Leonidas Stefanis

List of Publications by Year in descending order

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66343 36028 10,190 164 42 97 citations h-index g-index papers 169 169 169 13010 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Impaired Degradation of Mutant α-Synuclein by Chaperone-Mediated Autophagy. Science, 2004, 305, 1292-1295.	12.6	1,762
2	Â-Synuclein in Parkinson's Disease. Cold Spring Harbor Perspectives in Medicine, 2012, 2, a009399-a009399.	6.2	958
3	Cell-Produced α-Synuclein Is Secreted in a Calcium-Dependent Manner by Exosomes and Impacts Neuronal Survival. Journal of Neuroscience, 2010, 30, 6838-6851.	3.6	913
4	Wild Type α-Synuclein Is Degraded by Chaperone-mediated Autophagy and Macroautophagy in Neuronal Cells. Journal of Biological Chemistry, 2008, 283, 23542-23556.	3.4	553
5	Expression of A53T Mutant But Not Wild-Type α-Synuclein in PC12 Cells Induces Alterations of the Ubiquitin-Dependent Degradation System, Loss of Dopamine Release, and Autophagic Cell Death. Journal of Neuroscience, 2001, 21, 9549-9560.	3.6	540
6	Pathological roles of α-synuclein in neurological disorders. Lancet Neurology, The, 2011, 10, 1015-1025.	10.2	328
7	Abberant α-Synuclein Confers Toxicity to Neurons in Part through Inhibition of Chaperone-Mediated Autophagy. PLoS ONE, 2009, 4, e5515.	2.5	304
8	Proteasomal inhibition leads to formation of ubiquitin/αâ€synucleinâ€immunoreactive inclusions in PC12 cells. Journal of Neurochemistry, 2001, 78, 899-908.	3.9	253
9	Alpha-synuclein and Protein Degradation Systems: a Reciprocal Relationship. Molecular Neurobiology, 2013, 47, 537-551.	4.0	222
10	Autophagy and <scp>A</scp> lphaâ€ <scp>S</scp> ynuclein: <scp>R</scp> elevance to <scp>P</scp> arkinson's <scp>D</scp> isease and <scp>R</scp> elated <scp>S</scp> ynucleopathies. Movement Disorders, 2016, 31, 178-192.	3.9	216
11	Cell-produced α-synuclein oligomers are targeted to, and impair, the 26S proteasome. Neurobiology of Aging, 2010, 31, 953-968.	3.1	185
12	Boosting chaperone-mediated autophagy in vivo mitigates \hat{l}_{\pm} -synuclein-induced neurodegeneration. Brain, 2013, 136, 2130-2146.	7.6	175
13	Motor and Nonmotor Features of Carriers of the p.A53T Alphaâ€Synuclein Mutation: A Longitudinal Study. Movement Disorders, 2016, 31, 1226-1230.	3.9	134
14	Mediterranean diet adherence is related to reduced probability of prodromal Parkinson's disease. Movement Disorders, 2019, 34, 48-57.	3.9	134
15	Defective synaptic connectivity and axonal neuropathology in a human iPSC-based model of familial Parkinson's disease. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E3679-E3688.	7.1	122
16	Proteasomal Inhibition-Induced Inclusion Formation and Death in Cortical Neurons Require Transcription and Ubiquitination. Molecular and Cellular Neurosciences, 2002, 21, 223-238.	2.2	118
17	How is alphaâ€synuclein cleared from the cell?. Journal of Neurochemistry, 2019, 150, 577-590.	3.9	113
18	The protective role of AMP-activated protein kinase in alpha-synuclein neurotoxicity in vitro. Neurobiology of Disease, 2014, 63, 1-11.	4.4	97

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19	Inducible overâ€expression of wild type αâ€synuclein in human neuronal cells leads to caspaseâ€dependent nonâ€apoptotic death. Journal of Neurochemistry, 2009, 109, 1348-1362.	3.9	96
20	Neurobiology of α-Synuclein. Molecular Neurobiology, 2004, 30, 001-022.	4.0	95
