

Conor McClenaghan

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

908
citations

516710

16
h-index

677142

22
g-index

25
all docs

25
docs citations

25
times ranked

1091
citing authors

#	ARTICLE	IF	CITATIONS
1	ATP-sensitive potassium channels in zebrafish cardiac and vascular smooth muscle. <i>Journal of Physiology</i> , 2022, 600, 299-312.	2.9	6
2	Bridging Personal and Population in Excitability Diseases: Will Studies of Rare Diseases Bring Generalizable Mechanisms From Monogenic Channelopathies?. <i>Function</i> , 2022, 3, zqab072.	2.3	0
3	Isolation of Cardiac and Vascular Smooth Muscle Cells from Adult, Juvenile, Larval and Embryonic Zebrafish for Electrophysiological Studies. <i>Journal of Visualized Experiments</i> , 2022, , .	0.3	0
4	Complex consequences of Cantu syndrome SUR2 variant R1154Q in genetically modified mice. <i>JCI Insight</i> , 2021, 6, .	5.0	11
5	Consequences of SUR2[A478V] Mutation in Skeletal Muscle of Murine Model of Cantu Syndrome. <i>Cells</i> , 2021, 10, 1791.	4.1	10
6	Coronavirus Proteins as Ion Channels: Current and Potential Research. <i>Frontiers in Immunology</i> , 2020, 11, 573339.	4.8	56
7	The Mechanism of High-Output Cardiac Hypertrophy Arising From Potassium Channel Gain-of-Function in Cantu Syndrome. <i>Function</i> , 2020, 1, zqaa004.	2.3	18
8	Pathophysiological Consequences of KATP Channel Overactivity and Pharmacological Response to Glibenclamide in Skeletal Muscle of a Murine Model of Cantu Syndrome. <i>Frontiers in Pharmacology</i> , 2020, 11, 604885.	3.5	19
9	Glibenclamide reverses cardiovascular abnormalities of Cantu syndrome driven by KATP channel overactivity. <i>Journal of Clinical Investigation</i> , 2020, 130, 1116-1121.	8.2	40
10	ABCC9-related Intellectual disability Myopathy Syndrome is a KATP channelopathy with loss-of-function mutations in ABCC9. <i>Nature Communications</i> , 2019, 10, 4457.	12.8	31
11	Pulmonary Hypertension and ATP-Sensitive Potassium Channels. <i>Hypertension</i> , 2019, 74, 14-22.	2.7	24
12	Beta-cell excitability and excitability-driven diabetes in adult Zebrafish islets. <i>Physiological Reports</i> , 2019, 7, e14101.	1.7	8
13	Glibenclamide treatment in a Cantu syndrome patient with a pathogenic ABCC9 gain-of-function variant: Initial experience. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 1585-1590.	1.2	30
14	Cantu syndrome: Findings from 74 patients in the International Cantu Syndrome Registry. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2019, 181, 658-681.	1.6	50
15	Cantu syndrome-associated SUR2 (ABCC9) mutations in distinct structural domains result in KATP channel gain-of-function by differential mechanisms. <i>Journal of Biological Chemistry</i> , 2018, 293, 2041-2052.	3.4	34
16	Loss-of-Function <i>ABCC8</i> Mutations in Pulmonary Arterial Hypertension. <i>Circulation Genomic and Precision Medicine</i> , 2018, 11, e002087.	3.6	62
17	Cardiovascular consequences of KATP overactivity in Cantu syndrome. <i>JCI Insight</i> , 2018, 3, .	5.0	44
18	Bilayer-Mediated Structural Transitions Control Mechanosensitivity of the TREK-2 K2P Channel. <i>Structure</i> , 2017, 25, 708-718.e2.	3.3	64

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19	Conserved functional consequences of disease-associated mutations in the slide helix of Kir6.1 and Kir6.2 subunits of the ATP-sensitive potassium channel. <i>Journal of Biological Chemistry</i> , 2017, 292, 17387-17398.	3.4	31
20	Polymodal activation of the TREK-2 K2P channel produces structurally distinct open states. <i>Journal of General Physiology</i> , 2016, 147, 497-505.	1.9	65
21	K2P channel gating mechanisms revealed by structures of TREK-2 and a complex with Prozac. <i>Science</i> , 2015, 347, 1256-1259.	12.6	255
22	TRPA1 Agonist Activity of Probenecid Desensitizes Channel Responses: Consequences for Screening. <i>Assay and Drug Development Technologies</i> , 2012, 10, 533-541.	1.2	10
23	Increased prokineticin 2 expression in gut inflammation: role in visceral pain and intestinal ion transport. <i>Neurogastroenterology and Motility</i> , 2012, 24, 65.	3.0	33
24	GPR39 Is Coupled to TMEM16A in Intestinal Fibroblast-Like Cells. <i>PLoS ONE</i> , 2012, 7, e47686.	2.5	7