DISEASE:
Hyperkalemic periodic paralysis

NAME:	Hyperkalemic periodic paralysis
DESCRIPTION:	A rare muscle disorder characterized by episodic attacks of muscle weakness associated with an increase in serum potassium concentration.
ORPHACODE:	682
SYNONYMS:	Adynamia episodica hereditaria Familial hyperPP Familial hyperkalemic periodic paralysis Gamstorp disease Gamstorp episodic adynamy HYPP HyperKPP HyperPP Hyperkalemic PP Primary hyperPP Primary hyperkalemic periodic paralysis

XREF(S):	Orphanet OMIM MeSH MeSH ICD-10
ANALYTE(S):	SCN4A
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Periodic paralysis (myotonia) / Paramyotonia congenita (SCN4A gene)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- sodium voltage-gated channel alpha subunit 4