

Daniela Pietrobon

List of Publications by Year in descending order

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47
papers

6,923
citations

101543

36
h-index

223800

46
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49
docs citations

49
times ranked

4838
citing authors

#	ARTICLE	IF	CITATIONS
1	A Cacna1a Knockin Migraine Mouse Model with Increased Susceptibility to Cortical Spreading Depression. <i>Neuron</i> , 2004, 41, 701-710.	8.1	595
2	Pathophysiology of Migraine. <i>Annual Review of Physiology</i> , 2013, 75, 365-391.	13.1	523
3	Neurobiology of migraine. <i>Nature Reviews Neuroscience</i> , 2003, 4, 386-398.	10.2	498
4	Chaos and commotion in the wake of cortical spreading depression and spreading depolarizations. <i>Nature Reviews Neuroscience</i> , 2014, 15, 379-393.	10.2	318
5	Enhanced Excitatory Transmission at Cortical Synapses as the Basis for Facilitated Spreading Depression in CaV2.1 Knockin Migraine Mice. <i>Neuron</i> , 2009, 61, 762-773.	8.1	292
6	Functional Consequences of Mutations in the Human α_1A Calcium Channel Subunit Linked to Familial Hemiplegic Migraine. <i>Journal of Neuroscience</i> , 1999, 19, 1610-1619.	3.6	242
7	Familial hemiplegic migraine mutations increase Ca ²⁺ influx through single human CaV2.1 channels and decrease maximal CaV2.1 current density in neurons. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2002, 99, 13284-13289.	7.1	240
8	Inherited Neuronal Ion Channelopathies: New Windows on Complex Neurological Diseases. <i>Journal of Neuroscience</i> , 2008, 28, 11768-11777.	3.6	225
9	Migraine: a disorder of brain excitatory-inhibitory balance?. <i>Trends in Neurosciences</i> , 2012, 35, 507-520.	8.6	219
10	High cortical spreading depression susceptibility and migraine-associated symptoms in Ca _v 2.1 S218L mice. <i>Annals of Neurology</i> , 2010, 67, 85-98.	5.3	206
11	Direct measurement of proton transfer rates to a group controlling the dihydropyridine-sensitive Ca ²⁺ channel. <i>Nature</i> , 1987, 329, 243-246.	27.8	200
12	Novel mechanism of voltage-dependent gating in L-type calcium channels. <i>Nature</i> , 1990, 346, 651-655.	27.8	196
13	Structural and functional aspects of calcium homeostasis in eukaryotic cells. <i>FEBS Journal</i> , 1990, 193, 599-622.	0.2	196
14	Calcium Channels and Channelopathies of the Central Nervous System. <i>Molecular Neurobiology</i> , 2002, 25, 031-050.	4.0	190
15	Effect of Funiculosin and Antimycin A on the Redox-Driven H ⁺ Pumps in Mitochondria: on the Nature of "Leaks". <i>FEBS Journal</i> , 1981, 117, 389-394.	0.2	186
16	CaV2.1 channelopathies. <i>Pflugers Archiv European Journal of Physiology</i> , 2010, 460, 375-393.	2.8	184
17	Dystonia and cerebellar atrophy in Cacna1a null mice lacking P/Q calcium channel activity. <i>FASEB Journal</i> , 2001, 15, 1288-1290.	0.5	182
18	Increased Susceptibility to Cortical Spreading Depression in the Mouse Model of Familial Hemiplegic Migraine Type 2. <i>PLoS Genetics</i> , 2011, 7, e1002129.	3.5	179

