

## 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