

Kewei Wang

List of Publications by Year in descending order

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66
papers

1,929
citations

236925

25
h-index

276875

41
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67
all docs

67
docs citations

67
times ranked

2427
citing authors

#	ARTICLE	IF	CITATIONS
1	Exome Sequencing Reveals Mutations in TRPV3 as a Cause of Olmsted Syndrome. <i>American Journal of Human Genetics</i> , 2012, 90, 558-564.	6.2	300
2	Inhibition of Ca ²⁺ -activated Cl ⁻ channel ANO1/TMEM16A expression suppresses tumor growth and invasiveness in human prostate carcinoma. <i>Cancer Letters</i> , 2012, 326, 41-51.	7.2	158
3	Inhibition of Calcium-Activated Chloride Channel ANO1/TMEM16A Suppresses Tumor Growth and Invasion in Human Lung Cancer. <i>PLoS ONE</i> , 2015, 10, e0136584.	2.5	101
4	The current agonists and positive allosteric modulators of α_7 nAChR for CNS indications in clinical trials. <i>Acta Pharmaceutica Sinica B</i> , 2017, 7, 611-622.	12.0	87
5	Inhibition of calcium-activated chloride channel ANO1 suppresses proliferation and induces apoptosis of epithelium originated cancer cells. <i>Oncotarget</i> , 2016, 7, 78619-78630.	1.8	65
6	The fate of medications evaluated for ischemic stroke pharmacotherapy over the period 1995-2015. <i>Acta Pharmaceutica Sinica B</i> , 2016, 6, 522-530.	12.0	64
7	Heteromeric Heat-sensitive Transient Receptor Potential Channels Exhibit Distinct Temperature and Chemical Response. <i>Journal of Biological Chemistry</i> , 2012, 287, 7279-7288.	3.4	63
8	Honokiol protects brain against ischemia-reperfusion injury in rats through disrupting PSD95-nNOS interaction. <i>Brain Research</i> , 2013, 1491, 204-212.	2.2	53
9	Intracellular Proton-mediated Activation of TRPV3 Channels Accounts for the Exfoliation Effect of α -Hydroxyl Acids on Keratinocytes. <i>Journal of Biological Chemistry</i> , 2012, 287, 25905-25916.	3.4	50
10	Inhibition of ANO1/TMEM16A induces apoptosis in human prostate carcinoma cells by activating TNF- α signaling. <i>Cell Death and Disease</i> , 2018, 9, 703.	6.3	50
11	Antipruritic Effect of Natural Coumarin Osthole through Selective Inhibition of Thermosensitive TRPV3 Channel in the Skin. <i>Molecular Pharmacology</i> , 2018, 94, 1164-1173.	2.3	45
12	The Ca ²⁺ -Permeable Cation Transient Receptor Potential TRPV3 Channel: An Emerging Pivotal Target for Itch and Skin Diseases. <i>Molecular Pharmacology</i> , 2017, 92, 193-200.	2.3	40
13	Inhibition of Ca ²⁺ -activated chloride channel ANO1 suppresses ovarian cancer through inactivating PI3K/Akt signaling. <i>International Journal of Cancer</i> , 2019, 144, 2215-2226.	5.1	40
14	Deficiency of anti-inflammatory cytokine IL-4 leads to neural hyperexcitability and aggravates cerebral ischemia-reperfusion injury. <i>Acta Pharmaceutica Sinica B</i> , 2020, 10, 1634-1645.	12.0	39
15	Capsaicin enhances the antitumor activity of sorafenib in hepatocellular carcinoma cells and mouse xenograft tumors through increased ERK signaling. <i>Acta Pharmacologica Sinica</i> , 2018, 39, 438-448.	6.1	38
16	A pivotal role for the activation of TRPV3 channel in itch sensations induced by the natural skin sensitizer carvacrol. <i>Acta Pharmacologica Sinica</i> , 2018, 39, 331-335.	6.1	37
17	Pharmacological Inhibition of the Temperature-Sensitive and Ca ²⁺ -Permeable Transient Receptor Potential Vanilloid TRPV3 Channel by Natural Forsythoside B Attenuates Pruritus and Cytotoxicity of Keratinocytes. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2019, 368, 21-31.	2.5	36
18	Synergistic antitumor activity of sorafenib and artesunate in hepatocellular carcinoma cells. <i>Acta Pharmacologica Sinica</i> , 2020, 41, 1609-1620.	6.1	36

