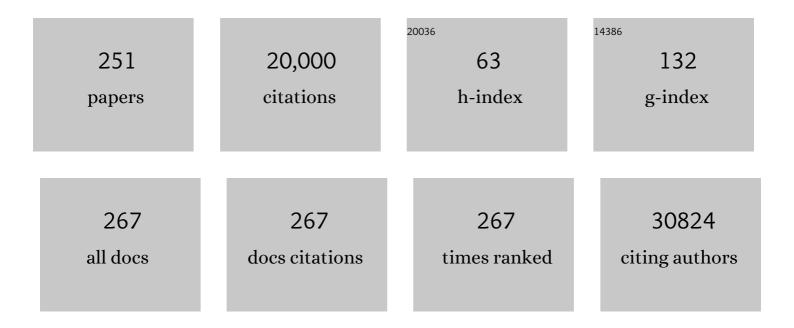
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Seizure prevalence in neurodegenerative diseases—a study of autopsy proven cases. European Journal of Neurology, 2022, 29, 12-18.	1.7	6
2	One-year outcome of brain injured patients undergoing early neurological rehabilitation: a prospective observational study. BMC Neurology, 2022, 22, 30.	0.8	8
3	Innovative therapeutic concepts of progressive multifocal leukoencephalopathy. Journal of Neurology, 2022, 269, 2403-2413.	1.8	12
4	Neuropsychiatric Symptoms in Parkinson's Disease Patients Are Associated with Reduced Health-Related Quality of Life and Increased Caregiver Burden. Brain Sciences, 2022, 12, 89.	1.1	17
5	Neurological management and work-up of neurotoxicity associated with CAR T cell therapy. Neurological Research and Practice, 2022, 4, 1.	1.0	9
6	Impact of Partial Volume Correction on [18F]GE-180 PET Quantification in Subcortical Brain Regions of Patients with Corticobasal Syndrome. Brain Sciences, 2022, 12, 204.	1.1	2
7	Longitudinal changes of early motor and cognitive symptoms in progressive supranuclear palsy: the OxQUIP study. BMJ Neurology Open, 2022, 4, e000214.	0.7	5
8	Safety, Pharmacokinetics, and Pharmacodynamics of Oral Venglustat in Patients with Parkinson's Disease and a GBA Mutation: Results from Part 1 of the Randomized, Double-Blinded, Placebo-Controlled MOVES-PD Trial. Journal of Parkinson's Disease, 2022, 12, 557-570.	1.5	34
9	Tau deposition patterns are associated with functional connectivity in primary tauopathies. Nature Communications, 2022, 13, 1362.	5.8	34
10	Reduction in Volume of Nucleus Basalis of Meynert Is Specific to Parkinson's Disease and Progressive Supranuclear Palsy but Not to Multiple System Atrophy. Frontiers in Aging Neuroscience, 2022, 14, 851788.	1.7	7
11	The Movement Disorder Society Criteria for the Diagnosis of Multiple System Atrophy. Movement Disorders, 2022, 37, 1131-1148.	2.2	222
12	Transcriptome and Proteome Analysis in LUHMES Cells Overexpressing Alpha-Synuclein. Frontiers in Neurology, 2022, 13, 787059.	1.1	9
13	Binding Stability of Antibody—α-Synuclein Complexes Predicts the Protective Efficacy of Anti-α-synuclein Antibodies. Molecular Neurobiology, 2022, 59, 3980-3995.	1.9	3
14	A new paradigm for diagnosis of neurodegenerative diseases: peripheral exosomes of brain origin. Translational Neurodegeneration, 2022, 11, 28.	3.6	37
15	Inferring the sequence of brain volume changes in progressive supranuclear palsy using MRI. Brain Communications, 2022, 4, .	1.5	1
16	Comparative analysis of machine learning algorithms for multi-syndrome classification of neurodegenerative syndromes. Alzheimer's Research and Therapy, 2022, 14, 62.	3.0	9
17	GBA-associated PD: chances and obstacles for targeted treatment strategies. Journal of Neural Transmission, 2022, 129, 1219-1233.	1.4	22
18	In Vivo Assessment of Neuroinflammation in <scp>4â€Repeat</scp> Tauopathies. Movement Disorders, 2021, 36, 883-894.	2.2	37

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19	Genetic determinants of survival in progressive supranuclear palsy: a genome-wide association study. Lancet Neurology, The, 2021, 20, 107-116.	4.9	62
20	The influence of the CRS-R score on functional outcome in patients with severe brain injury receiving early rehabilitation. BMC Neurology, 2021, 21, 44.	0.8	13
