

**ANALYTE:  
ACTN2**

<b>NAME:</b>	actinin alpha 2
<b>SYMBOL:</b>	ACTN2
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Ensembl</a> <a href="#">Genatlas</a> <a href="#">HGNC</a> <a href="#">OMIM</a> <a href="#">Reactome</a> <a href="#">SwissProt</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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### 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 structural heart defects \(gene panel\)](#)
- [Dilated Cardiomyopathy \(Gene panel\)](#)
- [Heart / Cardio disorders / Cardiopathy \(gene panel\)](#)
- [Hypertrophic cardiomyopathy \(gene panel\)](#)
- [Myopathy \(gene panel\)](#)
- [Neuromuscular disorders \(232 genes \(= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia\)\)](#)
- [Neuromuscular 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 \(Atrial fibrillation / Brugada syndrome / Catech. polymorphic ventricular tachycardia / Early repolarisation syndrome / Idiopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome\) \(gene panel\)](#)

### Related Diseases

- [Familial isolated dilated cardiomyopathy](#)
- [NON RARE IN EUROPE: Familial isolated hypertrophic cardiomyopathy](#)

### Related Gene Panels

- Cardiomyopathy (genepanel) - UZA
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Congenital structural heart defects - UGent
- Dilated Cardiomyopathy (79 genes) - IPG
- Hypertrophic cardiomyopathy (75 genes) - IPG
- Myopathy (genepanel) - UZA
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA
- Primary cardiac arrhythmias (113 genes) - VUB
- cardiopathy panel - UGent

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