Opis choroby *

Definicja

A rare, genetic keratinization disorder which is classically characterized by keratotic papules, acral pits, and acral wart-like lesions that can be associated with a trigger, and may occur anywhere on the body (including mucosal surfaces). Extracutaneous manifestations may include, nail anomalies, blepharitis, dry eye, neuropsychiatric illness and, recurrent parotid gland obstruction and xerostomia.

Dane

Klasyfikacja
Choroba

Synonimy
Darier-White disease
Choroba Dariera i White
Rogowacenie pęcherzykowe
Keratosis follicularis

Kod ORPHA
218

Kod OMIM
124200

Kod ICD10
Q82.8

Kod ICD11
EC20.2

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*Źródło

orphanet