

Dennis W Dickson

List of Publications by Year in descending order

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Version: 2024-02-01

834
papers

111,633
citations

¹⁷⁹
152
h-index

²⁸⁶
294
g-index

899
all docs

899
docs citations

899
times ranked

59675
citing authors

#	ARTICLE	IF	CITATIONS
1	Deep learning-based model for diagnosing Alzheimer's disease and tauopathies. <i>Neuropathology and Applied Neurobiology</i> , 2022, 48, .	1.8	33
2	Relationship Between ¹⁸ F-Flortaucipir Uptake and Histologic Lesion Types in 4-Repeat Tauopathies. <i>Journal of Nuclear Medicine</i> , 2022, 63, 931-935.	2.8	9
3	Genetic evaluation of dementia with Lewy bodies implicates distinct disease subgroups. <i>Brain</i> , 2022, 145, 1757-1762.	3.7	17
4	Autopsy Validation of Progressive Supranuclear Palsy-Predominant Speech/Language Disorder Criteria. <i>Movement Disorders</i> , 2022, 37, 213-218.	2.2	6
5	Genome-wide association study and functional validation implicates JADE1 in tauopathy. <i>Acta Neuropathologica</i> , 2022, 143, 33-53.	3.9	19
6	Alzheimer's disease and progressive supranuclear palsy share similar transcriptomic changes in distinct brain regions. <i>Journal of Clinical Investigation</i> , 2022, 132, .	3.9	13
7	Concurrent tau pathologies in frontotemporal lobar degeneration with TDP43 pathology. <i>Neuropathology and Applied Neurobiology</i> , 2022, 48, .	1.8	9
8	The temporal onset of the core features in dementia with Lewy bodies. <i>Alzheimer's and Dementia</i> , 2022, 18, 591-601.	0.4	19
9	Clinical Deep Phenotyping of <i>ABCA7</i> Mutation Carriers. <i>Neurology: Genetics</i> , 2022, 8, e655.	0.9	4
10	Neuropathological Findings of <i>CSF1R</i> -Related Leukoencephalopathy After Long-Term Immunosuppressive Therapy. <i>Movement Disorders</i> , 2022, 37, 439-440.	2.2	8
11	Diffuse Lewy body disease presenting as Parkinson's disease with progressive aphasia. <i>Neuropathology</i> , 2022, 42, 82-89.	0.7	4
12	Asymmetrical Primary Lateral Sclerosis Presenting as Corticobasal Syndrome. <i>Journal of Neuropathology and Experimental Neurology</i> , 2022, 81, 154-156.	0.9	2
13	TDP-43-associated atrophy in brains with and without frontotemporal lobar degeneration. <i>NeuroImage: Clinical</i> , 2022, 34, 102954.	1.4	3
14	Longitudinal atrophy in prodromal dementia with Lewy bodies points to cholinergic degeneration. <i>Brain Communications</i> , 2022, 4, fcac013.	1.5	15
15	Neuropathology of <i>McLeod</i> Syndrome. <i>Movement Disorders</i> , 2022, 37, 644-646.	2.2	5
16	TDP-43 represses cryptic exon inclusion in the FTD-ALS gene <i>UNC13A</i> . <i>Nature</i> , 2022, 603, 124-130.	13.7	193
17	Proximity proteomics of <i>C9orf72</i> dipeptide repeat proteins identifies molecular chaperones as modifiers of poly-GA aggregation. <i>Acta Neuropathologica Communications</i> , 2022, 10, 22.	2.4	22
18	Homotypic fibrillization of <i>TMEM106B</i> across diverse neurodegenerative diseases. <i>Cell</i> , 2022, 185, 1346-1355.e15.	13.5	70

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19	Amyloid fibrils in FTLN-TDP are composed of TMEM106B and not TDP-43. <i>Nature</i> , 2022, 605, 304-309.	13.7	85
20	Clinical and pathological characteristics of later onset multiple system atrophy. <i>Journal of Neurology</i> , 2022, 269, 4310-4321.	1.8	8
21	Plasma PolyQ-ATXN3 Levels Associate With Cerebellar Degeneration and Behavioral Abnormalities in a New AAV-Based SCA3 Mouse Model. <i>Frontiers in Cell and Developmental Biology</i> , 2022, 10, 863089.	1.8	5
22	Shared brain transcriptomic signature in TDP-43 type A FTLN patients with or without <i>GRN</i> mutations. <i>Brain</i> , 2022, 145, 2472-2485.	3.7	6
23	TREM2 interacts with TDP-43 and mediates microglial neuroprotection against TDP-43-related neurodegeneration. <i>Nature Neuroscience</i> , 2022, 25, 26-38.	7.1	52
24	Brainstem Biomarkers of Clinical Variant and Pathology in Progressive Supranuclear Palsy. <i>Movement Disorders</i> , 2022, 37, 702-712.	2.2	14
25	Manifestations of Alzheimer's disease genetic risk in the blood are evident in a multiomic analysis in healthy adults aged 18 to 90. <i>Scientific Reports</i> , 2022, 12, 6117.	1.6	12
26	APOE4 exacerbates β -synuclein seeding activity and contributes to neurotoxicity in Alzheimer's disease with Lewy body pathology. <i>Acta Neuropathologica</i> , 2022, 143, 641-662.	3.9	24
27	Histologic lesion type correlates of magnetic resonance imaging biomarkers in four-repeat tauopathies. <i>Brain Communications</i> , 2022, 4, .	1.5	5
28	Frequency and distribution of TAR DNA-binding protein 43 (TDP-43) pathology increase linearly with age in a large cohort of older adults with and without dementia. <i>Acta Neuropathologica</i> , 2022, 144, 159-160.	3.9	14
29	Neuropathology of Parkinson's disease after focused ultrasound thalamotomy. <i>Npj Parkinson's Disease</i> , 2022, 8, 59.	2.5	5
30	Diffusion tractography of superior cerebellar peduncle and dentatorubrothalamic tracts in two autopsy confirmed progressive supranuclear palsy variants: Richardson syndrome and the speech-language variant. <i>NeuroImage: Clinical</i> , 2022, 35, 103030.	1.4	8
31	Old age amyotrophic lateral sclerosis and limbic TDP-43 pathology. <i>Brain Pathology</i> , 2022, 32, .	2.1	6
32	Frequency of LATE neuropathologic change across the spectrum of Alzheimer's disease neuropathology: combined data from 13 community-based or population-based autopsy cohorts. <i>Acta Neuropathologica</i> , 2022, 144, 27-44.	3.9	67
33	<i>GRN</i> Mutations Are Associated with Lewy Body Dementia. <i>Movement Disorders</i> , 2022, 37, 1943-1948.	2.2	5
34	Mitochondrial genomic variation in dementia with Lewy bodies: association with disease risk and neuropathological measures. <i>Acta Neuropathologica Communications</i> , 2022, 10, .	2.4	0
35	SARS-CoV-2 Brain Regional Detection, Histopathology, Gene Expression, and Immunomodulatory Changes in Decedents with COVID-19. <i>Journal of Neuro pathology and Experimental Neurology</i> , 2022, 81, 666-695.	0.9	22
36	Sensitive ELISA-based detection method for the mitophagy marker p-S65-Ub in human cells, autopsy brain, and blood samples. <i>Autophagy</i> , 2021, 17, 2613-2628.	4.3	29

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37	Mitophagy alterations in Alzheimer's disease are associated with granulovacuolar degeneration and early tau pathology. <i>Alzheimer's and Dementia</i> , 2021, 17, 417-430.	0.4	34
38	Apoptotic Neuron-Derived Histone Amyloid Fibrils Induce α -Synuclein Aggregation. <i>Molecular Neurobiology</i> , 2021, 58, 867-876.	1.9	1
39	Lewy Body Disease is a Contributor to Logopenic Progressive Aphasia Phenotype. <i>Annals of Neurology</i> , 2021, 89, 520-533.	2.8	21
40	Genetic determinants of survival in progressive supranuclear palsy: a genome-wide association study. <i>Lancet Neurology</i> , The, 2021, 20, 107-116.	4.9	62
41	Early Selective Vulnerability of the CA2 Hippocampal Subfield in Primary Age-Related Tauopathy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 102-111.	0.9	35
42	Enrichment of Phosphorylated Tau (Thr181) and Functionally Interacting Molecules in Chronic Traumatic Encephalopathy Brain-derived Extracellular Vesicles. , 2021, 12, 1376.		3
43	Frequency of spinocerebellar ataxia mutations in patients with multiple system atrophy. <i>Clinical Autonomic Research</i> , 2021, 31, 117-125.	1.4	10
44	Progressive Supranuclear Palsy and Corticobasal Degeneration. <i>Advances in Experimental Medicine and Biology</i> , 2021, 1281, 151-176.	0.8	10
45	The Second NINDS/NIBIB Consensus Meeting to Define Neuropathological Criteria for the Diagnosis of Chronic Traumatic Encephalopathy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 210-219.	0.9	111
46	Association of Mitochondrial DNA Genomic Variation With Risk of Pick Disease. <i>Neurology</i> , 2021, 96, e1755-e1760.	1.5	1
47	Latent trait modeling of tau neuropathology in progressive supranuclear palsy. <i>Acta Neuropathologica</i> , 2021, 141, 667-680.	3.9	5
48	Deep Learning-Based Image Classification in Differentiating Tufted Astrocytes, Astrocytic Plaques, and Neuritic Plaques. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 306-312.	0.9	21
49	Genome sequencing analysis identifies new loci associated with Lewy body dementia and provides insights into its genetic architecture. <i>Nature Genetics</i> , 2021, 53, 294-303.	9.4	198
50	Underlying pathology identified after 20 years of disease course in two cases of slowly progressive frontotemporal dementia syndromes. <i>Neurocase</i> , 2021, 27, 212-222.	0.2	4
51	The AD tau core spontaneously self-assembles and recruits full-length tau to filaments. <i>Cell Reports</i> , 2021, 34, 108843.	2.9	30
52	TAR DNA-Binding Protein 43 Is Associated with Rate of Memory, Functional and Global Cognitive Decline in the Decade Prior to Death. <i>Journal of Alzheimer's Disease</i> , 2021, 80, 683-693.	1.2	7
53	Loss of Tmem106b leads to cerebellum Purkinje cell death and motor deficits. <i>Brain Pathology</i> , 2021, 31, e12945.	2.1	8
54	Investigating ELOVL7 coding variants in multiple system atrophy. <i>Neuroscience Letters</i> , 2021, 749, 135723.	1.0	2

