

Mirella Memmi

List of Publications by Year in descending order

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Version: 2024-02-01

28
papers

5,522
citations

361045

20
h-index

525886

27
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30
all docs

30
docs citations

30
times ranked

4306
citing authors

#	ARTICLE	IF	CITATIONS
1	Outcomes of Patients With Catecholaminergic Polymorphic Ventricular Tachycardia Treated With β -Blockers. <i>JAMA Cardiology</i> , 2022, 7, 504.	3.0	26
2	Identification of a SCN5A founder mutation causing sudden death, Brugada syndrome, and conduction blocks in Southern Italy. <i>Heart Rhythm</i> , 2021, 18, 1698-1706.	0.3	2
3	Association of Hydroxychloroquine With QTc Interval in Patients With COVID-19. <i>Circulation</i> , 2020, 142, 513-515.	1.6	31
4	Natural History and Risk Stratification in Andersen-Tawil Syndrome Type 1. <i>Journal of the American College of Cardiology</i> , 2020, 75, 1772-1784.	1.2	44
5	Mutation site-specific risk profile in patients with Type 1 Long QT Syndrome. <i>European Heart Journal</i> , 2020, 41, .	1.0	0
6	Role of CACNA1C variants in Brugada syndrome: clinical aspects and genetic testing strategies. <i>European Heart Journal</i> , 2020, 41, .	1.0	0
7	Efficacy and Limitations of Quinidine in Patients With Brugada Syndrome. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2019, 12, .	2.1	14
8	Interplay Between Genetic Substrate, QTc Duration, and Arrhythmia Risk in Patients With Long QT Syndrome. <i>Journal of the American College of Cardiology</i> , 2018, 71, 1663-1671.	1.2	137
9	CardioVAI: An automatic implementation of ACMG-AMP variant interpretation guidelines in the diagnosis of cardiovascular diseases. <i>Human Mutation</i> , 2018, 39, 1835-1846.	1.1	31
10	Challenges in Molecular Diagnostics of Channelopathies in the Next-Generation Sequencing Era: Less Is More?. <i>Frontiers in Cardiovascular Medicine</i> , 2016, 3, 29.	1.1	8
11	Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2016, 68, 2540-2550.	1.2	148
12	Gene-Specific Therapy With Mexiletine Reduces Arrhythmic Events in Patients With Long QT Syndrome Type 3. <i>Journal of the American College of Cardiology</i> , 2016, 67, 1053-1058.	1.2	191
13	Novel Insight Into the Natural History of Short QT Syndrome. <i>Journal of the American College of Cardiology</i> , 2014, 63, 1300-1308.	1.2	191
14	Clinical utility gene card for: Catecholaminergic polymorphic ventricular tachycardia (CPVT). <i>European Journal of Human Genetics</i> , 2014, 22, 152-152.	1.4	18
15	Arrhythmogenesis in Catecholaminergic Polymorphic Ventricular Tachycardia. <i>Circulation Research</i> , 2006, 99, 292-298.	2.0	293
16	Clinical Phenotype and Functional Characterization of CASQ2 Mutations Associated With Catecholaminergic Polymorphic Ventricular Tachycardia. <i>Circulation</i> , 2006, 114, 1012-1019.	1.6	189
17	FKBP12.6 Deficiency and Defective Calcium Release Channel (Ryanodine Receptor) Function Linked to Exercise-Induced Sudden Cardiac Death. <i>Cell</i> , 2003, 113, 829-840.	13.5	683
18	Clinical and Molecular Characterization of Patients With Catecholaminergic Polymorphic Ventricular Tachycardia. <i>Circulation</i> , 2002, 106, 69-74.	1.6	1,103

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19	Involvement of the cardiac ryanodine receptor/calcium release channel in catecholaminergic polymorphic ventricular tachycardia. <i>Journal of Cellular Physiology</i> , 2002, 190, 1-6.	2.0	182
20	Long QT syndrome, Brugada syndrome, and conduction system disease are linked to a single sodium channel mutation. <i>Journal of Clinical Investigation</i> , 2002, 110, 1201-1209.	3.9	172
21	Mutations in the Cardiac Ryanodine Receptor Gene (<i>hRyR2</i>) Underlie Catecholaminergic Polymorphic Ventricular Tachycardia. <i>Circulation</i> , 2001, 103, 196-200.	1.6	1,291
22	Inherited Brugada and Long QT-3 Syndrome Mutations of a Single Residue of the Cardiac Sodium Channel Confer Distinct Channel and Clinical Phenotypes. <i>Journal of Biological Chemistry</i> , 2001, 276, 30623-30630.	1.6	181
23	Novel Arrhythmogenic Mechanism Revealed by a Long-QT Syndrome Mutation in the Cardiac Na ⁺ Channel. <i>Circulation Research</i> , 2001, 88, 740-745.	2.0	114
24	Mechanisms of <i>I_{Ks}</i> suppression in LQT1 mutants. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2000, 279, H3003-H3011.	1.5	68
25	Novel characteristics of a misprocessed mutant HERG channel linked to hereditary long QT syndrome. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2000, 279, H1748-H1756.	1.5	88
26	Brugada syndrome and sudden cardiac death in children. <i>Lancet</i> , The, 2000, 355, 808-809.	6.3	244
27	Identical de novo mutation at the D4F104S1 locus in monozygotic male twins affected by facioscapulohumeral muscular dystrophy (FSHD) with different clinical expression.. <i>Journal of Medical Genetics</i> , 1998, 35, 778-783.	1.5	29
28	Concentrations of l-dopa in plasma and plasma ultrafiltrates. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 1996, 14, 1043-1046.	1.4	17