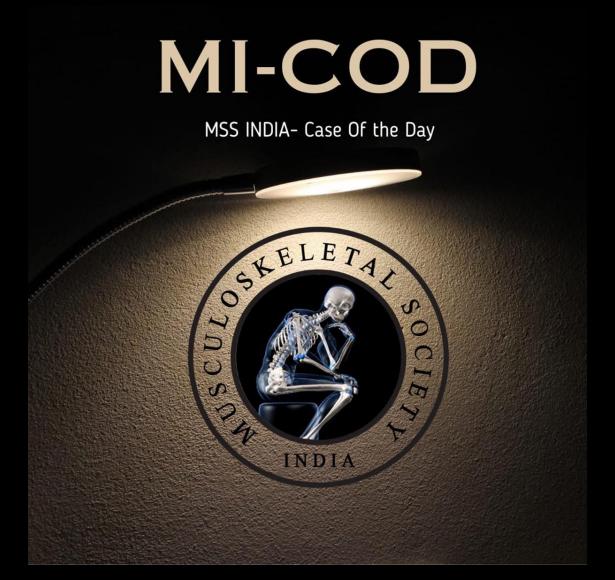
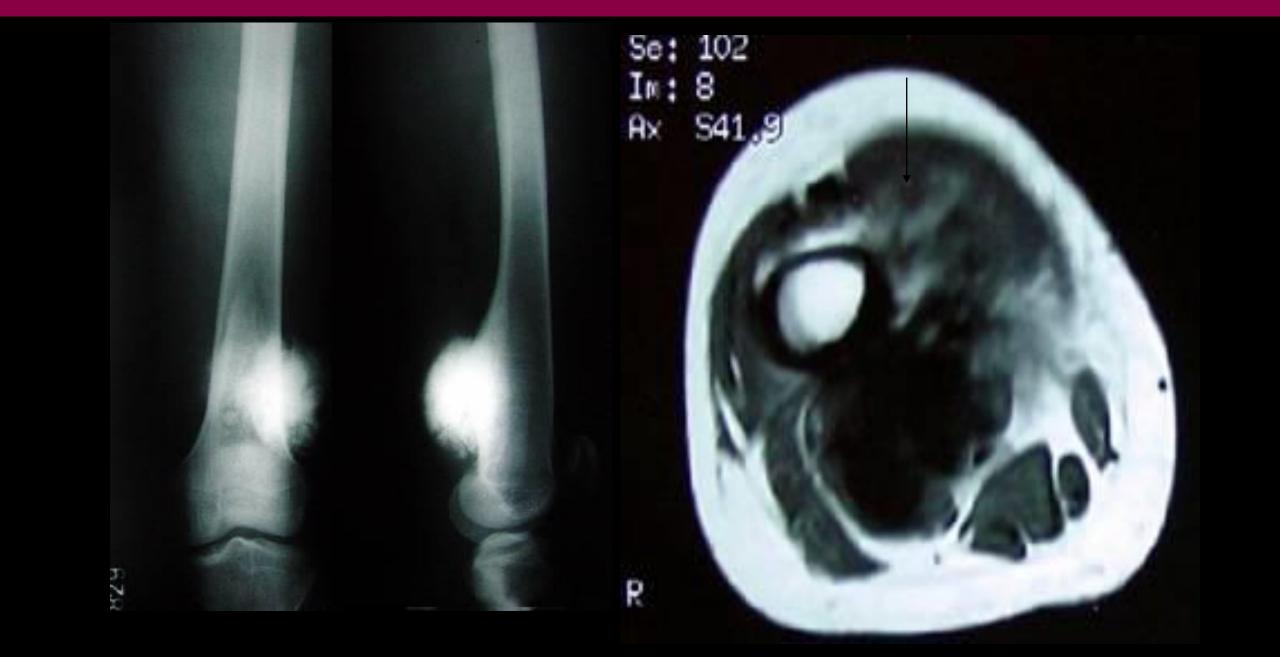
## MICOD – 24/06/2024 Case contributor – Dr. Nirmala Jambhekar

