

## Opis choroby \*

### Definicja

A rare ectodermal dysplasia syndrome characterized by hypotrichosis of scalp and eyebrows, finger syndactyly, intellectual disability and early eruption of teeth. Facial dysmorphism (i.e. round face with prominent forehead, cheeks and ears, and upward-slanting palpebral fissures), hypoplasia of median and distal phalanges, and kyphosis are additionally observed features. There have been no further descriptions in the literature since 1996.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Lopes-Marques de Faria syndrome

Zespół Lopesa i Marques de Faria

#### Kod ORPHA

2266

#### Kod OMIM

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#### Kod ICD10

Q82.4

#### Kod ICD11

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#### \*Źródło

orphanet