DISEASE:
Dilated cardiomyopathy with ataxia

NAME:	Dilated cardiomyopathy with ataxia
DESCRIPTION:	Dilated cardiomyopathy with ataxia (DCMA) is characterized by severe early onset (before the age of three years) dilated cardiomyopathy (DCM) with conduction defects (long QT syndrome), non-progressive cerebellar ataxia, testicular dysgenesis, and 3-methylglutaconic aciduria.
ORPHACODE:	66634
SYNONYMS:	3-methylglutaconic aciduria type 5 DCMA syndrome MGA5
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	DNAJC19
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- DnaJ heat shock protein family (Hsp40) member C19

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB