Francesc Cardellach

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Assessment of mitochondrial toxicity in newborns and infants with congenital cytomegalovirus infection treated with valganciclovir. Archives of Disease in Childhood, 2022, 107, 686-691.	1.0	Ο
2	Neuronal induction and bioenergetics characterization of human forearm adipose stem cells from Parkinson's disease patients and healthy controls. PLoS ONE, 2022, 17, e0265256.	1.1	0
3	Comment on Yeste et al. Polyphenols and IUCR Pregnancies: Intrauterine Growth Restriction and Hydroxytyrosol Affect the Development and Neurotransmitter Profile of the Hippocampus in a Pig Model. Antioxidants 2021, 10, 1505. Antioxidants, 2022, 11, 833.	2.2	0
4	Mitochondrial changes associated with viral infectious diseases in the paediatric population. Reviews in Medical Virology, 2021, 31, e2232.	3.9	3
5	A Mitocentric View of the Main Bacterial and Parasitic Infectious Diseases in the Pediatric Population. International Journal of Molecular Sciences, 2021, 22, 3272.	1.8	3
6	The 3-Year Effect of the Mediterranean Diet Intervention on Inflammatory Biomarkers Related to Cardiovascular Disease. Biomedicines, 2021, 9, 862.	1.4	11
7	Nutrition, Bioenergetics, and Metabolic Syndrome. Nutrients, 2020, 12, 2785.	1.7	26
8	Disrupted Mitochondrial and Metabolic Plasticity Underlie Comorbidity between Age-Related and Degenerative Disorders as Parkinson Disease and Type 2 Diabetes Mellitus. Antioxidants, 2020, 9, 1063.	2.2	8
9	Mitochondrial Dysfunction: A Common Hallmark Underlying Comorbidity between sIBM and Other Degenerative and Age-Related Diseases. Journal of Clinical Medicine, 2020, 9, 1446.	1.0	4
10	Mitochondrial Toxicogenomics for Antiretroviral Management: HIV Post-exposure Prophylaxis in Uninfected Patients. Frontiers in Genetics, 2020, 11, 497.	1.1	13
11	Mitochondrial implications in human pregnancies with intrauterine growth restriction and associated cardiac remodelling. Journal of Cellular and Molecular Medicine, 2019, 23, 3962-3973.	1.6	19
12	Mitochondrial and autophagic alterations in skin fibroblasts from Parkinson disease patients with Parkin mutations. Aging, 2019, 11, 3750-3767.	1.4	25
13	GBA mutation promotes early mitochondrial dysfunction in 3D neurosphere models. Aging, 2019, 11, 10338-10355.	1.4	15
14	Transcriptional alterations in skin fibroblasts from Parkinson's disease patients with parkin mutations. Neurobiology of Aging, 2018, 65, 206-216.	1.5	13
15	Exhaustion of mitochondrial and autophagic reserve may contribute to the development of LRRK2 G2019S -Parkinson's disease. Journal of Translational Medicine, 2018, 16, 160.	1.8	22
16	Medicina interna y enfermedades raras. Transición niño-adulto. Arbor, 2018, 194, 460.	0.1	2
17	Mitochondrial toxicity and caspase activation in HIV pregnant women. Journal of Cellular and Molecular Medicine, 2017, 21, 26-34.	1.6	5
18	Mitochondrial DNA disturbances and deregulated expression of oxidative phosphorylation and mitochondrial fusion proteins in sporadic inclusion body myositis. Clinical Science, 2016, 130, 1741-1751.	1.8	33

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19	Clinical and genetic characterization of the autoinflammatory diseases diagnosed in an adult reference center. Autoimmunity Reviews, 2016, 15, 9-15.	2.5	62
20	BACE-1, PS-1 and sAPPβ Levels Are Increased in Plasma from Sporadic Inclusion Body Myositis Patients: Surrogate Biomarkers among Inflammatory Myopathies. Molecular Medicine, 2015, 21, 817-823.	1.9	12
21	Selective Elimination of Mitochondrial Mutations in the Germline by Genome Editing. Cell, 2015, 161, 459-469.	13.5	245
22	Partial Immunological and Mitochondrial Recovery after Reducing Didanosine doses in Patients on Didanosine and Tenofovir-Based Regimens. Antiviral Therapy, 2008, 13, 231-240.	0.6	5
23	Mitochondrial DNA Depletion in Oocytes of HIV-Infected Antiretroviral-Treated Infertile Women. Antiviral Therapy, 2008, 13, 833-838.	0.6	34
24	Metabolic and Mitochondrial Effects of Switching Antiretroviral-Experienced Patients to Enfuvirtide, Tenofovir and Saquinavir/Ritonavir. Antiviral Therapy, 2006, 11, 625-630.	0.6	12
25	Mitochondrial Studies in Haart-Related Lipodystrophy: From Experimental Hypothesis to Clinical Findings. Antiviral Therapy, 2005, 10, 73-81.	0.6	22
26	<i>In Vivo</i> Effects of Highly Active Antiretroviral Therapies Containing the Protease Inhibitor Nelfinavir on Mitochondrially Driven Apoptosis. Antiviral Therapy, 2005, 10, 945-951.	0.6	17
27	Mitochondrial Effects of Antiretroviral Therapies in Asymptomatic Patients. Antiviral Therapy, 2004, 9, 47-55.	0.6	65
28	Effect of Smoking Cessation on Mitochondrial Respiratory Chain Function. Journal of Toxicology: Clinical Toxicology, 2003, 41, 223-228.	1.5	32
29	Mitochondrial Dna Depletion and Respiratory Chain Enzyme Deficiencies are Present in Peripheral Blood Mononuclear Cells of HIV-Infected Patients with Haart-Related Lipodystrophy. Antiviral Therapy, 2003, 8, 333-338.	0.6	67
30	Small-vessel vasculitis surrounding a spared temporal artery: Clinical and pathologic findings in a series of twenty-eight patients. Arthritis and Rheumatism, 2001, 44, 1387-1395.	6.7	105
31	Heart Mitochondrial Respiratory Chain Complexes Are Functionally Unaffected in Heavy Ethanol Drinkers Without Cardiomyopathy. Alcoholism: Clinical and Experimental Research, 2000, 24, 859-864.	1.4	1
32	Mitochondrial Cytochrome c Oxidase Inhibition during Acute Carbon Monoxide Poisoning. Basic and Clinical Pharmacology and Toxicology, 1998, 82, 199-202.	0.0	85
33	Biochemical and molecular effects of chronic haloperidol administration on brain and muscle mitochondria of rats. Journal of Neuroscience Research, 1998, 53, 475-481.	1.3	30
34	Biochemical parameters for the diagnosis of mitochondrial respiratory chain deficiency in humans, and their lack of age-related changes. Biochemical Journal, 1998, 329, 249-254.	1.7	98
35	Inherited susceptibility to several cancers but absence of linkage between dysplastic nevus syndrome and CDKN2A in a melanoma family with a mutation in the CDKN2A (P16INK4A) gene. Human Genetics, 1997, 101, 359-364.	1.8	58
36	Sporadic heteroplasmic single 5.5 Kb mitochondrial DNA deletion associated with cerebellar ataxia, hypogonadotropic hypogonadism, choroidal dystrophy, and mitochondrial respiratory chain complex I deficiency. Human Mutation, 1997, 10, 212-216.	1.1	16