Opis choroby *

Definicja

A rare vascular anomaly characterized by congenital, solitary or grouped, red-to-purple plaques which may bleed and enlarge over time. The lesions show a predilection for the lower extremities. Histological examination reveals numerous dilated, congested capillaries and venules in the papillary dermis, often with a deep dermal component, and with increased density of variably congested capillaries and venules also in the subcutaneous tissue. The overlying epidermis displays prominent acanthosis, papillomatosis, hyperkeratosis, parakeratosis, and crusting.

Dane

Klasyfikacja
Choroba

Kod ORPHA
464318

Kod OMIM
-

Kod ICD10
D18.0

Kod ICD11
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*Źródło

orphanet