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 Synonimy

Zespół wad wrodzonych Idiopathic multicentric osteolysis with or without

nephropathy

Idiopatyczna wielocentryczna osteoliza z lub bez

nefropatii

 Kod ORPHA
 Kod OMIM
 Kod ICD10

 2774
 166300
 M89.5

Kod ICD11

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*Źródło

orphanet