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55	Machine learning-based decision tree classifier for the diagnosis of progressive supranuclear palsy and corticobasal degeneration. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 931-941.	1.8	22
56	Long-read targeted sequencing uncovers clinicopathological associations for <i>C9orf72</i> -linked diseases. <i>Brain</i> , 2021, 144, 1082-1088.	3.7	17
57	Transcriptomic analysis to identify genes associated with selective hippocampal vulnerability in Alzheimer's disease. <i>Nature Communications</i> , 2021, 12, 2311.	5.8	44
58	Cerebral Amyloid Angiopathy Burden and Cerebral Microbleeds: Pathological Evidence for Distinct Phenotypes. <i>Journal of Alzheimer's Disease</i> , 2021, 81, 113-122.	1.2	8
59	MRI quantitative susceptibility mapping of the substantia nigra as an early biomarker for Lewy body disease. <i>Journal of Neuroimaging</i> , 2021, 31, 1020-1027.	1.0	13
60	Genome-wide analysis identifies a novel LINC-PINT splice variant associated with vascular amyloid pathology in Alzheimer's disease. <i>Acta Neuropathologica Communications</i> , 2021, 9, 93.	2.4	9
61	Old age genetically confirmed frontotemporal lobar degeneration with TDP43 has limbic predominant TDP43 deposition. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 1050-1059.	1.8	10
62	Tau isoforms are differentially expressed across the hippocampus in chronic traumatic encephalopathy and Alzheimer's disease. <i>Acta Neuropathologica Communications</i> , 2021, 9, 86.	2.4	38
63	Clinical, Imaging, and Pathologic Characteristics of Patients With Right vs Left Hemisphere-Predominant Logopenic Progressive Aphasia. <i>Neurology</i> , 2021, 97, e523-e534.	1.5	4
64	A molecular pathology, neurobiology, biochemical, genetic and neuroimaging study of progressive apraxia of speech. <i>Nature Communications</i> , 2021, 12, 3452.	5.8	34
65	Analysis of intraoperative human brain tissue transcriptome reveals putative risk genes and altered molecular pathways in glioma-related seizures. <i>Epilepsy Research</i> , 2021, 173, 106618.	0.8	7
66	Cerebral Microvascular Erdheim-Chester Disease: A Perivascular Hematopoietic Vasculopathy. <i>Cerebrovascular Diseases</i> , 2021, 50, 746-751.	0.8	4
67	Nuclear accumulation of CHMP7 initiates nuclear pore complex injury and subsequent TDP-43 dysfunction in sporadic and familial ALS. <i>Science Translational Medicine</i> , 2021, 13, .	5.8	68
68	Predictors of cognitive impairment in primary age-related tauopathy: an autopsy study. <i>Acta Neuropathologica Communications</i> , 2021, 9, 134.	2.4	32
69	Clinical, pathological and genetic characteristics of Perry disease—new cases and literature review. <i>European Journal of Neurology</i> , 2021, 28, 4010-4021.	1.7	10
70	Clinical features of autopsy-confirmed multiple system atrophy in the Mayo Clinic Florida brain bank. <i>Parkinsonism and Related Disorders</i> , 2021, 89, 155-161.	1.1	12
71	Cellular and pathological heterogeneity of primary tauopathies. <i>Molecular Neurodegeneration</i> , 2021, 16, 57.	4.4	85
72	Apolipoprotein E regulates lipid metabolism and β -synuclein pathology in human iPSC-derived cerebral organoids. <i>Acta Neuropathologica</i> , 2021, 142, 807-825.	3.9	25

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73	Microglial lysosome dysfunction contributes to white matter pathology and TDP-43 proteinopathy in GRN-associated FTD. <i>Cell Reports</i> , 2021, 36, 109581.	2.9	33
74	Cerebral Amyloid Angiopathy Pathology and Its Association With Amyloid- β^2 PET Signal. <i>Neurology</i> , 2021, 97, e1799-e1808.	1.5	10
75	<i>APOE3</i> -Jacksonville (V236E) variant reduces self-aggregation and risk of dementia. <i>Science Translational Medicine</i> , 2021, 13, eabc9375.	5.8	37
76	Neuropathology of progressive supranuclear palsy after treatment with tilavonemab. <i>Lancet Neurology</i> , 2021, 20, 786-787.	4.9	9
77	Hematologic Emergencies in the Postoperative Neurointensive Care Unit Setting: Illustrative Case Series and Differential Diagnosis. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2021, 30, 106019.	0.7	1
78	Nanoparticles With Affinity for α -Synuclein Sequester α -Synuclein to Form Toxic Aggregates in Neurons With Endolysosomal Impairment. <i>Frontiers in Molecular Neuroscience</i> , 2021, 14, 738535.	1.4	2
79	TSC1 loss increases risk for tauopathy by inducing tau acetylation and preventing tau clearance via chaperone-mediated autophagy. <i>Science Advances</i> , 2021, 7, eabg3897.	4.7	27
80	Capgras syndrome in dementia with Lewy bodies: a possible association of severe cortical Lewy body pathology. <i>Neurologia i Neurochirurgia Polska</i> , 2021, , .	0.6	2
81	AD-linked R47H- <i>TREM2</i> mutation induces disease-enhancing microglial states via AKT hyperactivation. <i>Science Translational Medicine</i> , 2021, 13, eabe3947.	5.8	55
82	HDAC6 Interacts With Poly (GA) and Modulates its Accumulation in c9FTD/ALS. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 809942.	1.8	4
83	TDP-43 Pathology in Alzheimer's Disease. <i>Molecular Neurodegeneration</i> , 2021, 16, 84.	4.4	92
84	Neuropathology and molecular diagnosis of Synucleinopathies. <i>Molecular Neurodegeneration</i> , 2021, 16, 83.	4.4	101
85	Aberrant Accumulation of BRCA1 in Alzheimer Disease and Other Tauopathies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 22-33.	0.9	18
86	Association between contact sports participation and chronic traumatic encephalopathy: a retrospective cohort study. <i>Brain Pathology</i> , 2020, 30, 63-74.	2.1	66
87	4-Repeat tau seeds and templating subtypes as brain and CSF biomarkers of frontotemporal lobar degeneration. <i>Acta Neuropathologica</i> , 2020, 139, 63-77.	3.9	89
88	Selective Vulnerability of the Nucleus Basalis of Meynert Among Neuropathologic Subtypes of Alzheimer Disease. <i>JAMA Neurology</i> , 2020, 77, 225.	4.5	50
89	Tau-positron emission tomography correlates with neuropathology findings. <i>Alzheimer's and Dementia</i> , 2020, 16, 561-571.	0.4	113
90	β^2 -Amyloid PET and neuropathology in dementia with Lewy bodies. <i>Neurology</i> , 2020, 94, e282-e291.	1.5	65

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91	Novel monoclonal antibodies targeting the RRM2 domain of human TDP-43 protein. <i>Neuroscience Letters</i> , 2020, 738, 135353.	1.0	3
92	Deciphering cellular transcriptional alterations in Alzheimer's disease brains. <i>Molecular Neurodegeneration</i> , 2020, 15, 38.	4.4	42
93	Association of mitochondrial genomic background with risk of Multiple System Atrophy. <i>Parkinsonism and Related Disorders</i> , 2020, 81, 200-204.	1.1	4
94	Astrocyte-derived clusterin suppresses amyloid formation in vivo. <i>Molecular Neurodegeneration</i> , 2020, 15, 71.	4.4	26
95	Clusterin ameliorates tau pathology in vivo by inhibiting fibril formation. <i>Acta Neuropathologica Communications</i> , 2020, 8, 210.	2.4	24
96	Neuronal intranuclear inclusion disease is genetically heterogeneous. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 1716-1725.	1.7	38
97	GBA variation and susceptibility to multiple system atrophy. <i>Parkinsonism and Related Disorders</i> , 2020, 77, 64-69.	1.1	12
98	Letter to the editor, "Movement disorders rounds: A case of missing pathology in a patient with LRRK2 Parkinson's disease". <i>Parkinsonism and Related Disorders</i> , 2020, 79, 130.	1.1	0
99	Association of ABI3 and PLCG2 missense variants with disease risk and neuropathology in Lewy body disease and progressive supranuclear palsy. <i>Acta Neuropathologica Communications</i> , 2020, 8, 172.	2.4	8
100	Orthostatic hypotension preceding dementia with Lewy bodies by over 15 years: a clinicopathologic case report. <i>Clinical Autonomic Research</i> , 2020, 30, 575-577.	1.4	3
101	<i>C9orf72</i> poly(GR) aggregation induces TDP-43 proteinopathy. <i>Science Translational Medicine</i> , 2020, 12, .	5.8	115
102	Sensitivity and Specificity of Tau and Amyloid β Positron Emission Tomography in Frontotemporal Lobar Degeneration. <i>Annals of Neurology</i> , 2020, 88, 1009-1022.	2.8	32
103	Trans-synaptic and retrograde axonal spread of Lewy pathology following pre-formed fibril injection in an in vivo A53T alpha-synuclein mouse model of synucleinopathy. <i>Acta Neuropathologica Communications</i> , 2020, 8, 150.	2.4	36
104	Associations of mitochondrial genomic variation with corticobasal degeneration, progressive supranuclear palsy, and neuropathological tau measures. <i>Acta Neuropathologica Communications</i> , 2020, 8, 162.	2.4	9
105	Tau and apolipoprotein E modulate cerebrovascular tight junction integrity independent of cerebral amyloid angiopathy in Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2020, 16, 1372-1383.	0.4	34
106	MAPT subhaplotypes in corticobasal degeneration: assessing associations with disease risk, severity of tau pathology, and clinical features. <i>Acta Neuropathologica Communications</i> , 2020, 8, 218.	2.4	8
107	Protein contributions to brain atrophy acceleration in Alzheimer's disease and primary age-related tauopathy. <i>Brain</i> , 2020, 143, 3463-3476.	3.7	45
108	Cerebrovascular pathology and misdiagnosis of multiple system atrophy: An autopsy study. <i>Parkinsonism and Related Disorders</i> , 2020, 75, 34-40.	1.1	8

