Can therefore consider use of conservative measures eg protection in sling, immobilisation for fractures etc

#### **Proximal humeral reconstruction**



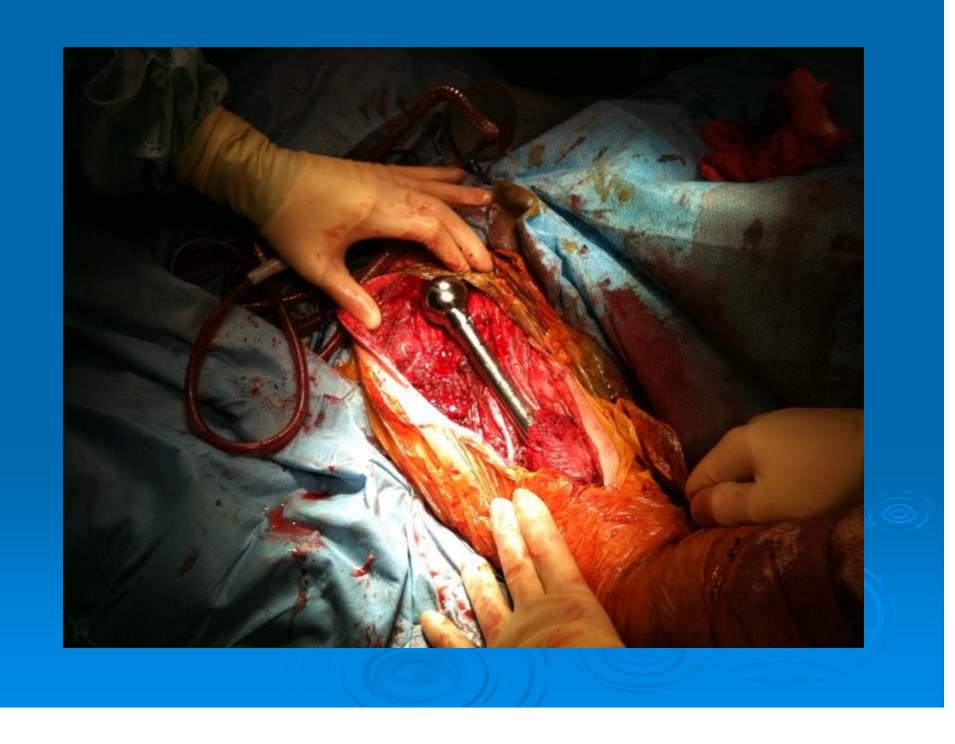
84 yo malePathological fracture1. Most likely diagnosis?2. What next?











## Reconstruction



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### Acknowledgements

#### MCQs adapted from:

 Review questions in Orthopaedics, Wright JM et al Lippincott Williams and Wilkins

Please look at www.Sarcoma.org for further information. There is a link to a free online text book (on the left hand side of the main page) from which the majority of the in information for this talk was taken and a lot of the pictures (which I unfortunately cannot put in to the uploaded version) also came form this resource