DISEASE:
Autosomal dominant cerebellar ataxia-deafness-narcolepsy syndrome

NAME:	Autosomal dominant cerebellar ataxia-deafness-narcolepsy syndrome
DESCRIPTION:	A rare polymorphic disorder, subtype of autosomal dominant cerebellar ataxia type 1 (ADCA type 1), characterized by ataxia, sensorineural deafness and narcolepsy with cataplexy and dementia.
ORPHACODE:	314404
SYNONYMS:	ADCA-DN syndrome Autosomal dominant cerebellar ataxia-hearing loss-narcolepsy syndrome
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	DNMT1
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RELATED CONTENT

Related Genetic Tests

- Hereditary Spastic Paraplegia (gene panel)
- Neuropathy (gene panel)

Related Laboratories

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

Related Analytes

- DNA methyltransferase 1

Related Gene Panels

- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Neuropathy (148 genes) - IPG