

Bo Sun

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

393
citations

840585

11
h-index

794469

19
g-index

21
all docs

21
docs citations

21
times ranked

649
citing authors

#	ARTICLE	IF	CITATIONS
1	Cardiac ryanodine receptor calcium release deficiency syndrome. <i>Science Translational Medicine</i> , 2021, 13, .	5.8	68
2	Limiting RyR2 Open Time Prevents Alzheimer's Disease-Related Neuronal Hyperactivity and Memory Loss but Not $\text{A}\beta$ -Amyloid Accumulation. <i>Cell Reports</i> , 2020, 32, 108169.	2.9	41
3	De novo ITPR1 variants are a recurrent cause of early-onset ataxia, acting via loss of channel function. <i>European Journal of Human Genetics</i> , 2018, 26, 1623-1634.	1.4	32
4	Suppression of ryanodine receptor function prolongs Ca^{2+} release refractoriness and promotes cardiac alternans in intact hearts. <i>Biochemical Journal</i> , 2016, 473, 3951-3964.	1.7	28
5	Generation and Characterization of a Mouse Model Harboring the Exon-3 Deletion in the Cardiac Ryanodine Receptor. <i>PLoS ONE</i> , 2014, 9, e95615.	1.1	27
6	The cardiac ryanodine receptor, but not sarcoplasmic reticulum Ca^{2+} -ATPase, is a major determinant of Ca^{2+} alternans in intact mouse hearts. <i>Journal of Biological Chemistry</i> , 2018, 293, 13650-13661.	1.6	27
7	Enhanced Cytosolic Ca^{2+} Activation Underlies a Common Defect of Central Domain Cardiac Ryanodine Receptor Mutations Linked to Arrhythmias. <i>Journal of Biological Chemistry</i> , 2016, 291, 24528-24537.	1.6	22
8	Limiting RyR2 open time prevents Alzheimer's disease-related deficits in the 3xTg-AD mouse model. <i>Journal of Neuroscience Research</i> , 2021, 99, 2906-2921.	1.3	18
9	Human RyR2 (Ryanodine Receptor 2) Loss-of-Function Mutations. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2021, 14, e010013.	2.1	18
10	Ca^{2+} -CaM Dependent Inactivation of RyR2 Underlies Ca^{2+} Alternans in Intact Heart. <i>Circulation Research</i> , 2021, 128, e63-e83.	2.0	17
11	Increased RyR2 open probability induces neuronal hyperactivity and memory loss with or without Alzheimer's disease-causing gene mutations. <i>Alzheimer's and Dementia</i> , 2022, 18, 2088-2098.	0.4	16
12	The central domain of cardiac ryanodine receptor governs channel activation, regulation, and stability. <i>Journal of Biological Chemistry</i> , 2020, 295, 15622-15635.	1.6	13
13	CPVT-associated cardiac ryanodine receptor mutation G357S with reduced penetrance impairs Ca^{2+} release termination and diminishes protein expression. <i>PLoS ONE</i> , 2017, 12, e0184177.	1.1	12
14	Subcellular localization of hippocampal ryanodine receptor 2 and its role in neuronal excitability and memory. <i>Communications Biology</i> , 2022, 5, 183.	2.0	12
15	Reduced expression of cardiac ryanodine receptor protects against stress-induced ventricular tachyarrhythmia, but increases the susceptibility to cardiac alternans. <i>Biochemical Journal</i> , 2018, 475, 169-183.	1.7	8
16	RyR2 disease mutations at the C-terminal domain intersubunit interface alter closed-state stability and channel activation. <i>Journal of Biological Chemistry</i> , 2021, 297, 100808.	1.6	7
17	A gain-of-function mutation in the ITPR1 gating domain causes male infertility in mice. <i>Journal of Cellular Physiology</i> , 2022, 237, 3305-3316.	2.0	7
18	The Cytoplasmic Region of Inner Helix S6 Is an Important Determinant of Cardiac Ryanodine Receptor Channel Gating. <i>Journal of Biological Chemistry</i> , 2016, 291, 26024-26034.	1.6	6

#	ARTICLE	IF	CITATIONS
19	Genetically and pharmacologically limiting RyR2 open time prevents neuronal hyperactivity of hippocampal CA1 neurons in brain slices of 5xFAD mice. <i>Neuroscience Letters</i> , 2021, 758, 136011.	1.0	6
20	Thioredoxin α 1 regulates calcium homeostasis in MPP ⁺ /MPTP-induced Parkinson's disease models. <i>European Journal of Neuroscience</i> , 2021, 54, 4827-4837.	1.2	4
21	Exhaled nitric oxide in neonates with or without hypoxemic respiratory failure. <i>World Journal of Emergency Medicine</i> , 2011, 2, 195.	0.5	4