Unconventional intronic splice site mutation in SCN5A channelopathy

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Citation Report

#	Article	IF	Citations
1	The Brugada syndrome. Current Opinion in Cardiology, 2007, 22, 163-170.	1.8	28
4	Gene mutations, atrial fibrillation, and the elusive cigar. Clinical Pharmacology and Therapeutics, 2007, 81, 26-28.	4.7	3
5	Non-SCN5A Related Brugada Syndromes: Verification of Normal Splicing and Trafficking of SCN5A Without Exonic Mutations. Annals of Human Genetics, 2007, 71, 8-17.	0.8	7
6	SCN5A channelopathies â€“ An update on mutations and mechanisms. Progress in Biophysics and Molecular Biology, 2008, 98, 120-136.	2.9	130
7	The Year in Review of Clinical Cardiac Electrophysiology. Journal of the American College of Cardiology, 2008, 51, 2075-2081.	2.8	4
8	Novel mRNA isoforms of the sodium channels Nav1.2, Nav1.3 and Nav1.7 encode predicted two-domain, truncated proteins. Neuroscience, 2008, 155, 797-808.	2.3	11
9	A splice site mutation in hERG leads to cryptic splicing in human long QT syndrome. Journal of Molecular and Cellular Cardiology, 2008, 44, 502-509.	1.9	20
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13	A novel 5â€² splice site mutation of SCN5A associated with Brugada syndrome resulting in multiple cryptic transcripts. International Journal of Cardiology, 2012, 158, 441-443.	1.7	2
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17	Targeted Next-Generation Sequencing of 51 Genes Involved in Primary Electrical Disease. Journal of Molecular Diagnostics, 2017, 19, 445-459.	2.8	15
18	Prolonged Right Ventricular Ejection Delay in Brugada Syndrome Depends on the Type of SCN5A Variantã€€â€• Electromechanical Coupling Through Tissue Velocity Imaging as a Bridge Between Genotyping and Phenotyping â€•. Circulation Journal, 2018, 82, 53-61.	1.6	6
19	Identification and functional characterization of mutations in LPL gene causing severe hypertriglyceridaemia and acute pancreatitis. Journal of Cellular and Molecular Medicine, 2020, 24, 1286-1299.	3.6	52
20	Compound Heterozygous SCN5A Mutations in Severe Sodium Channelopathy With Brugada Syndrome: A Case Report. Frontiers in Cardiovascular Medicine, 2020, 7, 117.	2.4	3
21	Clinical characterization of the first Belgian SCN5A founder mutation cohort. Europace, 2021, 23, 918-927.	1.7	3

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22	Novel SCN5A variants identified in a group of Iranian Brugada syndrome patients. Functional and Integrative Genomics, 2021, 21, 331-340.	3.5	1
23	Inherited Arrhythmias: Brugada Syndrome and Early Repolarisation Syndrome. , 2018, , 437-480.		0
24	Genomic and Non-Genomic Regulatory Mechanisms of the Cardiac Sodium Channel in Cardiac Arrhythmias. International Journal of Molecular Sciences, 2022, 23, 1381.	4.1	10
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