Dziedziczna poikilodermia z akrokeratozą
Kod Orpha: 2907 Kod OMIM:

Opis choroby *

Definicja
A rare hereditary poikiloderma characterized by infantile onset of vesicopustule formation on hands and feet and widespread eczematoid dermatitis (both spontaneously resolving during childhood), as well as gradually developing diffuse poikiloderma with striate and reticulate atrophy (excluding the face, scalp, and ears), and development of keratotic papules on hands, feet, elbows, and knees, beginning in early childhood. There have been no further descriptions in the literature since 1981.

Dane

Klasyfikacja
Choroba

Synonimy
Weary syndrome
Wrodzona poikilodermia z pęcherzami, typu Weary

Kod ORPHA
2907

Kod OMIM
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Kod ICD10
Q82.8

Kod ICD11
LD2B

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.