21	A multi-centre clinico-genetic analysis of the VPS35 gene in Parkinson disease indicates reduced penetrance for disease-associated variants. Journal of Medical Genetics, 2012, 49, 721-726.	3.2	94
22	Impairment of chaperone-mediated autophagy induces dopaminergic neurodegeneration in rats. Autophagy, 2016, 12, 2230-2247.	9.1	87
23	Autophagic pathways in Parkinson disease and related disorders. Expert Reviews in Molecular Medicine, 2011, 13, e8.	3.9	84
24	Synuclein-1 is selectively up-regulated in response to nerve growth factor treatment in PC12 cells. Journal of Neurochemistry, 2001, 76, 1165-1176.	3.9	80
25	Genetics of Parkinson's Disease. International Review of Neurobiology, 2017, 132, 197-231.	2.0	76
26	Modulation of β-glucocerebrosidase increases α-synuclein secretion and exosome release in mouse models of Parkinson's disease. Human Molecular Genetics, 2018, 27, 1696-1710.	2.9	75
27	Alpha-synuclein research: defining strategic moves in the battle against Parkinson's disease. Npj Parkinson's Disease, 2021, 7, 65.	5.3	74
28	Serotonergic pathology and disease burden in the premotor and motor phase of A53T \hat{l}_{\pm} -synuclein parkinsonism: a cross-sectional study. Lancet Neurology, The, 2019, 18, 748-759.	10.2	70
29	ADCY5-related movement disorders: Frequency, disease course and phenotypic variability in a cohort of paediatric patients. Parkinsonism and Related Disorders, 2017, 41, 37-43.	2.2	67
30	Long Non-coding RNAs Associated With Neurodegeneration-Linked Genes Are Reduced in Parkinson's Disease Patients. Frontiers in Cellular Neuroscience, 2019, 13, 58.	3.7	63
31	Plasma alpha-synuclein levels in patients with Parkinson's disease: a systematic review and meta-analysis. Neurological Sciences, 2019, 40, 929-938.	1.9	59
32	Targeting intracellular and extracellular alpha-synuclein as a therapeutic strategy in Parkinson's disease and other synucleinopathies. Expert Opinion on Therapeutic Targets, 2012, 16, 421-432.	3.4	58
33	Evidence of an association between the scavenger receptor class B member 2 gene and Parkinson's disease. Movement Disorders, 2012, 27, 400-405.	3.9	56
34	Chaperone mediated autophagy in aging: Starve to prosper. Ageing Research Reviews, 2016, 32, 13-21.	10.9	55
35	A Novel SNCA A30G Mutation Causes Familial Parkinson $\hat{\mathbb{E}}^{1}\!\!/_{4}$ s Disease. Movement Disorders, 2021, 36, 1624-1633.	3.9	54
36	Lysosomal alterations in peripheral blood mononuclear cells of Parkinson's disease patients. Movement Disorders, 2015, 30, 1830-1834.	3.9	53

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37	Chaperone mediated autophagy to the rescue: A new-fangled target for the treatment of neurodegenerative diseases. Molecular and Cellular Neurosciences, 2015, 66, 29-36.	2.2	53
38	A novel pathway for transcriptional regulation of αâ€synuclein. FASEB Journal, 2007, 21, 596-607.	0.5	50
39	Progressive striatonigral degenerationÂin a transgenic mouse model of multiple system atrophy: translational implications for interventional therapies. Acta Neuropathologica Communications, 2018, 6, 2.	5.2	50
40	Functional dissection of the αâ€synuclein promoter: transcriptional regulation by ZSCAN21 and ZNF219. Journal of Neurochemistry, 2009, 110, 1479-1490.	3.9	49
41	\hat{l}^2 -Glucocerebrosidase gene mutations in two cohorts of Greek patients with sporadic Parkinson's disease. Molecular Genetics and Metabolism, 2011, 104, 149-152.	1.1	47
42	Loss of \hat{l}^2 -Glucocerebrosidase Activity Does Not Affect Alpha-Synuclein Levels or Lysosomal Function in Neuronal Cells. PLoS ONE, 2013, 8, e60674.	2.5	47
