

Opis choroby *

Definicja

A rare genetic skin disease characterized by the triad of congenital scleroatrophy predominantly of the hands with sclerodactyly, palmoplantar keratoderma, and nail changes (consisting of hypoplasia, ridging, clubbing, and white discoloration). Additional features include palmar hypohidrosis and a high susceptibility to early-onset squamous cell carcinoma of affected skin areas.

Dane

Klasyfikacja

Choroba

Synonimy

Palmoplantar hyperkeratosis-sclerodactyly syndrome

Scleroatrophic syndrome

Zespół hiperkeratoza dłoniowo-podeszwowa - sklerodaktylia

Zespół Hurieza

Palmoplantar keratoderma-sclerodactyly syndrome

Scleroatrophic syndrome

Sclerotylosis

Kod ORPHA

384

Kod OMIM

181600

Kod ICD10

Q82.8

Kod ICD11

EC20.30

*Źródło

orphanet