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109	Loss of homeostatic microglial phenotype in CSF1R-related Leukoencephalopathy. <i>Acta Neuropathologica Communications</i> , 2020, 8, 72.	2.4	42
110	Utility of FDG-PET in diagnosis of Alzheimer-related TDP-43 proteinopathy. <i>Neurology</i> , 2020, 95, e23-e34.	1.5	27
111	Crystal structure of a conformational antibody that binds tau oligomers and inhibits pathological seeding by extracts from donors with Alzheimer's disease. <i>Journal of Biological Chemistry</i> , 2020, 295, 10662-10676.	1.6	21
112	Loss of TMEM106B leads to myelination deficits: implications for frontotemporal dementia treatment strategies. <i>Brain</i> , 2020, 143, 1905-1919.	3.7	44
113	Clinical and pathologic features of cognitive-predominant corticobasal degeneration. <i>Neurology</i> , 2020, 95, e35-e45.	1.5	9
114	Subtypes of dementia with Lewy bodies are associated with $\hat{\alpha}$ -synuclein and tau distribution. <i>Neurology</i> , 2020, 95, e155-e165.	1.5	47
115	Confirmation of ¹²³ I-FP-CIT SPECT Quantification Methods in Dementia with Lewy Bodies and Other Neurodegenerative Disorders. <i>Journal of Nuclear Medicine</i> , 2020, 61, 1628-1635.	2.8	18
116	Association of <i>Tripartite Motif Containing 11</i> rs564309 With Tau Pathology in Progressive Supranuclear Palsy. <i>Movement Disorders</i> , 2020, 35, 890-894.	2.2	6
117	Microglial Homeostasis Requires Balanced CSF-1/CSF-2 Receptor Signaling. <i>Cell Reports</i> , 2020, 30, 3004-3019.e5.	2.9	53
118	Cathepsin D regulates cerebral A β ²⁴² /40 ratios via differential degradation of A β ²⁴² and A β ²⁴⁰ . <i>Alzheimer's Research and Therapy</i> , 2020, 12, 80.	3.0	36
119	¹⁸ F-fluorodeoxyglucose positron emission tomography in dementia with Lewy bodies. <i>Brain Communications</i> , 2020, 2, fcaa040.	1.5	17
120	APOE4 exacerbates $\hat{\alpha}$ -synuclein pathology and related toxicity independent of amyloid. <i>Science Translational Medicine</i> , 2020, 12, .	5.8	90
121	Generation and Characterization of Novel Monoclonal Antibodies Targeting $\hat{\alpha}$ p62/sequestosome-1 Across Human Neurodegenerative Diseases. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 407-418.	0.9	8
122	Brain volume and flortaucipir analysis of progressive supranuclear palsy clinical variants. <i>NeuroImage: Clinical</i> , 2020, 25, 102152.	1.4	46
123	Pathology of Proven Corticobasal Degeneration Presenting as Richardson's Syndrome. <i>Movement Disorders Clinical Practice</i> , 2020, 7, 267-272.	0.8	6
124	Effect Modifiers of TDP-43-Associated Hippocampal Atrophy Rates in Patients with Alzheimer's Disease Neuropathological Changes. <i>Journal of Alzheimer's Disease</i> , 2020, 73, 1511-1523.	1.2	14
125	TDP-43 is associated with a reduced likelihood of rendering a clinical diagnosis of dementia with Lewy bodies in autopsy-confirmed cases of transitional/diffuse Lewy body disease. <i>Journal of Neurology</i> , 2020, 267, 1444-1453.	1.8	4
126	Elevated methylation levels, reduced expression levels, and frequent contractions in a clinical cohort of C9orf72 expansion carriers. <i>Molecular Neurodegeneration</i> , 2020, 15, 7.	4.4	34

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127	Exceptionally low likelihood of Alzheimer's dementia in APOE2 homozygotes from a 5,000-person neuropathological study. <i>Nature Communications</i> , 2020, 11, 667.	5.8	246
128	Posttranslational Modifications Mediate the Structural Diversity of Tauopathy Strains. <i>Cell</i> , 2020, 180, 633-644.e12.	13.5	300
129	Large-scale proteomic analysis of Alzheimer's disease brain and cerebrospinal fluid reveals early changes in energy metabolism associated with microglia and astrocyte activation. <i>Nature Medicine</i> , 2020, 26, 769-780.	15.2	547
130	Clinicopathologic and genetic features of multiple system atrophy with Lewy body disease. <i>Brain Pathology</i> , 2020, 30, 766-778.	2.1	19
131	Association between transactive response DNA-binding protein of 43 kDa type and cognitive resilience to Alzheimer's disease: a case-control study. <i>Neurobiology of Aging</i> , 2020, 92, 92-97.	1.5	13
132	Analysis of τ -synuclein species enriched from cerebral cortex of humans with sporadic dementia with Lewy bodies. <i>Brain Communications</i> , 2020, 2, fcaa010.	1.5	21
133	Pick's disease: clinicopathologic characterization of 21 cases. <i>Journal of Neurology</i> , 2020, 267, 2697-2704.	1.8	17
134	LRP10 variants in progressive supranuclear palsy. <i>Neurobiology of Aging</i> , 2020, 94, 311.e5-311.e10.	1.5	6
135	Chimeric Peptide Species Contribute to Divergent Dipeptide Repeat Pathology in c9ALS/FTD and SCA36. <i>Neuron</i> , 2020, 107, 292-305.e6.	3.8	51
136	Truncated stathmin-2 is a marker of TDP-43 pathology in frontotemporal dementia. <i>Journal of Clinical Investigation</i> , 2020, 130, 6080-6092.	3.9	117
137	Analysis of neurodegenerative disease-causing genes in dementia with Lewy bodies. <i>Acta Neuropathologica Communications</i> , 2020, 8, 5.	2.4	27
138	Longitudinal anatomic, functional, and molecular characterization of Pick disease phenotypes. <i>Neurology</i> , 2020, 95, e3190-e3202.	1.5	13
139	Loss of Tmem106b exacerbates τ FTLD pathologies and causes motor deficits in progranulin-deficient mice. <i>EMBO Reports</i> , 2020, 21, e50197.	2.0	35
140	Neuronal intermediate filament inclusion disease may be incorrectly classified as a subtype of FTLD-FUS. <i>Free Neuropathology</i> , 2020, 1, .	2.4	0
141	"Minimal change" multiple system atrophy with limbic-predominant τ -synuclein pathology. <i>Acta Neuropathologica</i> , 2019, 137, 167-169.	3.9	11
142	Clinicopathologic subtype of Alzheimer's disease presenting as corticobasal syndrome. <i>Alzheimer's and Dementia</i> , 2019, 15, 1218-1228.	0.4	34
143	Dipeptide repeat (DPR) pathology in the skeletal muscle of ALS patients with C9ORF72 repeat expansion. <i>Acta Neuropathologica</i> , 2019, 138, 667-670.	3.9	32
144	Reply: LATE to the PART-y. <i>Brain</i> , 2019, 142, e48-e48.	3.7	11

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145	The neuropathological diagnosis of Alzheimer's disease. <i>Molecular Neurodegeneration</i> , 2019, 14, 32.	4.4	1,497
146	C9orf72 intermediate repeats are associated with corticobasal degeneration, increased C9orf72 expression and disruption of autophagy. <i>Acta Neuropathologica</i> , 2019, 138, 795-811.	3.9	50
147	Coexistence of Progressive Supranuclear Palsy With Pontocerebellar Atrophy and Myotonic Dystrophy Type 1. <i>Journal of Neuropathology and Experimental Neurology</i> , 2019, 78, 756-762.	0.9	3
148	C-terminal and full length TDP-43 specie differ according to FTLTDP lesion type but not genetic mutation. <i>Acta Neuropathologica Communications</i> , 2019, 7, 100.	2.4	11
149	Evaluation of Associations of Alzheimer's Disease Risk Variants that Are Highly Expressed in Microglia with Neuropathological Outcome Measures. <i>Journal of Alzheimer's Disease</i> , 2019, 70, 659-666.	1.2	6
150	Extensive transcriptomic study emphasizes importance of vesicular transport in C9orf72 expansion carriers. <i>Acta Neuropathologica Communications</i> , 2019, 7, 150.	2.4	40
151	Microglia in frontotemporal lobar degeneration with progranulin or C9ORF72 mutations. <i>Annals of Clinical and Translational Neurology</i> , 2019, 6, 1782-1796.	1.7	20
152	Structure-based inhibitors halt prion-like seeding by Alzheimer's disease and tauopathy derived brain tissue samples. <i>Journal of Biological Chemistry</i> , 2019, 294, 16451-16464.	1.6	51
153	Cerebrovascular pathology presenting as corticobasal syndrome: An autopsy case series of cerebrovascular CBS. <i>Parkinsonism and Related Disorders</i> , 2019, 68, 79-84.	1.1	14
154	Neuropathologic basis of frontotemporal dementia in progressive supranuclear palsy. <i>Movement Disorders</i> , 2019, 34, 1655-1662.	2.2	14
155	Progressive supranuclear palsy is not associated with neurogenic orthostatic hypotension. <i>Neurology</i> , 2019, 93, e1339-e1347.	1.5	16
156	PET-detectable tau pathology correlates with long-term neuropsychiatric outcomes in patients with traumatic brain injury. <i>Brain</i> , 2019, 142, 3265-3279.	3.7	54
157	Antemortem volume loss mirrors TDP-43 staging in older adults with non-frontotemporal lobar degeneration. <i>Brain</i> , 2019, 142, 3621-3635.	3.7	37
158	The influence of tau, amyloid, alpha-synuclein, TDP-43, and vascular pathology in clinically normal elderly individuals. <i>Neurobiology of Aging</i> , 2019, 77, 26-36.	1.5	51
159	Enhanced phosphorylation of T153 in soluble tau is a defining biochemical feature of the A152T tau risk variant. <i>Acta Neuropathologica Communications</i> , 2019, 7, 10.	2.4	3
160	A nonsynonymous mutation in PLCG2 reduces the risk of Alzheimer's disease, dementia with Lewy bodies and frontotemporal dementia, and increases the likelihood of longevity. <i>Acta Neuropathologica</i> , 2019, 138, 237-250.	3.9	87
161	Association of MAPT H1 subhaplotypes with neuropathology of lewy body disease. <i>Movement Disorders</i> , 2019, 34, 1325-1332.	2.2	15
162	CNS small vessel disease. <i>Neurology</i> , 2019, 92, 1146-1156.	1.5	343

