Marie A Bogoyevitch

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2272403/publications.pdf

Version: 2024-02-01

60 papers

4,285 citations

230014 27 h-index 58 g-index

61 all docs

61 does citations

61 times ranked

6645 citing authors

#	Article	IF	CITATIONS
1	Bimolecular Fluorescence Complementation: Quantitative Analysis of In Cell Interaction of Nuclear Transporter Importin α with Cargo Proteins. Methods in Molecular Biology, 2022, 2502, 215-233.	0.4	1
2	Impact of Respiratory Syncytial Virus Infection on Host Functions: Implications for Antiviral Strategies. Physiological Reviews, 2020, 100, 1527-1594.	13.1	30
3	The broad spectrum antiviral ivermectin targets the host nuclear transport importin $\hat{l}\pm\hat{l}^21$ heterodimer. Antiviral Research, 2020, 177, 104760.	1.9	255
4	The ataxin-1 interactome reveals direct connection with multiple disrupted nuclear transport pathways. Nature Communications, 2020, 11, 3343.	5 . 8	15
5	Nuclear bodies formed by polyQ-ataxin-1 protein are liquid RNA/protein droplets with tunable dynamics. Scientific Reports, 2020, 10, 1557.	1.6	15
6	Subversion of Host Cell Mitochondria by RSV to Favor Virus Production is Dependent on Inhibition of Mitochondrial Complex I and ROS Generation. Cells, 2019, 8, 1417.	1.8	28
7	Oligonucleotide-directed STAT3 alternative splicing switch drives anti-tumorigenic outcomes in MCF10 human breast cancer cells. Biochemical and Biophysical Research Communications, 2019, 513, 1076-1082.	1.0	6
8	Pathogenic E2K mutation of doublecortin X (DCX) alters microtubule stabilisation and actin filament association. Biochemical and Biophysical Research Communications, 2019, 513, 540-545.	1.0	1
9	Doublecortin X (DCX) serine 28 phosphorylation is a regulatory switch, modulating association of DCX with microtubules and actin filaments. Biochimica Et Biophysica Acta - Molecular Cell Research, 2019, 1866, 638-649.	1.9	9
10	Respiratory syncytial virus co-opts host mitochondrial function to favour infectious virus production. ELife, 2019, 8, .	2.8	47
11	TrawlerWeb: an online de novo motif discovery tool for next-generation sequencing datasets. BMC Genomics, 2018, 19, 238.	1.2	12
12	Complementary proteomics strategies capture an ataxin-1 interactome in Neuro-2a cells. Scientific Data, 2018, 5, 180262.	2.4	8
13	c-Jun N-terminal kinase activity is required for efficient respiratory syncytial virus production. Biochemical and Biophysical Research Communications, 2017, 483, 64-68.	1.0	7
14	Mitochondrial protein p32/HAPB1/gC1qR/C1qbp is required for efficient respiratory syncytial virus production. Biochemical and Biophysical Research Communications, 2017, 489, 460-465.	1.0	25
15	Dynamic microtubule association of Doublecortin X (DCX) is regulated by its C-terminus. Scientific Reports, 2017, 7, 5245.	1.6	15
16	Quantifying the dynamics of the oligomeric transcription factor STAT3 by pair correlation of molecular brightness. Nature Communications, 2016, 7, 11047.	5 . 8	28
17	JNK Signaling: Regulation and Functions Based on Complex Protein-Protein Partnerships. Microbiology and Molecular Biology Reviews, 2016, 80, 793-835.	2.9	348
18	Aurora A phosphorylation of WD40-repeat protein 62 in mitotic spindle regulation. Cell Cycle, 2016, 15, 413-424.	1.3	26

