Jeremy E Chojnacki

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	A Novel Pharmacologic Inhibitor of the NLRP3 Inflammasome Limits Myocardial Injury After Ischemia–Reperfusion in the Mouse. Journal of Cardiovascular Pharmacology, 2014, 63, 316-322.	1.9	215
2	Inhibition of the NLRP3 inflammasome limits the inflammatory injury following myocardial ischemia–reperfusion in the mouse. International Journal of Cardiology, 2016, 209, 215-220.	1.7	173
3	NLRP3 Inflammasome Inhibitor Ameliorates Amyloid Pathology in a Mouse Model of Alzheimer's Disease. Molecular Neurobiology, 2018, 55, 1977-1987.	4.0	153
4	Pharmacologic Inhibition of the NLRP3 Inflammasome Preserves Cardiac Function After Ischemic and Nonischemic Injury in the Mouse. Journal of Cardiovascular Pharmacology, 2015, 66, 1-8.	1.9	128
5	Development and Characterization of a Hydroxyl-Sulfonamide Analogue, 5-Chloro- <i>N</i> -[2-(4-hydroxysulfamoyl-phenyl)-ethyl]-2-methoxy-benzamide, as a Novel NLRP3 Inflammasome Inhibitor for Potential Treatment of Multiple Sclerosis. ACS Chemical Neuroscience, 2017. 8, 2194-2201.	3.5	77
6	Bivalent Ligand Containing Curcumin and Cholesterol as a Fluorescence Probe for Aβ Plaques in Alzheimer's Disease. ACS Chemical Neuroscience, 2012, 3, 141-146.	3.5	70
7	Discovery of 5-(4-Hydroxyphenyl)-3-oxo-pentanoic Acid [2-(5-Methoxy-1H-indol-3-yl)-ethyl]-amide as a Neuroprotectant for Alzheimer's Disease by Hybridization of Curcumin and Melatonin. ACS Chemical Neuroscience, 2014, 5, 690-699.	3.5	66
8	Curcumin/Melatonin Hybrid 5-(4-Hydroxy-phenyl)-3-oxo-pentanoic Acid [2-(5-Methoxy-1 <i>H</i> -indol-3-yl)-ethyl]-amide Ameliorates AD-Like Pathology in the APP/PS1 Mouse Model. ACS Chemical Neuroscience, 2015, 6, 1393-1399.	3.5	51
9	Design and biological characterization of hybrid compounds of curcumin and thalidomide for multiple myeloma. Organic and Biomolecular Chemistry, 2013, 11, 4757.	2.8	47
10	BF3·OEt2-promoted concise synthesis of difluoroboron-derivatized curcumins from aldehydes and 2,4-pentanedione. Tetrahedron Letters, 2013, 54, 2070-2073.	1.4	45
11	Dissecting the sequence determinants for dephosphorylation by the catalytic subunits of phosphatases PP1 and PP2A. Nature Communications, 2020, 11, 3583.	12.8	38
12	Bivalent ligands incorporating curcumin and diosgenin as multifunctional compounds against Alzheimer's disease. Bioorganic and Medicinal Chemistry, 2015, 23, 7324-7331.	3.0	29
13	Bivalent Compound 17MN Exerts Neuroprotection through Interaction at Multiple Sites in a Cellular Model of Alzheimer's Disease. Journal of Alzheimer's Disease, 2015, 47, 1021-1033.	2.6	14
14	Mechanistic Insight of Bivalent Compound 21MO as Potential Neuroprotectant for Alzheimer's Disease. Molecules, 2016, 21, 412.	3.8	9
15	Structural and mechanistic insights into the interaction of the circadian transcription factor BMAL1 with the KIX domain of the CREB-binding protein. Journal of Biological Chemistry, 2019, 294, 16604-16619.	3.4	9
16	PLDMS: Phosphopeptide Library Dephosphorylation Followed by Mass Spectrometry Analysis to Determine the Specificity of Phosphatases for Dephosphorylation Site Sequences. Methods in Molecular Biology, 2022, , 43-64.	0.9	1
17	P1-400: DEVELOPMENT OF CURCUMIN/MELATONIN HYBRIDS AND THEIR POTENTIAL APPLICATION IN AD. , 2014, 10, P460-P460.		0

¹⁸ P4-170: Mechanistic studies of a bivalent compound containing curcumin and a membrane anchorage as a neuroprotectant in mc65 cell model. , 2015, 11, P844-P844.

#	Article	IF	CITATIONS
19	P1-305: Design and evaluation of fluorescent probes to elucidate the mechanism of curcumin/melatonin hybrids for Alzheimer's disease. , 2015, 11, P473-P473.		0