

Francisca Diaz

List of Publications by Year in descending order

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Version: 2024-02-01

50
papers

4,502
citations

159585

30
h-index

182427

51
g-index

54
all docs

54
docs citations

54
times ranked

7278
citing authors

#	ARTICLE	IF	CITATIONS
1	Nicotine Exposure Along with Oral Contraceptive Treatment in Female Rats Exacerbates Post-cerebral Ischemic Hypoperfusion Potentially via Altered Histamine Metabolism. <i>Translational Stroke Research</i> , 2021, 12, 817-828.	4.2	8
2	Simultaneous nicotine and oral contraceptive exposure alters brain energy metabolism and exacerbates ischemic stroke injury in female rats. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2021, 41, 793-804.	4.3	13
3	Cardiomyopathic mutations in essential light chain reveal mechanisms regulating the super relaxed state of myosin. <i>Journal of General Physiology</i> , 2021, 153, .	1.9	14
4	Enhanced glycolysis and GSK3 inactivation promote brain metabolic adaptations following neuronal mitochondrial stress. <i>Human Molecular Genetics</i> , 2021, , .	2.9	0
5	ATAD3A has a scaffolding role regulating mitochondria inner membrane structure and protein assembly. <i>Cell Reports</i> , 2021, 37, 110139.	6.4	34
6	Respiratory supercomplexes act as a platform for complex III-mediated maturation of human mitochondrial complexes I and IV. <i>EMBO Journal</i> , 2020, 39, e102817.	7.8	102
7	Hypoxia Promotes Mitochondrial Complex I Abundance via HIF-1 α in Complex III and Complex IV Deficient Cells. <i>Cells</i> , 2020, 9, 2197.	4.1	6
8	Methods and models for functional studies on mtDNA mutations. , 2020, , 305-349.		1
9	Mitochondrial disease disrupts hepatic allostasis and lowers the threshold for immune-mediated liver toxicity. <i>Molecular Metabolism</i> , 2020, 37, 100981.	6.5	8
10	Impaired mitophagy links mitochondrial disease to epithelial stress in methylmalonyl-CoA mutase deficiency. <i>Nature Communications</i> , 2020, 11, 970.	12.8	65
11	Myopathy reversion in mice after restauration of mitochondrial complex I. <i>EMBO Molecular Medicine</i> , 2020, 12, e10674.	6.9	29
12	Metformin delays neurological symptom onset in a mouse model of neuronal complex I deficiency. <i>JCI Insight</i> , 2020, 5, .	5.0	8
13	Photobiomodulation enhancement of cell proliferation at 660nm does not require cytochrome c oxidase. <i>Journal of Photochemistry and Photobiology B: Biology</i> , 2019, 194, 71-75.	3.8	51
14	Ablation of Cytochrome c in Adult Forebrain Neurons Impairs Oxidative Phosphorylation Without Detectable Apoptosis. <i>Molecular Neurobiology</i> , 2019, 56, 3722-3735.	4.0	9
15	Overexpression of PGC-1 α in aging muscle enhances a subset of young-like molecular patterns. <i>Aging Cell</i> , 2018, 17, e12707.	6.7	57
16	The Organization of Mitochondrial Supercomplexes is Modulated by Oxidative Stress In Vivo in Mouse Models of Mitochondrial Encephalopathy. <i>International Journal of Molecular Sciences</i> , 2018, 19, 1582.	4.1	16
17	ATAD3 controls mitochondrial cristae structure, influencing mtDNA replication and cholesterol levels in muscle. <i>Journal of Cell Science</i> , 2018, 131, .	2.0	68
18	Cytochrome c Oxidase Activity Is a Metabolic Checkpoint that Regulates Cell Fate Decisions During T Cell Activation and Differentiation. <i>Cell Metabolism</i> , 2017, 25, 1254-1268.e7.	16.2	125