#	Article	IF	CITATIONS
19	Opposing roles for JNK and Aurora A in regulating WD40-Repeat Protein 62 association with spindle microtubules. Journal of Cell Science, 2015, 128, 527-40.	1.2	41
20	Dual role of Src kinase in governing neuronal survival. Brain Research, 2015, 1594, 1-14.	1.1	15
21	Hyperosmotic stress sustains cytokine-stimulated phosphorylation of STAT3, but slows its nuclear trafficking and impairs STAT3-dependent transcription. Cellular Signalling, 2014, 26, 815-824.	1.7	5
22	Differences in c-Jun N-terminal kinase recognition and phosphorylation of closely related stathmin-family members. Biochemical and Biophysical Research Communications, 2014, 446, 248-254.	1.0	17
23	Intracellular mobility and nuclear trafficking of the stress-activated kinase JNK1 are impeded by hyperosmotic stress. Biochimica Et Biophysica Acta - Molecular Cell Research, 2014, 1843, 253-264.	1.9	10
24	The JNK1/JNK3 interactome – Contributions by the JNK3 unique N-terminus and JNK common docking site residues. Biochemical and Biophysical Research Communications, 2014, 453, 576-581.	1.0	10
25	Oxidative stress impairs multiple regulatory events to drive persistent cytokine-stimulated STAT3 phosphorylation. Biochimica Et Biophysica Acta - Molecular Cell Research, 2014, 1843, 483-494.	1.9	31
26	Identification and characterization of bi-thiazole-2,2′-diamines as kinase inhibitory scaffolds. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2013, 1834, 1077-1088.	1.1	8
27	A novel retro-inverso peptide is a preferential JNK substrate-competitive inhibitor. International Journal of Biochemistry and Cell Biology, 2013, 45, 1939-1950.	1.2	8
28	p32 protein levels are integral to mitochondrial and endoplasmic reticulum morphology, cell metabolism and survival. Biochemical Journal, 2013, 453, 381-391.	1.7	61
29	Characterization of a microtubuleâ€associated protein, doublecortin (DCX), as a substrate of câ€jun Nâ€terminal Kinases (JNKs). FASEB Journal, 2013, 27, 1042.3.	0.2	0
30	Selective STAT3-α or -β expression reveals spliceform-specific phosphorylation kinetics, nuclear retention and distinct gene expression outcomes. Biochemical Journal, 2012, 447, 125-136.	1.7	48
31	WD40-repeat protein 62 is a JNK-phosphorylated spindle pole protein required for spindle maintenance and timely mitotic progression Journal of Cell Science, 2012, 125, 5096-109.	1.2	69
32	Characterization of a novel JNK (c-Jun N-terminal kinase) inhibitory peptide. Biochemical Journal, 2011, 434, 399-413.	1.7	27
33	C-Jun N-terminal kinase controls TDP-43 accumulation in stress granules induced by oxidative stress. Molecular Neurodegeneration, 2011, 6, 57.	4.4	103
34	c-Jun N-terminal kinase (JNK) signaling: Recent advances and challenges. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2010, 1804, 463-475.	1.1	257
35	c-Jun N-terminal Kinase Phosphorylation of Stathmin Confers Protection against Cellular Stress. Journal of Biological Chemistry, 2010, 285, 29001-29013.	1.6	30
36	Inhibitors of c-Jun N-terminal kinases—JuNK no more?. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2008, 1784, 76-93.	1.1	114