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19	Synthesis and biological activities of indolizine derivatives as alpha-7 nAChR agonists. <i>European Journal of Medicinal Chemistry</i> , 2016, 115, 94-108.	5.5	35
20	Comparison of Gating Properties and Use-Dependent Block of Nav1.5 and Nav1.7 Channels by Anti-Arrhythmics Mexiletine and Lidocaine. <i>PLoS ONE</i> , 2015, 10, e0128653.	2.5	35
21	Effect of magnolol on cerebral injury and blood brain barrier dysfunction induced by ischemia-reperfusion in vivo and in vitro. <i>Metabolic Brain Disease</i> , 2017, 32, 1109-1118.	2.9	34
22	The Ca ²⁺ -activated chloride channel ANO1/TMEM16A: An emerging therapeutic target for epithelium-originated diseases?. <i>Acta Pharmaceutica Sinica B</i> , 2021, 11, 1412-1433.	12.0	34
23	Inhibition of the Warm Temperature-Activated Ca ²⁺ -Permeable Transient Receptor Potential Vanilloid TRPV3 Channel Attenuates Atopic Dermatitis. <i>Molecular Pharmacology</i> , 2019, 96, 393-400.	2.3	33
24	Elevated Expression of Acid-Sensing Ion Channel 3 Inhibits Epilepsy via Activation of Interneurons. <i>Molecular Neurobiology</i> , 2016, 53, 485-498.	4.0	30
25	Exploiting the Diversity of Ion Channels: Modulation of Ion Channels for Therapeutic Indications. <i>Handbook of Experimental Pharmacology</i> , 2019, 260, 187-205.	1.8	27
26	Visceral Hyperalgesia Induced by Forebrain-Specific Suppression of Native Kv7/KCNQ/M-Current in Mice. <i>Molecular Pain</i> , 2011, 7, 1744-8069-7-84.	2.1	23
27	The role of Piezo1 in conventional aqueous humor outflow dynamics. <i>IScience</i> , 2021, 24, 102042.	4.1	23
28	The Mpro structure-based modifications of ebsele derivatives for improved antiviral activity against SARS-CoV-2 virus. <i>Bioorganic Chemistry</i> , 2021, 117, 105455.	4.1	22
29	Electrophysiological and pharmacological characterization of a novel and potent neuronal Kv7 channel opener SCR2682 for antiepilepsy. <i>FASEB Journal</i> , 2019, 33, 9154-9166.	0.5	21
30	Selective Activation of Nociceptor TRPV1 Channel and Reversal of Inflammatory Pain in Mice by a Novel Coumarin Derivative Muralatin L from <i>Murraya alata</i> . <i>Journal of Biological Chemistry</i> , 2016, 291, 640-651.	3.4	20
31	Inhibition of temperature-sensitive TRPV3 channel by two natural isochlorogenic acid isomers for alleviation of dermatitis and chronic pruritus. <i>Acta Pharmaceutica Sinica B</i> , 2022, 12, 723-734.	12.0	19
32	Anti-pruritic and anti-inflammatory effects of natural verbascoside through selective inhibition of temperature-sensitive Ca ²⁺ -permeable TRPV3 channel. <i>Journal of Dermatological Science</i> , 2020, 97, 229-231.	1.9	16
33	Design and Synthesis of Novel Positive Allosteric Modulators of $\alpha 7$ Nicotinic Acetylcholine Receptors with the Ability To Rescue Auditory Gating Deficit in Mice. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 159-173.	6.4	13
34	Detection of Lipase Activity in Cells by a Fluorescent Probe Based on Formation of Self-Assembled Micelles. <i>IScience</i> , 2020, 23, 101294.	4.1	13
35	Discovery of 4-arylthiophene-3-carboxylic acid as inhibitor of ANO1 and its effect as analgesic agent. <i>Acta Pharmaceutica Sinica B</i> , 2021, 11, 1947-1964.	12.0	13
36	Pharmacological Activation of Thermo-Sensitive Transient Receptor Potential Vanilloid 3 Channels Inhibits Hair Growth by Inducing Cell Death of Hair Follicle Outer Root Sheath. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2019, 370, 299-307.	2.5	12