21	The "zig-zag―sign in progressive supranuclear palsy – The slowness of vertical saccades was the clue. Parkinsonism and Related Disorders, 2021, 83, 6-7.	1.1	Ο
22	Auditory Stimulation Modulates Resting-State Functional Connectivity in Unresponsive Wakefulness Syndrome Patients. Frontiers in Neuroscience, 2021, 15, 554194.	1.4	7
23	One Year Trajectory of Caregiver Burden in Parkinson's Disease and Analysis of Gender-Specific Aspects. Brain Sciences, 2021, 11, 295.	1.1	19
24	Outcomes of <scp>SARS oVâ€2</scp> Infections in Patients with Neurodegenerative Diseases in the <scp>LEOSS</scp> Cohort. Movement Disorders, 2021, 36, 791-793.	2.2	13
25	Safety and efficacy of tilavonemab in progressive supranuclear palsy: a phase 2, randomised, placebo-controlled trial. Lancet Neurology, The, 2021, 20, 182-192.	4.9	74
26	First symptom guides diagnosis and prognosis in neurodegenerative diseases—a retrospective study of autopsy proven cases. European Journal of Neurology, 2021, 28, 1801-1811.	1.7	11
27	Genotype–Phenotype Relations for the Atypical Parkinsonism Genes: MDSGene Systematic Review. Movement Disorders, 2021, 36, 1499-1510.	2.2	22
28	Evidence for pathogenicity of variant ATM Val1729Leu in a family with ataxia telangiectasia. Neurogenetics, 2021, 22, 143-147.	0.7	2
29	Comprehensive miRNome-Wide Profiling in a Neuronal Cell Model of Synucleinopathy Implies Involvement of Cell Cycle Genes. Frontiers in Cell and Developmental Biology, 2021, 9, 561086.	1.8	9
30	Clinical Features Observed in General Practice Associated With the Subsequent Diagnosis of Progressive Supranuclear Palsy. Frontiers in Neurology, 2021, 12, 637176.	1.1	9
31	Prothrombotic immune thrombocytopenia after COVID-19 vaccination. Blood, 2021, 138, 350-353.	0.6	145
32	DescribePSP and ProPSP: German Multicenter Networks for Standardized Prospective Collection of Clinical Data, Imaging Data, and Biomaterials of Patients With Progressive Supranuclear Palsy. Frontiers in Neurology, 2021, 12, 644064.	1.1	3
33	Validation of the Parkinson's Disease Caregiver Burden Questionnaire in Progressive Supranuclear Palsy. Parkinson's Disease, 2021, 2021, 1-7.	0.6	3
34	Cortical [<scp>¹⁸F</scp>] <scp>PI</scp> â€2620 Binding Differentiates Corticobasal Syndrome Subtypes. Movement Disorders, 2021, 36, 2104-2115.	2.2	46
35	Allogeneic BK Virus-Specific T-Cell Treatment in 2 Patients With Progressive Multifocal Leukoencephalopathy. Neurology: Neuroimmunology and NeuroInflammation, 2021, 8, e1020.	3.1	19
36	Binding characteristics of [¹⁸ F]PI-2620 distinguish the clinically predicted tau isoform in different tauopathies by PET. Journal of Cerebral Blood Flow and Metabolism, 2021, 41, 2957-2972.	2.4	30

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37	Feasibility of short imaging protocols for [18F]PI-2620 tau-PET in progressive supranuclear palsy. European Journal of Nuclear Medicine and Molecular Imaging, 2021, 48, 3872-3885.	3.3	22
38	Dual-Phase β-Amyloid PET Captures Neuronal Injury and Amyloidosis in Corticobasal Syndrome. Frontiers in Aging Neuroscience, 2021, 13, 661284.	1.7	13
39	Impact of TSPO Receptor Polymorphism on [18F]GE-180 Binding in Healthy Brain and Pseudo-Reference Regions of Neurooncological and Neurodegenerative Disorders. Life, 2021, 11, 484.	1.1	11
40	Clinical Features of Patients With Progressive Supranuclear Palsy in an US Insurance Claims Database. Frontiers in Neurology, 2021, 12, 571800.	1.1	14
