

Opis choroby *

Definicja

A rare, genetic, acrokeratoderma disease characterized by multiple, symmetrical, asymptomatic, skin-colored (rarely, brownish), flat-topped, wart-like papules located on the dorsal aspects of the hands and feet (occasionally found on other parts of the body, such as knees, elbows and forearms), typically associated with palmoplantar punctate keratosis and variable nail involvement (including leukonychia, thickening, ridging, longitudinal striations and splitting). Histology reveals undulating hyperkeratosis, papillomatosis, hypergranulosis, and acanthosis, creating a characteristic 'church spire' appearance, with no acantholysis nor dyskeratosis associated.

Dane

Klasyfikacja

Choroba

Synonimy

AKV of Hopf

AKV of Hopf

Kod ORPHA

79151

Kod OMIM

101900

Kod ICD10

Q82.8

Kod ICD11

EC20.Y

[*Źródło](#)

orphanet