

## 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