DISEASE:
HANAC syndrome

NAME:	HANAC syndrome
DESCRIPTION:	A rare multisystemic disease characterized by small-vessel brain disease, cerebral aneurysm, and extracerebral findings involving the kidney, muscle, and small vessels of the eye.
ORPHACODE:	73229
SYNONYMS:	Autosomal dominant familial hematuria-retinal arteriolar tortuosity-contractures syndrome Hereditary angiopathy-nephropathy-aneurysms-muscle cramps syndrome
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	COL4A1
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Porencephaly / Hemorrhagic stroke / Cerebral small vessel disease / Idiopathic cerebral white matter lesions / HANAC / Isolated retinal arteriolar tortuosity

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Gent

Related Analytes

- collagen type IV alpha 1 chain

Related Gene Panels

- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Porencephaly; Hemorrhagic stroke; Cerebral small vessel disease; Idiopathic cerebral white matter lesions; HANAC; Isolated retinal arteriolar tortuosity - UGent