DISEASE:
Hereditary sensory and autonomic neuropathy type 2

NAME:	Hereditary sensory and autonomic neuropathy type 2
DESCRIPTION:	A rare hereditary sensory and autonomic neuropathy characterized by profound and universal sensory loss involving large and small fiber nerves.
ORPHACODE:	970
SYNONYMS:	Autosomal recessive sensory radicular neuropathy HSAN2 Hereditary sensory and autonomic neuropathy type II Neurogenic acroosteolysis
XREF(S):	Orphanet OMIM OMIM OMIM OMIM ICD-10
ANALYTE(S):	SCN9A WNK1 RETREG1 KIF1A

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RELATED CONTENT

Related Genetic Tests

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

Related Laboratories

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

Related Analytes

- kinesin family member 1A
- reticulophagy regulator 1
- sodium voltage-gated channel alpha subunit 9
- WNK lysine deficient protein kinase 1

Related Gene Panels

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