DISEASE:
Familial cerebral saccular aneurysm

NAME:	Familial cerebral saccular aneurysm
DESCRIPTION:	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.
ORPHACODE:	231160
SYNONYMS:	Familial berry aneurysm Familial intracranial saccular aneurysm

XREF(S):	Orphanet OMIM ICD-10 OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM
ANALYTE(S):	ANGPTL6 COL3A1 ENG TGFBR3 THSD1
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- Centre de Génétique Médicale UCL
- Centrum Menselijke Erfelijkheid - KUL

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- transforming growth factor beta receptor 3
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