

Tally M Largent-Milnes

List of Publications by Year in descending order

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Version: 2024-02-01

66
papers

1,634
citations

257450

24
h-index

330143

37
g-index

69
all docs

69
docs citations

69
times ranked

2003
citing authors

#	ARTICLE	IF	CITATIONS
1	Regulation of mitochondrial fission by GIPC-mediated Drp1 retrograde transport. <i>Molecular Biology of the Cell</i> , 2022, 33, mbcE21060286.	2.1	10
2	Î²IV-spectrin as a stalk cell-intrinsic regulator of VEGF signaling. <i>Nature Communications</i> , 2022, 13, 1326.	12.8	11
3	Extracellular Alterations in pH and K ⁺ Modify the Murine Brain Endothelial Cell Total and Phospho-Proteome. <i>Pharmaceutics</i> , 2022, 14, 1469.	4.5	1
4	Cannabinoid-2 Agonism with AM2301 Mitigates Morphine-Induced Respiratory Depression. <i>Cannabis and Cannabinoid Research</i> , 2021, 6, 401-412.	2.9	8
5	An underrepresented majority: A systematic review utilizing allodynic criteria to examine the present scarcity of discrete animal models for episodic migraine. <i>Cephalalgia</i> , 2021, 41, 404-416.	3.9	4
6	The Effects of Repeated Morphine Treatment on the Endogenous Cannabinoid System in the Ventral Tegmental Area. <i>Frontiers in Pharmacology</i> , 2021, 12, 632757.	3.5	8
7	Green Light Antinociceptive and Reversal of Thermal and Mechanical Hypersensitivity Effects Rely on Endogenous Opioid System Stimulation. <i>Journal of Pain</i> , 2021, 22, 1646-1656.	1.4	11
8	Sex hormones regulate NHE1 functional expression and brain endothelial proteome to control paracellular integrity of the blood endothelial barrier. <i>Brain Research</i> , 2021, 1763, 147448.	2.2	4
9	Analgesic Potential of Terpenes Derived from <i>Cannabis sativa</i> . <i>Pharmacological Reviews</i> , 2021, 73, 1269-1297.	16.0	25
10	Sex differences in the expression of the endocannabinoid system within V1M cortex and PAG of Sprague Dawley rats. <i>Biology of Sex Differences</i> , 2021, 12, 60.	4.1	23
11	Brain Penetrant, but not Peripherally Restricted, Synthetic Cannabinoid 1 Receptor Agonists Promote Morphine-Mediated Respiratory Depression. <i>Cannabis and Cannabinoid Research</i> , 2021, . .	2.9	5
12	Glial neuroimmune signaling in opioid reward. <i>Brain Research Bulletin</i> , 2020, 155, 102-111.	3.0	33
13	Heat shock protein 90 inhibitors block the antinociceptive effects of opioids in mouse chemotherapy-induced neuropathy and cancer bone pain models. <i>Pain</i> , 2020, 161, 1798-1807.	4.2	8
14	Acute visceral pain relief mediated by A3AR agonists in rats: involvement of N-type voltage-gated calcium channels. <i>Pain</i> , 2020, 161, 2179-2190.	4.2	21
15	Activation of sphingosine-1-phosphate receptor subtype 1 in the central nervous system contributes to morphine-induced hyperalgesia and antinociceptive tolerance in rodents. <i>Pain</i> , 2020, 161, 2107-2118.	4.2	19
16	Editorial: Novel Molecular Targets for the Treatment of Pain. <i>Frontiers in Molecular Neuroscience</i> , 2020, 13, 625714.	2.9	0
17	Functional NHE1 expression is critical to blood brain barrier integrity and sumatriptan blood to brain uptake. <i>PLoS ONE</i> , 2020, 15, e0227463.	2.5	8
18	Synthesis and Structure-Activity Relationships of 5-aryl-14-alkoxy-pyridomorphinans: Identification of a ¼ Opioid Receptor Agonist/¼ Opioid Receptor Antagonist Ligand with Systemic Antinociceptive Activity and Diminished Opioid Side Effects. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 7663-7694.	6.4	21

