Giuseppe De Michele

List of Publications by Year
in descending order

27	2,625	516710 16	526287 27
papers	citations	h-index	g-index

27	27	27	3223
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Optical coherence tomography angiography findings in Huntingtonâ€™s disease. Neurological Sciences, 2021, 42, 995-1001.	1.9	10
2	Spinocerebellar ataxia type 48: last but not least. Neurological Sciences, 2021, 42, 1577-1577.	1.9	1
3	NGS in Hereditary Ataxia: When Rare Becomes Frequent. International Journal of Molecular Sciences, 2021, 22, 8490.	4.1	12
4	STUB1 â€Related Ataxias: A Challenging Diagnosis. Movement Disorders Clinical Practice, 2020, 7, 733-734.	1.5	4
5	Spinocerebellar ataxia type 48: last but not least. Neurological Sciences, 2020, 41, 2423-2432.	1.9	31
6	PERK-Mediated Unfolded Protein Response Activation and Oxidative Stress in PARK20 Fibroblasts. Frontiers in Neuroscience, 2019, 13, 673.	2.8	38
7	Spinocerebellar ataxia 48 presenting with ataxia associated with cognitive, psychiatric, and extrapyramidal features: A report of two Italian families. Parkinsonism and Related Disorders, 2019, 65, 91-96.	2.2	43
8	LRP10 variants in Parkinson's disease and dementia with Lewy bodies in the South-West of the Netherlands. Parkinsonism and Related Disorders, 2019, 65, 243-247.	2.2	14
9	Longitudinal study of a cohort of MSA-C patients in South Italy: survival and clinical features. Neurological Sciences, 2019, 40, 2105-2109.	1.9	11
10	Alteration of endosomal trafficking is associated with early-onset parkinsonism caused by SYNJ1 mutations. Cell Death and Disease, 2018, 9, 385.	6.3	48
11	Peculiar combinations of individually non-pathogenic missense mitochondrial DNA variants cause low penetrance Leberâ€™s hereditary optic neuropathy. PLoS Genetics, 2018, 14, e1007210.	3.5	47
12	Myocardial 123I-metaiodobenzylguanidine scintigraphy in patients with homozygous and heterozygous parkin mutations. Journal of Nuclear Cardiology, 2017, 24, 103-107.	2.1	10
13	A new model to study neurodegeneration in ataxia oculomotor apraxia type 2. Human Molecular Genetics, 2015, 24, 5759-5774.	2.9	34
14	PARK20 caused by SYNJ1 homozygous Arg258Gln mutation in a new Italian family. Neurogenetics, 2014, 15, 183-188.	1.4	107
15	Mutation of senataxin alters disease-specific transcriptional networks in patients with ataxia with oculomotor apraxia type 2. Human Molecular Genetics, 2014, 23, 4758-4769.	2.9	43
16	Is serum uric acid related to non-motor symptoms in de-novo Parkinson's disease patients?. Parkinsonism and Related Disorders, 2014, 20, 772-775.	2.2	32
17	Other autosomal recessive and childhood ataxias. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2012, 103, 343-357.	1.8	10
18	Neurophysiological evaluation of motor corticospinal pathways by TMS in idiopathic early-onset Parkinsonâ€™s disease. Clinical Neurophysiology, 2011, 122, 546-549.	1.5	17

#	Article	IF	Citations
19	Benign hereditary chorea: Clinical and neuroimaging features in an Italian family. Movement Disorders, 2010, 25, 1491-1495.	3.9	32
20	Genetic screening for LRRK2 gene G2019S mutation in Parkinson's disease patients from Southern Italy. Parkinsonism and Related Disorders, 2009, 15, 242-244.	2.2	16
21	A novel mutation in GCH-1 gene in a case of dopa-responsive dystonia. Journal of Neurology, 2007, 254, 1133-1134.	3.6	2
22	Neurophysiological evidence of corticospinal tract abnormality in patients with Parkin mutations. Journal of Neurology, 2006, 253, 275-279.	3.6	29
23	Suppression of myoclonus in SCA2 by piracetam. Movement Disorders, 2006, 21, 116-118.	3.9	14
24	Characterization of nigrostriatal dysfunction in spinocerebellar ataxia 17. Movement Disorders, 2006, 21, 872-875.	3.9	22
25	How much phenotypic variation can be attributed toparkingenotype?. Annals of Neurology, 2003, 54, 176-185.	5.3	271
26	Parkin mutations are frequent in patients with isolated earlyâ€onset parkinsonism. Brain, 2003, 126, 1271-1278.	7.6	279
27	Association between Early-Onset Parkinson's Disease and Mutations in theParkinGene. New England Journal of Medicine, 2000, 342, 1560-1567.	27.0	1,448