41	Relationship Satisfaction in People with Parkinson's Disease and Their Caregivers: A Cross-Sectional Observational Study. Brain Sciences, 2021, 11, 822.	1.1	11
42	Non-invasive and high-throughput interrogation of exon-specific isoform expression. Nature Cell Biology, 2021, 23, 652-663.	4.6	11
43	Differential expression of gut miRNAs in idiopathic Parkinson's disease. Parkinsonism and Related Disorders, 2021, 88, 46-50.	1.1	8
44	Atypical pantothenate kinase-associated neurodegeneration with variable phenotypes in an Egyptian family. Heliyon, 2021, 7, e07469.	1.4	0
45	Superiority of Formalin-Fixed Paraffin-Embedded Brain Tissue for in vitro Assessment of Progressive Supranuclear Palsy Tau Pathology With [18F]PI-2620. Frontiers in Neurology, 2021, 12, 684523.	1.1	11
46	Does the Anti‶au Strategy in Progressive Supranuclear Palsy Need to Be Reconsidered? <scp>No</scp> . Movement Disorders Clinical Practice, 2021, 8, 1038-1040.	0.8	5
47	<scp>COVID</scp> â€19 Vaccineâ€Associated Cerebral Venous Thrombosis in Germany. Annals of Neurology, 2021, 90, 627-639.	2.8	122
48	Evolving concepts in progressive supranuclear palsy and other 4-repeat tauopathies. Nature Reviews Neurology, 2021, 17, 601-620.	4.9	41
49	Frequency and Characterization of Movement Disorders in Anti-IgLON5 Disease. Neurology, 2021, 97, .	1.5	50
50	iPS Cell-Based Model for MAPT Haplotype as a Risk Factor for Human Tauopathies Identifies No Major Differences in TAU Expression. Frontiers in Cell and Developmental Biology, 2021, 9, 726866.	1.8	4
51	Safety and efficacy of anti-tau monoclonal antibody gosuranemab in progressive supranuclear palsy: a phase 2, randomized, placebo-controlled trial. Nature Medicine, 2021, 27, 1451-1457.	15.2	63
52	Alpha-Synuclein defects autophagy by impairing SNAP29-mediated autophagosome-lysosome fusion. Cell Death and Disease, 2021, 12, 854.	2.7	39
53	Neurological symptoms and complications in predominantly hospitalized COVIDâ€19 patients: Results of the European multinational Lean European Open Survey on SARSâ€Infected Patients (LEOSS). European Journal of Neurology, 2021, 28, 3925-3937.	1.7	25
54	Treatment of upper limb spasticity with inhibitory repetitive transcranial magnetic stimulation: A randomized placebo-controlled trial. NeuroRehabilitation, 2021, 49, 425-434.	0.5	14

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55	Neuropathology of progressive supranuclear palsy after treatment with tilavonemab – Author's reply. Lancet Neurology, The, 2021, 20, 787-788.	4.9	3
56	PD-1-inhibitor pembrolizumab for treatment of progressive multifocal leukoencephalopathy. Therapeutic Advances in Neurological Disorders, 2021, 14, 175628642199368.	1.5	9
57	A Modified Progressive Supranuclear Palsy Rating Scale. Movement Disorders, 2021, 36, 1203-1215.	2.2	13
58	Reply to: "Application of the <scp>mPSPRS</scp> to the Salerno Cohort― Movement Disorders, 2021, 36, 2451-2452.	2.2	0
59	Microglial activation states drive glucose uptake and FDG-PET alterations in neurodegenerative diseases. Science Translational Medicine, 2021, 13, eabe5640.	5.8	108
60	Cerebral Microstructural Alterations in Patients With Early Parkinson's Disease Detected With Quantitative Magnetic Resonance Measurements. Frontiers in Aging Neuroscience, 2021, 13, 763331.	1.7	5
61	Analysis of Transition of Patients with Parkinson's Disease into Institutional Care: A Retrospective Pilot Study. Brain Sciences, 2021, 11, 1470.	1.1	9
