

# 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

---

[\\*Źródło](#)

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.