Opis choroby *

Definicja

An extremely rare familial bone deformity described only in Japanese patients to date. The deformity is bilateral in nearly half of patients (with bilateral involvement, the condition is symmetrical) and sometimes causes ulnar nerve palsy or cubitus varus.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
3383

Kod OMIM
191000

Kod ICD10
Q74.0

Kod ICD11
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*Źródło

orphanet