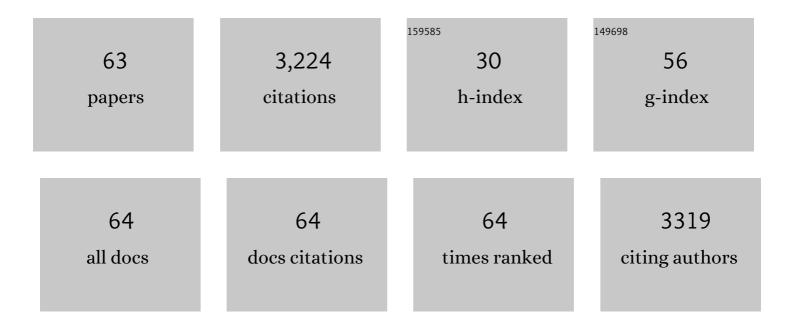
Steve Mcgaraughty

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Discovery of (R)-(3-fluoropyrrolidin-1-yl)(6-((5-(trifluoromethyl)pyridin-2-yl)oxy)quinolin-2-yl)methanone (ABBV-318) and analogs as small molecule Nav1.7/ Nav1.8 blockers for the treatment of pain. Bioorganic and Medicinal Chemistry, 2022, 63, 116743.	3.0	0
2	Characterization of psoriasiform dermatitis induced by systemic injection of interleukinâ€⊋3 minicircles in mice. Journal of Dermatology, 2019, 46, 482-497.	1.2	12
3	Monocytes/Macrophages play a pathogenic role in IL-23 mediated psoriasis-like skin inflammation. Scientific Reports, 2019, 9, 5310.	3.3	42
4	<scp>IL</scp> â€36 receptor antagonistic antibodies inhibit inflammatory responses in preclinical models of psoriasiform dermatitis. Experimental Dermatology, 2019, 28, 113-120.	2.9	29
5	Characterization and comparison of rat monosodium iodoacetate and medial meniscal tear models of osteoarthritic pain. Journal of Orthopaedic Research, 2018, 36, 2109-2117.	2.3	12
6	Short-term oral gavage administration of adenine induces a model of fibrotic kidney disease in rats. Journal of Pharmacological and Toxicological Methods, 2018, 94, 34-43.	0.7	10
7	Losartan improves renal function and pathology in obese ZSF-1 rats. Journal of Basic and Clinical Physiology and Pharmacology, 2018, 29, 281-290.	1.3	11
8	Characterization and pharmacological modulation of noci-responsive deep dorsal horn neurons across diverse rat models of pathological pain. Journal of Neurophysiology, 2018, 120, 1893-1905.	1.8	6
9	TRPV3 modulates nociceptive signaling through peripheral and supraspinal sites in rats. Journal of Neurophysiology, 2017, 118, 904-916.	1.8	9
10	Longitudinal Changes in Measured Glomerular Filtration Rate, Renal Fibrosis and Biomarkers in a Rat Model of Type 2 Diabetic Nephropathy. American Journal of Nephrology, 2016, 44, 339-353.	3.1	33
11	Mechanistic Insights Into the Analgesic Efficacy of A-1264087, a Novel Neuronal Ca2+ Channel Blocker That Reduces Nociception in Rat Preclinical Pain Models. Journal of Pain, 2014, 15, 387.e1-387.e14.	1.4	10
12	A mixed Ca2+ channel blocker, A-1264087, utilizes peripheral and spinal mechanisms to inhibit spinal nociceptive transmission in a rat model of neuropathic pain. Journal of Neurophysiology, 2014, 111, 394-404.	1.8	10
13	Characterization of the triazine, T4, a representative from a novel series of CaV2 inhibitors with strong state-dependence, poor use-dependence, and distinctively fast kinetics. European Journal of Pharmacology, 2014, 745, 234-242.	3.5	4
14	A peripherally acting, selective T-type calcium channel blocker, ABT-639, effectively reduces nociceptive and neuropathic pain in rats. Biochemical Pharmacology, 2014, 89, 536-544.	4.4	58
15	Development and Validation of a Medium-Throughput Electrophysiological Assay for KCNQ2/3 Channel Openers Using QPatch HT. Assay and Drug Development Technologies, 2013, 11, 17-24.	1.2	2
16	Disturbances in slow-wave sleep are induced by models of bilateral inflammation, neuropathic, and postoperative pain, but not osteoarthritic pain in rats. Pain, 2013, 154, 1092-1102.	4.2	38
17	Antagonism of Supraspinal Histamine H ₃ Receptors Modulates Spinal Neuronal Activity in Neuropathic Rats. Journal of Pharmacology and Experimental Therapeutics, 2012, 343, 13-20.	2.5	31