43	Endogenous oligodendroglial alpha-synuclein and TPPP/p25α orchestrate alpha-synuclein pathology in experimental multiple system atrophy models. Acta Neuropathologica, 2019, 138, 415-441.	7.7	45
44	Lack of p53 delays apoptosis, but increases ubiquitinated inclusions, in proteasomal inhibitor-treated cultured cortical neurons. Molecular and Cellular Neurosciences, 2003, 24, 430-441.	2.2	43
45	Circulating Brainâ€Enriched MicroRNAs for Detection and Discrimination of Idiopathic and Genetic Parkinson's Disease. Movement Disorders, 2020, 35, 457-467.	3.9	43
46	Regulation of \hat{l}_{\pm} -synuclein by bFGF in cultured ventral midbrain dopaminergic neurons. Journal of Neurochemistry, 2003, 84, 803-813.	3.9	39
47	Differentially Expressed Circular <scp>RNAs</scp> in Peripheral Blood Mononuclear Cells of Patients with Parkinson's Disease. Movement Disorders, 2021, 36, 1170-1179.	3.9	38
48	Validation of differentially expressed brainâ€enriched microRNAs in the plasma of PD patients. Annals of Clinical and Translational Neurology, 2020, 7, 1594-1607.	3.7	36
49	Alpha-synuclein dimerization in erythrocytes of patients with genetic and non-genetic forms of Parkinson's Disease. Neuroscience Letters, 2018, 672, 145-149.	2.1	35
50	Selective neuroprotective effects of the S18Y polymorphic variant of UCH-L1 in the dopaminergic system. Human Molecular Genetics, 2012, 21, 874-889.	2.9	34
51	Salivary alpha-synuclein as a biomarker for Parkinson's disease: a systematic review. Journal of Neural Transmission, 2019, 126, 1373-1382.	2.8	34
52	Higher probability of prodromal Parkinson disease is related to lower cognitive performance. Neurology, 2019, 92, e2261-e2272.	1.1	34
53	Reduced serum immunoglobulin G concentrations in multiple sclerosis: prevalence and association with disease-modifying therapy and disease course. Therapeutic Advances in Neurological Disorders, 2019, 12, 175628641987834.	3.5	31
54	Selective cognitive impairment and hyposmia in p.A53T <i>SNCA</i> PD vs typical PD. Neurology, 2018, 90, e864-e869.	1.1	28

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55	Frontotemporal dementia as the presenting phenotype of p.A53T mutation carriers in the alpha-synuclein gene. Parkinsonism and Related Disorders, 2017, 35, 82-87.	2.2	27
56	Organochlorine pesticide levels in Greek patients with Parkinson's disease. Toxicology Reports, 2020, 7, 596-601.	3.3	27
57	In silico modeling of the effects of alpha-synuclein oligomerization on dopaminergic neuronal homeostasis. BMC Systems Biology, 2014, 8, 54.	3.0	26
58	Corticobasal degeneration and corticobasal syndrome: A review. Clinical Parkinsonism & Related Disorders, 2019, 1, 66-71.	0.9	26
59	Increased dimerization of alpha-synuclein in erythrocytes in Gaucher disease and aging. Neuroscience Letters, 2012, 528, 205-209.	2.1	24
60	Isolated delusional syndrome in Parkinson's Disease. Parkinsonism and Related Disorders, 2010, 16, 550-552.	2.2	23
61	Distinct alphaâ€Synuclein species induced by seeding are selectively cleared by the Lysosome or the Proteasome in neuronally differentiated SHâ€SY5Y cells. Journal of Neurochemistry, 2021, 156, 880-896.	3.9	22
62	The relationship between environmental factors and different Parkinson's disease subtypes in Greece: Data analysis of the Hellenic Biobank of Parkinson's disease. Parkinsonism and Related Disorders, 2019, 67, 105-112.	2.2	21
63	Autophagy dysfunction in peripheral blood mononuclear cells of Parkinson's disease patients. Neuroscience Letters, 2019, 704, 112-115.	2.1	21