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19	DAGL \pm Inhibition as a Non-invasive and Translational Model of Episodic Headache. <i>Frontiers in Pharmacology</i> , 2020, 11, 615028.	3.5	11
20	Chronic Morphine-Induced Changes in Signaling at the A ₃ Adenosine Receptor Contribute to Morphine-Induced Hyperalgesia, Tolerance, and Withdrawal. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2020, 374, 331-341.	2.5	30
21	Title is missing!. , 2020, 15, e0227463.		0
22	Title is missing!. , 2020, 15, e0227463.		0
23	Title is missing!. , 2020, 15, e0227463.		0
24	Title is missing!. , 2020, 15, e0227463.		0
25	A Novel Angiotensin-(1-7) Glycosylated Mas Receptor Agonist for Treating Vascular Cognitive Impairment and Inflammation-Related Memory Dysfunction. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2019, 369, 9-25.	2.5	47
26	Sphingosine-1-phosphate receptor 1 activation in astrocytes contributes to neuropathic pain. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 10557-10562.	7.1	76
27	Animal Models for the Study of Bone-Derived Pain. <i>Methods in Molecular Biology</i> , 2019, 1914, 391-407.	0.9	6
28	Cdk5-mediated CRMP2 phosphorylation is necessary and sufficient for peripheral neuropathic pain. <i>Neurobiology of Pain (Cambridge, Mass)</i> , 2019, 5, 100022.	2.5	46
29	A Kappa Opioid Receptor Agonist Blocks Bone Cancer Pain Without Altering Bone Loss, Tumor Size, or Cancer Cell Proliferation in a Mouse Model of Cancer-Induced Bone Pain. <i>Journal of Pain</i> , 2018, 19, 612-625.	1.4	19
30	Continuous remote ischemic conditioning attenuates cognitive and motor deficits from moderate traumatic brain injury. <i>Journal of Trauma and Acute Care Surgery</i> , 2018, 85, 48-53.	2.1	11
31	Loss of Blood-Brain Barrier Integrity in a KCl-Induced Model of Episodic Headache Enhances CNS Drug Delivery. <i>ENeuro</i> , 2018, 5, ENEURO.0116-18.2018.	1.9	26
32	Peripherally restricted cannabinoid 1 receptor agonist as a novel analgesic in cancer-induced bone pain. <i>Pain</i> , 2018, 159, 1814-1823.	4.2	29
33	Chronic morphine exposure potentiates p-glycoprotein trafficking from nuclear reservoirs in cortical rat brain microvessels. <i>PLoS ONE</i> , 2018, 13, e0192340.	2.5	15
34	Remote ischemic conditioning preserves cognition and motor coordination in a mouse model of traumatic brain injury. <i>Journal of Trauma and Acute Care Surgery</i> , 2017, 83, 1074-1081.	2.1	19
35	Long-lasting antinociceptive effects of green light in acute and chronic pain in rats. <i>Pain</i> , 2017, 158, 347-360.	4.2	81
36	Targeting the S1P/S1PR1 axis mitigates cancer-induced bone pain and neuroinflammation. <i>Pain</i> , 2017, 158, 1733-1742.	4.2	55