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163	Disproportionately enlarged subarachnoid-space hydrocephalus (DESH) in normal pressure hydrocephalus misinterpreted as atrophy: autopsy and radiological evidence. <i>Neurocase</i> , 2019, 25, 151-155.	0.2	8
164	Neuroimaging correlates with neuropathologic schemes in neurodegenerative disease. <i>Alzheimer's and Dementia</i> , 2019, 15, 927-939.	0.4	48
165	Brain atrophy in primary age-related tauopathy is linked to transactive response DNA-binding protein of 43 kDa. <i>Alzheimer's and Dementia</i> , 2019, 15, 799-806.	0.4	14
166	Limbic-predominant age-related TDP-43 encephalopathy (LATE): consensus working group report. <i>Brain</i> , 2019, 142, 1503-1527.	3.7	873
167	X-Linked Lymphoproliferative Syndrome Presenting as Adult-Onset Multi-Infarct Dementia. <i>Journal of Neuropathology and Experimental Neurology</i> , 2019, 78, 460-466.	0.9	6
168	Association of <i>MAPT</i> Subhaplotypes With Risk of Progressive Supranuclear Palsy and Severity of Tau Pathology. <i>JAMA Neurology</i> , 2019, 76, 710.	4.5	39
169	Mixed Alzheimer's and Lewy-related Pathology Can Cause Corticobasal Syndrome with Visual Hallucinations. <i>Internal Medicine</i> , 2019, 58, 1813-1813.	0.3	0
170	Tau exhibits unique seeding properties in globular glial tauopathy. <i>Acta Neuropathologica Communications</i> , 2019, 7, 36.	2.4	28
171	ADAR2 mislocalization and widespread RNA editing aberrations in C9orf72-mediated ALS/FTD. <i>Acta Neuropathologica</i> , 2019, 138, 49-65.	3.9	48
172	In vivo binding of a tau imaging probe, [¹¹ C]PBB3, in patients with progressive supranuclear palsy. <i>Movement Disorders</i> , 2019, 34, 744-754.	2.2	36
173	Heritability and genetic variance of dementia with Lewy bodies. <i>Neurobiology of Disease</i> , 2019, 127, 492-501.	2.1	29
174	Ethnoracial differences in Alzheimer's disease from the FLorida Autopsied Multi-Ethnic (FLAME) cohort. <i>Alzheimer's and Dementia</i> , 2019, 15, 635-643.	0.4	29
175	Prominent auditory deficits in primary progressive aphasia: A case study. <i>Cortex</i> , 2019, 117, 396-406.	1.1	14
176	Sensitivity and Specificity of Diagnostic Criteria for Progressive Supranuclear Palsy. <i>Movement Disorders</i> , 2019, 34, 1144-1153.	2.2	98
177	Genome-wide analyses as part of the international FTLTDP whole-genome sequencing consortium reveals novel disease risk factors and increases support for immune dysfunction in FTLTDP. <i>Acta Neuropathologica</i> , 2019, 137, 879-899.	3.9	90
178	Selective loss of cortical endothelial tight junction proteins during Alzheimer's disease progression. <i>Brain</i> , 2019, 142, 1077-1092.	3.7	120
179	Aberrant deposition of stress granule-resident proteins linked to C9orf72-associated TDP-43 proteinopathy. <i>Molecular Neurodegeneration</i> , 2019, 14, 9.	4.4	111
180	Genetic meta-analysis of diagnosed Alzheimer's disease identifies new risk loci and implicates AÎ ² , tau, immunity and lipid processing. <i>Nature Genetics</i> , 2019, 51, 414-430.	9.4	1,962

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181	rAAV-based brain slice culture models of Alzheimer's and Parkinson's disease inclusion pathologies. <i>Journal of Experimental Medicine</i> , 2019, 216, 539-555.	4.2	48
182	Heterochromatin anomalies and double-stranded RNA accumulation underlie <i>C9orf72</i> poly(PR) toxicity. <i>Science</i> , 2019, 363, .	6.0	181
183	Subventricular glial nodules in neurofibromatosis 1 with craniofacial dysmorphism and occipital meningoencephalocele. <i>ENeurologicalSci</i> , 2019, 17, 100213.	0.5	1
184	Pathological, imaging and genetic characteristics support the existence of distinct TDP-43 types in non-FTLD brains. <i>Acta Neuropathologica</i> , 2019, 137, 227-238.	3.9	65
185	A comprehensive screening of copy number variability in dementia with Lewy bodies. <i>Neurobiology of Aging</i> , 2019, 75, 223.e1-223.e10.	1.5	13
186	APOE4-mediated amyloid- β^2 pathology depends on its neuronal receptor LRP1. <i>Journal of Clinical Investigation</i> , 2019, 129, 1272-1277.	3.9	96
187	Pathological analysis of ErbB family and NRG-1 protein in progressive supranuclear palsy. <i>The Journal of Kansai Medical University</i> , 2019, 70, 13-17.	0.3	0
188	Target-enriched sequencing of chromosome 17q21.31 in sporadic tauopathies reveals no candidate variants. <i>Neurobiology of Aging</i> , 2018, 66, 177.e7-177.e10.	1.5	1
189	Daytime sleepiness in dementia with Lewy bodies is associated with neuronal depletion of the nucleus basalis of Meynert. <i>Parkinsonism and Related Disorders</i> , 2018, 50, 99-103.	1.1	22
190	Multiple system atrophy and apolipoprotein E. <i>Movement Disorders</i> , 2018, 33, 647-650.	2.2	15
191	Mitotic defects lead to neuronal aneuploidy and apoptosis in frontotemporal lobar degeneration caused by MAPT mutations. <i>Molecular Biology of the Cell</i> , 2018, 29, 575-586.	0.9	36
192	Relationships between lewy and tau pathologies in 375 consecutive non-Alzheimer's olfactory bulbs. <i>Movement Disorders</i> , 2018, 33, 333-334.	2.2	1
193	TDP-43 pathology disrupts nuclear pore complexes and nucleocytoplasmic transport in ALS/FTD. <i>Nature Neuroscience</i> , 2018, 21, 228-239.	7.1	404
194	Potential genetic modifiers of disease risk and age at onset in patients with frontotemporal lobar degeneration and GRN mutations: a genome-wide association study. <i>Lancet Neurology</i> , The, 2018, 17, 548-558.	4.9	97
195	FDG-PET in tau-negative amnesic dementia resembles that of autopsy-proven hippocampal sclerosis. <i>Brain</i> , 2018, 141, 1201-1217.	3.7	67
196	Recent advances in neuropathology, biomarkers and therapeutic approach of multiple system atrophy. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018, 89, 175-184.	0.9	94
197	Establishing diagnostic criteria for Perry syndrome. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018, 89, 482-487.	0.9	40
198	Investigating the genetic architecture of dementia with Lewy bodies: a two-stage genome-wide association study. <i>Lancet Neurology</i> , The, 2018, 17, 64-74.	4.9	195

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199	Association Between Microinfarcts and Blood Pressure Trajectories. <i>JAMA Neurology</i> , 2018, 75, 212.	4.5	15
200	The limbic and neocortical contribution of α -synuclein, tau, and amyloid β to disease duration in dementia with Lewy bodies. <i>Alzheimer's and Dementia</i> , 2018, 14, 330-339.	0.4	69
201	Conserved brain myelination networks are altered in Alzheimer's and other neurodegenerative diseases. <i>Alzheimer's and Dementia</i> , 2018, 14, 352-366.	0.4	116
202	Parkinson's disease: experimental models and reality. <i>Acta Neuropathologica</i> , 2018, 135, 13-32.	3.9	89
203	Neuropathology of Parkinson disease. <i>Parkinsonism and Related Disorders</i> , 2018, 46, S30-S33.	1.1	363
204	Association study between multiple system atrophy and TREM2 p.R47H. <i>Neurology: Genetics</i> , 2018, 4, e257.	0.9	9
205	<i>CSF1R</i> -related leukoencephalopathy. <i>Neurology</i> , 2018, 91, 1092-1104.	1.5	126
206	Tangential Flow Filtration for Highly Efficient Concentration of Extracellular Vesicles from Large Volumes of Fluid. <i>Cells</i> , 2018, 7, 273.	1.8	262
207	A <i>C6orf10/LOC101929163</i> locus is associated with age of onset in <i>C9orf72</i> carriers. <i>Brain</i> , 2018, 141, 2895-2907.	3.7	39
208	Association of Apolipoprotein E ϵ 4 With Transactive Response DNA-Binding Protein 43. <i>JAMA Neurology</i> , 2018, 75, 1347.	4.5	60
209	APOE ϵ 2 is associated with increased tau pathology in primary tauopathy. <i>Nature Communications</i> , 2018, 9, 4388.	5.8	100
210	<i>ABI3</i> and <i>PLCG2</i> missense variants as risk factors for neurodegenerative diseases in Caucasians and African Americans. <i>Molecular Neurodegeneration</i> , 2018, 13, 53.	4.4	75
211	Sex and age interact to determine clinicopathologic differences in Alzheimer's disease. <i>Acta Neuropathologica</i> , 2018, 136, 873-885.	3.9	69
212	TLR5 decoy receptor as a novel anti-amyloid therapeutic for Alzheimer's disease. <i>Journal of Experimental Medicine</i> , 2018, 215, 2247-2264.	4.2	50
213	Age- and disease-dependent increase of the mitophagy marker phospho-ubiquitin in normal aging and Lewy body disease. <i>Autophagy</i> , 2018, 14, 1404-1418.	4.3	87
214	Poly(GR) impairs protein translation and stress granule dynamics in <i>C9orf72</i> -associated frontotemporal dementia and amyotrophic lateral sclerosis. <i>Nature Medicine</i> , 2018, 24, 1136-1142.	15.2	241
215	Loss of <i>Tmem106b</i> is unable to ameliorate frontotemporal dementia-like phenotypes in an AAV mouse model of <i>C9ORF72</i> -repeat induced toxicity. <i>Acta Neuropathologica Communications</i> , 2018, 6, 42.	2.4	20
216	Replication of progressive supranuclear palsy genome-wide association study identifies <i>SLCO1A2</i> and <i>DUSP10</i> as new susceptibility loci. <i>Molecular Neurodegeneration</i> , 2018, 13, 37.	4.4	54

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217	Epigenome-wide DNA methylation profiling in Progressive Supranuclear Palsy reveals major changes at DLX1. <i>Nature Communications</i> , 2018, 9, 2929.	5.8	20
218	Poly-GR dipeptide repeat polymers correlate with neurodegeneration and Clinicopathological subtypes in C9ORF72-related brain disease. <i>Acta Neuropathologica Communications</i> , 2018, 6, 63.	2.4	79
219	TDP-43 and Alzheimer's Disease Pathologic Subtype in Non-Amnesic Alzheimer's Disease Dementia. <i>Journal of Alzheimer's Disease</i> , 2018, 64, 1227-1233.	1.2	20
220	TMEM106B haplotypes have distinct gene expression patterns in aged brain. <i>Molecular Neurodegeneration</i> , 2018, 13, 35.	4.4	30
221	Dipeptide repeat proteins activate a heat shock response found in C9ORF72-ALS/FTLD patients. <i>Acta Neuropathologica Communications</i> , 2018, 6, 55.	2.4	24
222	Genome-wide pleiotropy analysis of neuropathological traits related to Alzheimer's disease. <i>Alzheimer's Research and Therapy</i> , 2018, 10, 22.	3.0	27
223	Divergent brain gene expression patterns associate with distinct cell-specific tau neuropathology traits in progressive supranuclear palsy. <i>Acta Neuropathologica</i> , 2018, 136, 709-727.	3.9	47
224	Long-read sequencing across the C9orf72 GGGGCC repeat expansion: implications for clinical use and genetic discovery efforts in human disease. <i>Molecular Neurodegeneration</i> , 2018, 13, 46.	4.4	111
225	APOE ϵ 4 is associated with severity of Lewy body pathology independent of Alzheimer pathology. <i>Neurology</i> , 2018, 91, e1182-e1195.	1.5	122
226	Identification and functional characterization of novel mutations including frameshift mutation in exon 4 of CSF1R in patients with adult-onset leukoencephalopathy with axonal spheroids and pigmented glia. <i>Journal of Neurology</i> , 2018, 265, 2415-2424.	1.8	23
227	Diffuse Lewy body disease manifesting as corticobasal syndrome. <i>Neurology</i> , 2018, 91, e268-e279.	1.5	37
228	Corticobasal degeneration with TDP-43 pathology presenting with progressive supranuclear palsy syndrome: a distinct clinicopathologic subtype. <i>Acta Neuropathologica</i> , 2018, 136, 389-404.	3.9	59
229	Duration and Pathologic Correlates of Lewy Body Disease. <i>JAMA Neurology</i> , 2017, 74, 310.	4.5	48
230	Pathology of Neurodegenerative Diseases. <i>Cold Spring Harbor Perspectives in Biology</i> , 2017, 9, a028035.	2.3	865
231	ABCA7 loss-of-function variants, expression, and neurologic disease risk. <i>Neurology: Genetics</i> , 2017, 3, e126.	0.9	26
232	Tau aggregation influences cognition and hippocampal atrophy in the absence of beta-amyloid: a clinico-imaging-pathological study of primary age-related tauopathy (PART). <i>Acta Neuropathologica</i> , 2017, 133, 705-715.	3.9	125
233	Frontotemporal dementia with the V337M MAPT mutation. <i>Neurology</i> , 2017, 88, 758-766.	1.5	76
234	Shared genetic risk between corticobasal degeneration, progressive supranuclear palsy, and frontotemporal dementia. <i>Acta Neuropathologica</i> , 2017, 133, 825-837.	3.9	90

