Opis choroby *

Definicja

A rare bone disease characterized by bone resorption affecting the distal phalanx, most commonly the terminal tuft, in the absence of a known cause. Patients present with shortening of the affected fingers or toes, associated with nail abnormalities (dystrophic or hypertrophic nails) and skin changes (such as ulceration or pigment anomalies).

Dane

Klasyfikacja
Choroba

Synonimy
Idiopathic phalangeal acroosteolysis
Idiopathic phalangeal acroosteolysis

Kod ORPHA
444316

Kod OMIM
-

Kod ICD10
M89.5

Kod ICD11
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*Źródło

orphanet