

# Samantha J Perez-Miller

## List of Publications by Year in descending order

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Version: 2024-02-01

30  
papers

1,210  
citations

430874

18  
h-index

477307

29  
g-index

35  
all docs

35  
docs citations

35  
times ranked

1708  
citing authors

#	ARTICLE	IF	CITATIONS
1	The small molecule compound C65780 alleviates pain by stabilizing voltage-gated sodium channels in the inactivated and slowly-recovering state. <i>Neuropharmacology</i> , 2022, , 109057.	4.1	0
2	Stereospecific Effects of Benzimidazolonepiperidine Compounds on T-Type Ca <sup>2+</sup> Channels and Pain. <i>ACS Chemical Neuroscience</i> , 2022, 13, 2035-2047.	3.5	4
3	Comparison of quinazoline and benzoylpyrazoline chemotypes targeting the CaV <sub>1</sub> - $\hat{I}^2$ interaction as antagonists of the N-type CaV <sub>2.2</sub> channel. <i>Channels</i> , 2021, 15, 128-135.	2.8	4
4	Novel Compounds Targeting Neuropilin Receptor 1 with Potential To Interfere with SARS-CoV-2 Virus Entry. <i>ACS Chemical Neuroscience</i> , 2021, 12, 1299-1312.	3.5	30
5	Antihypertensive drug treatment and susceptibility to SARS-CoV-2 infection in human PSC-derived cardiomyocytes and primary endothelial cells. <i>Stem Cell Reports</i> , 2021, 16, 2459-2472.	4.8	11
6	SARS-CoV-2 spike protein co-opts VEGF-A/neuropilin-1 receptor signaling to induce analgesia. <i>Pain</i> , 2021, 162, 243-252.	4.2	119
7	Selective targeting of Nav1.7 via inhibition of the CRMP2-Ubc9 interaction reduces pain in rodents. <i>Science Translational Medicine</i> , 2021, 13, eabh1314.	12.4	23
8	Druggability of CRMP2 for Neurodegenerative Diseases. <i>ACS Chemical Neuroscience</i> , 2020, 11, 2492-2505.	3.5	13
9	The role of cyclin-dependent kinase 5 in neuropathic pain. <i>Pain</i> , 2020, 161, 2674-2689.	4.2	20
10	A modulator of the low-voltage-activated T-type calcium channel that reverses HIV glycoprotein 120-, paclitaxel-, and spinal nerve ligation-induced peripheral neuropathies. <i>Pain</i> , 2020, 161, 2551-2570.	4.2	12
11	The Natural Flavonoid Naringenin Elicits Analgesia through Inhibition of Nav1.8 Voltage-Gated Sodium Channels. <i>ACS Chemical Neuroscience</i> , 2019, 10, 4834-4846.	3.5	20
12	Small Molecule Targeting TDP-43's RNA Recognition Motifs Reduces Locomotor Defects in a <i>Drosophila</i> Model of Amyotrophic Lateral Sclerosis (ALS). <i>ACS Chemical Biology</i> , 2019, 14, 2006-2013.	3.4	45
13	Structural Insights Into TDP-43 and Effects of Post-translational Modifications. <i>Frontiers in Molecular Neuroscience</i> , 2019, 12, 301.	2.9	86
14	Targeting the CaV <sub>1</sub> - $\hat{I}^2$ interaction yields an antagonist of the N-type CaV <sub>2.2</sub> channel with broad antinociceptive efficacy. <i>Pain</i> , 2019, 160, 1644-1661.	4.2	30
15	Remodeling the interactions between TDP43 and RNA for development of therapeutics for ALS. <i>FASEB Journal</i> , 2019, 33, 670.1.	0.5	0
16	Homology-guided mutational analysis reveals the functional requirements for antinociceptive specificity of collapsin response mediator protein 2-derived peptides. <i>British Journal of Pharmacology</i> , 2018, 175, 2244-2260.	5.4	40
17	A Chemical Biology Approach to Model Pontocerebellar Hypoplasia Type 1B (PCH1B). <i>ACS Chemical Biology</i> , 2018, 13, 3000-3010.	3.4	9
18	Chemical shift perturbation mapping of the Ubc9-CRMP2 interface identifies a pocket in CRMP2 amenable for allosteric modulation of Nav1.7 channels. <i>Channels</i> , 2018, 12, 219-227.	2.8	17

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19	Inhibition of the Ubc9 E2 SUMO-conjugating enzymeâ€‘CRMP2 interaction decreases Nav1.7 currents and reverses experimental neuropathic pain. <i>Pain</i> , 2018, 159, 2115-2127.	4.2	49
20	A single structurally conserved SUMOylation site in CRMP2 controls Nav1.7 function. <i>Channels</i> , 2017, 11, 316-328.	2.8	34
21	(S)-Lacosamide Binding to Collapsin Response Mediator Protein 2 (CRMP2) Regulates CaV2.2 Activity by Subverting Its Phosphorylation by Cdk5. <i>Molecular Neurobiology</i> , 2016, 53, 1959-1976.	4.0	50
22	Catalytic contribution of threonine 244 in human ALDH2. <i>Chemico-Biological Interactions</i> , 2013, 202, 32-40.	4.0	4
23	Expression and purification of functional human glycogen synthase-1 (hGYS1) in insect cells. <i>Protein Expression and Purification</i> , 2013, 90, 78-83.	1.3	12
24	Discovery of a Novel Class of Covalent Inhibitor for Aldehyde Dehydrogenases. <i>Journal of Biological Chemistry</i> , 2011, 286, 43486-43494.	3.4	65
25	High resolution X-ray structures of mouse major urinary protein nasal isoform in complex with pheromones. <i>Protein Science</i> , 2010, 19, 1469-1479.	7.6	23
26	Alda-1 is an agonist and chemical chaperone for the common human aldehyde dehydrogenase 2 variant. <i>Nature Structural and Molecular Biology</i> , 2010, 17, 159-164.	8.2	193
27	Light Chain C-Terminal Region Reinforces the Stability of Clathrin Heavy Chain Trimers. <i>Traffic</i> , 2007, 8, 1101-1110.	2.7	31
28	Coenzyme Isomerization Is Integral to Catalysis in Aldehyde Dehydrogenaseâ€‘â€‘. <i>Biochemistry</i> , 2003, 42, 7100-7109.	2.5	152
29	Three-dimensional structures of the three human class I alcohol dehydrogenases. <i>Protein Science</i> , 2001, 10, 697-706.	7.6	65
30	Order and disorder in mitochondrial aldehyde dehydrogenase. <i>Chemico-Biological Interactions</i> , 2001, 130-132, 3-14.	4.0	23