18	An Automated Electrophysiological Assay for Differentiating CaV2.2 Inhibitors Based on State Dependence and Kinetics. Assay and Drug Development Technologies, 2012, 10, 542-550.	1.2	7

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19	TRPA1 in Drug Discovery. Methods in Pharmacology and Toxicology, 2012, , 43-59.	0.2	4
20	TRPV1 antagonist, Aâ€889425, inhibits mechanotransmission in a subclass of rat primary afferent neurons following peripheral inflammation. Synapse, 2012, 66, 187-195.	1.2	17
21	Spontaneous firing and evoked responses of spinal nociceptive neurons are attenuated by blockade of P2X3 and P2X2/3 receptors in inflamed rats. Journal of Neuroscience Research, 2012, 90, 1597-1606.	2.9	27
22	Oral and cutaneous thermosensory profile of selective TRPV1 inhibition by ABT-102 in a randomized healthy volunteer trial. Pain, 2011, 152, 1192-1200.	4.2	102
23	TRPV1-related modulation of spinal neuronal activity and behavior in a rat model of osteoarthritic pain. Brain Research, 2011, 1369, 158-166.	2.2	45
24	Coexpression and activation of TRPV1 suppress the activity of the KCNQ2/3 channel. Journal of General Physiology, 2011, 138, 341-352.	1.9	26
25	TRPA1 Modulation of Spontaneous and Mechanically Evoked Firing of Spinal Neurons in Uninjured, Osteoarthritic, and Inflamed Rats. Molecular Pain, 2010, 6, 1744-8069-6-14.	2.1	150
26	Characterization of Fasudil in Preclinical Models of Pain. Journal of Pain, 2010, 11, 941-949.	1.4	17
27	Antagonism of TRPV1 receptors indirectly modulates activity of thermoregulatory neurons in the medial preoptic area of rats. Brain Research, 2009, 1268, 58-67.	2.2	30
28	Pharmacological MRI in awake rats predicts selective binding of α ₄ β ₂ nicotinic receptors. Synapse, 2008, 62, 159-168.	1.2	24
29	A Selective Na _v 1.8 Sodium Channel Blocker, A-803467		

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37	Pharmacological and functional magnetic resonance imaging techniques in CNS drug discovery. Expert Opinion on Drug Discovery, 2006, 1, 211-224.	5.0	4
38	5-(3-Bromophenyl)-7-(6-morpholin-4-ylpyridin-3-yl)pyrido[2,3-d]pyrimidin-4-ylamine: structure–activity relationships of 7-substituted heteroaryl analogs as non-nucleoside adenosine kinase inhibitors. Bioorganic and Medicinal Chemistry, 2005, 13, 3705-3720.	3.0	49
39	Endogenous opioid mechanisms partially mediate P2X3 /P2X2/3 -related antinociception in rat models of inflammatory and chemogenic pain but not neuropathic pain. British Journal of Pharmacology, 2005, 146, 180-188.	5.4	35
40	Anticonvulsant and Antinociceptive Actions of Novel Adenosine Kinase Inhibitors. Current Topics in Medicinal Chemistry, 2005, 5, 43-58.	2.1	100
41	Lesions of the periaqueductal gray disrupt input to the rostral ventromedial medulla following microinjections of morphine into the medial or basolateral nuclei of the amygdala. Brain Research, 2004, 1009, 223-227.	2.2	53
42	Increased WDR spontaneous activity and receptive field size in rats following a neuropathic or inflammatory injury: implications for mechanical sensitivity. Neuroscience Letters, 2004, 372, 123-126.	2.1	43
43	Distinct neurochemical mechanisms are activated following administration of different P2X receptor agonists into the hindpaw of a rat. Brain Research, 2003, 965, 187-193.	2.2	35
44	Adenosine kinase inhibitors: polar 7-Substitutent of pyridopyrimidine derivatives improving their locomotor selectivity. Bioorganic and Medicinal Chemistry Letters, 2003, 13, 3041-3044.	2.2	38