62	Patient Safety in a Box: Implementation and Evaluation of the Emergency Box in Geriatric and Parkinson Patients. Journal of Clinical Medicine, 2021, 10, 5618.	1.0	1
63	Associations between sex, body mass index, and the individual microglial response in Alzheimer's disease. Alzheimer's and Dementia, 2021, 17, .	0.4	0
64	Can SARS-CoV-2 Infection Lead to Neurodegeneration and Parkinson's Disease?. Brain Sciences, 2021, 11, 1654.	1.1	22
65	Feasibility of short imaging protocols for [¹⁸ F]Plâ€2620 tauâ€PET in progressive supranuclear palsy. Alzheimer's and Dementia, 2021, 17, .	0.4	0
66	Validation of the Movement Disorder Society Criteria for the Diagnosis of 4â€Repeat Tauopathies. Movement Disorders, 2020, 35, 171-176.	2.2	37
67	Disease-modifying strategies in primary tauopathies. Neuropharmacology, 2020, 167, 107842.	2.0	7
68	Loss of fragile X mental retardation protein precedes Lewy pathology in Parkinson's disease. Acta Neuropathologica, 2020, 139, 319-345.	3.9	17
69	Video-tutorial for the Movement Disorder Society criteria for progressive supranuclear palsy. Parkinsonism and Related Disorders, 2020, 78, 200-203.	1.1	8
70	How specific are non-motor symptoms in the prodrome of Parkinson's disease compared to other movement disorders?. Parkinsonism and Related Disorders, 2020, 81, 213-218.	1.1	8
71	Alexithymia Is Associated with Reduced Quality of Life and Increased Caregiver Burden in Parkinson's Disease. Brain Sciences, 2020, 10, 401.	1.1	18
72	Longitudinal TSPO expression in tau transgenic P301S mice predicts increased tau accumulation and deteriorated spatial learning. Journal of Neuroinflammation, 2020, 17, 208.	3.1	19

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73	Reply to: "Brief Clinical Rating Scales Should Not Be Overlooked― Movement Disorders, 2020, 35, 1886-1886.	2.2	3
74	Basic Fibroblast Growth Factor 2-Induced Proteome Changes Endorse Lewy Body Pathology in Hippocampal Neurons. IScience, 2020, 23, 101349.	1.9	4
75	Postinfectious Onset of Myasthenia Gravis in a COVID-19 Patient. Frontiers in Neurology, 2020, 11, 576153.	1.1	64
76	FGF2 Affects Parkinson's Disease-Associated Molecular Networks Through Exosomal Rab8b/Rab31. Frontiers in Genetics, 2020, 11, 572058.	1.1	12
77	Brain Morphological Alterations Are Detected in Earlyâ€6tage Parkinson's Disease with MRI Morphometry. Journal of Neuroimaging, 2020, 30, 786-792.	1.0	8
78	Can Autonomic Testing and Imaging Contribute to the Early Diagnosis of Multiple System Atrophy? A Systematic Review and Recommendations by the <scp>Movement Disorder Society</scp> Multiple System Atrophy Study Group. Movement Disorders Clinical Practice, 2020, 7, 750-762.	0.8	31
79	Clinical Conditions "Suggestive of Progressive Supranuclear Palsyâ€â€"Diagnostic Performance. Movement Disorders, 2020, 35, 2301-2313.	2.2	22
80	Assessment of ¹⁸ F-PI-2620 as a Biomarker in Progressive Supranuclear Palsy. JAMA Neurology, 2020, 77, 1408.	4.5	145
81	Hospitalization Rates and Comorbidities in Patients with Progressive Supranuclear Palsy in Germany from 2010 to 2017. Journal of Clinical Medicine, 2020, 9, 2454.	1.0	3
82	Glial activation is moderated by sex in response to amyloidosis but not to tau pathology in mouse models of neurodegenerative diseases. Journal of Neuroinflammation, 2020, 17, 374.	3.1	28
83	Microglial activation in vivo is moderated by sex in response to amyloidosis but not to tau pathology in mouse models of Alzheimer's disease. Alzheimer's and Dementia, 2020, 16, e039574.	0.4	Ο
