

Opis choroby *

Definicja

A rare genetic vascular anomaly characterized by the presence of angiomatous lesions affecting the skin, brain, and spinal cord. Lesions of the central nervous system have a marked tendency to bleed. There have been no further descriptions in the literature since 1988.

Dane

Klasyfikacja

Choroba

Kod ORPHA

1062

Kod OMIM

106070

Kod ICD10

D18.0

Kod ICD11

-

[*Źródło](#)

orphanet