Opis choroby *

Definicja

A rare ectodermal dysplasia syndrome characterized by a variably severe clinical picture comprising dry, thin skin, onychodysplasia, trichodysplasia, and dental abnormalities (such as hypodontia, microdontia, and persistence of deciduous teeth). There have been no further descriptions in the literature since 1990.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
1660

Kod OMIM
125640

Kod ICD10
Q82.4

Kod ICD11
LD27.0Y

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*Źródło

orphanet