

## Opis choroby \*

### Definicja

A rare genetic neurovascular malformation characterized by sac-like bulging of cerebral arteries due to weakening of the endothelial layer. Familial occurrence is suspected when two or more affected first- to third-degree relatives are present in a family. Aneurysms may remain asymptomatic throughout life, or rupture and thereby cause potentially life-threatening subarachnoid hemorrhage. Patients with familial cerebral saccular aneurysm are more likely to develop more than one brain aneurysm, are at greater risk of rupture, and tend to have poorer outcome after rupture than patients with sporadic cerebral aneurysms.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Familial berry aneurysm  
Rodzinny tętniak w kształcie jagody  
Rodzinny wewnątrzczaszkowy tętniak  
woreczkowaty  
Familial intracranial saccular aneurysm

#### Kod ORPHA

231160

#### Kod OMIM

614252

#### Kod ICD10

I67.1

#### Kod ICD11

8B22.6

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#### [\\*Źródło](#)

orphanet