64	Mendelian Randomisation Study of Smoking, Alcohol, and Coffee Drinking in Relation to Parkinson's Disease. Journal of Parkinson's Disease, 2022, 12, 267-282.	2.8	21
65	Neuropsychiatric symptoms and α-Synuclein profile of patients with Parkinson's disease dementia, dementia with Lewy bodies and Alzheimer's disease. Journal of Neurology, 2018, 265, 2295-2301.	3.6	20
66	Ageâ€dependent variation of female preponderance across different phenotypes of multiple sclerosis: A retrospective crossâ€sectional study. CNS Neuroscience and Therapeutics, 2019, 25, 527-531.	3.9	19
67	Medical cannabis as an alternative therapeutics for Parkinsons' disease: Systematic review. Complementary Therapies in Clinical Practice, 2020, 39, 101154.	1.7	19
68	Complex Effects of the ZSCAN21 Transcription Factor on Transcriptional Regulation of α-Synuclein in Primary Neuronal Cultures and in Vivo. Journal of Biological Chemistry, 2016, 291, 8756-8772.	3.4	18
69	Clinical rating scale for pantothenate kinaseâ€associated neurodegeneration: A pilot study. Movement Disorders, 2017, 32, 1620-1630.	3.9	18
70	Phenotypic Characteristics in GBA-Associated Parkinson's Disease: A Study in a Greek Population. Journal of Parkinson's Disease, 2018, 8, 101-105.	2.8	18
71	Midbrain morphology in idiopathic normal pressure hydrocephalus: A progressive supranuclear palsy mimic. Acta Neurologica Scandinavica, 2020, 141, 328-334.	2.1	18
72	Resistance of naturally secreted αâ€synuclein to proteolysis. FASEB Journal, 2014, 28, 3146-3158.	0.5	18

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73	Motor function and the probability of prodromal Parkinson's disease in older adults. Movement Disorders, 2019, 34, 1345-1353.	3.9	16
74	Frontotemporal dementia spectrum: first genetic screen in a Greek cohort. Neurobiology of Aging, 2019, 75, 224.e1-224.e8.	3.1	16
75	Frailty and Prodromal Parkinson's Disease: Results From the HELIAD Study. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2021, 76, 622-629.	3.6	16
76	Activation of FADD-Dependent Neuronal Death Pathways as a Predictor of Pathogenicity for LRRK2 Mutations. PLoS ONE, 2016, 11, e0166053.	2.5	16
77	A Prospective Validation of the Updated Movement Disorders Society Research Criteria for Prodromal Parkinson's Disease. Movement Disorders, 2020, 35, 1802-1809.	3.9	15
78	Recurrent Fulminant Tumefactive Demyelination With Marburg-Like Features and Atypical Presentation: Therapeutic Dilemmas and Review of Literature. Frontiers in Neurology, 2020, 11, 536.	2.4	15
79	Endogenous Levels of Alpha-Synuclein Modulate Seeding and Aggregation in Cultured Cells. Molecular Neurobiology, 2022, 59, 1273-1284.	4.0	15
80	Dairy Intake and Parkinson's Disease: A Mendelian Randomization Study. Movement Disorders, 2022, 37, 857-864.	3.9	15
81	Regulation of αâ€synuclein expression in cultured cortical neurons. Journal of Neurochemistry, 2011, 117, 275-285.	3.9	14
82	Cerebrospinal Fluid α-Synuclein Species in Cognitive and Movements Disorders. Brain Sciences, 2021, 11, 119.	2.3	14
83	\hat{l}_{\pm} -Synuclein dimerization in erythrocytes of Gaucher disease patients: correlation with lipid abnormalities and oxidative stress. Neuroscience Letters, 2016, 613, 1-5.	2.1	13
84	Identification of a prospective early motor progression cluster of Parkinson's disease: Data from the PPMI study. Journal of the Neurological Sciences, 2018, 387, 103-108.	0.6	13
