Alessia Indrieri

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3508658/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Linear Skin Defects with Multiple Congenital Anomalies (LSDMCA): An Unconventional Mitochondrial Disorder. Genes, 2021, 12, 263.	2.4	8
2	Integrated Genomics Identifies miR-181/TFAM Pathway as a Critical Driver of Drug Resistance in Melanoma. International Journal of Molecular Sciences, 2021, 22, 1801.	4.1	20
3	The Role of MicroRNAs in Mitochondria-Mediated Eye Diseases. Frontiers in Cell and Developmental Biology, 2021, 9, 653522.	3.7	9
4	Correction of oxidative stress enhances enzyme replacement therapy in Pompe disease. EMBO Molecular Medicine, 2021, 13, e14434.	6.9	13
5	Mutation-Independent Therapies for Retinal Diseases: Focus on Gene-Based Approaches. Frontiers in Neuroscience, 2020, 14, 588234.	2.8	9
6	74P Targeting mitochondria as a novel therapeutic strategy in biliary tract cancer. Annals of Oncology, 2020, 31, S269.	1.2	0
7	Dopamine, Alpha-Synuclein, and Mitochondrial Dysfunctions in Parkinsonian Eyes. Frontiers in Neuroscience, 2020, 14, 567129.	2.8	31
8	α-synuclein overexpression in the retina leads to vision impairment and degeneration of dopaminergic amacrine cells. Scientific Reports, 2020, 10, 9619.	3.3	27
9	The Pervasive Role of the miR-181 Family in Development, Neurodegeneration, and Cancer. International Journal of Molecular Sciences, 2020, 21, 2092.	4.1	93
10	The HOPS complex subunit VPS39 controls ciliogenesis through autophagy. Human Molecular Genetics, 2020, 29, 1018-1029.	2.9	16
11	COVID-19: High-JAKing of the Inflammatory "Flight―by Ruxolitinib to Avoid the Cytokine Storm. Frontiers in Oncology, 2020, 10, 599502.	2.8	9
12	miRâ€181a/b downregulation exerts a protective action on mitochondrial disease models. EMBO Molecular Medicine, 2019, 11, .	6.9	58
13	Activation of Autophagy, Observed in Liver Tissues From Patients With Wilson Disease and From ATP7B-Deficient Animals, Protects Hepatocytes From Copper-Induced Apoptosis. Gastroenterology, 2019, 156, 1173-1189.e5.	1.3	150
14	Synthetic long non-coding RNAs [SINEUPs] rescue defective gene expression in vivo. Scientific Reports, 2016, 6, 27315.	3.3	37
15	Microphthalmia With Linear Skin Lesions (MLS) Syndrome: An Unconventional Mitochondrial Disorder. , 2016, , 1449-1451.		1
16	Metabolic Regulation of the Ultradian Oscillator Hes1 by Reactive Oxygen Species. Journal of Molecular Biology, 2015, 427, 1887-1902.	4.2	11
17	The impairment of HCCS leads to MLS syndrome by activating a nonâ€canonical cell death pathway in the brain and eyes. EMBO Molecular Medicine, 2014, 6, 849-849.	6.9	0
18	The impairment of HCCS leads to MLS syndrome by activating a non anonical cell death pathway in the brain and eyes. EMBO Molecular Medicine, 2013, 5, 280-293.	6.9	33

#	Article	IF	CITATIONS
19	Mutations in COX7B Cause Microphthalmia with Linear Skin Lesions, an Unconventional Mitochondrial Disease. American Journal of Human Genetics, 2012, 91, 942-949.	6.2	104
20	Drug Repurposing to Target the Apoptosome in MAPKi-Resistant Melanoma. SSRN Electronic Journal, 0, , .	0.4	1