

Giulia Franco

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7458672/publications.pdf>

Version: 2024-02-01

17
papers

146
citations

1307594

7
h-index

1281871

11
g-index

17
all docs

17
docs citations

17
times ranked

318
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutational analysis of COQ2 in patients with MSA in Italy. <i>Neurobiology of Aging</i> , 2016, 45, 213.e1-213.e2.	3.1	25
2	Disruption of Mitochondrial Homeostasis: The Role of PINK1 in Parkinson's Disease. <i>Cells</i> , 2021, 10, 3022.	4.1	21
3	Progressive Encephalomyelitis with Rigidity and Myoclonus Associated With Anti-GlyR Antibodies and Hodgkin's Lymphoma: A Case Report. <i>Frontiers in Neurology</i> , 2017, 8, 401.	2.4	17
4	Selective inversion recovery quantitative magnetization transfer imaging: Toward a 3 T clinical application in multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2020, 26, 457-467.	3.0	16
5	A de novo C19orf12 heterozygous mutation in a patient with MPAN. <i>Parkinsonism and Related Disorders</i> , 2018, 48, 109-111.	2.2	15
6	Mutations in TMEM230 are rare in autosomal dominant Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2017, 39, 87-88.	2.2	11
7	Unravelling Genetic Factors Underlying Corticobasal Syndrome: A Systematic Review. <i>Cells</i> , 2021, 10, 171.	4.1	8
8	Dysautonomia in Parkinson's Disease: Impact of Glucocerebrosidase Gene Mutations on Cardiovascular Autonomic Control. <i>Frontiers in Neuroscience</i> , 2022, 16, 842498.	2.8	6
9	Validation of the Italian version of the PSP Quality of Life questionnaire. <i>Neurological Sciences</i> , 2019, 40, 2587-2594.	1.9	5
10	Validation of the Italian version of carers' quality-of-life questionnaire for parkinsonism (PQoL). <i>Journal of Parkinsonism and Related Disorders</i> , 2019, 10, 101-105.	1.9	5
11	Screening of LRP10 mutations in Parkinson's disease patients from Italy. <i>Parkinsonism and Related Disorders</i> , 2021, 89, 17-21.	2.2	5
12	Parkinsonism in diseases predominantly presenting with dystonia. <i>International Review of Neurobiology</i> , 2019, 149, 307-326.	2.0	4
13	A case report of late-onset cerebellar ataxia associated with a rare p.R342W TGM6 (SCA35) mutation. <i>BMC Neurology</i> , 2020, 20, 408.	1.8	3
14	Microscopic Polyangiitis With Selective Involvement of Central and Peripheral Nervous System: A Case Report. <i>Frontiers in Neurology</i> , 2020, 11, 269.	2.4	3
15	Parkinsonism and ataxia. <i>Journal of the Neurological Sciences</i> , 2021, , 120020.	0.6	2
16	Late-onset leukoencephalopathy in a patient with recessive EARS2 mutations. <i>Neurology: Genetics</i> , 2020, 6, e488.	1.9	0
17	Expanding the genotypic and phenotypic spectrum of Beta-protein-associated neurodegeneration. <i>European Journal of Neurology</i> , 2021, 28, e25-e27.	3.3	0