Opis choroby *

Definicja

A rare skin disease characterized by widespread cutaneous telangiectases usually ﬁrst 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
-

Kod ICD10
L98.8

Kod ICD11
EF20.4

---

*Źródło

orphanet