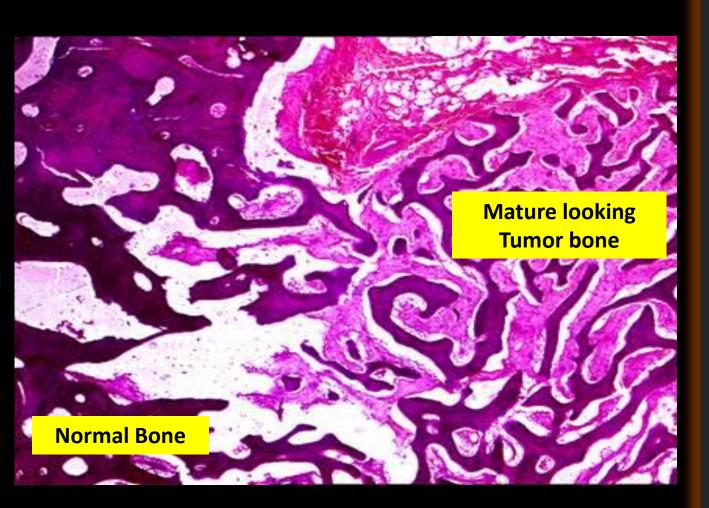


### Parosteal Osteosarcoma



#### *Kumar VS, Barwar N, Khan SA. Surface osteosarcomas: Diagnosis, treatment and outcome. Indian J Orthop. 2014 May*

### Parosteal Osteosarcoma



- Surface osteosarcomas are a rare form of osteosarcomas (OS) ~ 3-6% of all OS & occur at later age compared to conventional OS
- Three major groups of surface osteosarcomas
  - ✓ Parosteal OS (low grade & most common)
  - Periosteal OS (intermediate grade)
  - ✓ High grade surface OS
- Parosteal & periosteal OS are distinct clinical entities and it is important to identify the clinico radiological differences between the two types
- Classical site is lower end of the femur followed by the upper end of tibia and upper end of humerus
- Periosteal variant affects the tibia more commonly than the parosteal variety

Charecteristic	Parosteal osteosarcoma	Periosteal osteosarcoma	High grade surface osteosarcoma
Grade	Usually low grade	Intermediate grade	High grade
Age of affection	Between 20 and 30 years	Between 18 and 20 years	Between 20 and 30 years
Gender	More common in females	Common in males (as compared to parosteal type)	Common in males
Site of affection	Posterior aspect of the distal femur is the commonest site	Metaphysio-diaphyseal portion of the tibia	Mid femur, distal femur, mid tibia (in order)
Radiological picture	Sclerotic lesion over the surface of bone with thickening of the cortex and presence of a periosteal line between the tumor and the normal bone (string sign)	Broad based soft tissue mass with destruction of underlying bone with perpendicular periosteal reaction going into the soft tissue mass	Lesion attached to the host bone with a broad base with dense to moderate mineralization with a fluffy, immature appearance noted prominently at the base of the lesion. Lucent zone and perpendicular periosteal reaction are absent
Histopathology	Characterized by being well differentiated with malignant osteoid formation and a spindle cell stroma. Most tumors are Grade 1 lesions	Shows lobulated islands of malignant cartilage and areas of moderately high grade spindle cells located peripherally	Shows areas of malignant spindle cells with a high degree of cellular atypia (Grade 3 or 4 lesion) and variable amount of osteoid formation
Treatment	Minimal role of neo-adjuvant chemotherapy. Upfront surgery is the standard of care. Dedifferentiated tumors are high grade lesions that need to be treated like conventional osteosarcoma	Role of chemotherapy is controversial. Wide excision is the preferred local therapy	Neo-adjuvant chemotherapy followed by surgery

# Thank you