

**ANALYTE:**  
**CACNA1H**

<b>NAME:</b>	calcium voltage-gated channel subunit alpha1 H
<b>SYMBOL:</b>	CACNA1H
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	Cav3.2
<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>	26 Oct 2023 - 23:49

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## RELATED CONTENT

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### Related Genetic Tests

- [Early onset epileptic encephalopathy \(gene panel - 845 genes\)](#)
- [Epilepsy gene panel](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Nephropathies, hereditary \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
- [Neuromuscular disorders \(232 genes \(= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia\)\)](#)
- [Neuromuscular disorders \(gene panel\)](#)
- [Tubulopathy \(gene panel\)](#)

### Related Diseases

- [Childhood absence epilepsy](#)
- [Familial hyperaldosteronism type IV](#)

### Related Gene Panels

- [Early onset epileptic encephalopathy \(845 genes\) - ULB](#)
- [Epilepsy gene panel - VUB](#)
- [Intellectual disability & Epilepsy - UGent](#)
- [Intellectual disability \(gene panel\)](#)
- [Nephropathies, hereditary \(219 genes\) - KUL](#)

- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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