#	ARTICLE	IF	CITATIONS
235	Systems biology approach to late-onset Alzheimer's disease genome-wide association study identifies novel candidate genes validated using brain expression data and <i>Caenorhabditis elegans</i> experiments. , 2017, 13, 1133-1142.		40
236	Î±-synuclein astrogliopathy: A possible specific feature in Î±-synucleinopathy. <i>Neuropathology</i> , 2017, 37, 379-381.	0.7	5
237	Transethnic genome-wide scan identifies novel Alzheimer's disease loci. <i>Alzheimer's and Dementia</i> , 2017, 13, 727-738.	0.4	166
238	Association Between Vascular Pathology and Rate of Cognitive Decline Independent of Alzheimer's Disease Pathology. <i>Journal of the American Geriatrics Society</i> , 2017, 65, 1836-1841.	1.3	15
239	Clinical diagnosis of progressive supranuclear palsy: The movement disorder society criteria. <i>Movement Disorders</i> , 2017, 32, 853-864.	2.2	1,402
240	Fluorescence and autoradiographic evaluation of tau PET ligand PBB3 to Î±-synuclein pathology. <i>Movement Disorders</i> , 2017, 32, 884-892.	2.2	55
241	In-depth clinico-pathological examination of RNA foci in a large cohort of C9ORF72 expansion carriers. <i>Acta Neuropathologica</i> , 2017, 134, 255-269.	3.9	76
242	DCTN1-related neurodegeneration: Perry syndrome and beyond. <i>Parkinsonism and Related Disorders</i> , 2017, 41, 14-24.	1.1	62
243	Diagnosis and management of dementia with Lewy bodies. <i>Neurology</i> , 2017, 89, 88-100.	1.5	2,805
244	White-matter integrity on DTI and the pathologic staging of Alzheimer's disease. <i>Neurobiology of Aging</i> , 2017, 56, 172-179.	1.5	158
245	Multisite Assessment of Aging-Related Tau Astrogliopathy (ARTAG). <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 605-619.	0.9	38
246	Increased cytoplasmic TDP-43 reduces global protein synthesis by interacting with RACK1 on polyribosomes. <i>Human Molecular Genetics</i> , 2017, 26, 1407-1418.	1.4	78
247	Poly(GP) proteins are a useful pharmacodynamic marker for <i>C9ORF72</i> -associated amyotrophic lateral sclerosis. <i>Science Translational Medicine</i> , 2017, 9, .	5.8	179
248	Distribution and characteristics of transactive response DNA binding protein 43 kDa pathology in progressive supranuclear palsy. <i>Movement Disorders</i> , 2017, 32, 246-255.	2.2	46
249	Brain tau deposition linked to systemic causes of death in normal elderly. <i>Neurobiology of Aging</i> , 2017, 50, 163-166.	1.5	2
250	Histones facilitate Î±-synuclein aggregation during neuronal apoptosis. <i>Acta Neuropathologica</i> , 2017, 133, 547-558.	3.9	20
251	A candidate regulatory variant at the <i>TREM</i> gene cluster associates with decreased Alzheimer's disease risk and increased <i>TREML1</i> and <i>TREM2</i> brain gene expression. <i>Alzheimer's and Dementia</i> , 2017, 13, 663-673.	0.4	48
252	Cognitive impairment in progressive supranuclear palsy is associated with tau burden. <i>Movement Disorders</i> , 2017, 32, 1772-1779.	2.2	46

#	ARTICLE	IF	CITATIONS
253	Regional analysis and genetic association of nigrostriatal degeneration in Lewy body disease. <i>Movement Disorders</i> , 2017, 32, 1584-1593.	2.2	15
254	Parkinson's disease susceptibility variants and severity of Lewy body pathology. <i>Parkinsonism and Related Disorders</i> , 2017, 44, 79-84.	1.1	17
255	DCTN1 variation in pathologically-confirmed PSP and CBD tauopathy. <i>Parkinsonism and Related Disorders</i> , 2017, 44, 151-153.	1.1	3
256	Impaired endo-lysosomal membrane integrity accelerates the seeding progression of α -synuclein aggregates. <i>Scientific Reports</i> , 2017, 7, 7690.	1.6	73
257	Rates of hippocampal atrophy and presence of post-mortem TDP-43 in patients with Alzheimer's disease: a longitudinal retrospective study. <i>Lancet Neurology</i> , The, 2017, 16, 917-924.	4.9	159
258	Reply re: "Profile of cognitive impairment and underlying pathology in multiple system atrophy". <i>Movement Disorders</i> , 2017, 32, 1339-1340.	2.2	3
259	Reduced orexin immunoreactivity in Perry syndrome and multiple system atrophy. <i>Parkinsonism and Related Disorders</i> , 2017, 42, 85-89.	1.1	9
260	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. <i>Nature Genetics</i> , 2017, 49, 1373-1384.	9.4	783
261	An acetylation-phosphorylation switch that regulates tau aggregation propensity and function. <i>Journal of Biological Chemistry</i> , 2017, 292, 15277-15286.	1.6	100
262	TIA1 Mutations in Amyotrophic Lateral Sclerosis and Frontotemporal Dementia Promote Phase Separation and Alter Stress Granule Dynamics. <i>Neuron</i> , 2017, 95, 808-816.e9.	3.8	493
263	Conserved DNA methylation combined with differential frontal cortex and cerebellar expression distinguishes C9orf72-associated and sporadic ALS, and implicates SERPINA1 in disease. <i>Acta Neuropathologica</i> , 2017, 134, 715-728.	3.9	40
264	Alzheimer's Disease-Related Dementias Summit 2016: National research priorities. <i>Neurology</i> , 2017, 89, 2381-2391.	1.5	109
265	Distinct binding of PET ligands PBB3 and AV-1451 to tau fibril strains in neurodegenerative tauopathies. <i>Brain</i> , 2017, 140, aww339.	3.7	153
266	Repetitive element transcripts are elevated in the brain of C9orf72 ALS/FTLD patients. <i>Human Molecular Genetics</i> , 2017, 26, 3421-3431.	1.4	101
267	Loss of clusterin shifts amyloid deposition to the cerebrovasculature via disruption of perivascular drainage pathways. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E6962-E6971.	3.3	96
268	Perry Syndrome: A Distinctive Type of TDP-43 Proteinopathy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 676-682.	0.9	50
269	Neonatal AAV delivery of alpha-synuclein induces pathology in the adult mouse brain. <i>Acta Neuropathologica Communications</i> , 2017, 5, 51.	2.4	24
270	Abnormal expression of homeobox genes and transthyretin in C9ORF72 expansion carriers. <i>Neurology: Genetics</i> , 2017, 3, e161.	0.9	12

#	ARTICLE	IF	CITATIONS
271	Profile of cognitive impairment and underlying pathology in multiple system atrophy. <i>Movement Disorders</i> , 2017, 32, 405-413.	2.2	95
272	Study of <i>LRRK2</i> variation in tauopathy: Progressive supranuclear palsy and corticobasal degeneration. <i>Movement Disorders</i> , 2017, 32, 115-123.	2.2	48
273	FTDP τ 17 with Pick body-like inclusions associated with a novel tau mutation, p.E372G. <i>Brain Pathology</i> , 2017, 27, 612-626.	2.1	11
274	An investigation of cerebrovascular lesions in dementia with Lewy bodies compared to Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2017, 13, 257-266.	0.4	41
275	Analysis of C9orf72 repeat expansions in a large international cohort of dementia with Lewy bodies. <i>Neurobiology of Aging</i> , 2017, 49, 214.e13-214.e15.	1.5	12
276	Distinct spatiotemporal accumulation of N-truncated and full-length amyloid- β 42 in Alzheimer's disease. <i>Brain</i> , 2017, 140, 3301-3316.	3.7	14
277	Brain calcifications and <i>PCDH12</i> variants. <i>Neurology: Genetics</i> , 2017, 3, e166.	0.9	15
278	The lysosomal protein cathepsin L is a progranulin protease. <i>Molecular Neurodegeneration</i> , 2017, 12, 55.	4.4	81
279	Linkage, whole genome sequence, and biological data implicate variants in <i>RAB10</i> in Alzheimer's disease resilience. <i>Genome Medicine</i> , 2017, 9, 100.	3.6	67
280	Clinical and neuropathological features of ALS/FTD with <i>TIA1</i> mutations. <i>Acta Neuropathologica Communications</i> , 2017, 5, 96.	2.4	38
281	p62 Pathology Model in the Rat Substantia Nigra with Filamentous Inclusions and Progressive Neurodegeneration. <i>PLoS ONE</i> , 2017, 12, e0169291.	1.1	15
282	Globular Glial Tauopathy Presenting as Semantic Variant Primary Progressive Aphasia. <i>JAMA Neurology</i> , 2016, 73, 123.	4.5	21
283	Cerebellar ataxia in progressive supranuclear palsy: An autopsy study of PSP τ C. <i>Movement Disorders</i> , 2016, 31, 653-662.	2.2	60
284	<i>MAPT</i> haplotype diversity in multiple system atrophy. <i>Parkinsonism and Related Disorders</i> , 2016, 30, 40-45.	1.1	23
285	Human whole genome genotype and transcriptome data for Alzheimer's and other neurodegenerative diseases. <i>Scientific Data</i> , 2016, 3, 160089.	2.4	361
286	Genetic risk factors for the posterior cortical atrophy variant of Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2016, 12, 862-871.	0.4	93
287	<i>RAB39B</i> gene mutations are not a common cause of Parkinson's disease or dementia with Lewy bodies. <i>Neurobiology of Aging</i> , 2016, 45, 107-108.	1.5	21
288	Tremor in progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2016, 27, 93-97.	1.1	17

