

Robert Dumaine

List of Publications by Year in descending order

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Version: 2024-02-01

42
papers

4,229
citations

201575

27
h-index

276775

41
g-index

44
all docs

44
docs citations

44
times ranked

3159
citing authors

#	ARTICLE	IF	CITATIONS
1	The neuronal potassium current I _A is a potential target for pain during chronic inflammation. <i>Physiological Reports</i> , 2021, 9, e14975.	0.7	2
2	In utero exposure to nicotine abolishes the postnatal response of the cardiac sodium current to isoproterenol in newborn rabbit atrium. <i>Heart Rhythm</i> , 2019, 16, 494-501.	0.3	2
3	Apelin-13 Regulates Vasopressin-Induced Aquaporin-2 Expression and Trafficking in Kidney Collecting Duct Cells. <i>Cellular Physiology and Biochemistry</i> , 2019, 53, 687-700.	1.1	24
4	A Systematic Exploration of Macrocyclization in Apelin-13: Impact on Binding, Signaling, Stability, and Cardiovascular Effects. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 2266-2277.	2.9	30
5	A novel three base-pair deletion in domain two of the cardiac sodium channel causes Brugada syndrome. <i>Journal of Electrocardiology</i> , 2018, 51, 667-673.	0.4	1
6	The hypotensive effect of activated apelin receptor is correlated with β^2 -arrestin recruitment. <i>Pharmacological Research</i> , 2018, 131, 7-16.	3.1	23
7	In-utero exposure to nicotine alters the development of the rabbit cardiac conduction system and provides a potential mechanism for sudden infant death syndrome. <i>Archives of Toxicology</i> , 2017, 91, 3947-3960.	1.9	12
8	Apelin Compared With Dobutamine Exerts Cardioprotection and Extends Survival in a Rat Model of Endotoxin-Induced Myocardial Dysfunction*. <i>Critical Care Medicine</i> , 2017, 45, e391-e398.	0.4	30
9	Structure-activity relationship of novel macrocyclic biased apelin receptor agonists. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 449-458.	1.5	27
10	ELABELA Improves Cardio-Renal Outcome in Fatal Experimental Septic Shock. <i>Critical Care Medicine</i> , 2017, 45, e1139-e1148.	0.4	49
11	Discovery and Structure-Activity Relationship of a Bioactive Fragment of ELABELA that Modulates Vascular and Cardiac Functions. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 2962-2972.	2.9	100
12	A Brugada syndrome proband with compound heterozygote <i>SCN5A</i> mutations identified from a Chinese family in Singapore. <i>Europace</i> , 2016, 18, 897-904.	0.7	16
13	C-Terminal Modifications of Apelin-13 Significantly Change Ligand Binding, Receptor Signaling, and Hypotensive Action. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 2431-2440.	2.9	48
14	Prolongation of Action Potential Duration and QT Interval During Epilepsy Linked to Increased Contribution of Neuronal Sodium Channels to Cardiac Late Na ⁺ Current. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2015, 8, 912-920.	2.1	48
15	Functional up-regulation of Nav1.8 sodium channel in A β afferent fibers subjected to chronic peripheral inflammation. <i>Journal of Neuroinflammation</i> , 2014, 11, 45.	3.1	43
16	About half of the late sodium current in cardiac myocytes from dog ventricle is due to non-cardiac-type Na ⁺ channels. <i>Journal of Molecular and Cellular Cardiology</i> , 2012, 53, 593-598.	0.9	45
17	Modulation of canine cardiac sodium current by Apelin. <i>Journal of Molecular and Cellular Cardiology</i> , 2010, 48, 694-701.	0.9	27
18	Larger dispersion of INa in female dog ventricle as a mechanism for gender-specific incidence of cardiac arrhythmias. <i>Cardiovascular Research</i> , 2009, 81, 82-89.	1.8	43