85	Levodopa-induced skin disorders in patients with Parkinson disease: a systematic literature review approach. Acta Neurologica Belgica, 2019, 119, 325-336.	1.1	13
86	High discriminatory ability of peripheral and CFSF biomarkers in Lewy body diseases. Journal of Neural Transmission, 2020, 127, 311-322.	2.8	13
87	Cerebrospinal fluid biomarker profiling in corticobasal degeneration: Application of the AT(N) and other classification systems. Parkinsonism and Related Disorders, 2021, 82, 44-49.	2.2	13
88	Decreased levels of alphaâ€synuclein in cerebrospinal fluid of patients with clinically isolated syndrome and multiple sclerosis. Journal of Neurochemistry, 2015, 134, 748-755.	3.9	12
89	Peripheral alpha-synuclein levels in patients with genetic and non-genetic forms of Parkinson's disease. Parkinsonism and Related Disorders, 2020, 73, 35-40.	2.2	12
90	Serum Uric Acid Level as a Biomarker in Idiopathic and Genetic (p.A53T Alpha-Synuclein Carriers) Parkinson's Disease: Data from the PPMI Study. Journal of Parkinson's Disease, 2020, 10, 481-487.	2.8	12

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91	Nurr1 repression mediates cardinal features of Parkinson's disease in α-synuclein transgenic mice. Human Molecular Genetics, 2021, 30, 1469-1483.	2.9	12
92	Distinct profiles of LRRK2 activation and Rab GTPase phosphorylation in clinical samples from different PD cohorts. Npj Parkinson's Disease, 2022, 8, .	5.3	12
93	Severe dysphagia as the presenting symptom of Wernicke-Korsakoff syndrome in a non-alcoholic man. Neurological Sciences, 2008, 29, 45-46.	1.9	11
94	Impairment of chaperone-mediated autophagy affects neuronal homeostasis through altered expression of DJ-1 and CRMP-2 proteins. Molecular and Cellular Neurosciences, 2019, 95, 1-12.	2.2	11
95	Psychosisâ€Like Behavior and Hyperdopaminergic Dysregulation in Human α â€Synuclein <scp>BAC</scp> Transgenic Rats. Movement Disorders, 2021, 36, 716-728.	3.9	11
96	Biallelic <scp><i>RFC1</i></scp> pentanucleotide repeat expansions in Greek patients with lateâ€onset ataxia. Clinical Genetics, 2021, 100, 90-94.	2.0	11
97	Can We Treat Neurodegenerative Proteinopathies by Enhancing Protein Degradation?. Movement Disorders, 2022, 37, 1346-1359.	3.9	11
98	The different faces of the p. A53T alpha-synuclein mutation: A screening of Greek patients with parkinsonism and/or dementia. Neuroscience Letters, 2018, 672, 136-139.	2.1	10
99	MOG antibody-associated demyelinating disease mimicking typical multiple sclerosis: A case for expanding anti-MOG testing?. Multiple Sclerosis and Related Disorders, 2019, 33, 67-69.	2.0	10
100	Autophagy mediates the clearance of oligodendroglial SNCA/alpha-synuclein and TPPP/p25A in multiple system atrophy models. Autophagy, 2022, 18, 2104-2133.	9.1	10
101	Duration of paroxysmal atrial fibrillation in cryptogenic stroke is not associated with stroke severity and early outcomes. Journal of the Neurological Sciences, 2017, 376, 191-195.	0.6	9
102	123Iâ€FPâ€CIT SPECT [(123) Iâ€2βâ€carbomethoxyâ€3βâ€(4â€iodophenyl)â€Nâ€(3â€fluoropropyl) nortropane emission computed tomography] Imaging in a p.A53T αâ€synuclein Parkinson's disease cohort versus Parkinson's disease. Movement Disorders, 2018, 33, 1734-1739.	single pho	oton 9
103	Clinical, neuropsychological and imaging characteristics of Alzheimer's disease patients presenting as corticobasal syndrome. Journal of the Neurological Sciences, 2019, 398, 142-147.	0.6	9