84	18 Fâ€Plâ€2620 tauâ€PET in corticobasal syndrome (ActiGliA cohort). Alzheimer's and Dementia, 2020, 16, e041469.	0.4	1
85	Microglial activation and brain networks in Alzheimer's disease: The ActiGliA cohort study. Alzheimer's and Dementia, 2020, 16, e043265.	0.4	0
86	Reply to: †̃Letter to the Editor on "Copathology Progressive Supranuclear Palsy: Does It Matter?â€â€™. Movement Disorders, 2020, 35, 2126-2126.	2.2	2
87	Distribution patterns of tau pathology in progressive supranuclear palsy. Acta Neuropathologica, 2020, 140, 99-119.	3.9	210
88	Copathology in Progressive Supranuclear Palsy: Does It Matter?. Movement Disorders, 2020, 35, 984-993.	2.2	48
89	Author response: Use of β2-adrenoreceptor agonist and antagonist drugs and risk of Parkinson disease. Neurology, 2020, 94, 899-899.	1.5	0
90	Longitudinal correlation between neurofilament light chain and UMSARS in Multiple System Atrophy. Clinical Neurology and Neurosurgery, 2020, 195, 105924.	0.6	3

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91	Fibroblast Growth Factor 2â€Mediated Regulation of Neuronal Exosome Release Depends on VAMP3/Cellubrevin in Hippocampal Neurons. Advanced Science, 2020, 7, 1902372.	5.6	33
92	Mindfulness and Psychological Flexibility are Inversely Associated with Caregiver Burden in Parkinson's Disease. Brain Sciences, 2020, 10, 111.	1.1	16
93	Consensus-Based Recommendations for Advance Directives of People with Parkinson's Disease in Regard to Typical Complications by German Movement Disorder Specialists. Journal of Clinical Medicine, 2020, 9, 449.	1.0	7
94	The Progressive Supranuclear Palsy Clinical Deficits Scale. Movement Disorders, 2020, 35, 650-661.	2.2	31
95	β-adrenoreceptors and the risk of Parkinson's disease. Lancet Neurology, The, 2020, 19, 247-254.	4.9	49
96	Alpha-synuclein fragments trigger distinct aggregation pathways. Cell Death and Disease, 2020, 11, 84.	2.7	19
97	Rare Variants in Specific Lysosomal Genes Are Associated With Parkinson's Disease. Movement Disorders, 2020, 35, 1245-1248.	2.2	37
98	Early-phase [18F]PI-2620 tau-PET imaging as a surrogate marker of neuronal injury. European Journal of Nuclear Medicine and Molecular Imaging, 2020, 47, 2911-2922.	3.3	36
99	Looking into the prediagnostic phase of progressive supranuclear palsy. Parkinsonism and Related Disorders, 2020, 74, 74-75.	1.1	1
100	Private variants in PRKN are associated with late-onset Parkinson's disease. Parkinsonism and Related Disorders, 2020, 75, 24-26.	1.1	4
101	A call for a global COVID-19 Neuro Research Coalition. Lancet Neurology, The, 2020, 19, 482-484.	4.9	22
102	<scp><i>LRP1</i></scp> : A Novel Mediator of Tau Uptake. Movement Disorders, 2020, 35, 1136-1136.	2.2	1
103	Towards a consensus on developmental regression. Neuroscience and Biobehavioral Reviews, 2019, 107, 3-5.	2.9	14
104	One decade ago, one decade ahead in progressive supranuclear palsy. Movement Disorders, 2019, 34, 1284-1293.	2.2	12
105	Tau links developmental to neurodegenerative diseases. Neuroscience and Biobehavioral Reviews, 2019, 104, 26-27.	2.9	1
106	Safety and efficacy of epigallocatechin gallate in multiple system atrophy (PROMESA): a randomised, double-blind, placebo-controlled trial. Lancet Neurology, The, 2019, 18, 724-735.	4.9	79
107	Neuronal precursor cells with dopaminergic commitment in the rostral migratory stream of the mouse. Scientific Reports, 2019, 9, 13359.	1.6	12
