Opis choroby *

Definicja

A rare, genetic, ectodermal dysplasia characterized by a widespread, early-onset, reticulate hyperpigmentation that persists throughout life, mild, diffuse non-cicatricial alopecia, and onychodystrophy. There are no dental anomalies. Patients may also present with adermatoglyphia, palmoplantar hyperkeratosis, acral dorsal blistering, and hypohidrosis or hyperhidrosis.

Dane

Klasyfikacja
Choroba

Kod ORPHA
86920

Kod OMIM
125595

Kod ICD10
Q82.4

Kod ICD11
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*Źródło

orphanet