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19	Lidocaine-Induced Brugada Syndrome Phenotype Linked to a Novel Double Mutation in the Cardiac Sodium Channel. <i>Circulation Research</i> , 2008, 103, 396-404.	2.0	45
20	Comparison of K ⁺ currents in cardiac Purkinje cells isolated from rabbit and dog. <i>Journal of Molecular and Cellular Cardiology</i> , 2007, 42, 378-389.	0.9	60
21	Genetic and biophysical basis for bupivacaine-induced ST segment elevation and VT/VF. Anesthesia unmasked Brugada syndrome. <i>Heart Rhythm</i> , 2006, 3, 1074-1078.	0.3	53
22	P6-4. <i>Heart Rhythm</i> , 2006, 3, S302.	0.3	1
23	Identification of Two Domains Involved in the Assembly of Transient Receptor Potential Canonical Channels*. <i>Journal of Biological Chemistry</i> , 2006, 281, 30356-30364.	1.6	50
24	Compound Heterozygous Mutations P336L and I1660V in the Human Cardiac Sodium Channel Associated With the Brugada Syndrome. <i>Circulation</i> , 2006, 114, 2026-2033.	1.6	102
25	Expression pattern of neuronal and skeletal muscle voltage-gated Na ⁺ -channels in the developing mouse heart. <i>Journal of Physiology</i> , 2005, 564, 683-696.	1.3	95
26	Further Insights into the Effect of Quinidine in Short QT Syndrome Caused by a Mutation in HERG. <i>Journal of Cardiovascular Electrophysiology</i> , 2005, 16, 54-58.	0.8	189
27	Short QT syndrome. <i>Cmaj</i> , 2005, 173, 1349-1354.	0.9	64
28	Lidocaine-induced Brugada syndrome phenotype linked to a novel double mutation in the cardiac sodium channel. <i>Heart Rhythm</i> , 2005, 2, S294.	0.3	0
29	Cryptic 5' splice site activation in SCN5A associated with Brugada syndrome. <i>Journal of Molecular and Cellular Cardiology</i> , 2005, 38, 555-560.	0.9	51
30	Genetic and biophysical basis for bupivacaine-induced ST segment elevation and VT/VF. Anesthesia-mediated acquired Brugada syndrome. <i>Heart Rhythm</i> , 2005, 2, S49.	0.3	0
31	Sudden Death Associated With Short-QT Syndrome Linked to Mutations in HERG. <i>Circulation</i> , 2004, 109, 30-35.	1.6	804
32	Divergent action potential morphologies reveal nonequilibrium properties of human cardiac Na channels. <i>Cardiovascular Research</i> , 2004, 64, 477-487.	1.8	14
33	Phenotypic Characterization of a Large European Family with Brugada Syndrome Displaying a Sudden Unexpected Death Syndrome Mutation in SCN5A. <i>Journal of Cardiovascular Electrophysiology</i> , 2004, 15, 64-69.	0.8	50
34	Value of Electrocardiographic Parameters and Ajmaline Test in the Diagnosis of Brugada Syndrome Caused by SCN5A Mutations. <i>Circulation</i> , 2004, 110, 3023-3027.	1.6	163
35	Molecular mechanisms underlying the long QT syndrome. <i>Current Opinion in Cardiology</i> , 2002, 17, 36-42.	0.8	32
36	Genetic and biophysical basis of sudden unexplained nocturnal death syndrome (SUNDS), a disease allelic to Brugada syndrome. <i>Human Molecular Genetics</i> , 2002, 11, 337-345.	1.4	334

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37	Novel mutations in domain I of SCN5A cause Brugada syndrome. <i>Molecular Genetics and Metabolism</i> , 2002, 75, 317-324.	0.5	61
38	A Molecular Link between the Sudden Infant Death Syndrome and the Long-QT Syndrome. <i>New England Journal of Medicine</i> , 2000, 343, 262-267.	13.9	340
39	Ionic Mechanisms Responsible for the Electrocardiographic Phenotype of the Brugada Syndrome Are Temperature Dependent. <i>Circulation Research</i> , 1999, 85, 803-809.	2.0	557
40	Actions of dolasetron and its major metabolite on guinea-pig papillary muscle fibres and the β -subunit of human heart sodium channels expressed in <i>Xenopus</i> oocytes. <i>Drug Development Research</i> , 1996, 37, 223-230.	1.4	2
41	HERG, a Primary Human Ventricular Target of the Nonsedating Antihistamine Terfenadine. <i>Circulation</i> , 1996, 94, 817-823.	1.6	262
42	Multiple Mechanisms of Na ⁺ Channel-Linked Long-QT Syndrome. <i>Circulation Research</i> , 1996, 78, 916-924.	2.0	285