

## Opis choroby \*

### Definicja

A syndromic genetic deafness characterized by erythrokeratoderma, hypotrichosis, nail dystrophy and sensorineural hearing loss. Erythema, recurrent skin infections and mucositis have also been associated.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Hypotrichosis-hearing loss syndrome

Hypotrichosis-hearing loss syndrome

#### Kod ORPHA

330029

#### Kod OMIM

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#### Kod ICD10

H90.5

#### Kod ICD11

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#### \*Źródło

orphanet