## Opis choroby *

Definicja

A rare genetic cerebral small vessel disease characterized by isolated marked tortuosity of second-order and third-order retinal arteries with normal first-order arteries and venous system, typically located in the macular and peripapillary area and developing during childhood or early adulthood. The disease may be asymptomatic, although most patients present variable degrees of transient vision loss due to retinal hemorrhage following physical exertion or minor trauma.

Dane

**Klasyfikacja**
Choroba

Synonimy
Familial isolated retinal arterial tortuosity
Krętość tętnic siatkówki
Krwotok w siatkówce z krętością naczyń
Retinal arteriolar tortuosity
Retinal hemorrhage with vascular tortuosity
Tortuosity of retinal arteries

**Kod ORPHA**
75326

**Kod OMIM**
180000

**Kod ICD10**
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**Kod ICD11**
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*Źródło

orphanet