DISEASE:
Andersen-Tawil syndrome

NAME:	Andersen-Tawil syndrome
DESCRIPTION:	A rare disorder characterized by periodic muscle paralysis, prolongation of the QT interval with a variety of ventricular arrhythmias (leading to predisposition to sudden cardiac death) and characteristic physical features: short stature, scoliosis, low-set ears, hypertelorism, broad nasal root, micrognathia, clinodactyly, brachydactyly and syndactyly.
ORPHACODE:	37553
SYNONYMS:	Andersen syndrome LQT7 Long QT syndrome type 7
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	KCNJ2 KCNJ5
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RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Cardiopathies, hereditary (gene panel)
- Primary cardiac arrhythmias (Atrial fibrillation / Brugada syndome / Catech. polymorphic ventricular tachycardia / Early repolaristion syndrome / Ideopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome) (gene pane)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- potassium inwardly rectifying channel subfamily J member 2
- potassium inwardly rectifying channel subfamily J member 5

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Primary cardiac arrhythmias (113 genes) - VUB