Opis choroby *

Definicja

A rare neurovascular malformation characterized by a unilateral, direct communication between the arterial and venous system in the retina via abnormal, enlarged vessels, but without interposed capillaries. The inferotemporal vasculature is most commonly affected. Patients may be asymptomatic or present with variable degrees of visual loss. Local vascular complications include vascular occlusions or retinal or vitreous hemorrhages. The anomaly may occur in isolation or as part of Wyburn-Mason syndrome, in which intracranial (usually ipsilateral) arteriovenous malformations are present.

Dane

Klasyfikacja
Wada morfologiczna

Synonimy
Congenital arteriovenous anastomoses of the retina
Wrodzone anastomozy tętniczo-żylne w siatkówce
Congenital arteriovenous communication of the retina
Congenital retinal arteriovenous anastomoses

Kod ORPHA
353334

Kod OMIM
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Kod ICD10
Q14.1

Kod ICD11
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*Źródło

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