DISEASE:
Hereditary sensory neuropathy-deafness-dementia syndrome

NAME:	Hereditary sensory neuropathy-deafness-dementia syndrome
DESCRIPTION:	A rare genetic neurological disorder characterized by sensorineural hearing loss, sensory neuropathy, behavioral abnormalities, and dementia. Occurrence of seizures has also been reported. Age of onset is between adolescence and adulthood. The disease is progressive, with fatal outcome typically in the fifth to sixth decade.
ORPHACODE:	456318
SYNONYMS:	HSAN1E HSN1E Hereditary sensory neuropathy-sensorineural hearing loss-dementia syndrome
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	DNMT1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Neuropathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- DNA methyltransferase 1

Related Gene Panels

- Neuropathy (148 genes) - IPG