

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

*Źródło

orphanet