## 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** 

**Kod ICD10** 

\_

D48.0

Kod ICD11

2F9B

## \*Źródło

orphanet