

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

-

Kod ICD10

D18.0

Kod ICD11

2E81.2Y

*Źródło

orphanet