Genochondromatoza typu 2
Kod Orpha: 93398 Kod OMIM: 137360

Opis choroby *

Definicja
Genochondromatosis type 2 is a rare genetic bone development disorder characterized by normal clavicles and symmetrical, generalized metaphyseal enchondromas, particularly in the distal femur, proximal humerus, and bones of the wrists, hands, and feet. Lesions regress later in life with growth cartilage obliteration. Clinical examination is normal and the course of the disease is benign.
Dane

Klasyfikacja
Choroba

Kod ORPHA
93398

Kod OMIM
137360

Kod ICD10
Q78.4

Kod ICD11
LD24.2Y

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.