

## Opis choroby \*

### Definicja

A rare genetic skin disease characterized by infantile onset of diffuse alopecia, abnormal skin pigmentation (hypo- and hyperpigmented macules of the trunk and face and areas of reticular hypo- and hyperpigmentation of the extremities), palmoplantar keratoderma, and nail dystrophy. Patients develop recurrent spinocellular carcinomas later in life. Brittle teeth resulting in early loss of dentition have also been described.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

447961

#### Kod OMIM

618373

#### Kod ICD10

D04.8

#### Kod ICD11

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### \*Źródło

orphanet