DISEASE:
Hereditary sensory and autonomic neuropathy type 1

NAME:	Hereditary sensory and autonomic neuropathy type 1
DESCRIPTION:	Hereditary sensory neuropathy type I (HSN I) is a slowly progressive neurological disorder characterised by prominent predominantly distal sensory loss, autonomic disturbances, autosomal dominant inheritance, and juvenile or adulthood disease onset.
ORPHACODE:	36386
SYNONYMS:	HSAN1 Hereditary sensory and autonomic neuropathy type I
XREF(S):	Orphanet OMIM OMIM OMIM OMIM ICD-10
ANALYTE(S):	ATL1 SPTLC1 SPTLC2 ATL3
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RELATED CONTENT

Related Genetic Tests

- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Neuropathy (gene panel)
- Spastic Paraplegia (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- atlastin GTPase 1
- atlastin GTPase 3
- serine palmitoyltransferase long chain base subunit 1
- serine palmitoyltransferase long chain base subunit 2

Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- Neuropathy (148 genes) - IPG
- Spastic Paraplegia (89 genes) - IPG

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