Dziedziczna poikilodermia stwardniająca, typ Weary'ego
Kod Orpha: 221039 Kod OMIM: 173700

Opis choroby *

Definicja
A rare genetic skin disease characterized by generalized poikiloderma with marked accentuation in flexural regions and on extensor surfaces, sclerosis of palms and soles, and linear and reticulated hyperkeratotic and sclerotic bands in the axilla and the antecubital and popliteal fossae. Subcutaneous calcification, finger clubbing, Raynaud phenomenon, and cardiac abnormalities (such as severe aortic stenosis) have also been reported.
Dane

Klasyfikacja
Choroba

Kod ORPHA
221039

Kod OMIM
173700

Kod ICD10
Q82.8

Kod ICD11
EC10

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.