108	PET Imaging of Astrogliosis and Tau Facilitates Diagnosis of Parkinsonian Syndromes. Frontiers in Aging Neuroscience, 2019, 11, 249.	1.7	30

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109	Use of β2-adrenoreceptor agonist and antagonist drugs and risk of Parkinson disease. Neurology, 2019, 93, e135-e142.	1.5	29
110	Four-repeat tauopathies. Progress in Neurobiology, 2019, 180, 101644.	2.8	141
111	A critique of the second consensus criteria for multiple system atrophy. Movement Disorders, 2019, 34, 975-984.	2.2	73
112	Clinical, pathophysiological and genetic features of motor symptoms in autosomal dominant Alzheimer's disease. Brain, 2019, 142, 1429-1440.	3.7	36
113	How to apply the movement disorder society criteria for diagnosis of progressive supranuclear palsy. Movement Disorders, 2019, 34, 1228-1232.	2.2	93
114	Safety and Tolerability of Pharmacotherapies for Parkinson's Disease in Geriatric Patients. Drugs and Aging, 2019, 36, 511-530.	1.3	38
115	Neuroimaging biomarkers for clinical trials in atypical parkinsonian disorders: Proposal for a Neuroimaging Biomarker Utility System. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2019, 11, 301-309.	1.2	30
116	Multiple molecular pathways stimulating macroautophagy protect from alpha-synuclein-induced toxicity in human neurons. Neuropharmacology, 2019, 149, 13-26.	2.0	14
117	Unbiased Screens for Modifiers of Alpha-Synuclein Toxicity. Current Neurology and Neuroscience Reports, 2019, 19, 8.	2.0	8
118	Mitochondrial damage by α-synuclein causes cell death in human dopaminergic neurons. Cell Death and Disease, 2019, 10, 865.	2.7	112
119	Classification of atypical parkinsonism per pathology versus phenotype. International Review of Neurobiology, 2019, 149, 37-47.	0.9	10
120	Genetic mimics of the non-genetic atypical parkinsonian disorders – the â€~atypical' atypical. International Review of Neurobiology, 2019, 149, 327-351.	0.9	8
121	Progressive supranuclear palsy. International Review of Neurobiology, 2019, 149, 49-86.	0.9	19
122	Severity dependent distribution of impairments in PSP and CBS: Interactive visualizations. Parkinsonism and Related Disorders, 2019, 60, 138-145.	1.1	7
123	Pearls & Oy-sters: Ocular motor apraxia as essential differential diagnosis to supranuclear gaze palsy. Neurology, 2018, 90, 482-485.	1.5	10
124	CXCR4 involvement in neurodegenerative diseases. Translational Psychiatry, 2018, 8, 73.	2.4	66
125	Selective Genetic Overlap Between Amyotrophic Lateral Sclerosis and Diseases of the Frontotemporal Dementia Spectrum. JAMA Neurology, 2018, 75, 860.	4.5	79
126	Is it Useful to Classify Progressive Supranuclear Palsy and Corticobasal Degeneration as Different Disorders? No. Movement Disorders Clinical Practice, 2018, 5, 141-144.	0.8	28

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127	Recommendations of the Global Multiple System Atrophy Research Roadmap Meeting. Neurology, 2018, 90, 74-82.	1.5	23
128	Symptomatic therapy of multiple system atrophy. Autonomic Neuroscience: Basic and Clinical, 2018, 211, 26-30.	1.4	18
129	K-variant BCHE and pesticide exposure: Gene-environment interactions in a case–control study of Parkinson's disease in Egypt. Scientific Reports, 2018, 8, 16525.	1.6	21
130	Epigenome-wide DNA methylation profiling in Progressive Supranuclear Palsy reveals major changes at DLX1. Nature Communications, 2018, 9, 2929.	5.8	20
