DISEASE:
Autosomal dominant Charcot-Marie-Tooth disease type 2 due to TFG mutation

NAME:	Autosomal dominant Charcot-Marie-Tooth disease type 2 due to TFG mutation
DESCRIPTION:	A rare, axonal hereditary motor and sensory neuropathy characterized by adult onset of slowly progressive distal muscle weakness and atrophy, decreased deep tendon reflexes of lower limbs, and mild distal sensory loss leading to gait difficulties in most patients.
ORPHACODE:	435819
SYNONYMS:	CMT2 due to TFG mutation
XREF(S):	Orphanet
ANALYTE(S):	TFG
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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