#	Article	IF	Citations
37	Changes in the Transcriptional Profile of Cardiac Myocytes Following Green Fluorescent Protein Expression. DNA and Cell Biology, 2007, 26, 727-736.	0.9	12
38	Gene expression profiling reveals complex changes following MEK-EE expression in cardiac myocytes. International Journal of Biochemistry and Cell Biology, 2007, 39, 349-365.	1.2	3
39	Necrotic death of neurons following an excitotoxic insult is prevented by a peptide inhibitor of c-jun N-terminal kinase. Journal of Neurochemistry, 2007, 102, 65-76.	2.1	31
40	A new paradigm for protein kinase inhibition: blocking phosphorylation without directly targeting ATP binding. Drug Discovery Today, 2007, 12, 622-633.	3.2	170
41	Contrasting actions of prolonged mitogen-activated protein kinase activation on cell survival. Biochemical and Biophysical Research Communications, 2006, 345, 843-850.	1.0	4
42	The isoform-specific functions of the c-Jun N-terminal Kinases (JNKs): differences revealed by gene targeting. BioEssays, 2006, 28, 923-934.	1.2	166
43	Uses for JNK: the Many and Varied Substrates of the c-Jun N-Terminal Kinases. Microbiology and Molecular Biology Reviews, 2006, 70, 1061-1095.	2.9	488
44	Peptide inhibitors of protein kinasesâ€"discovery, characterisation and use. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2005, 1754, 79-99.	1.1	61
45	Therapeutic promise of JNK ATP-noncompetitive inhibitors. Trends in Molecular Medicine, 2005, 11, 232-239.	3.5	41
46	Reverse Two-hybrid Screening Identifies Residues of JNK Required for Interaction with the Kinase Interaction Motif of JNK-interacting Protein-1. Journal of Biological Chemistry, 2004, 279, 43178-43189.	1.6	25
47	Contribution of the Membrane-distal Tyrosine in Intracellular Signaling by the Granulocyte Colony-stimulating Factor Receptor. Journal of Biological Chemistry, 2004, 279, 326-340.	1.6	14
48	The Critical Features and the Mechanism of Inhibition of a Kinase Interaction Motif-based Peptide Inhibitor of JNK. Journal of Biological Chemistry, 2004, 279, 36327-36338.	1.6	54
49	An update on the cardiac effects of erythropoietin cardioprotection by erythropoietin and the lessons learnt from studies in neuroprotection. Cardiovascular Research, 2004, 63, 208-216.	1.8	121
50	Counting on mitogen-activated protein kinasesâ€"ERKs 3, 4, 5, 6, 7 and 8. Cellular Signalling, 2004, 16, 1345-1354.	1.7	118
51	Targeting the JNK MAPK cascade for inhibition: basic science and therapeutic potential. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2004, 1697, 89-101.	1.1	231
52	Identification of the Critical Features of a Small Peptide Inhibitor of JNK Activity. Journal of Biological Chemistry, 2002, 277, 10987-10997.	1.6	189
53	Crossing the Membrane: Nonviral and Viral Delivery Methods for Use In Vitro and In Vivo. DNA and Cell Biology, 2002, 21, 855-856.	0.9	1
54	Taking the Cell by Stealth or Storm? Protein Transduction Domains (PTDs) as Versatile Vectors for Delivery. DNA and Cell Biology, 2002, 21, 879-894.	0.9	38

#	Article	IF	CITATIONS
55	Adrenergic receptor stimulation of the mitogen-activated protein kinase cascade and cardiac hypertrophy. Biochemical Journal, 1996, 314, 115-121.	1.7	158
56	Stimulation of the Stress-Activated Mitogen-Activated Protein Kinase Subfamilies in Perfused Heart. Circulation Research, 1996, 79, 162-173.	2.0	462
57	Endothelin-1, phorbol esters and phenylephrine stimulate MAP kinase activities in ventricular cardiomyocytes. FEBS Letters, 1993, 317, 271-275.	1.3	160
58	Mitogen-activated protein (MAP) kinase stimulation by phorbol esters and external load in the isolated perfused heart. Biochemical Society Transactions, 1993, 21, 356S-356S.	1.6	3
59	Acidic fibroblast growth factor or endothelin-1 stimulate the MAP kinase cascade in cardiac myocytes. Biochemical Society Transactions, 1993, 21, 358S-358S.	1.6	4
60	Effects of catecholamines on protein synthesis and cyclic AMP concentrations in the isolated working heart. Biochemical Society Transactions, 1991, 19, 276S-276S.	1.6	0