DISEASE:
Charcot-Marie-Tooth disease type 2P

NAME:	Charcot-Marie-Tooth disease type 2P
DESCRIPTION:	Charcot-Marie-Tooth disease type 2P is a rare, genetic, axonal hereditary motor and sensory neuropathy disorder characterized by adulthood-onset of slowly progressive, occasionally asymmetrical, distal muscle weakness and atrophy (predominantly in the lower limbs), pan-modal sensory loss, muscle cramping in extremities and/or trunk, pes cavus and absent or reduced deep tendon reflexes. Gait anomalies and variable autonomic disturbances, such as erectile dysfunction and urinary urgency, may be associated.
ORPHACODE:	300319
SYNONYMS:	CMT2P
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	LRSAM1
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RELATED CONTENT

Related Genetic Tests

- Neuropathy (gene panel)

Related Laboratories

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

Related Analytes

- leucine rich repeat and sterile alpha motif containing 1

Related Gene Panels

- Neuropathy (148 genes) - IPG