45	Effects of A-317491, a novel and selective P2X3 /P2X2/3 receptor antagonist, on neuropathic, inflammatory and chemogenic nociception following intrathecal and intraplantar administration. British Journal of Pharmacology, 2003, 140, 1381-1388.	5.4	164
46	Capsaicin Infused Into the PAG Affects Rat Tail Flick Responses to Noxious Heat and Alters Neuronal Firing in the RVM. Journal of Neurophysiology, 2003, 90, 2702-2710.	1.8	108
47	A-317491, a novel potent and selective non-nucleotide antagonist of P2X3 and P2X2/3 receptors, reduces chronic inflammatory and neuropathic pain in the rat. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 17179-17184.	7.1	440
48	Analgesic and anti-inflammatory effects of A-286501, a novel orally active adenosine kinase inhibitor. Pain, 2002, 96, 107-118.	4.2	50
49	Microinjection of morphine into various amygdaloid nuclei differentially affects nociceptive responsiveness and RVM neuronal activity. Pain, 2002, 96, 153-162.	4.2	93
50	Comparison of the ability of adenosine kinase inhibitors and adenosine receptor agonists to attenuate thermal hyperalgesia and reduce motor performance in rats. Pharmacology Biochemistry and Behavior, 2002, 73, 573-581.	2.9	32
51	Structure–activity studies of 5-substituted pyridopyrimidines as adenosine kinase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 83-86.	2.2	30
52	Pyridopyrimidine analogues as novel adenosine kinase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 2071-2074.	2.2	33
53	A-134974: a novel adenosine kinase inhibitor, relieves tactile allodynia via spinal sites of action in peripheral nerve injured rats. Brain Research, 2001, 905, 104-110.	2.2	33
54	Recent Developments in the Discovery of Novel Adenosine Kinase Inhibitors: Mechanism of Action and Therapeutic Potential. CNS Neuroscience & Therapeutics, 2001, 7, 415-432.	4.0	35

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55	Design of Adenosine Kinase Inhibitors from the NMR-Based Screening of Fragments. Journal of Medicinal Chemistry, 2000, 43, 4781-4786.	6.4	81
56	The effects of strychnine, bicuculline, and ketamine on `immersion-inhibited' dorsal horn convergent neurons in intact and spinalized rats. Brain Research, 1998, 784, 63-70.	2.2	9
57	Pain-Modulating Neurons and Behavioral State. , 1998, , .		0
58	Relationship between mechano-receptive fields of dorsal horn convergent neurons and the response to noxious immersion of the ipsilateral hindpaw in rats. Pain, 1997, 70, 133-140.	4.2	13
59	Effects of Noxious Hindpaw Immersion on Evoked and Spontaneous Firing of Contralateral Convergent Dorsal Horn Neurons in Both Intact and Spinalized Rats. Brain Research Bulletin, 1997, 43, 263-267.	3.0	12
60	A correlogram analysis of the activity in the rostral ventromedial medulla of awake rats and in rats anesthetized with ketamine or pentobarbital following the administration of morphine. Experimental Brain Research, 1995, 106, 283-90.	1.5	4
61	Simultaneous multi- and single-unit recordings in the rostral ventromedial medulla of ketamine-anaesthetized rats, and the cross-correlogram analysis of their interactions. Experimental Brain Research, 1993, 92, 489-94.	1.5	14
62	Two distinct unit activity responses to morphine in the rostral ventromedial medulla of awake rats. Brain Research, 1993, 604, 331-333.	2.2	22
63	Investigating the role of anaesthetics on the rostral ventromedial medulla: implications for a GABAergic link between ON and OFF cells. Neuroscience Letters, 1993, 149, 119-122.	2.1	9