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	Dysautonomia in Parkinsonâ€™s Disease: Impact of Glucocerebrosidase Gene Mutations on Cardiovascular Autonomic Control. Frontiers in Neuroscience, 2022, 16, 842498.	2.8	6
2	Expanding the genotypic and phenotypic spectrum of Betaâ€propeller poteinâ€associated neurodegeneration. European Journal of Neurology, 2021, 28, e25-e27.	3.3	0
3	Unravelling Genetic Factors Underlying Corticobasal Syndrome: A Systematic Review. Cells, 2021, 10, 171.	4.1	8
4	Screening of LRP10 mutations in Parkinson's disease patients from Italy. Parkinsonism and Related Disorders, 2021, 89, 17-21.	2.2	5
5	Parkinsonism and ataxia. Journal of the Neurological Sciences, 2021, , 120020.	0.6	2
6	Disruption of Mitochondrial Homeostasis: The Role of PINK1 in Parkinsonâ€™s Disease. Cells, 2021, 10, 3022.	4.1	21
7	Selective inversion recovery quantitative magnetization transfer imaging: Toward a 3 T clinical application in multiple sclerosis. Multiple Sclerosis Journal, 2020, 26, 457-467.	3.0	16
8	Late-onset leukoencephalopathy in a patient with recessive EARS2 mutations. Neurology: Genetics, 2020, 6, e488.	1.9	0
9	A case report of late-onset cerebellar ataxia associated with a rare p.R342W TGM6 (SCA35) mutation. BMC Neurology, 2020, 20, 408.	1.8	3
10	Microscopic Polyangiitis With Selective Involvement of Central and Peripheral Nervous System: A Case Report. Frontiers in Neurology, 2020, 11, 269.	2.4	3
11	Validation of the Italian version of the PSP Quality of Life questionnaire. Neurological Sciences, 2019, 40, 2587-2594.	1.9	5
12	Validation of the Italian version of carersâ€™ quality-of-life questionnaire for parkinsonism (PQoL) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50	1.9	5
13	Parkinsonism in diseases predominantly presenting with dystonia. International Review of Neurobiology, 2019, 149, 307-326.	2.0	4
14	A de novo C19orf12 heterozygous mutation in a patient with MPAN. Parkinsonism and Related Disorders, 2018, 48, 109-111.	2.2	15
15	Mutations in TMEM230 are rare in autosomal dominant Parkinson's disease. Parkinsonism and Related Disorders, 2017, 39, 87-88.	2.2	11
16	Progressive Encephalomyelitis with Rigidity and Myoclonus Associated With Anti-GlyR Antibodies and Hodgkinâ€™s Lymphoma: A Case Report. Frontiers in Neurology, 2017, 8, 401.	2.4	17
17	Mutational analysis of COQ2 in patients with MSA in Italy. Neurobiology of Aging, 2016, 45, 213.e1-213.e2.	3.1	25