Opis choroby *

Definicja

A very rare syndrome characterized by progressive loss of bone, usually the capsal and tarsal bones, resulting in deformity and disability, as well as chronic renal failure in many cases. The bone and renal disorders are sometimes associated with intellectual deficit and facial abnormalities.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Idiopathic multicentric osteolysis with or without nephropathy
Idiopatyczna wielocentryczna osteoliza z lub bez nefropatii

Kod ORPHA
2774

Kod OMIM
166300

Kod ICD10
M89.5

Kod ICD11
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*Źródło

orphanet