

Valentina Melzi

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

181
citations

1307594

7
h-index

1474206

9
g-index

11
all docs

11
docs citations

11
times ranked

418
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular analysis of SMARD1 patient-derived cells demonstrates that nonsense-mediated mRNA decay is impaired. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2022, 93, 908-910.	1.9	3
2	Insights into the identification of a molecular signature for amyotrophic lateral sclerosis exploiting integrated microRNA profiling of iPSC-derived motor neurons and exosomes. <i>Cellular and Molecular Life Sciences</i> , 2022, 79, 189.	5.4	12
3	Stathmins and Motor Neuron Diseases: Pathophysiology and Therapeutic Targets. <i>Biomedicines</i> , 2022, 10, 711.	3.2	9
4	Generation of two hiPSC lines (UMILi027-A and UMILi028-A) from early and late-onset Congenital Central hypoventilation Syndrome (CCHS) patients carrying a polyalanine expansion mutation in the PHOX2B gene. <i>Stem Cell Research</i> , 2022, 61, 102781.	0.7	0
5	Expanding the genotypic and phenotypic spectrum of Beta-amyloid-associated neurodegeneration. <i>European Journal of Neurology</i> , 2021, 28, e25-e27.	3.3	0
6	Key role of SMN/SYNERGIC and RNA-Motif 7 in spinal muscular atrophy: RNA-Seq and motif analysis of human motor neurons. <i>Brain</i> , 2019, 142, 276-294.	7.6	31
7	A de novo C19orf12 heterozygous mutation in a patient with MPAN. <i>Parkinsonism and Related Disorders</i> , 2018, 48, 109-111.	2.2	15
8	MicroRNA expression analysis identifies a subset of downregulated miRNAs in ALS motor neuron progenitors. <i>Scientific Reports</i> , 2018, 8, 10105.	3.3	53
9	Genome-wide RNA-seq of iPSC-derived motor neurons indicates selective cytoskeletal perturbation in Brown-Vialetto disease that is partially rescued by riboflavin. <i>Scientific Reports</i> , 2017, 7, 46271.	3.3	22
10	Mutations in TMEM230 are rare in autosomal dominant Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2017, 39, 87-88.	2.2	11
11	Mutational analysis of COQ2 in patients with MSA in Italy. <i>Neurobiology of Aging</i> , 2016, 45, 213.e1-213.e2.	3.1	25