131	Variation at the <i>TRIM11</i> locus modifies progressive supranuclear palsy phenotype. Annals of Neurology, 2018, 84, 485-496.	2.8	37
132	Exosomal secretion of α-synuclein as protective mechanism after upstream blockage of macroautophagy. Cell Death and Disease, 2018, 9, 757.	2.7	117
133	Progressive supranuclear palsy and multiple system atrophy: clinicopathological concepts and therapeutic challenges. Current Opinion in Neurology, 2018, 31, 448-454.	1.8	19
134	New classification of tauopathies. Revue Neurologique, 2018, 174, 664-668.	0.6	39
135	Immune-related genetic enrichment in frontotemporal dementia: An analysis of genome-wide association studies. PLoS Medicine, 2018, 15, e1002487.	3.9	111
136	Manual MRI morphometry in Parkinsonian syndromes. Movement Disorders, 2017, 32, 778-782.	2.2	67
137	<scp>PERK</scp> activation mitigates tau pathology <i>inÂvitro</i> and <i>inÂvivo</i> . EMBO Molecular Medicine, 2017, 9, 371-384.	3.3	93
138	Shared genetic risk between corticobasal degeneration, progressive supranuclear palsy, and frontotemporal dementia. Acta Neuropathologica, 2017, 133, 825-837.	3.9	90
139	Differentiation of atypical Parkinson syndromes. Journal of Neural Transmission, 2017, 124, 997-1004.	1.4	30
140	Multiple System Atrophy. , 2017, , 183-192.		2
141	Which ante mortem clinical features predict progressive supranuclear palsy pathology?. Movement Disorders, 2017, 32, 995-1005.	2.2	121
142	Radiological biomarkers for diagnosis in PSP: Where are we and where do we need to be?. Movement Disorders, 2017, 32, 955-971.	2.2	179
143	Clinical diagnosis of progressive supranuclear palsy: The movement disorder society criteria. Movement Disorders, 2017, 32, 853-864.	2.2	1,402
144	Longitudinal magnetic resonance imaging in progressive supranuclear palsy: A new combined score for clinical trials. Movement Disorders, 2017, 32, 842-852.	2.2	52

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145	Reply to: MRI measures of brainstem in parkinsonian syndromes: Where we stand and where we need to go. Movement Disorders, 2017, 32, 1261-1262.	2.2	1
146	Advances in progressive supranuclear palsy: new diagnostic criteria, biomarkers, and therapeutic approaches. Lancet Neurology, The, 2017, 16, 552-563.	4.9	303
147	Tau Diagnostics and Clinical Studies. Journal of Molecular Neuroscience, 2017, 63, 123-130.	1.1	11
148	Protective efficacy of phosphodiesterase-1 inhibition against alpha-synuclein toxicity revealed by compound screening in LUHMES cells. Scientific Reports, 2017, 7, 11469.	1.6	52
149	c.207C>G mutation in sepiapterin reductase causes autosomal dominant dopa-responsive dystonia. Neurology: Genetics, 2017, 3, e197.	0.9	10
150	Drug Safety Analysis in a Real-Life Cohort of Parkinson's Disease Patients with Polypharmacy. CNS Drugs, 2017, 31, 1093-1102.	2.7	15
151	Preclinical Analysis of Fetal Human Mesencephalic Neural Progenitor Cell Lines: Characterization and Safety In Vitro and In Vivo. Stem Cells Translational Medicine, 2017, 6, 576-588.	1.6	11
152	[18F]-THK5351 PET Correlates with Topology and Symptom Severity in Progressive Supranuclear Palsy. Frontiers in Aging Neuroscience, 2017, 9, 440.	1.7	58
153	The Differential Diagnosis and Treatment of Atypical Parkinsonism. Deutsches Ärzteblatt International, 2016, 113, 61-9.	0.6	135
154	Differential effects of social and physical environmental enrichment on brain plasticity, cognition, and ultrasonic communication in rats. Journal of Comparative Neurology, 2016, 524, 1586-1607.	0.9	122