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19	Familial hemiplegic migraine. <i>Neurotherapeutics</i> , 2007, 4, 274-284.	4.4	171
20	Migraine: New Molecular Mechanisms. <i>Neuroscientist</i> , 2005, 11, 373-386.	3.5	168
21	α ₁ Subunits Form the Pore of Three Cerebellar R-Type Calcium Channels with Different Pharmacological and Permeation Properties. <i>Journal of Neuroscience</i> , 2000, 20, 171-178.	3.6	162
22	Functional Diversity of P-Type and R-Type Calcium Channels in Rat Cerebellar Neurons. <i>Journal of Neuroscience</i> , 1996, 16, 6353-6363.	3.6	160
23	Function and dysfunction of synaptic calcium channels: insights from mouse models. <i>Current Opinion in Neurobiology</i> , 2005, 15, 257-265.	4.2	158
24	Complete Loss of P/Q Calcium Channel Activity Caused by a CACNA1A Missense Mutation Carried by Patients with Episodic Ataxia Type 2. <i>American Journal of Human Genetics</i> , 2001, 68, 759-764.	6.2	147
25	A Systems Neuroscience Approach to Migraine. <i>Neuron</i> , 2018, 97, 1004-1021.	8.1	134
26	Specific Kinetic Alterations of Human CaV2.1 Calcium Channels Produced by Mutation S218L Causing Familial Hemiplegic Migraine and Delayed Cerebral Edema and Coma after Minor Head Trauma. <i>Journal of Biological Chemistry</i> , 2005, 280, 17678-17686.	3.4	123
27	Defective glutamate and K ⁺ clearance by cortical astrocytes in familial hemiplegic migraine type 2. <i>EMBO Molecular Medicine</i> , 2016, 8, 967-986.	6.9	110
28	Presynaptic R-Type Calcium Channels Contribute to Fast Excitatory Synaptic Transmission in the Rat Hippocampus. <i>Journal of Neuroscience</i> , 2001, 21, 8715-8721.	3.6	103
29	Calcium channels and migraine. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2013, 1828, 1655-1665.	2.6	78
30	Diagnostic and therapeutic aspects of hemiplegic migraine. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020, 91, 764-771.	1.9	66
31	Insights into migraine mechanisms and CaV2.1 calcium channel function from mouse models of familial hemiplegic migraine. <i>Journal of Physiology</i> , 2010, 588, 1871-1878.	2.9	54
32	Mechanism underlying unaltered cortical inhibitory synaptic transmission in contrast with enhanced excitatory transmission in CaV2.1 knockin migraine mice. <i>Neurobiology of Disease</i> , 2014, 69, 225-234.	4.4	50
33	Non-canonical glutamate signaling in a genetic model of migraine with aura. <i>Neuron</i> , 2021, 109, 611-628.e8.	8.1	41
34	Abnormal cortical synaptic transmission in CaV2.1 knockin mice with the S218L missense mutation which causes a severe familial hemiplegic migraine syndrome in humans. <i>Frontiers in Cellular Neuroscience</i> , 2015, 9, 8.	3.7	40
35	Modal Gating of Human CaV2.1 (P/Q-type) Calcium Channels. <i>Journal of General Physiology</i> , 2004, 124, 445-461.	1.9	38
36	The differential expression of low-threshold K ⁺ currents generates distinct firing patterns in different subtypes of adult mouse trigeminal ganglion neurones. <i>Journal of Physiology</i> , 2008, 586, 5101-5118.	2.9	38

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37	Role of different voltage-gated Ca ²⁺ channels in cortical spreading depression: Specific requirement of P/Q-type Ca ²⁺ channels. <i>Channels</i> , 2011, 5, 110-114.	2.8	37
38	Heterogeneity of Astrocytic and Neuronal GLT-1 at Cortical Excitatory Synapses, as Revealed by its Colocalization With Na ⁺ /K ⁺ -ATPase \pm Isoforms. <i>Cerebral Cortex</i> , 2019, 29, 3331-3350.	2.9	37
39	Genetic mouse models of migraine. <i>Journal of Headache and Pain</i> , 2019, 20, 79.	6.0	31
40	Astrocyte dysfunction increases cortical dendritic excitability and promotes cranial pain in familial migraine. <i>Science Advances</i> , 2020, 6, eaaz1584.	10.3	23
41	Biological science of headache channels. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2010, 97, 73-83.	1.8	22
42	Enhanced Thalamocortical Synaptic Transmission and Dysregulation of the Excitatoryâ€“Inhibitory Balance at the Thalamocortical Feedforward Inhibitory Microcircuit in a Genetic Mouse Model of Migraine. <i>Journal of Neuroscience</i> , 2019, 39, 9841-9851.	3.6	18
43	Ion channels in migraine disorders. <i>Current Opinion in Physiology</i> , 2018, 2, 98-108.	1.8	16
44	Specific activation of GluN1-N2B NMDA receptors underlies facilitation of cortical spreading depression in a genetic mouse model of migraine with reduced astrocytic glutamate clearance. <i>Neurobiology of Disease</i> , 2021, 156, 105419.	4.4	14
45	Differential effect of FHM2 mutation on synaptic plasticity in distinct hippocampal regions. <i>Cephalalgia</i> , 2019, 39, 1333-1338.	3.9	8
46	Cortical spreading depression and familial hemiplegic migraine 2015. <i>Journal of Headache and Pain</i> , 2015, 16, A20.	6.0	3
47	Cav2.1 Channels and Migraine. , 2014, , 3-25.		0