

Lishuang Cao

List of Publications by Year in descending order

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Version: 2024-02-01

20
papers

1,900
citations

567281

15
h-index

752698

20
g-index

21
all docs

21
docs citations

21
times ranked

3060
citing authors

#	ARTICLE	IF	CITATIONS
1	Combined small-molecule inhibition accelerates developmental timing and converts human pluripotent stem cells into nociceptors. <i>Nature Biotechnology</i> , 2012, 30, 715-720.	17.5	515
2	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
3	Molecular and functional variation in iPSC-derived sensory neurons. <i>Nature Genetics</i> , 2018, 50, 54-61.	21.4	191
4	Pharmacological reversal of a pain phenotype in iPSC-derived sensory neurons and patients with inherited erythromelalgia. <i>Science Translational Medicine</i> , 2016, 8, 335ra56.	12.4	154
5	An intracellular P2X receptor required for osmoregulation in <i>Dictyostelium discoideum</i> . <i>Nature</i> , 2007, 448, 200-203.	27.8	130
6	Characterizing Human Stem Cell-derived Sensory Neurons at the Single-cell Level Reveals Their Ion Channel Expression and Utility in Pain Research. <i>Molecular Therapy</i> , 2014, 22, 1530-1543.	8.2	127
7	The biophysical and molecular basis of TRPV1 proton gating. <i>EMBO Journal</i> , 2011, 30, 994-1002.	7.8	93
8	Permeation Properties of a P2X Receptor in the Green Algae <i>Ostreococcus tauri</i> . <i>Journal of Biological Chemistry</i> , 2008, 283, 15122-15126.	3.4	67
9	Influence of the N Terminus on the Biophysical Properties and Pharmacology of TREK1 Potassium Channels. <i>Molecular Pharmacology</i> , 2014, 85, 671-681.	2.3	52
10	Thr339-to-Serine Substitution in Rat P2X2 Receptor Second Transmembrane Domain Causes Constitutive Opening and Indicates a Gating Role for Lys308. <i>Journal of Neuroscience</i> , 2007, 27, 12916-12923.	3.6	51
11	Pharmacology in translation: the preclinical and early clinical profile of the novel $\frac{1}{2}$ / $\frac{3}{3}$ functionally selective GABA _A receptor positive allosteric modulator PF06372865. <i>British Journal of Pharmacology</i> , 2018, 175, 708-725.	5.4	49
12	Polar Residues in the Second Transmembrane Domain of the Rat P2X2 Receptor That Affect Spontaneous Gating, Unitary Conductance, and Rectification. <i>Journal of Neuroscience</i> , 2009, 29, 14257-14264.	3.6	46
13	P2X receptor channels show threefold symmetry in ionic charge selectivity and unitary conductance. <i>Nature Neuroscience</i> , 2011, 14, 17-18.	14.8	45
14	GLX530159, a novel, selective, mechanosensitive two-pore domain potassium (K _{2P}) channel opener, reduces rat dorsal root ganglion neuron excitability. <i>British Journal of Pharmacology</i> , 2018, 175, 2272-2283.	5.4	40
15	The Chimeric Approach Reveals That Differences in the TRPV1 Pore Domain Determine Species-specific Sensitivity to Block of Heat Activation. <i>Journal of Biological Chemistry</i> , 2011, 286, 39663-39672.	3.4	31
16	Rescue of functional $\frac{1}{5}$ F508-CFTR channels by co-expression with truncated CFTR constructs in COS-1 cells. <i>FEBS Letters</i> , 2003, 554, 173-178.	2.8	15
17	Chronic exposure to EGF affects trafficking and function of ENaC channel in cystic fibrosis cells. <i>Biochemical and Biophysical Research Communications</i> , 2005, 331, 503-511.	2.1	12
18	Investigation of the structure activity relationship of flufenamic acid derivatives at the human TREK channel K _{2P} 18.1. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 4919-4924.	2.2	11

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19	Functional analysis of CFTR chloride channel activity in cells with elevated MDR1 expression. <i>Biochemical and Biophysical Research Communications</i> , 2003, 304, 248-252.	2.1	6
20	Role of the domain encompassing Arg304-Ile328 in rat P2X2 receptor conformation revealed by alterations in complex glycosylation at Asn298. <i>Biochemical Journal</i> , 2008, 416, 137-143.	3.7	6