

Rosa Pasquariello

List of Publications by Year in descending order

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Version: 2024-02-01

14
papers

141
citations

1307594

7
h-index

1199594

12
g-index

14
all docs

14
docs citations

14
times ranked

371
citing authors

#	ARTICLE	IF	CITATIONS
1	Leigh-like neuroimaging features associated with new biallelic mutations in OPA1. <i>European Journal of Paediatric Neurology</i> , 2017, 21, 671-677.	1.6	25
2	Neuroanatomical correlates of childhood apraxia of speech: A connectomic approach. <i>NeuroImage: Clinical</i> , 2016, 12, 894-901.	2.7	18
3	Behavioral and neurobiological correlates of childhood apraxia of speech in Italian children. <i>Brain and Language</i> , 2015, 150, 177-185.	1.6	17
4	Structural brain damage and visual disorders in children with cerebral palsy due to periventricular leukomalacia. <i>NeuroImage: Clinical</i> , 2020, 28, 102430.	2.7	17
5	TMEM5-associated dystroglycanopathy presenting with CMD and mild limb-girdle muscle involvement. <i>Neuromuscular Disorders</i> , 2016, 26, 459-461.	0.6	15
6	Corticopontocerebellar Connectivity Disruption in Congenital Hemiplegia. <i>Neurorehabilitation and Neural Repair</i> , 2015, 29, 858-866.	2.9	13
7	Expanding the clinical and genetic heterogeneity of SPAX5. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 595-601.	3.7	11
8	Diffusion Tractography Biomarkers of Pediatric Cerebellar Hypoplasia/Atrophy: Preliminary Results Using Constrained Spherical Deconvolution. <i>American Journal of Neuroradiology</i> , 2016, 37, 917-923.	2.4	8
9	Teaching NeuroImages: Leigh-like features expand the picture of PMPCA-related disorders. <i>Neurology</i> , 2019, 92, e168-e169.	1.1	6
10	Longitudinal follow up of a boy affected by Pol III-related leukodystrophy: a detailed phenotype description. <i>BMC Medical Genetics</i> , 2015, 16, 53.	2.1	5
11	Relapsing-Remitting Course of Cystic Leukoencephalopathy. <i>Pediatric Neurology</i> , 2018, 89, 63-65.	2.1	4
12	Phenotypic Definition and Genotype-Phenotype Correlates in PMPCA-Related Disease. <i>Applied Sciences (Switzerland)</i> , 2021, 11, 748.	2.5	1
13	Reconsidering NMIHBA Core Features: Macrocephaly Is Not a So Unusual Sign in PRUNE1-Related Encephalopathy. <i>Journal of Pediatric Neurology</i> , 2021, 19, 116-123.	0.2	1
14	Correlating Neuroimaging and CNVs Data: 7 Years of Cytogenomic Microarray Analysis on Patients Affected by Neurodevelopmental Disorders. <i>Journal of Pediatric Genetics</i> , 2021, 10, 292-299.	0.7	0