Opis choroby *

Definicja

A rare bone tumor characterized by a benign lesion composed of lobules of spindle shaped or stellate cells and an abundant myxoid or chondroid matrix. The tumor may occur in almost any osseous location but is most common in long bones, in particular the proximal tibia and the distal femur. Pain is the most common presenting symptom. Prognosis is excellent even in cases with local recurrence.

Dane

Klasyfikacja
Choroba

Kod ORPHA
404507

Kod OMIM
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Kod ICD10
D16.9

Kod ICD11
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*Źródło

orphanet