

## **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