

ANALYTE:
ABCC9

NAME:	ATP binding cassette subfamily C member 9
SYMBOL:	ABCC9
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CMD1O SUR2 sulfonylurea receptor 2
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
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RELATED CONTENT

Related Genetic Tests

- [Cardiomyopathy, hereditary \(gene panel\)](#)
- [Cardiomyopathy: hypertrophic cardiomyopathy, dilated cardiomyopathy, restrictive cardiomyopathy, left ventricular non-compaction cardiomyopathy, arrhythmogenic right ventricular cardiomyopathy \(gene panel\)](#)
- [Cardiopathies, hereditary \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Dilated Cardiomyopathy \(Gene panel\)](#)
- [Heart / Cardio disorders / Cardiopathy \(gene panel\)](#)
- [Inherited cardiac arrhythmia \(gene panel\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
- [Primary Electrical disorders / Brugada syndrome / Long QT syndrome \(LQT\) / Short QT syndrome \(SQT\) / Arrhythmogenic right ventricular cardiomyopathy \(ARVC\) / Catecholaminergic polymorphic ventricular tachycardia \(CPVT\) \(gene panel\)](#)
- [Primary cardiac arrhythmias \(Atrial fibrillation / Brugada syndrome / Catech. polymorphic ventricular tachycardia / Early repolarisation syndrome / Idiopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome\) \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)

Related Diseases

- [Acromegaly facial appearance syndrome](#)

- Brugada syndrome
- Cantú syndrome
- Familial atrial fibrillation
- Familial isolated dilated cardiomyopathy
- Hypertrichosis-acromegaloid facial appearance syndrome

Related Gene Panels

- Cardiomyopathy (genepanel) - UZA
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Congenital malformation (1721 genes) - ULB
- Dilated Cardiomyopathy (79 genes) - IPG
- Inherited cardiac arrhythmia (25 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA
- Primary cardiac arrhythmias (113 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- cardiopathy panel - UGent

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