Opis choroby *

Definicja

A rare, genetic, ectodermal dysplasia syndrome characterized by dental abnormalities (primarily agenesis of the permanent and deciduous teeth with cone-shaped incisors and canines), onychodysplasia, palmoplantar hyperkeratosis, dry skin and, more variably, hypotrichosis, and sweat gland dysfunction (hyper- or hypohidrosis).

Dane

Klasyfikacja
Choroba

Synonimy
OODD
OODD

Kod ORPHA
2721

Kod OMIM
257980

Kod ICD10
Q82.4

Kod ICD11
LD27.0Y

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*Źródło

orphanet