Opis choroby *

Definicja

A rare, genetic, chronic, recurrent, slowly progressive, epidermal disease characterized by small, sterile, pustular eruptions, involving the nails and surrounding skin of the fingers and/or toes, which coalesce and burst, leaving erythematous, atrophic skin where new pustules develop. Onychodystrophy is frequently associated and anonychia and osteolysis are reported in severe cases. Local expansion (to involve the hands, forearms and/or feet) and involvement of mucosal surfaces (e.g. conjunctiva, tongue, urethra) may be observed.

Dane

Klasyfikacja
Choroba

Kod ORPHA
163931

Kod OMIM
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Kod ICD10
L40.2

Kod ICD11
EA90.41

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*Źródło

orphanet