Alessia Indrieri

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3508658/publications.pdf

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20 papers	630 citations	840728 11 h-index	940516 16 g-index
22 all docs	22 docs citations	22 times ranked	1051 citing authors

#	Article	IF	Citations
1	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
2	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
3	The Pervasive Role of the miR-181 Family in Development, Neurodegeneration, and Cancer. International Journal of Molecular Sciences, 2020, 21, 2092.	4.1	93
4	miRâ€181a/b downregulation exerts a protective action on mitochondrial disease models. EMBO Molecular Medicine, 2019, 11, .	6.9	58
5	Synthetic long non-coding RNAs [SINEUPs] rescue defective gene expression in vivo. Scientific Reports, 2016, 6, 27315.	3.3	37
6	The impairment of HCCS leads to MLS syndrome by activating a nonâ€canonical cell death pathway in the brain and eyes. EMBO Molecular Medicine, 2013, 5, 280-293.	6.9	33
7	Dopamine, Alpha-Synuclein, and Mitochondrial Dysfunctions in Parkinsonian Eyes. Frontiers in Neuroscience, 2020, 14, 567129.	2.8	31
8	Î±-synuclein overexpression in the retina leads to vision impairment and degeneration of dopaminergic amacrine cells. Scientific Reports, 2020, 10, 9619.	3.3	27
9	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
10	The HOPS complex subunit VPS39 controls ciliogenesis through autophagy. Human Molecular Genetics, 2020, 29, 1018-1029.	2.9	16
11	Correction of oxidative stress enhances enzyme replacement therapy in Pompe disease. EMBO Molecular Medicine, 2021, 13, e14434.	6.9	13
12	Metabolic Regulation of the Ultradian Oscillator Hes1 by Reactive Oxygen Species. Journal of Molecular Biology, 2015, 427, 1887-1902.	4.2	11
13	Mutation-Independent Therapies for Retinal Diseases: Focus on Gene-Based Approaches. Frontiers in Neuroscience, 2020, 14, 588234.	2.8	9
14	COVID-19: High-JAKing of the Inflammatory â€œFlightâ€•by Ruxolitinib to Avoid the Cytokine Storm. Frontiers in Oncology, 2020, 10, 599502.	2.8	9
15	The Role of MicroRNAs in Mitochondria-Mediated Eye Diseases. Frontiers in Cell and Developmental Biology, 2021, 9, 653522.	3.7	9
16	Linear Skin Defects with Multiple Congenital Anomalies (LSDMCA): An Unconventional Mitochondrial Disorder. Genes, 2021, 12, 263.	2.4	8
17	Drug Repurposing to Target the Apoptosome in MAPKi-Resistant Melanoma. SSRN Electronic Journal, 0, , .	0.4	1
18	Microphthalmia With Linear Skin Lesions (MLS) Syndrome: An Unconventional Mitochondrial Disorder. , 2016, , 1449-1451.		1

#	Article	IF	Citations
19	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
20	74P Targeting mitochondria as a novel therapeutic strategy in biliary tract cancer. Annals of Oncology, 2020, 31, S269.	1.2	0