Opis choroby *

Definicja

A rare congenital hemangioma characterized by a superficial, red to violaceous lesion with overlying telangiectasia and a surrounding pale halo, which initially behaves like a rapidly involuting congenital hemangioma, beginning to involute shortly after birth. Involution is then aborted, and a residual tumor virtually indistinguishable from non-involuting congenital hemangioma remains. This lesion grows proportionally with the child and does not regress.

Dane

Klasyfikacja
Choroba

Kod ORPHA
458785

Kod OMIM
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Kod ICD10
D18.0

Kod ICD11
2E81.2Y

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*Źródło

orphanet