Opis choroby *

Definicja

A rare ectodermal dysplasia syndrome characterized by the association of amelogenesis imperfecta and trichodysplasia with symmetrical pits in the cuticles of hair shafts. There have been no further descriptions in the literature since 1993.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
79129

Kod OMIM
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Kod ICD10
Q82.4

Kod ICD11
LD27.0Y

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*Źródło

orphanet