DISEASE:
Cerebral autosomal dominant arteriopathy-subcortical infarcts-leukoencephalopathy

NAME:	Cerebral autosomal dominant arteriopathy-subcortical infarcts-leukoencephalopathy
DESCRIPTION:	CADASIL (Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy) is a hereditary cerebrovascular disorder characterized by mid-adult onset of recurrent subcortical ischemic stroke and cognitive impairment progressing to dementia in addition to migraines with aura and mood disturbances seen in about a third of patients.
ORPHACODE:	136
SYNONYMS:	CADASIL Hereditary multi-infarct dementia
XREF(S):	Orphanet OMIM MeSH MedDRA ICD-10
ANALYTE(S):	NOTCH3
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Source URL: http://gentest.healthdata.be/disease/525

RELATED CONTENT

Related Genetic Tests

- Cadasil (exons of EGFL domains (2 - 24))
- Neurodegeneration (gene panel)

Related Laboratories

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

Related Analytes

- notch receptor 3

Related Gene Panels

- Neurodegeneration (99 genes) - IPG