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19	Respiration-Deficient Astrocytes Survive As Glycolytic Cells <i>In Vivo</i> . <i>Journal of Neuroscience</i> , 2017, 37, 4231-4242.	3.6	97
20	GSNOR Deficiency Enhances <i>In Situ</i> Skeletal Muscle Strength, Fatigue Resistance, and RyR1 S-Nitrosylation Without Impacting Mitochondrial Content and Activity. <i>Antioxidants and Redox Signaling</i> , 2017, 26, 165-181.	5.4	18
21	The CoQH ₂ /CoQ Ratio Serves as a Sensor of Respiratory Chain Efficiency. <i>Cell Reports</i> , 2016, 15, 197-209.	6.4	215
22	Pioglitazone ameliorates the phenotype of a novel Parkinson's disease mouse model by reducing neuroinflammation. <i>Molecular Neurodegeneration</i> , 2016, 11, 25.	10.8	57
23	Enhanced Transcriptional Activity and Mitochondrial Localization of STAT3 Co-induce Axon Regrowth in the Adult Central Nervous System. <i>Cell Reports</i> , 2016, 15, 398-410.	6.4	91
24	Sustained AMPK activation improves muscle function in a mitochondrial myopathy mouse model by promoting muscle fiber regeneration. <i>Human Molecular Genetics</i> , 2016, 25, 3178-3191.	2.9	23
25	Mitochondrial Diseases Part I: Mouse models of OXPHOS deficiencies caused by defects in respiratory complex subunits or assembly factors. <i>Mitochondrion</i> , 2015, 21, 76-91.	3.4	36
26	The Mitochondrial Metallochaperone SCO1 Is Required to Sustain Expression of the High-Affinity Copper Transporter CTR1 and Preserve Copper Homeostasis. <i>Cell Reports</i> , 2015, 10, 933-943.	6.4	37
27	Mitochondrial Diseases Part II: Mouse models of OXPHOS deficiencies caused by defects in regulatory factors and other components required for mitochondrial function. <i>Mitochondrion</i> , 2015, 22, 96-118.	3.4	23
28	Mitochondrial Diseases Part III: Therapeutic interventions in mouse models of OXPHOS deficiencies. <i>Mitochondrion</i> , 2015, 23, 71-80.	3.4	10
29	Periodic Estrogen Receptor-Beta Activation: A Novel Approach to Prevent Ischemic Brain Damage. <i>Neurochemical Research</i> , 2015, 40, 2009-2017.	3.3	15
30	Partial complex I deficiency due to the CNS conditional ablation of Ndufa5 results in a mild chronic encephalopathy but no increase in oxidative damage. <i>Human Molecular Genetics</i> , 2014, 23, 1399-1412.	2.9	33
31	Oral Contraceptives and Nicotine Synergistically Exacerbate Cerebral Ischemic Injury in the Female Brain. <i>Translational Stroke Research</i> , 2013, 4, 402-412.	4.2	11
32	Cells Lacking Rieske Iron-Sulfur Protein Have a Reactive Oxygen Species-Associated Decrease in Respiratory Complexes I and IV. <i>Molecular and Cellular Biology</i> , 2012, 32, 415-429.	2.3	107
33	A defect in the mitochondrial complex III, but not complex IV, triggers early ROS-dependent damage in defined brain regions. <i>Human Molecular Genetics</i> , 2012, 21, 5066-5077.	2.9	81
34	Glycolytic oligodendrocytes maintain myelin and long-term axonal integrity. <i>Nature</i> , 2012, 485, 517-521.	27.8	1,120
35	Mitochondrial disorders caused by mutations in respiratory chain assembly factors. <i>Seminars in Fetal and Neonatal Medicine</i> , 2011, 16, 197-204.	2.3	51
36	Cytochrome c oxidase deficiency: Patients and animal models. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2010, 1802, 100-110.	3.8	113

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37	PGC-1 β induced expression partially compensates for respiratory chain defects in cells from patients with mitochondrial disorders. <i>Human Molecular Genetics</i> , 2009, 18, 1805-1812.	2.9	99
38	Endurance exercise is protective for mice with mitochondrial myopathy. <i>Journal of Applied Physiology</i> , 2009, 106, 1712-1719.	2.5	76
39	Mouse models of oxidative phosphorylation defects: Powerful tools to study the pathobiology of mitochondrial diseases. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2009, 1793, 171-180.	4.1	33
40	Evaluation of the Mitochondrial Respiratory Chain and Oxidative Phosphorylation System Using Blue Native Gel Electrophoresis. <i>Current Protocols in Human Genetics</i> , 2009, 63, Unit19.4.	3.5	49
41	Evaluation of the Mitochondrial Respiratory Chain and Oxidative Phosphorylation System Using Polarography and Spectrophotometric Enzyme Assays. <i>Current Protocols in Human Genetics</i> , 2009, 63, Unit19.3.	3.5	178
42	A 3 α UTR Modification of the Mitochondrial Rieske Iron Sulfur Protein in Mice Produces a Specific Skin Pigmentation Phenotype. <i>Journal of Investigative Dermatology</i> , 2008, 128, 2343-2345.	0.7	8
43	Mitochondrial biogenesis and turnover. <i>Cell Calcium</i> , 2008, 44, 24-35.	2.4	118
44	Cytochrome c oxidase deficiency in neurons decreases both oxidative stress and amyloid formation in a mouse model of Alzheimer's disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 14163-14168.	7.1	160
45	Role of Cytochrome c in Apoptosis: Increased Sensitivity to Tumor Necrosis Factor Alpha Is Associated with Respiratory Defects but Not with Lack of Cytochrome c Release. <i>Molecular and Cellular Biology</i> , 2007, 27, 1771-1783.	2.3	54
46	Cytochrome c Oxidase Is Required for the Assembly/Stability of Respiratory Complex I in Mouse Fibroblasts. <i>Molecular and Cellular Biology</i> , 2006, 26, 4872-4881.	2.3	213
47	Mice lacking COX10 in skeletal muscle recapitulate the phenotype of progressive mitochondrial myopathies associated with cytochrome c oxidase deficiency. <i>Human Molecular Genetics</i> , 2005, 14, 2737-2748.	2.9	145
48	Human mitochondrial DNA with large deletions repopulates organelles faster than full-length genomes under relaxed copy number control. <i>Nucleic Acids Research</i> , 2002, 30, 4626-4633.	14.5	139
49	An out-of-frame cytochrome b gene deletion from a patient with parkinsonism is associated with impaired complex III assembly and an increase in free radical production. <i>Annals of Neurology</i> , 2000, 48, 774-781.	5.3	126
50	An out-of-frame cytochrome b gene deletion from a patient with parkinsonism is associated with impaired complex III assembly and an increase in free radical production. <i>Annals of Neurology</i> , 2000, 48, 774-781.	5.3	2