Opis choroby *

Definicja

A rare bone sarcoma characterized by a usually benign space-occupying lesion, which is nevertheless locally aggressive and massively damaging to surrounding bone tissue. The tumor is composed of giant multinucleated cells (osteoclast-like cells), mononuclear macrophages, and mononuclear stromal cells which secrete pro-myeloid and pro-osteoclastic factors. Metastasis and malignant transformation are rare, but the recurrence rate is high.

Dane

Klasyfikacja
Choroba

Synonimy
GCT of bone
GCT kości
Osteoclastoma

Kod ORPHA
363976

Kod OMIM
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Kod ICD10
D48.0

Kod ICD11
2F9B

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*Źródło

orphanet