

Tatiana R. Rosenstock

List of Publications by Year in descending order

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Version: 2024-02-01

39
papers

1,471
citations

304368

22
h-index

315357

38
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43
all docs

43
docs citations

43
times ranked

2487
citing authors

#	ARTICLE	IF	CITATIONS
1	Analysis of Mitochondrial Dysfunction by Microplate Reader in hiPSC-Derived Neuronal Cell Models of Neurodegenerative Disorders. <i>Methods in Molecular Biology</i> , 2022, , 1-21.	0.4	2
2	Redox profiles of amyotrophic lateral sclerosis lymphoblasts with or without known SOD1 mutations. <i>European Journal of Clinical Investigation</i> , 2022, 52, e13798.	1.7	3
3	Oxygen Consumption Evaluation: An Important Indicator of Metabolic State, , and Cell Fate Along. <i>Methods in Molecular Biology</i> , 2021, 2240, 207-230.	0.4	2
4	Neonatal Rotenone Administration Induces Psychiatric Disorder-Like Behavior and Changes in Mitochondrial Biogenesis and Synaptic Proteins in Adulthood. <i>Molecular Neurobiology</i> , 2021, 58, 3015-3030.	1.9	5
5	Mitochondrial SIRT3 confers neuroprotection in Huntington's disease by regulation of oxidative challenges and mitochondrial dynamics. <i>Free Radical Biology and Medicine</i> , 2021, 163, 163-179.	1.3	42
6	Increased acute blood flow induced by the aqueous extract of <i>Euterpe oleracea</i> Mart. fruit pulp in rats in vivo is not related to the direct activation of endothelial cells. <i>Journal of Ethnopharmacology</i> , 2021, 271, 113885.	2.0	3
7	Haloperidol rescues the schizophrenia-like phenotype in adulthood after rotenone administration in neonatal rats. <i>Psychopharmacology</i> , 2021, 238, 2569-2585.	1.5	3
8	New Avenues for the Treatment of Huntington's Disease. <i>International Journal of Molecular Sciences</i> , 2021, 22, 8363.	1.8	55
9	Autophagy Dysfunction as a Phenotypic Readout in hiPSC-Derived Neuronal Cell Models of Neurodegenerative Diseases. <i>Methods in Molecular Biology</i> , 2021, , 103-136.	0.4	4
10	Human Induced Pluripotent Stem Cell Models of Neurodegenerative Disorders for Studying the Biomedical Implications of Autophagy. <i>Journal of Molecular Biology</i> , 2020, 432, 2754-2798.	2.0	15
11	Mitochondrial Dysfunction, Neurogenesis, and Epigenetics: Putative Implications for Amyotrophic Lateral Sclerosis Neurodegeneration and Treatment. <i>Frontiers in Neuroscience</i> , 2020, 14, 679.	1.4	38
12	Decreased Mitochondrial Function, Biogenesis, and Degradation in Peripheral Blood Mononuclear Cells from Amyotrophic Lateral Sclerosis Patients as a Potential Tool for Biomarker Research. <i>Molecular Neurobiology</i> , 2020, 57, 5084-5102.	1.9	20
13	S179. ADMINISTRATION OF ROTENONE DURING NEONATAL PERIOD INDUCES SCHIZOPHRENIA-LIKE BEHAVIOR. <i>Schizophrenia Bulletin</i> , 2020, 46, S106-S106.	2.3	0
14	Chemical Screening Approaches Enabling Drug Discovery of Autophagy Modulators for Biomedical Applications in Human Diseases. <i>Frontiers in Cell and Developmental Biology</i> , 2019, 7, 38.	1.8	37
15	Mitochondrial Dysfunction and Changes in High-Energy Compounds in Different Cellular Models Associated to Hypoxia: Implication to Schizophrenia. <i>Scientific Reports</i> , 2019, 9, 18049.	1.6	18
16	Metabolic Alteration and Amyotrophic Lateral Sclerosis Outcome: A Systematic Review. <i>Frontiers in Neurology</i> , 2019, 10, 1205.	1.1	17
17	In Vitro Screening Platforms for Identifying Autophagy Modulators in Mammalian Cells. <i>Methods in Molecular Biology</i> , 2019, 1880, 389-428.	0.4	14
18	Histone Deacetylase Inhibitors Protect Against Pyruvate Dehydrogenase Dysfunction in Huntington's Disease. <i>Journal of Neuroscience</i> , 2017, 37, 2776-2794.	1.7	50