104	Clinico-radiologic features and therapeutic strategies in tumefactive demyelination: a retrospective analysis of 50 consecutive cases. Therapeutic Advances in Neurological Disorders, 2021, 14, 175628642110065.	3.5	9
105	Expanding the Spectrum of <scp><i>AP5Z1â€</i></scp> Related Hereditary Spastic Paraplegia (<scp>HSPâ€6PG48</scp>): A Multicenter Study on a Rare Disease. Movement Disorders, 2021, 36, 1034-1038.	3.9	9
106	Serum uric acid level as a putative biomarker in Parkinson's disease patients carrying GBA1 mutations: 2-Year data from the PPMI study. Parkinsonism and Related Disorders, 2021, 84, 1-4.	2.2	9
107	Immunopathology of Tumefactive Demyelinating Lesions-From Idiopathic to Drug-Related Cases. Frontiers in Neurology, 2022, 13, 868525.	2.4	9
108	Potential Utility of Neurosonology in Paroxysmal Atrial Fibrillation Detection in Patients with Cryptogenic Stroke. Journal of Clinical Medicine, 2019, 8, 2002.	2.4	8

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109	Recurrent myelitis and asymptomatic hypophysitis in IgG4-related disease: case-based review. Rheumatology International, 2020, 40, 337-343.	3.0	8
110	HLA-DPB1*03 as Risk Allele and HLA-DPB1*04 as Protective Allele for Both Early- and Adult-Onset Multiple Sclerosis in a Hellenic Cohort. Brain Sciences, 2020, 10, 374.	2.3	8
111	Apathy: An underestimated feature in GBA and LRRK2 non-manifesting mutation carriers. Parkinsonism and Related Disorders, 2021, 91, 1-8.	2.2	8
112	\hat{l}_{\pm} -Synuclein Induces the GSK-3-Mediated Phosphorylation and Degradation of NURR1 and Loss of Dopaminergic Hallmarks. Molecular Neurobiology, 2021, 58, 6697-6711.	4.0	8
113	Parkinsonism and dementia. Journal of the Neurological Sciences, 2022, 433, 120015.	0.6	8
114	High content screening and proteomic analysis identify a kinase inhibitor that rescues pathological phenotypes in a patient-derived model of Parkinson's disease. Npj Parkinson's Disease, 2022, 8, 15.	5. 3	8
115	Evaluation of the interaction between LRRK2 and PARK16 loci in determining risk of Parkinson's disease: analysis of a large multicenter study. Neurobiology of Aging, 2017, 49, 217.e1-217.e4.	3.1	7
116	Three new case reports of Arteriovenous malformation-related Amyotrophic Lateral Sclerosis. Journal of the Neurological Sciences, 2018, 393, 58-62.	0.6	7
117	Autoimmune hemolytic anemia, demyelinating relapse, and AQP1 antibodies after alemtuzumab infusion. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, e711.	6.0	7
118	Elevated Serum α-Synuclein Levels in Huntington's Disease Patients. Neuroscience, 2020, 431, 34-39.	2.3	7
119	Driving and Alzheimer's dementia or mild cognitive impairment: a systematic review of the existing guidelines emphasizing on the neurologist's role. Neurological Sciences, 2021, 42, 4953-4963.	1.9	7
120	HLA-DRB1 differences in allelic distribution between familial and sporadic multiple sclerosis in a Hellenic cohort. Postgraduate Medicine, 2019, 131, 490-495.	2.0	6
121	HLA-DRB1 allele impact on pediatric multiple sclerosis in a Hellenic cohort. Multiple Sclerosis Journal - Experimental, Translational and Clinical, 2020, 6, 205521732090804.	1.0	6
122	Paraneoplastic basal ganglia encephalitis associated with anti-CV2/CRMP-5 and anti-Yo antibodies in a patient with non-small-cell lung cancer. Neurological Sciences, 2020, 41, 2649-2651.	1.9	6
123	REM sleep behavior disorder and other sleep abnormalities in p. A53T SNCA mutation carriers. Sleep, 2021, 44, .	1.1	6
124	Childhoodâ€Onset Chorea Caused by a Recurrent De Novo <i>DRD2</i> Variant. Movement Disorders, 2021, 36, 1472-1473.	3.9	6
