Rodzinny włókniak kostniejący
Kod Orpha: 435329 Kod OMIM: 137575

Opis choroby *

Definicja
A rare genetic bone disease characterized by multifocal, painless, benign fibrocemento-osseous lesions of the jaws which expand progressively and can cause severe facial deformity. It usually manifests at an early age and is often associated with abnormalities of the long bones and pathologic fractures. Radiologically, the lesions are of mixed radiopaque/radiolucent appearance. Incomplete surgical removal may lead to more rapid growth of the residual lesion.

Dane

Klasyfikacja
Choroba

Synonimy
Multiple ossifying fibroma
Familial Gigantiform cementoma
Multiple ossifying fibroma

Kod ORPHA
435329

Kod OMIM
137575

Kod ICD10
D16.4

Kod ICD11
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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.