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19	Selfish brain and selfish immune system interplay: A theoretical framework for metabolic comorbidities of mood disorders. <i>Neuroscience and Biobehavioral Reviews</i> , 2017, 72, 43-49.	2.9	25
20	Comparative Mitochondrial-Based Protective Effects of Resveratrol and Nicotinamide in Huntington's Disease Models. <i>Molecular Neurobiology</i> , 2017, 54, 5385-5399.	1.9	105
21	Dysregulation of autophagy as a common mechanism in lysosomal storage diseases. <i>Essays in Biochemistry</i> , 2017, 61, 733-749.	2.1	138
22	Gene Expression Control by Glucocorticoid Receptors during Innate Immune Responses. <i>Frontiers in Endocrinology</i> , 2016, 7, 31.	1.5	81
23	CD36 is expressed in a defined subpopulation of neurons in the olfactory epithelium. <i>Scientific Reports</i> , 2016, 6, 25507.	1.6	34
24	Insulin and IGF-1 regularize energy metabolites in neural cells expressing full-length mutant huntingtin. <i>Neuropeptides</i> , 2016, 58, 73-81.	0.9	28
25	Activation of IGF-1 and Insulin Signaling Pathways Ameliorate Mitochondrial Function and Energy Metabolism in Huntington's Disease Human Lymphoblasts. <i>Molecular Neurobiology</i> , 2015, 51, 331-348.	1.9	66
26	Deficits of pyruvate dehydrogenase and mitochondrial dysfunction in Huntington's disease - role of histone deacetylase inhibitors. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2014, 1837, e67.	0.5	0
27	Insulin and IGF-1 improve mitochondrial function in a PI-3K/Akt-dependent manner and reduce mitochondrial generation of reactive oxygen species in Huntington's disease knock-in striatal cells. <i>Free Radical Biology and Medicine</i> , 2014, 74, 129-144.	1.3	116
28	Current Therapeutic Advances in Patients and Experimental Models of Huntington's Disease. <i>Current Drug Targets</i> , 2014, 15, 313-334.	1.0	8
29	Glutathione redox cycle dysregulation in Huntington's disease knock-in striatal cells. <i>Free Radical Biology and Medicine</i> , 2012, 53, 1857-1867.	1.3	60
30	FK506 ameliorates cell death features in Huntington's disease striatal cell models. <i>Neurochemistry International</i> , 2011, 59, 600-609.	1.9	29
31	Glutamate-induced alterations in Ca^{2+} signaling are modulated by mitochondrial Ca^{2+} handling capacity in brain slices of R6/1 transgenic mice. <i>European Journal of Neuroscience</i> , 2010, 32, 60-70.	1.2	30
32	Mitochondrial-Associated Metabolic Changes and Neurodegeneration in Huntingtons Disease - from Clinical Features to the Bench. <i>Current Drug Targets</i> , 2010, 11, 1218-1236.	1.0	50
33	Calcium and cell death signaling in neurodegeneration and aging. <i>Anais Da Academia Brasileira De Ciencias</i> , 2009, 81, 467-475.	0.3	65
34	Old mice present increased levels of succinate dehydrogenase activity and lower vulnerability to dyskinetic effects of 3-nitropropionic acid. <i>Pharmacology Biochemistry and Behavior</i> , 2009, 91, 327-332.	1.3	14
35	Increase in bax expression and apoptosis are associated in Huntington's disease progression. <i>Neuroscience Letters</i> , 2008, 438, 59-63.	1.0	25
36	Chapter Two Evaluation of Some Cell Death Features by Real Time Real Space Microscopy. <i>Methods in Enzymology</i> , 2008, 442, 27-50.	0.4	5

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37	Bax affects intracellular Ca ²⁺ stores and induces Ca ²⁺ wave propagation. <i>Cell Death and Differentiation</i> , 2004, 11, 1265-1276.	5.0	42
38	Mitochondrial calcium, oxidative stress and apoptosis in a neurodegenerative disease model induced by 3-nitropropionic acid. <i>Journal of Neurochemistry</i> , 2004, 88, 1220-1228.	2.1	99
39	Mitochondria, calcium and pro-apoptotic proteins as mediators in cell death signaling. <i>Brazilian Journal of Medical and Biological Research</i> , 2003, 36, 183-190.	0.7	119