125	Fingolimod as a first- or second-line treatment in a mini-series of young Hellenic patients with adolescent-onset multiple sclerosis: focus on immunological data. Neurological Sciences, 2022, 43, 2641-2649.	1.9	6
126	Dietary Inflammatory Index score and prodromal Parkinson's disease incidence: The HELIAD study. Journal of Nutritional Biochemistry, 2022, 105, 108994.	4.2	6

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127	Magnetic resonance imaging reveals Creutzfeldt–Jakob disease in a patient with apparent dementia with Lewy bodies. Journal of the Neurological Sciences, 2014, 340, 130-132.	0.6	5
128	A homozygous GDAP2 loss-of-function variant in a patient with adult-onset cerebellar ataxia. Brain, 2020, 143, e49-e49.	7.6	5
129	DaTSCAN (123I-FP-CIT SPECT) imaging in early versus mid and late onset Parkinson's disease: Longitudinal data from the PPMI study. Parkinsonism and Related Disorders, 2020, 77, 36-42.	2.2	5
130	Late life psychotic features in prodromal Parkinson's disease. Parkinsonism and Related Disorders, 2021, 86, 67-73.	2.2	5
131	lgG4-related autoimmune manifestations in Alemtuzumab-treated multiple sclerosis patients. Journal of Neuroimmunology, 2021, 361, 577759.	2.3	5
132	Cortical involvement and leptomeningeal inflammation in myelin oligodendrocyte glycoprotein antibody disease: A three-dimensional fluid-attenuated inversion recovery MRI study. Multiple Sclerosis Journal, 2022, 28, 718-729.	3.0	4
133	Dopamine transporter SPECT imaging in corticobasal syndrome: A peak into the underlying pathology?. Acta Neurologica Scandinavica, 2022, 145, 762-769.	2.1	4
134	The Interaction between <scp><i>HLAâ€DRB1</i></scp> and Smoking in Parkinson's Disease Revisited. Movement Disorders, 2022, 37, 1929-1937.	3.9	4
135	Dopamine agonists and delusional jealousy in Parkinson's disease: A crossâ€sectional prevalence study. Movement Disorders, 2013, 28, 689-689.	3.9	3
136	Neuroimaging findings in Hunter disease. Journal of the Neurological Sciences, 2014, 342, 200-201.	0.6	3
137	Serum Uric Acid in LRRK2 Related Parkinson's Disease: Longitudinal Data from the PPMI Study. Journal of Parkinson's Disease, 2021, 11, 633-640.	2.8	3
138	Variant transthyretin amyloidosis (ATTRv) polyneuropathy in Greece: a broad overview with a focus on non-endemic unexplored regions of the country. Neuromuscular Disorders, 2021, 31, 1251-1258.	0.6	3
139	Effects of a structured dance program in Parkinson's disease. A Greek pilot study. Complementary Therapies in Clinical Practice, 2022, 46, 101528.	1.7	3
140	Natalizumab therapy in patients with pediatric-onset multiple sclerosis in Greece: clinical and immunological insights of time-long administration and future directions—a single-center retrospective observational study. Naunyn-Schmiedeberg's Archives of Pharmacology, 2022, 395, 933-943.	3.0	3
141	Lipid level alteration in human and cellular models of alpha synuclein mutations. Npj Parkinson's Disease, 2022, 8, 52.	5.3	3
142	Callosal Angle Sub-Score of the Radscale in Patients with Idiopathic Normal Pressure Hydrocephalus Is Associated with Positive Tap Test Response. Journal of Clinical Medicine, 2022, 11, 2898.	2.4	3
143	Resolution of unilateral obstructive hydrocephalus complicating expanding thrombosed basilar apex aneurysm after anticoagulation treatment. Journal of the Neurological Sciences, 2014, 341, 179-181.	0.6	2
144	Analysis of a founder mutation in the <i>TH</i> gene in a cohort of greek patients with Parkinson's disease. Movement Disorders, 2016, 31, 1753-1754.	3.9	2