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37	Different KChIPs Compete for Heteromultimeric Assembly with Pore-Forming Kv4 Subunits. <i>Biophysical Journal</i> , 2015, 108, 2658-2669.	0.5	11
38	Activity-induced spontaneous spikes in GABAergic neurons suppress seizure discharges: an implication of computational modeling. <i>Oncotarget</i> , 2017, 8, 32384-32397.	1.8	11
39	Visualizing TRPA1 in the Plasma Membrane for Rapidly Screening Optical Control Agonists via a Photochromic Ligand Based Fluorescent Probe. <i>Analytical Chemistry</i> , 2020, 92, 1934-1939.	6.5	10
40	Pharmacological Characterization of H05, a Novel Serotonin and Noradrenaline Reuptake Inhibitor with Moderate 5-HT _{2A} Antagonist Activity for the Treatment of Depression. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2018, 365, 624-635.	2.5	9
41	Activation of Neuronal Voltage-Gated Potassium Kv7/KCNQ/M-Current by a Novel Channel Opener SCR2682 for Alleviation of Chronic Pain. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2021, 377, 20-28.	2.5	9
42	Natural Piperine Improves Lipid Metabolic Profile of High-Fat Diet-Fed Mice by Upregulating SR-B1 and ABCG8 Transporters. <i>Journal of Natural Products</i> , 2021, 84, 373-381.	3.0	9
43	Photosensitive and Photoswitchable TRPA1 Agonists Optically Control Pain through Channel Desensitization. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 16282-16292.	6.4	9
44	Inhibition of intracellular proton-sensitive Ca ²⁺ -permeable TRPV3 channels protects against ischemic brain injury. <i>Acta Pharmaceutica Sinica B</i> , 2022, 12, 2330-2347.	12.0	9
45	Molecular determinants for the chemical activation of the warmth-sensitive TRPV3 channel by the natural monoterpene carvacrol. <i>Journal of Biological Chemistry</i> , 2022, , 101706.	3.4	9
46	Anticonvulsant effect of dipropofol by enhancing native GABA currents in cortical neurons in mice. <i>Journal of Neurophysiology</i> , 2018, 120, 1404-1414.	1.8	8
47	Design, synthesis and biological activities of piperidine-spirooxadiazole derivatives as $\alpha 7$ nicotinic receptor antagonists. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112774.	5.5	8
48	Identification of two natural coumarin enantiomers for selective inhibition of TRPV2 channels. <i>FASEB Journal</i> , 2020, 34, 12338-12353.	0.5	8
49	Inhibition of Nav1.7 channel by a novel blocker QLS-81 for alleviation of neuropathic pain. <i>Acta Pharmacologica Sinica</i> , 2021, 42, 1235-1247.	6.1	8
50	DIC/Oxyma [®] -based accelerated synthesis and oxidative folding studies of centipede toxin α -Tx. <i>Journal of Peptide Science</i> , 2022, 28, e3368.	1.4	8
51	Negative modulation of NMDA receptor channel function by DREAM/calsenilin/KChIP3 provides neuroprotection?. <i>Frontiers in Molecular Neuroscience</i> , 2012, 5, 39.	2.9	7
52	Discovery of fused heterocyclic carboxamide derivatives as novel $\alpha 7$ -nAChR agonists: Synthesis, preliminary SAR and biological evaluation. <i>European Journal of Medicinal Chemistry</i> , 2019, 182, 111618.	5.5	7
53	Synthesis and Biological Evaluation of Novel Triazine Derivatives as Positive Allosteric Modulators of $\alpha 7$ Nicotinic Acetylcholine Receptors. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 12379-12396.	6.4	6
54	Evodiamine Lowers Blood Lipids by Up-Regulating the PPAR γ /ABCG1 Pathway in High-Fat-Diet-Fed Mice. <i>Journal of Natural Products</i> , 2021, 84, 3110-3116.	3.0	6

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55	Efficient Chemical Synthesis and Oxidative Folding Studies of Scorpion Toxin Peptide WaTx. <i>Acta Chimica Sinica</i> , 2022, 80, 444.	1.4	6
56	Discovery, synthesis, and optimization of teixobactin, a novel antibiotic without detectable bacterial resistance. <i>Journal of Peptide Science</i> , 2022, 28, .	1.4	6
57	Interactions of KChIP4a and its mutants with Ca ²⁺ or Kv4.3 N-terminus by affinity capillary electrophoresis. <i>Analytical Biochemistry</i> , 2014, 449, 99-105.	2.4	5
58	The Tetramerization Domain Potentiates Kv4 Channel Function by Suppressing Closed-State Inactivation. <i>Biophysical Journal</i> , 2014, 107, 1090-1104.	0.5	5
59	Prefrontal inhibition of neuronal K _v 7 channels enhances prepulse inhibition of acoustic startle reflex and resistance to hypofrontality. <i>British Journal of Pharmacology</i> , 2020, 177, 4720-4733.	5.4	5
60	Optimization of 4-arylthiophene-3-carboxylic acid derivatives as inhibitors of ANO1: Lead optimization studies toward their analgesic efficacy for inflammatory pain. <i>European Journal of Medicinal Chemistry</i> , 2022, 237, 114413.	5.5	5
61	Selective activation of TRPA1 ion channels by nitrobenzene skin sensitizers DNFB and DNCB. <i>Journal of Biological Chemistry</i> , 2022, 298, 101555.	3.4	4
62	Involvement of TMEM16A/ANO1 upregulation in the oncogenesis of colorectal cancer. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2022, 1868, 166370.	3.8	4
63	Piezo2 downregulation via the Cre-lox system affects aqueous humor dynamics in mice. <i>Molecular Vision</i> , 2021, 27, 354-364.	1.1	3
64	Small molecule-driven direct reprogramming of Müller cells into bipolar-like cells. <i>Cell Proliferation</i> , 2022, 55, e13184.	5.3	3
65	Chemical conversion of nicotinamide into type I positive allosteric modulator of $\alpha 7$ nAChRs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 1928-1933.	2.2	2
66	Design and synthesis of novel β -aminoamides derivatives as Nav1.7 inhibitors for antinociception. <i>Chinese Chemical Letters</i> , 2022, 33, 1643-1646.	9.0	1