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37	Synergistic attenuation of chronic pain using mu opioid and cannabinoid receptor 2 agonists. <i>Neuropharmacology</i> , 2017, 116, 59-70.	4.1	70
38	17 β -Estradiol induces spreading depression and pain behavior in alert female rats. <i>Oncotarget</i> , 2017, 8, 114109-114122.	1.8	16
39	Effect of Centruroides Antivenom on Reversal of Methamphetamine-Induced Hyperkinesia and Hyperthermia in Rats. <i>Journal of Pharmaceutics & Pharmacology</i> , 2017, 5, 1-5.	0.5	0
40	Structure-Activity Relationships of [des-Arg ⁷]Dynorphin A Analogues at the μ Opioid Receptor. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 10291-10298.	6.4	11
41	The cystine/glutamate antiporter system xc ⁻ drives breast tumor cell glutamate release and cancer-induced bone pain. <i>Pain</i> , 2016, 157, 2605-2616.	4.2	32
42	Angiotensin-(1-7)/Mas receptor as an antinociceptive agent in cancer-induced bone pain. <i>Pain</i> , 2016, 157, 2709-2721.	4.2	46
43	Enkephalin analogues with N-phenyl-N-(piperidin-2-ylmethyl)propionamide derivatives: Synthesis and biological evaluations. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 222-227.	2.2	4
44	Use of Animal Models in Understanding Cancer-induced Bone Pain. <i>Cancer Growth and Metastasis</i> , 2015, 8s1, CGM.S21215.	3.5	39
45	A membrane-delimited N-myristoylated CRMP2 peptide aptamer inhibits CaV2.2 trafficking and reverses inflammatory and postoperative pain behaviors. <i>Pain</i> , 2015, 156, 1247-1264.	4.2	71
46	Temperature Differentially Facilitates Spontaneous but Not Evoked Glutamate Release from Cranial Visceral Primary Afferents. <i>PLoS ONE</i> , 2015, 10, e0127764.	2.5	9
47	Design and synthesis of novel bivalent ligands (MOR and DOR) by conjugation of enkephalin analogues with 4-anilidopiperidine derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 4683-4688.	2.2	10
48	Discovery of Novel Multifunctional Ligands with μ / κ Opioid Agonist/Neurokinin-1 (NK1) Antagonist Activities for the Treatment of Pain. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 8573-8583.	6.4	16
49	Discovery of 5-substituted tetrahydronaphthalen-2-yl-methyl with N-phenyl-N-(piperidin-4-yl)propionamide derivatives as potent opioid receptor ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 6185-6194.	3.0	2
50	Angiotensin(1-7) as an Antinociceptive Agent in Cancer-Induced Bone Pain. <i>FASEB Journal</i> , 2015, 29, 897.4.	0.5	0
51	External QX-314 inhibits evoked cranial primary afferent synaptic transmission independent of TRPV1. <i>Journal of Neurophysiology</i> , 2014, 112, 2697-2706.	1.8	14
52	Physiological temperatures drive glutamate release onto trigeminal superficial dorsal horn neurons. <i>Journal of Neurophysiology</i> , 2014, 111, 2222-2231.	1.8	12
53	Capsaicin-responsive corneal afferents do not contain TRPV1 at their central terminals in trigeminal nucleus caudalis in rats. <i>Journal of Chemical Neuroanatomy</i> , 2014, 61-62, 1-12.	2.1	23
54	Animal models for opioid addiction drug discovery. <i>Expert Opinion on Drug Discovery</i> , 2014, 9, 1345-1354.	5.0	12

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55	Truncation of the peptide sequence in bifunctional ligands with mu and delta opioid receptor agonist and neurokinin 1 receptor antagonist activities. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 4975-4978.	2.2	11
56	Disease modification of breast cancer-induced bone remodeling by cannabinoid 2 receptor agonists. <i>Journal of Bone and Mineral Research</i> , 2013, 28, 92-107.	2.8	64
57	Tachykinin NK1 receptor antagonist co-administration attenuates opioid withdrawal-mediated spinal microglia and astrocyte activation. <i>European Journal of Pharmacology</i> , 2012, 684, 64-70.	3.5	31
58	Repeated morphine treatment-mediated hyperalgesia, allodynia and spinal glial activation are blocked by co-administration of a selective cannabinoid receptor type-2 agonist. <i>Journal of Neuroimmunology</i> , 2012, 244, 23-31.	2.3	43
59	Activation of descending pain-facilitatory pathways from the rostral ventromedial medulla by cholecystokinin elicits release of prostaglandin-E2 in the spinal cord. <i>Pain</i> , 2012, 153, 86-94.	4.2	41
60	Discovery of a Potent and Efficacious Peptide Derivative for μ Opioid Agonist/Neurokinin 1 Antagonist Activity with a 2,6-Dimethyl-Tyrosine: In vitro, In vivo, and NMR-Based Structural Studies. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 2029-2038.	6.4	30
61	Novel peptide ligands with dual acting pharmacophores designed for the pathophysiology of neuropathic pain. <i>Brain Research</i> , 2011, 1395, 1-11.	2.2	32
62	Recently patented and promising ORL-1 ligands: where have we been and where are we going?. <i>Expert Opinion on Therapeutic Patents</i> , 2010, 20, 291-305.	5.0	32
63	A cannabinoid 2 receptor agonist attenuates bone cancer-induced pain and bone loss. <i>Life Sciences</i> , 2010, 86, 646-653.	4.3	71
64	Novel d-amino acid tetrapeptides produce potent antinociception by selectively acting at peripheral μ -opioid receptors. <i>European Journal of Pharmacology</i> , 2008, 583, 62-72.	3.5	88
65	Oxycodone Plus Ultra-Low-Dose Naltrexone Attenuates Neuropathic Pain and Associated μ -Opioid Receptor-Gs Coupling. <i>Journal of Pain</i> , 2008, 9, 700-713.	1.4	64
66	A Structure-Activity Relationship Study and Combinatorial Synthetic Approach of C-Terminal Modified Bifunctional Peptides That Are μ Opioid Receptor Agonists and Neurokinin 1 Receptor Antagonists. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 1369-1376.	6.4	48