155	l-DOPA-induced dyskinesia is associated with a deficient numerical downregulation of striatal tyrosine hydroxylase mRNA-expressing neurons. Neuroscience, 2016, 331, 120-133.	1.1	7
156	A Review of Treatment Options for Progressive Supranuclear Palsy. CNS Drugs, 2016, 30, 629-636.	2.7	36
157	Parkinson's disease: SNCA-, PARK2-, and LRRK2- targeting microRNAs elevated in cingulate gyrus. Parkinsonism and Related Disorders, 2016, 33, 115-121.	1.1	72
158	Chronic consumption of <i>Annona muricata</i> juice triggers and aggravates cerebral tau phosphorylation in wildâ€ŧype and <i><scp>MAPT</scp></i> transgenic mice. Journal of Neurochemistry, 2016, 139, 624-639.	2.1	26
159	Differentiation of neurodegenerative parkinsonian syndromes by volumetric magnetic resonance imaging analysis and support vector machine classification. Movement Disorders, 2016, 31, 1506-1517.	2.2	120
160	A genome-wide association study in multiple system atrophy. Neurology, 2016, 87, 1591-1598.	1.5	139
161	Current Treatment of Multiple System Atrophy. Current Treatment Options in Neurology, 2016, 18, 51.	0.7	9
162	microRNA profiling: increased expression of miR-147a and miR-518e in progressive supranuclear palsy (PSP). Neurogenetics, 2016, 17, 165-171.	0.7	20

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163	The PROMESA-protocol: progression rate of multiple system atrophy under EGCG supplementation as anti-aggregation-approach. Journal of Neural Transmission, 2016, 123, 439-445.	1.4	32
164	Power calculations and placebo effect for future clinical trials in progressive supranuclear palsy. Movement Disorders, 2016, 31, 742-747.	2.2	29
165	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). Autophagy, 2016, 12, 1-222.	4.3	4,701
166	Progressive supranuclear palsy: progression and survival. Journal of Neurology, 2016, 263, 380-389.	1.8	55
167	The phenotypic spectrum of progressive supranuclear palsy. Parkinsonism and Related Disorders, 2016, 22, S34-S36.	1.1	65
168	Psychosis in Parkinson's disease: identification, prevention and treatment. Journal of Neural Transmission, 2016, 123, 45-50.	1.4	39
169	Early Neurodegeneration in the Brain of a Child Without Functional PKR-like Endoplasmic Reticulum Kinase. Journal of Neuropathology and Experimental Neurology, 2015, 74, 850-857.	0.9	27
170	From a single nucleotide polymorphism to tau pathology: Appoptosin is the missing link. Movement Disorders, 2015, 30, 1871-1872.	2.2	2
171	Genome-wide association study of corticobasal degeneration identifies risk variants shared with progressive supranuclear palsy. Nature Communications, 2015, 6, 7247.	5.8	170
172	A new dopaminergic nigro-olfactory projection. Acta Neuropathologica, 2015, 130, 333-348.	3.9	89
173	Glucocerebrosidase deficiency and mitochondrial impairment in experimental Parkinson disease. Journal of the Neurological Sciences, 2015, 356, 129-136.	0.3	23
174	Improved preparation of nasal lavage fluid (NLF) as a noninvasive sample for proteomic biomarker discovery. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2015, 1854, 741-745.	1.1	6
175	Long-term treatment with I-DOPA or pramipexole affects adult neurogenesis and corresponding non-motor behavior in a mouse model of Parkinson's disease. Neuropharmacology, 2015, 95, 367-376.	2.0	51
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