DISEASE:
MME-related autosomal dominant Charcot Marie Tooth disease type 2

NAME:	MME-related autosomal dominant Charcot Marie Tooth disease type 2
DESCRIPTION:	A rare autosomal dominant hereditary axonal motor and sensory neuropathy characterized by adult onset of slowly progressive distal muscle weakness and atrophy, sensory impairment, and hyporeflexia beginning in the lower limbs. Progressive gait disturbance may lead to loss of independent ambulation in some patients at a higher age.
ORPHACODE:	497757
SYNONYMS:	MME-related autosomal dominant CMT2 MME-related autosomal dominant hereditary motor and sensory neuropathy type 2
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	MME
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Source URL: http://gentest.healthdata.be/disease/2655