DISEASE:
Charcot-Marie-Tooth disease type 4G

NAME:	Charcot-Marie-Tooth disease type 4G
DESCRIPTION:	Charcot-Marie-Tooth disease type 4G (CMT4G) is a subtype of Charcot-Marie-Tooth disease type 4 characterized by early childhood onset of progressive distal muscle weakness and atrophy, delayed motor development, prominent distal sensory impairment, areflexia, moderately reduced nerve conduction velocities, and foot and hand deformities in Balkan (Russe) Gypsies.
ORPHACODE:	99953
SYNONYMS:	CMT4G HMSNR Hereditary motor and sensory neuropathy, Russe Type
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	HK1
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RELATED CONTENT

Related Genetic Tests

- Neuropathy (gene panel)
- Peripheral neuropathy (gene panel)

Related Laboratories

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

Related Analytes

- hexokinase 1

Related Gene Panels

- Neuropathy (148 genes) - IPG
- Neuropathy (genepanel) - UZA