#	ARTICLE	IF	CITATIONS
289	Impact of sex and APOE4 on cerebral amyloid angiopathy in Alzheimer's disease. <i>Acta Neuropathologica</i> , 2016, 132, 225-234.	3.9	73
290	Genetic modification of H2AX renders mesenchymal stromal cell-derived dopamine neurons more resistant to DNA damage and subsequent apoptosis. <i>Cytotherapy</i> , 2016, 18, 1483-1492.	0.3	7
291	Arguing against the proposed definition changes of PD. <i>Movement Disorders</i> , 2016, 31, 1619-1622.	2.2	55
292	TYROBP genetic variants in early-onset Alzheimer's disease. <i>Neurobiology of Aging</i> , 2016, 48, 222.e9-222.e15.	1.5	69
293	Spt4 selectively regulates the expression of <i>C9orf72</i> sense and antisense mutant transcripts. <i>Science</i> , 2016, 353, 708-712.	6.0	116
294	TREM2 p.R47H substitution is not associated with dementia with Lewy bodies. <i>Neurology: Genetics</i> , 2016, 2, e85.	0.9	16
295	An MRI-Based Atlas for Correlation of Imaging and Pathologic Findings in Alzheimer's Disease. <i>Journal of Neuroimaging</i> , 2016, 26, 264-268.	1.0	3
296	Adult-onset cerebello-brainstem dominant form of X-linked adrenoleukodystrophy presenting as multiple system atrophy: case report and literature review. <i>Neuropathology</i> , 2016, 36, 64-76.	0.7	25
297	Association of <i>GBA</i> Mutations and the E326K Polymorphism With Motor and Cognitive Progression in Parkinson Disease. <i>JAMA Neurology</i> , 2016, 73, 1217.	4.5	185
298	Chronic Traumatic Encephalopathy Pathology in Multiple System Atrophy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2016, 75, 963-970.	0.9	54
299	Cerebral peduncle angle: Unreliable in differentiating progressive supranuclear palsy from other neurodegenerative diseases. <i>Parkinsonism and Related Disorders</i> , 2016, 32, 31-35.	1.1	5
300	An autoradiographic evaluation of AV-1451 Tau PET in dementia. <i>Acta Neuropathologica Communications</i> , 2016, 4, 58.	2.4	388
301	Prosaposin is a regulator of progranulin levels and oligomerization. <i>Nature Communications</i> , 2016, 7, 11992.	5.8	68
302	LRRK2 variation and dementia with Lewy bodies. <i>Parkinsonism and Related Disorders</i> , 2016, 31, 98-103.	1.1	30
303	[18F]AV-1451 tau-PET uptake does correlate with quantitatively measured 4R-tau burden in autopsy-confirmed corticobasal degeneration. <i>Acta Neuropathologica</i> , 2016, 132, 931-933.	3.9	116
304	A large-scale comparison of cortical thickness and volume methods for measuring Alzheimer's disease severity. <i>NeuroImage: Clinical</i> , 2016, 11, 802-812.	1.4	249
305	Juvenile onset Parkinsonism with <i>œpure nigral</i> degeneration and POLG1 mutation. <i>Parkinsonism and Related Disorders</i> , 2016, 30, 83-85.	1.1	9
306	CCNF mutations in amyotrophic lateral sclerosis and frontotemporal dementia. <i>Nature Communications</i> , 2016, 7, 11253.	5.8	174

#	ARTICLE	IF	CITATIONS
307	Expression and processing analyses of wild type and p.R47H TREM2 variant in Alzheimer's disease brains. <i>Molecular Neurodegeneration</i> , 2016, 11, 72.	4.4	55
308	microRNA profiling: increased expression of miR-147a and miR-518e in progressive supranuclear palsy (PSP). <i>Neurogenetics</i> , 2016, 17, 165-171.	0.7	20
309	<i>MAPT</i> haplotype H1G is associated with increased risk of dementia with Lewy bodies. <i>Alzheimer's and Dementia</i> , 2016, 12, 1297-1304.	0.4	32
310	Predicting Survival in Dementia With Lewy Bodies With Hippocampal Volumetry. <i>Movement Disorders</i> , 2016, 31, 989-994.	2.2	32
311	Neuropathologic differences by race from the National Alzheimer's Coordinating Center. <i>Alzheimer's and Dementia</i> , 2016, 12, 669-677.	0.4	75
312	Updated TDP-43 in Alzheimer's disease staging scheme. <i>Acta Neuropathologica</i> , 2016, 131, 571-585.	3.9	244
313	Genetic Disorders with Tau Pathology: A Review of the Literature and Report of Two Patients with Tauopathy and Positive Family Histories. <i>Neurodegenerative Diseases</i> , 2016, 16, 12-21.	0.8	35
314	Propagation of tau pathology: hypotheses, discoveries, and yet unresolved questions from experimental and human brain studies. <i>Acta Neuropathologica</i> , 2016, 131, 27-48.	3.9	147
315	TDP-43 functions within a network of hnRNP proteins to inhibit the production of a truncated human SORT1 receptor. <i>Human Molecular Genetics</i> , 2016, 25, 534-545.	1.4	70
316	Aging-related tau astroglialopathy (ARTAG): harmonized evaluation strategy. <i>Acta Neuropathologica</i> , 2016, 131, 87-102.	3.9	380
317	C9orf72 poly(GA) aggregates sequester and impair HR23 and nucleocytoplasmic transport proteins. <i>Nature Neuroscience</i> , 2016, 19, 668-677.	7.1	268
318	Proaggregant nuclear factor(s) trigger rapid formation of α -synuclein aggregates in apoptotic neurons. <i>Acta Neuropathologica</i> , 2016, 132, 77-91.	3.9	27
319	Plasma sphingolipid changes with autopsy-confirmed Lewy body or Alzheimer's pathology. <i>Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2016, 3, 43-50.	1.2	44
320	The first NINDS/NIBIB consensus meeting to define neuropathological criteria for the diagnosis of chronic traumatic encephalopathy. <i>Acta Neuropathologica</i> , 2016, 131, 75-86.	3.9	708
321	C9orf72 promoter hypermethylation is reduced while hydroxymethylation is acquired during reprogramming of ALS patient cells. <i>Experimental Neurology</i> , 2016, 277, 171-177.	2.0	21
322	Mixed tau and TDP-43 pathology in a patient with unclassifiable primary progressive aphasia. <i>Neurocase</i> , 2016, 22, 55-59.	0.2	11
323	Evaluating pathogenic dementia variants in posterior cortical atrophy. <i>Neurobiology of Aging</i> , 2016, 37, 38-44.	1.5	23
324	Tissue Transglutaminase and Its Product Isopeptide Are Increased in Alzheimer's Disease and APP ^{swe} /PS1 ^{dE9} Double Transgenic Mice Brains. <i>Molecular Neurobiology</i> , 2016, 53, 5066-5078.	1.9	31

#	ARTICLE	IF	CITATIONS
325	The presenilin 1 p.Gly206Ala mutation is a frequent cause of early-onset Alzheimer's disease in Hispanics in Florida. <i>American Journal of Neurodegenerative Disease</i> , 2016, 5, 94-101.	0.1	4
326	Transmission of Soluble and Insoluble α -Synuclein to Mice. <i>Journal of Neuro pathology and Experimental Neurology</i> , 2015, 74, 1158-1169.	0.9	14
327	(Patho)physiological relevance of α -PINK-dependent ubiquitin phosphorylation. <i>EMBO Reports</i> , 2015, 16, 1114-1130.	2.0	147
328	Genetically-controlled Vesicle-Associated Membrane Protein 1 expression may contribute to Alzheimer's pathophysiology and susceptibility. <i>Molecular Neurodegeneration</i> , 2015, 10, 18.	4.4	13
329	TAR DNA-binding protein 43 and pathological subtype of Alzheimer's disease impact clinical features. <i>Annals of Neurology</i> , 2015, 78, 697-709.	2.8	96
330	Frontotemporal dementia-associated N279K tau mutant disrupts subcellular vesicle trafficking and induces cellular stress in iPSC-derived neural stem cells. <i>Molecular Neurodegeneration</i> , 2015, 10, 46.	4.4	58
331	Jump from Pre-mutation to Pathologic Expansion in C9orf72. <i>American Journal of Human Genetics</i> , 2015, 96, 962-970.	2.6	50
332	Clinical Correlations With Lewy Body Pathology in <i>LRRK2</i> -Related Parkinson Disease. <i>JAMA Neurology</i> , 2015, 72, 100.	4.5	272
333	Rarity of the Alzheimer Disease "Protective <i>APP</i> A673T Variant in the United States. <i>JAMA Neurology</i> , 2015, 72, 209.	4.5	41
334	CHCHD2 and Parkinson's disease. <i>Lancet Neurology</i> , The, 2015, 14, 679.	4.9	16
335	A novel tau mutation, p.K317N, causes globular glial tauopathy. <i>Acta Neuropathologica</i> , 2015, 130, 199-214.	3.9	38
336	Mitochondrial targeting sequence variants of the <i>CHCHD2</i> gene are a risk for Lewy body disorders. <i>Neurology</i> , 2015, 85, 2016-2025.	1.5	51
337	Tau deposition drives neuropathological, inflammatory and behavioral abnormalities independently of neuronal loss in a novel mouse model. <i>Human Molecular Genetics</i> , 2015, 24, 6198-6212.	1.4	52
338	A Novel Tau Mutation in Exon 12, p.Q336H, Causes Hereditary Pick Disease. <i>Journal of Neuro pathology and Experimental Neurology</i> , 2015, 74, 1042-1052.	0.9	27
339	Chronic traumatic encephalopathy pathology in a neurodegenerative disorders brain bank. <i>Acta Neuropathologica</i> , 2015, 130, 877-889.	3.9	235
340	Three sib-pairs of autopsy-confirmed progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2015, 21, 101-105.	1.1	42
341	<i>PARK10</i> is a major locus for sporadic neuropathologically confirmed Parkinson disease. <i>Neurology</i> , 2015, 84, 972-980.	1.5	48
342	Clinical presentation of a patient with SLC20A2 and THAP1 deletions: Differential diagnosis of oromandibular dystonia. <i>Parkinsonism and Related Disorders</i> , 2015, 21, 329-331.	1.1	5

