

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

*Źródło

orphanet