DISEASE:
Autosomal dominant Charcot-Marie-Tooth disease type 2K

NAME:	Autosomal dominant Charcot-Marie-Tooth disease type 2K
DESCRIPTION:	An axonal Charcot-Marie-Tooth (CMT) peripheral sensorimotor polyneuropathy.
ORPHACODE:	99944
SYNONYMS:	CMT2K
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	GDAP1
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RELATED CONTENT

Related Genetic Tests

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

Related Laboratories

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

Related Analytes

- ganglioside induced differentiation associated protein 1

Related Gene Panels

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