Opis choroby *

Definicja

Hypotrichosis-osteolysis-periodontitis-palmoplantar keratoderma syndrome is an extremely rare ectodermal dysplasia syndrome characterized by hypotrichosis universalis with mild to severe scarring alopecia, acro-osteolysis, onychogryphosis, thin and tapered fingertips, periodontitis and caries leading to premature teeth loss, linear or reticular palmoplantar keratoderma and erythematous, scaling, psoriasis-like skin lesions on arms and legs. Lingua plicata and ventricular tachycardia have also been observed.

Dane

Klasyfikacja
Choroba

Synonimy
HOPP syndrome
Zespół hipotrichoza-osteoliza-zapalenie przyzębia-hiperkeratoza dłoniowo-podeszwowa
Zespół HOPP
Hypotrichosis-osteolysis-periodontitis-palmoplantar hyperkeratosis syndrome
Hypotrichosis-striate palmoplantar hyperkeratosis-acroosteolysis-periodontitis syndrome
Hypotrichosis-striate palmoplantar keratoderma-acroosteolysis-periodontitis syndrome

Kod ORPHA
307936

Kod OMIM
607658

Kod ICD10
Q82.8

Kod ICD11
EC20.31

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*Źródło

orphanet