

**DISEASE:**  
**Brugada syndrome**

<b>NAME:</b>	Brugada syndrome
<b>DESCRIPTION:</b>	A cardiac disorder characterized on electrocardiogram (ECG) by ST segment elevation with a coved aspect on the right precordial leads, and a clinical susceptibility to ventricular tachyarrhythmias and sudden death occurring in the absence of overt myocardial abnormalities.
<b>ORPHACODE:</b>	130
<b>SYNOMYS:</b>	Idiopathic ventricular fibrillation, Brugada type

XREF(S):	<a href="#">Orphanet</a> <a href="#">MedDRA</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">OMIM</a> <a href="#">MeSH</a> <a href="#">ICD-10</a>
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<b>ANALYTE(S):</b>	<u>HCN4</u> <u>ABCC9</u> <u>CACNB2</u> <u>CACNA1C</u> <u>CACNA2D1</u> <u>KCNE3</u> <u>GPD1L</u> <u>KCND3</u> <u>SCN3B</u> <u>TRPM4</u> <u>SCN10A</u> <u>SLMAP</u> <u>SCN1B</u> <u>AKAP9</u> <u>KCNE5</u> <u>RANGRF</u> <u>SCN5A</u> <u>SCNN1A</u> <u>SEMA3A</u> <u>SCN2B</u> <u>KCNJ8</u> <u>PKP2</u>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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### Related Genetic Tests

- [Brugada syndrome](#)
- [Cardiomyopathy, hereditary \(gene panel\)](#)
- [Cardiopathies, hereditary \(gene panel\)](#)
- [Long QT syndrome](#)
- [Primary cardiac arrhythmias \(Atrial fibrillation / Brugada syndrome / Catech. polymorphic ventricular tachycardia / Early repolarisation syndrome / Ideopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome\) \(gene panel\)](#)

### Related Laboratories

- [Centrum Medische Genetica - UZ Brussel VUB](#)
- [Centrum Medische Genetica - UZ Gent](#)
- [Centrum Menselijke Erfelijheid - KUL](#)

### Related Analytes

- [ATP binding cassette subfamily C member 9](#)
- [A-kinase anchoring protein 9](#)
- [calcium voltage-gated channel subunit alpha1 C](#)
- [calcium voltage-gated channel auxiliary subunit alpha2delta 1](#)
- [calcium voltage-gated channel auxiliary subunit beta 2](#)
- [glycerol-3-phosphate dehydrogenase 1 like](#)
- [hyperpolarization activated cyclic nucleotide gated potassium channel 4](#)
- [potassium voltage-gated channel subfamily D member 3](#)

- [potassium voltage-gated channel subfamily E regulatory subunit 3](#)
- [potassium voltage-gated channel subfamily E regulatory subunit 5](#)
- [potassium inwardly rectifying channel subfamily J member 8](#)
- [plakophilin 2](#)
- [RAN guanine nucleotide release factor](#)
- [sodium voltage-gated channel alpha subunit 10](#)
- [sodium voltage-gated channel beta subunit 1](#)
- [sodium voltage-gated channel beta subunit 2](#)
- [sodium voltage-gated channel beta subunit 3](#)
- [sodium voltage-gated channel alpha subunit 5](#)
- [sodium channel epithelial 1 subunit alpha](#)
- [semaphorin 3A](#)
- [sarcolemma associated protein](#)
- [transient receptor potential cation channel subfamily M member 4](#)

## Related Gene Panels

- [Brugada syndrome \(16 genes\) - VUB](#)
- [Cardiomyopathy, hereditary \(208 genes\) - VUB](#)
- [Cardiopathies, hereditary \(102 genes\) - KUL](#)
- [Primary cardiac arrhythmias \(113 genes\) - VUB](#)

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