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145	Quantifying apraxia and ophthalmokinetic abnormalities in patients with atypical Parkinsonism: A new way to differential diagnosis?. Parkinsonism and Related Disorders, 2019, 61, 39-44.	2.2	2
146	Area postrema involvement in chronic lymphocytic inflammation with pontine perivascular enhancement. Neurological Sciences, 2021, 42, 361-364.	1.9	2
147	CSF and Circulating NfL as Biomarkers for the Discrimination of Parkinson Disease From Atypical Parkinsonian Syndromes. Neurology: Clinical Practice, 2021, 11, e867-e875.	1.6	2
148	<scp><i>HINT1</i></scp> â€related neuropathy in Greek patients with <scp>Charcotâ€Marieâ€Tooth</scp> disease. Journal of the Peripheral Nervous System, 2021, 26, 444-448.	3.1	2
149	Co-occurrence between hereditary angioedema and multiple sclerosis: Therapeutic management of both diseases with fingolimod. Clinical Neurology and Neurosurgery, 2022, 216, 107222.	1.4	2
150	Brain Arteriovenous Malformation and Amyotrophic Lateral Sclerosis: a Review Based on Published Cases. SN Comprehensive Clinical Medicine, 2020, 2, 392-396.	0.6	1
151	Preserved eye movements in adults with spinal muscular atrophy. Muscle and Nerve, 2021, 63, 765-769.	2.2	1
152	The "hypointense substantia nigra―sign. A novel MRI marker of progressive supranuclear palsy. Journal of the Neurological Sciences, 2021, 421, 117286.	0.6	1
153	Odor Identification Testing Can Assist in the Clinical Distinction Between Psychiatric Disorders and Neurological/Neurodegenerative Disorders. Alzheimer Disease and Associated Disorders, 2021, 35, 244-249.	1.3	1
154	The factors associated with the presence of psychotic symptoms in the HELIAD Greek community study of older adults. Aging and Mental Health, 2021, , 1-10.	2.8	1
155	A double-hit in vivo model of GBA viral microRNA-mediated downregulation and human alpha-synuclein overexpression demonstrates nigrostriatal degeneration. Neurobiology of Disease, 2022, 163, 105612.	4.4	1
156	Lupus-like disease and progressive multifocal leukoencephalopathy following etanercept treatment: just a coincidence?. Clinical and Experimental Rheumatology, 2022, 40, 671-672.	0.8	1
157	Teaching Neuro <i>Images</i> : MRI-visible Virchow-Robin perivascular spaces in cerebral small-vessel disease. Neurology, 2014, 83, e119-20.	1.1	0
158	Response to correspondence: Testing for myelin oligodendrocyte glycoprotein antibody (MOG-lgG) in typical MS. Multiple Sclerosis and Related Disorders, 2019, 35, 156-157.	2.0	0
159	Ocular flutter as the cardinal feature of anti-GM2 rhombencephalitis. Neurological Sciences, 2021, 42, 3003-3005.	1.9	0
160	Reply to: "αâ€Synuclein (<scp><i>SNCA</i></scp>) <scp>A30G</scp> Mutation as a Cause of a Complex Phenotype Without Parkinsonism― Movement Disorders, 2021, 36, 2212-2213.	3.9	0
161	Effect of Apolipoprotein E4 on the Driving Behavior of Patients with Amnestic Mild Cognitive Impairment or Mild Alzheimer's Disease Dementia. Journal of Alzheimer's Disease, 2021, 84, 1005-1014.	2.6	О
162	Alpha-synuclein gene and Parkinson's disease. , 2020, , 19-34.		O

#	Article	lF	CITATIONS
163	Asymptomatic carriers of the p.A53T SNCA mutation: Data from the PPMI study. Parkinsonism and Related Disorders, 2022, 98, 72-74.	2.2	O
164	Lupus-like disease and progressive multifocal leukoencephalopathy following etanercept treatment: just a coincidence?. Clinical and Experimental Rheumatology, 2021, , .	0.8	0