

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