

Opis choroby *

Definicja

A rare skin disease characterized by widespread cutaneous telangiectases usually appearing on the lower limbs and slowly progressing upwards to involve the trunk and arms. The lesions can be diffuse, localized, macular, plaque-like, discrete, or confluent. Recurrent bleeding from the skin and mucous membranes is not a common feature. Likewise, co-existing epidermal or dermal abnormalities, like atrophy, depigmentation, or purpura, are absent. The condition is non-hereditary, and to establish the diagnosis, other primary and secondary telangiectases must be excluded.

Dane

Klasyfikacja

Choroba

Synonimy

GET

GET

Kod ORPHA

280774

Kod OMIM

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Kod ICD10

L98.8

Kod ICD11

EF20.4

*Źródło

orphanet