

Jing Guo

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5825119/publications.pdf>

Version: 2024-02-01

13
papers

621
citations

840776

11
h-index

1125743

13
g-index

13
all docs

13
docs citations

13
times ranked

592
citing authors

#	ARTICLE	IF	CITATIONS
1	Central Nervous System Targets: Glial Cell Mechanisms in Chronic Pain. <i>Neurotherapeutics</i> , 2020, 17, 846-860.	4.4	138
2	STING controls nociception via type I interferon signalling in sensory neurons. <i>Nature</i> , 2021, 591, 275-280.	27.8	107
3	miRNA-711 Binds and Activates TRPA1 Extracellularly to Evoke Acute and Chronic Pruritus. <i>Neuron</i> , 2018, 99, 449-463.e6.	8.1	79
4	Anti-PD-1 treatment impairs opioid antinociception in rodents and nonhuman primates. <i>Science Translational Medicine</i> , 2020, 12, .	12.4	54
5	HepaCAM controls astrocyte self-organization and coupling. <i>Neuron</i> , 2021, 109, 2427-2442.e10.	8.1	52
6	STING suppresses bone cancer pain via immune and neuronal modulation. <i>Nature Communications</i> , 2021, 12, 4558.	12.8	50
7	Central opioid receptors mediate morphine-induced itch and chronic itch via disinhibition. <i>Brain</i> , 2021, 144, 665-681.	7.6	45
8	Lysophospholipids Contribute to Oxaliplatin-Induced Acute Peripheral Pain. <i>Journal of Neuroscience</i> , 2020, 40, 9519-9532.	3.6	28
9	PD-1 Regulates GABAergic Neurotransmission and GABA-Mediated Analgesia and Anesthesia. <i>Science</i> , 2020, 23, 101570.	4.1	23
10	Repurposing cancer drugs identifies kenpauillone which ameliorates pathologic pain in preclinical models via normalization of inhibitory neurotransmission. <i>Nature Communications</i> , 2021, 12, 6208.	12.8	16
11	PDCD4 Knockdown Induces Senescence in Hepatoma Cells by Up-Regulating the p21 Expression. <i>Frontiers in Oncology</i> , 2018, 8, 661.	2.8	15
12	Degradation of the Tumor Suppressor PDCD4 Is Impaired by the Suppression of p62/SQSTM1 and Autophagy. <i>Cells</i> , 2020, 9, 218.	4.1	10
13	Identification and characterization of novel candidate compounds targeting μ - and δ -transmembrane μ -opioid receptor isoforms. <i>British Journal of Pharmacology</i> , 2021, 178, 2709-2726.	5.4	4