#	ARTICLE	IF	CITATIONS
343	Clinicopathologic and ¹¹ C-Pittsburgh compound B implications of Thal amyloid phase across the Alzheimer's disease spectrum. <i>Brain</i> , 2015, 138, 1370-1381.	3.7	270
344	A truncating SOD1 mutation, p.Gly141X, is associated with clinical and pathologic heterogeneity, including frontotemporal lobar degeneration. <i>Acta Neuropathologica</i> , 2015, 130, 145-157.	3.9	24
345	Genome-wide association study of corticobasal degeneration identifies risk variants shared with progressive supranuclear palsy. <i>Nature Communications</i> , 2015, 6, 7247.	5.8	170
346	When DLB, PD, and PSP masquerade as MSA. <i>Neurology</i> , 2015, 85, 404-412.	1.5	272
347	Distinct brain transcriptome profiles in C9orf72-associated and sporadic ALS. <i>Nature Neuroscience</i> , 2015, 18, 1175-1182.	7.1	330
348	Predicting amyloid status in corticobasal syndrome using modified clinical criteria, magnetic resonance imaging and fluorodeoxyglucose positron emission tomography. <i>Alzheimer's Research and Therapy</i> , 2015, 7, 8.	3.0	32
349	<i>C9ORF72</i> repeat expansions in mice cause TDP-43 pathology, neuronal loss, and behavioral deficits. <i>Science</i> , 2015, 348, 1151-1154.	6.0	332
350	PRKAR1B mutations are a rare cause of FUS negative neuronal intermediate filament inclusion disease. <i>Brain</i> , 2015, 138, e357-e357.	3.7	9
351	Atypical multiple system atrophy is a new subtype of frontotemporal lobar degeneration: frontotemporal lobar degeneration associated with τ -synuclein. <i>Acta Neuropathologica</i> , 2015, 130, 93-105.	3.9	65
352	PART, a distinct tauopathy, different from classical sporadic Alzheimer disease. <i>Acta Neuropathologica</i> , 2015, 129, 757-762.	3.9	139
353	ALS/FTD Mutation-Induced Phase Transition of FUS Liquid Droplets and Reversible Hydrogels into Irreversible Hydrogels Impairs RNP Granule Function. <i>Neuron</i> , 2015, 88, 678-690.	3.8	716
354	Novel clinical associations with specific C9ORF72 transcripts in patients with repeat expansions in C9ORF72. <i>Acta Neuropathologica</i> , 2015, 130, 863-876.	3.9	104
355	Intraneuronal amyloid- β^2 accumulation in basal forebrain cholinergic neurons: a marker of vulnerability, yet inversely related to neurodegeneration. <i>Brain</i> , 2015, 138, 1444-1445.	3.7	3
356	Concurrent neurodegenerative pathologies in periventricular nodular heterotopia. <i>Acta Neuropathologica</i> , 2015, 130, 895-897.	3.9	5
357	The <i>TMEM106B</i> locus and TDP-43 pathology in older persons without FTLD. <i>Neurology</i> , 2015, 85, 1354-1355.	1.5	14
358	Late-onset Alzheimer disease risk variants mark brain regulatory loci. <i>Neurology: Genetics</i> , 2015, 1, e15.	0.9	64
359	Cerebellar c9RAN proteins associate with clinical and neuropathological characteristics of C9ORF72 repeat expansion carriers. <i>Acta Neuropathologica</i> , 2015, 130, 559-573.	3.9	89
360	Role for the microtubule-associated protein tau variant p.A152T in risk of τ -synucleinopathies. <i>Neurology</i> , 2015, 85, 1680-1686.	1.5	31

#	ARTICLE	IF	CITATIONS
361	Mitochondrial ATP synthase activity is impaired by suppressed <i>O</i> -GlcNAcylation in Alzheimer's disease. <i>Human Molecular Genetics</i> , 2015, 24, 6492-6504.	1.4	74
362	C9orf72 BAC Transgenic Mice Display Typical Pathologic Features of ALS/FTD. <i>Neuron</i> , 2015, 88, 892-901.	3.8	249
363	Pattern of brain atrophy rates in autopsy-confirmed dementia with Lewy bodies. <i>Neurobiology of Aging</i> , 2015, 36, 452-461.	1.5	113
364	Hippocampal sclerosis in Lewy body disease is a TDP-43 proteinopathy similar to FTLTDP Type A. <i>Acta Neuropathologica</i> , 2015, 129, 53-64.	3.9	67
365	Exonic Re-Sequencing of the Chromosome 2q24.3 Parkinson's Disease Locus. <i>PLoS ONE</i> , 2015, 10, e0128586.	1.1	0
366	Relationships between typical histopathological hallmarks and the ferritin in the hippocampus from patients with Alzheimer's disease. <i>Acta Neurobiologiae Experimentalis</i> , 2015, 75, 391-8.	0.4	15
367	ER-mitochondria associations are regulated by the VAPB-PTPIP51 interaction and are disrupted by ALS/FTD-associated TDP-43. <i>Nature Communications</i> , 2014, 5, 3996.	5.8	463
368	Abnormal daytime sleepiness in dementia with Lewy bodies compared to Alzheimer's disease using the Multiple Sleep Latency Test. <i>Alzheimer's Research and Therapy</i> , 2014, 6, 76.	3.0	45
369	FUS is Phosphorylated by DNA-PK and Accumulates in the Cytoplasm after DNA Damage. <i>Journal of Neuroscience</i> , 2014, 34, 7802-7813.	1.7	129
370	Genetic analysis implicates APOE, SNCA and suggests lysosomal dysfunction in the etiology of dementia with Lewy bodies. <i>Human Molecular Genetics</i> , 2014, 23, 6139-6146.	1.4	178
371	Antemortem MRI findings associated with microinfarcts at autopsy. <i>Neurology</i> , 2014, 82, 1951-1958.	1.5	45
372	<i>LRRK2</i> exonic variants and risk of multiple system atrophy. <i>Neurology</i> , 2014, 83, 2256-2261.	1.5	46
373	Genome-Wide Association Meta-analysis of Neuropathologic Features of Alzheimer's Disease and Related Dementias. <i>PLoS Genetics</i> , 2014, 10, e1004606.	1.5	305
374	Quantitative characterization of brain β -amyloid using a joint PiB/FDG PET image histogram. , 2014, . .		0
375	Alterations in microRNA-124 and AMPA receptors contribute to social behavioral deficits in frontotemporal dementia. <i>Nature Medicine</i> , 2014, 20, 1444-1451.	15.2	165
376	Early Alzheimer's Disease Neuropathology Detected by Proton MR Spectroscopy. <i>Journal of Neuroscience</i> , 2014, 34, 16247-16255.	1.7	117
377	Aggregation-prone c9FTD/ALS poly(GA) RAN-translated proteins cause neurotoxicity by inducing ER stress. <i>Acta Neuropathologica</i> , 2014, 128, 505-524.	3.9	284
378	Genetic modifiers in carriers of repeat expansions in the C9ORF72 gene. <i>Molecular Neurodegeneration</i> , 2014, 9, 38.	4.4	63

#	ARTICLE	IF	CITATIONS
379	Association of MAPT haplotypes with Alzheimer's disease risk and MAPT brain gene expression levels. <i>Alzheimer's Research and Therapy</i> , 2014, 6, 39.	3.0	106
380	Effects of Multiple Genetic Loci on Age at Onset in Late-Onset Alzheimer Disease. <i>JAMA Neurology</i> , 2014, 71, 1394.	4.5	166
381	Expanded C9ORF72 Hexanucleotide Repeat in Depressive Pseudodementia. <i>JAMA Neurology</i> , 2014, 71, 775.	4.5	28
382	Dementia with Lewy bodies. <i>Neurology</i> , 2014, 83, 801-809.	1.5	143
383	Familial Progressive Supranuclear Palsy: A Literature Review. <i>Neurodegenerative Diseases</i> , 2014, 13, 180-182.	0.8	14
384	Regional proton magnetic resonance spectroscopy patterns in dementia with Lewy bodies. <i>Neurobiology of Aging</i> , 2014, 35, 1483-1490.	1.5	29
385	Effects of the C57BL/6 strain background on tauopathy progression in the rTg4510 mouse model. <i>Molecular Neurodegeneration</i> , 2014, 9, 8.	4.4	25
386	Novel mutation in MAPT exon 13 (p.N410H) causes corticobasal degeneration. <i>Acta Neuropathologica</i> , 2014, 127, 271-282.	3.9	66
387	TMEM106B protects C9ORF72 expansion carriers against frontotemporal dementia. <i>Acta Neuropathologica</i> , 2014, 127, 397-406.	3.9	133
388	Characterization of DNA hypermethylation in the cerebellum of c9FTD/ALS patients. <i>Brain Research</i> , 2014, 1584, 15-21.	1.1	70
389	Ribosomal Protein s15 Phosphorylation Mediates LRRK2 Neurodegeneration in Parkinson's Disease. <i>Cell</i> , 2014, 157, 472-485.	13.5	239
390	Severe amygdala dysfunction in a MAPT transgenic mouse model of frontotemporal dementia. <i>Neurobiology of Aging</i> , 2014, 35, 1769-1777.	1.5	48
391	SLC20A2 and THAP1 deletion in familial basal ganglia calcification with dystonia. <i>Neurogenetics</i> , 2014, 15, 23-30.	0.7	56
392	Convergence of pathology in dementia with Lewy bodies and Alzheimer's disease: a role for the novel interaction of alpha-synuclein and presenilin 1 in disease. <i>Brain</i> , 2014, 137, 1958-1970.	3.7	44
393	Genetic Screening and Functional Characterization of PDGFRB Mutations Associated with Basal Ganglia Calcification of Unknown Etiology. <i>Human Mutation</i> , 2014, 35, 964-971.	1.1	45
394	Ataxin-2 as potential disease modifier in C9ORF72 expansion carriers. <i>Neurobiology of Aging</i> , 2014, 35, 2421.e13-2421.e17.	1.5	74
395	Late-onset Alzheimer disease genetic variants in posterior cortical atrophy and posterior AD. <i>Neurology</i> , 2014, 82, 1455-1462.	1.5	51
396	Staging TDP-43 pathology in Alzheimer's disease. <i>Acta Neuropathologica</i> , 2014, 127, 441-450.	3.9	278

