

Pathology Summary

Benign Bone Tumours

Can be incidental findings or locally very aggressive

1. Osteochondroma
 - a. Can be associated with Hereditary Multiple Osteochondromatosis
 - b. Portion of the physis 'breaks' away and grows off the side of the bone
 - c. Cauliflower-like appearance
 - d. Histology: cellular cartilage
 - e. Malignant transformation can occur – growth in adulthood requires investigation!
2. Giant Cell Tumour
 - a. Most common ages 20-40 years
 - b. 1% exhibit metastatic behaviour
 - c. 20% of all benign bone tumours
 - d. Peri-articular location
 - e. Mainstay of treatment is surgery (some role for anti-osteoclastic agents)
3. Aneurysmal Bone Cyst
 - a. Similar pattern to GCT, but is a distinct entity
 - b. Fluid-fluid levels on MRI scans
 - c. Mainstay of treatment is surgical, other modalities such as embolization are also used

Malignant Bone Tumours

Require early referral to a subspecialist multidisciplinary centre

1. Osteosarcoma
 - a. Most common primary bone sarcoma
 - b. Peak in young patients (10-25 years) and older patients (30% are >40 years)
 - c. Histologic hallmark is malignant cells producing abnormal osteoid
 - d. Metastasize hematogenously, predominantly to lungs, bone
 - e. Mainstay of treatment is chemotherapy and surgery (cure rates reported between 60-75%)
2. Chondrosarcoma
 - a. Can be primary or arise secondarily from lesions such as enchondromas or osteochondromas
 - b. Wide variation in metastatic potential
 - c. Can occur in any bone
 - d. Mainstay of treatment is surgical
3. Ewing Family Tumour
 - a. Characterized by small round blue cells on histology that have specific translocations
 - b. Most common ages 5-30 years
 - c. Often have a large soft tissue mass associated
 - d. Chemotherapy, radiotherapy and surgery all have a role in management

Benign Soft Tissue Tumours

1. Biology
 - a. Grow slowly, do not metastasize
 - b. Well differentiated appearance similar to normal tissue
2. Some Subtypes
 - a. Lipoma
 - b. Hemangioma
 - c. Schwannoma
 - d. Fibromatosis

Etc ...

Malignant Soft Tissue Tumours

1. Biology
 - a. Appear to recapitulate different tissue types
 - b. Histology shows usual features of malignancy (hypercellularity, pleomorphism, atypia and abnormal mitoses)
2. Some Subtypes
 - a. Liposarcoma
 - i. Several variants with differing behaviours
 - ii. Often spread lymphatically as well as hematogenously
 - b. Synovial Sarcoma
 - i. Most common young adult soft tissue sarcoma
 - ii. Misnomer – does not occur in the synovium!
 - iii. Has a specific translocation
 - c. Pleomorphic Undifferentiated Sarcoma
 - i. “The tumour previously known as MFH”
 - ii. High grade
 - iii. More common in older people

Metastatic Tumours

Tumours that have spread from elsewhere to the skeleton or soft tissues
Increasing problem

- a. Skeleton is the third most common site of metastasis
- b. 70% of bone metastases are painful
- c. Many different treatment options