Opis choroby *

Definicja

A rare subtype of cerebrofacial arteriovenous metameric syndrome characterized by unilateral arteriovenous malformations involving the hypothalamus and nasal region (medial prosencephalic group). The condition manifests in childhood. Common presenting signs and symptoms are progressive neurological deficit, hemorrhage, and cosmetic complaints like facial asymmetry.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
CAMS1
CAMS1

Kod ORPHA
141194

Kod OMIM
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Kod ICD10
Q28.2

Kod ICD11
LA90.3Y

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*Źródło

orphanet