OSTEOGENIC SARCOMA

Hsin-Nung Shih M.D. PROFESSOR DIVISION OF JOINT RECONSTRUCTION DEPARTMENT OF ORTHOPEADIC CHANG GUNG MEMORIAL HOSPITAL

CHANG GUNG UNIVERSITY,COLLEGE OF MEDICINE TAIWAN

Osteogenic Sarcoma

Second most common primary malignant bone tumor Variable in its radiologic and morphologic presentation **CODE** may cause diagnostic confusion or mistaking it for a benign tumor























Osteogenic Sarcoma Capsule Summary

Incidence

Age

Signs Skeletal distribution 15% of primary bone tumors

15-25 Y/O (85%<30Yrs)

pain, swelling, pathologic fracture

- 54% knee rarely in spine, ribs and phalanges
- ✤ 90% metaphyseal
- 9% diaphyseal

Radiologic features

Gross pathology

Histology



Osteogenic Sarcoma Modern Classification

I. –	Primary, high-grade,	75%
	intramedullary OGS	
П.	Multifocal OGS	1-2%
Ш.	Secondary intramedullary OGS	5-7%
IV.	Solitary, low-grade,	4-5%
	Intramedullary	
V.	Intracortical OGS	0.2%
VI.	OGS of the bones of the jaw	6%
VII.	Juxtacortical OGS	7-10%
J.M.	Mirra 1989	

Osteogenic Sarcoma Serologic Findings: Alk-P

Alk-P (<50%) Initial</p>
After treatment
prognostic indicators



Osteogenic Sarcoma Differential Diagnosis

Callus Osteoblastoma Pseudomalignant osteoblastoma Aneurysmal bone cyst Chondroblastoma Giant cell tumor Ewing's sarcoma Chondrosarcoma Mesenchymal chondrosarcoma Fibrosarcoma



Osteogenic Sarcoma Clinical work-up and management

Systemic approach Pre-op bone scan Pre-op CT scan or MRI Biopsy Pre-op chemotherapy Radical surgery Post-op chemotherapy



Principle of OGS Treatment

Pre-op evaluation Plain X-ray: local + chest □ Chest CT □ Local: CT scan or MRI Bone scan Biopsy Neoadjuvant chemotherapy Radical surgery Amputation or Limb-salvage surgery Post-operative chemotherapy Other treatment

Osteogenic Sarcoma Clinical Course

High-grade biologic malignancy
 85% lung metastasis

 (diagnosis ± surgical intervention)
 Die within 2 yrs without
 chemotherapy (± Radiotherapy)



Osteogenic Sarcoma Consideration of Limb Salvage

Age
Staging
Location
Sizing

Grading
Biopsy wound
Pathologic fracture
Reconstructive material

***Technique Demand**



Orthopedic Oncology

Local control of non-metastatic
Classic high-grade osteosarcoma

Local therapy20%+ chemotherapy70-90%

disease-free survival > 5 yrs 50-70%

1970-1990 Rosen G.



Orthopedic Oncology

Limb-salvage vs. Amputation Osteosarcoma N=227, distal femur *Local recurrence Similarity *Survival rates No difference *Indications

MA SIMON, HJ MANKIN 1986, JBJS























Resection Arthrodesis of the Knee for Osteosarcoma: An Alternative When Mobile Joint Reconstruction Is Not Feasible

Hsin-Nung Shih, MD; Lih-Yuann Shih, MD

(Chang Gung Med J 2005;28:411-20)

Background:

Wide resection and mobile joint reconstruction are preferable for treating an osteosarcoma around the knee. In certain situations, resection arthrodesis or an amputation is suggested.

Methods:

he past decade, 86 patients with an osteosarcoma around the knee were treated surgically in our institution. Wide resection and endoprosthetic reconstruction were performed in 35 patients, resection arthrodesis was performed in 36 patients, and an amputation was performed in 15 patients. The oncological and functional results were compared. Special attention was paid to the indications, techniques, and complications of patients receiving resection arthrodesis.



Results:

Extensive tumor involvement was the main reason, followed by inappropriate previous treatment, for precluding mobile joint reconstruction. The local recurrence rates were similar among the 3 groups (11.4% for the endoprosthetic group, 11.1% for the arthrodesis group, and 6.7% for the amputation group). The 5-year survival rate was 39% for the arthrodesis group, which was significantly lower than that of the endoprosthetic group (60%, p = 0.040), although it was higher than that of the amputation group (13%, p = 0.056). Major complications were found in 7 patients receiving resection arthrodesis (7/24, 29%), and these included nonunion, infection, and allograft fracture. Functional results for the arthrodesis patients were inferior to those of the endoprosthetic patients, but most patients were grateful for preservation of the limb despite certain handicaps.

Conclusions:

The importance of early and proper planning of treatment cannot be overstressed when treating osteosarcomas. Resection arthrodesis offers a durable reconstruction alternative to amputation in a special group of patients when extensive resection precludes mobile joint reconstruction.































Transient neurological disturbances induced by the chemotherapy of high-dose methotrexate for osteogenic sarcoma

Klu, Mee-Chou; Liaw, Chuang-Chl; Yang, Tsai-Shen; Lai, Gi-Ming; Hsl, Shin-Nun; Lu, Chin-Song

Anti-Cancer Drugs 1994,5,p.480-482

Temporary neurologic abnormalities were observed In one out of 23 patients undergoing chemotherapy with high-dose methotrexate (HD-MTX) for osteogenic sarcoma. This patient developed sequential symptoms including alternative hemlparesis, dysarthria and altered consciousness 5 days after the second course of HD-MTX (8 gm/m2 by 6 h continuous Infusion) with leucov-orin rescue. Laboratory evaluations disclosed normal electrolytes, hemograms and non-toxic serum MTX levels at the onset of the symptoms. Computed tomography of the brain was normal but electroencephalography showed focal theta and delta slow waves over the right temporalparietal-occlpital area. The neurological symptoms resolved completely within 72 h.

Synchronous multifocal osteosarcoma: report of one case.

Acta Paediatr Taiwan. 2006 May-Jun;47(3):146-9.

Tsai MH, Yang CP, Jaing TH, Shih HN.

Synchronous multifocal osteosarcoma (SMOS), defined as more than one bone lesion at presentation, is a rare variant form of osteosarcoma. The onset is usually in childhood or early adolescence without pulmonary metastasis. The prognosis has been dismal. Whether SMOS represents a true multicentic origin or merely **bone-to-bone metastases remains controversial.** Here, we report a case of SMOS in a 10-year-old girl, with the dominant primary sclerotic tumor arising from the right distal femur. Despite aggressive chemotherapy and limb salvage surgery, she died of progressive multiple axial skeletal and symmetrical metaphyseal long bone diseases within one year after diagnosis. No pulmonary metastasis was found before she died.



Biochemical Marker of Bone Metabolism

Markers of bone formation	Markers of bone resorption	
Serum	Urine	Serum
Alkaline phosphatase (ALP)	Calcium	Cross-linked Carboxyterminal telopeptide type I collagen (ICTP)
Bone-specific alkaline phosphatase	Hydroxyproline	
Osteocalcin	Pyridinoline and Deoxypyridinoline	Other non-collagenous proteins (?)
Procollagen I C-terminal extension peptide (PICP)	Cross-linked aminoterminal telopeptide type I collagen (INTP)	
Other non-collagenous proteins (?)		



Osteogenic Sarcoma Treatment

A Team Work





CHANG-GUNG MEMORIAL HOSPITAL LINKOU MEDICAL CENTER TAIWAN