#	ARTICLE	IF	CITATIONS
397	A familial form of parkinsonism, dementia, and motor neuron disease: A longitudinal study. <i>Parkinsonism and Related Disorders</i> , 2014, 20, 1129-1134.	1.1	6
398	Primary age-related tauopathy (PART): a common pathology associated with human aging. <i>Acta Neuropathologica</i> , 2014, 128, 755-766.	3.9	1,060
399	Recommendations of the Alzheimer's Disease-Related Dementias Conference. <i>Neurology</i> , 2014, 83, 851-860.	1.5	103
400	Clinicopathologic assessment and imaging of tauopathies in neurodegenerative dementias. <i>Alzheimer's Research and Therapy</i> , 2014, 6, 1.	3.0	156
401	Is pathological aging a successful resistance against amyloid-beta or preclinical Alzheimer's disease?. <i>Alzheimer's Research and Therapy</i> , 2014, 6, 24.	3.0	35
402	FDG-PET in pathologically confirmed spontaneous 4R-tauopathy variants. <i>Journal of Neurology</i> , 2014, 261, 710-716.	1.8	60
403	TDP-43 is a key player in the clinical features associated with Alzheimer's disease. <i>Acta Neuropathologica</i> , 2014, 127, 811-824.	3.9	336
404	Concurrent variably protease-sensitive prionopathy and amyotrophic lateral sclerosis. <i>Acta Neuropathologica</i> , 2014, 128, 313-315.	3.9	9
405	TDP-43 in Alzheimer's disease is not associated with clinical FTD or Parkinsonism. <i>Journal of Neurology</i> , 2014, 261, 1344-1348.	1.8	22
406	Frontotemporal dementia and its subtypes: a genome-wide association study. <i>Lancet Neurology</i> , The, 2014, 13, 686-699.	4.9	302
407	Regional distribution of synaptic markers and APP correlate with distinct clinicopathological features in sporadic and familial Alzheimer's disease. <i>Brain</i> , 2014, 137, 1533-1549.	3.7	100
408	Age-related decline in white matter integrity in a mouse model of tauopathy: an <i>in vivo</i> diffusion tensor magnetic resonance imaging study. <i>Neurobiology of Aging</i> , 2014, 35, 1364-1374.	1.5	58
409	Absence of <i>C9ORF72</i> expanded or intermediate repeats in autopsy-confirmed Parkinson's disease. <i>Movement Disorders</i> , 2014, 29, 827-830.	2.2	24
410	Differential clinicopathologic and genetic features of late-onset amnesic dementias. <i>Acta Neuropathologica</i> , 2014, 128, 411-421.	3.9	119
411	Genome-wide association interaction analysis for Alzheimer's disease. <i>Neurobiology of Aging</i> , 2014, 35, 2436-2443.	1.5	61
412	ApoE variant p.V236E is associated with markedly reduced risk of Alzheimer's disease. <i>Molecular Neurodegeneration</i> , 2014, 9, 11.	4.4	57
413	Clinical, positron emission tomography, and pathological studies of DNAJC13 p.N855S Parkinsonism. <i>Movement Disorders</i> , 2014, 29, 1684-1687.	2.2	20
414	Divergent Phenotypes in Mutant TDP-43 Transgenic Mice Highlight Potential Confounds in TDP-43 Transgenic Modeling. <i>PLoS ONE</i> , 2014, 9, e86513.	1.1	23

#	ARTICLE	IF	CITATIONS
415	TREM2 in neurodegeneration: evidence for association of the p.R47H variant with frontotemporal dementia and Parkinson's disease. <i>Molecular Neurodegeneration</i> , 2013, 8, 19.	4.4	323
416	Normal cognition in transgenic BRI2- Δ^2 mice. <i>Molecular Neurodegeneration</i> , 2013, 8, 15.	4.4	74
417	Population-specific frequencies for <i>LRRK2</i> susceptibility variants in the genetic epidemiology of Parkinson's disease (GEO-PD) consortium. <i>Movement Disorders</i> , 2013, 28, 1740-1744.	2.2	30
418	Robust cytoplasmic accumulation of phosphorylated TDP-43 in transgenic models of tauopathy. <i>Acta Neuropathologica</i> , 2013, 126, 39-50.	3.9	24
419	LRRK2 phosphorylates novel tau epitopes and promotes tauopathy. <i>Acta Neuropathologica</i> , 2013, 126, 809-827.	3.9	85
420	Globular glial tauopathies (GGT): consensus recommendations. <i>Acta Neuropathologica</i> , 2013, 126, 537-544.	3.9	168
421	Nonamnesic mild cognitive impairment progresses to dementia with Lewy bodies. <i>Neurology</i> , 2013, 81, 2032-2038.	1.5	191
422	Association between repeat sizes and clinical and pathological characteristics in carriers of C9ORF72 repeat expansions (Xpansize-72): a cross-sectional cohort study. <i>Lancet Neurology</i> , The, 2013, 12, 978-988.	4.9	232
423	Endogenous Tau Aggregates in Oligodendrocytes of rTg4510 Mice Induced by Human P301L Tau. <i>Journal of Alzheimer's Disease</i> , 2013, 38, 589-600.	1.2	11
424	Novel A18T and pA29S substitutions in α -synuclein may be associated with sporadic Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 1057-1060.	1.1	63
425	MRI and pathology of REM sleep behavior disorder in dementia with Lewy bodies. <i>Neurology</i> , 2013, 81, 1681-1689.	1.5	58
426	Frontal asymmetry in behavioral variant frontotemporal dementia: clinicoimaging and pathogenetic correlates. <i>Neurobiology of Aging</i> , 2013, 34, 636-639.	1.5	54
427	Tau pathology in frontotemporal lobar degeneration with C9ORF72 hexanucleotide repeat expansion. <i>Acta Neuropathologica</i> , 2013, 125, 289-302.	3.9	87
428	Unconventional Translation of C9ORF72 GGGGCC Expansion Generates Insoluble Polypeptides Specific to c9FTD/ALS. <i>Neuron</i> , 2013, 77, 639-646.	3.8	962
429	Neurocognitive speed associates with frontotemporal lobar degeneration TDP-43 subtypes. <i>Journal of Clinical Neuroscience</i> , 2013, 20, 1737-1741.	0.8	1
430	Pallidonigrolyusian atrophy associated with p.A152T variant in MAPT. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 838-841.	1.1	9
431	Parkinsonian features in hereditary diffuse leukoencephalopathy with spheroids (HDLS) and CSF1R mutations. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 869-877.	1.1	119
432	Mutations in protein N-arginine methyltransferases are not the cause of FTL-D-FUS. <i>Neurobiology of Aging</i> , 2013, 34, 2235.e11-2235.e13.	1.5	13

#	ARTICLE	IF	CITATIONS
433	Quantitative neurofibrillary tangle density and brain volumetric MRI analyses in Alzheimer's disease presenting as logopenic progressive aphasia. <i>Brain and Language</i> , 2013, 127, 127-134.	0.8	53
434	TARDBP mutations in Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2013, 19, 312-315.	1.1	49
435	Sequence variants in eukaryotic translation initiation factor 4-gamma (eIF4G1) are associated with Lewy body dementia. <i>Acta Neuropathologica</i> , 2013, 125, 425-438.	3.9	20
436	Corticobasal degeneration with olivopontocerebellar atrophy and TDP-43 pathology: an unusual clinicopathologic variant of CBD. <i>Acta Neuropathologica</i> , 2013, 125, 741-752.	3.9	40
437	Criteria for the diagnosis of corticobasal degeneration. <i>Neurology</i> , 2013, 80, 496-503.	1.5	1,445
438	Adenosine monophosphate-activated protein kinase overactivation leads to accumulation of α -synuclein oligomers and decrease of neurites. <i>Neurobiology of Aging</i> , 2013, 34, 1504-1515.	1.5	82
439	Brain regional correlation of amyloid- β with synapses and apolipoprotein E in non-demented individuals: potential mechanisms underlying regional vulnerability to amyloid- β accumulation. <i>Acta Neuropathologica</i> , 2013, 125, 535-547.	3.9	51
440	Antisense transcripts of the expanded C9ORF72 hexanucleotide repeat form nuclear RNA foci and undergo repeat-associated non-ATG translation in c9FTD/ALS. <i>Acta Neuropathologica</i> , 2013, 126, 829-844.	3.9	506
441	Reduced C9orf72 gene expression in c9FTD/ALS is caused by histone trimethylation, an epigenetic event detectable in blood. <i>Acta Neuropathologica</i> , 2013, 126, 895-905.	3.9	263
442	Progressive amnesic dementia, hippocampal sclerosis, and mutation in C9ORF72. <i>Acta Neuropathologica</i> , 2013, 126, 545-554.	3.9	30
443	Corticospinal tract degeneration associated with TDP-43 type C pathology and semantic dementia. <i>Brain</i> , 2013, 136, 455-470.	3.7	37
444	C9ORF72 repeat expansions in cases with previously identified pathogenic mutations. <i>Neurology</i> , 2013, 81, 1332-1341.	1.5	84
445	Progressive Supranuclear Palsy: High-Field-Strength MR Microscopy in the Human Substantia Nigra and Globus Pallidus. <i>Radiology</i> , 2013, 266, 280-288.	3.6	26
446	Similarities between familial and sporadic autopsy-proven progressive supranuclear palsy. <i>Neurology</i> , 2013, 80, 2076-2078.	1.5	31
447	CSF1R mutations link POLD and HDLS as a single disease entity. <i>Neurology</i> , 2013, 80, 1033-1040.	1.5	136
448	Clinical and electrophysiologic variability in amyotrophic lateral sclerosis within a kindred harboring the C9ORF72 repeat expansion. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2013, 14, 132-137.	1.1	7
449	Atp13a2-deficient mice exhibit neuronal ceroid lipofuscinosis, limited α -synuclein accumulation and age-dependent sensorimotor deficits. <i>Human Molecular Genetics</i> , 2013, 22, 2067-2082.	1.4	124
450	Profilin-1 mutations are rare in patients with amyotrophic lateral sclerosis and frontotemporal dementia. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2013, 14, 463-469.	1.1	25

#	ARTICLE	IF	CITATIONS
451	Risk factors for dementia with Lewy bodies. <i>Neurology</i> , 2013, 81, 833-840.	1.5	136
452	Clinicopathologic variability of the <i>GRN</i> A9D mutation, including amyotrophic lateral sclerosis. <i>Neurology</i> , 2013, 80, 1771-1777.	1.5	24
453	Polysomnographic Findings in Dementia With Lewy Bodies. <i>Neurologist</i> , 2013, 19, 1-6.	0.4	75
454	The ALS disease-associated mutant TDP-43 impairs mitochondrial dynamics and function in motor neurons. <i>Human Molecular Genetics</i> , 2013, 22, 4706-4719.	1.4	251
455	<i>TMEM106B</i> p.T185S regulates <i>TMEM106B</i> protein levels: implications for frontotemporal dementia. <i>Journal of Neurochemistry</i> , 2013, 126, 781-791.	2.1	87
456	Linking Protective GAB2 Variants, Increased Cortical GAB2 Expression and Decreased Alzheimer's Disease Pathology. <i>PLoS ONE</i> , 2013, 8, e64802.	1.1	13
457	Neuropathology of parkinsonism. , 2013, , 239-257.		0
458	Diversity of pathological features other than Lewy bodies in familial Parkinson's disease due to SNCA mutations. <i>American Journal of Neurodegenerative Disease</i> , 2013, 2, 266-75.	0.1	19
459	Brain Expression Genome-Wide Association Study (eGWAS) Identifies Human Disease-Associated Variants. <i>PLoS Genetics</i> , 2012, 8, e1002707.	1.5	225
460	Neuroimaging signatures of frontotemporal dementia genetics: C9ORF72, tau, progranulin and sporadics. <i>Brain</i> , 2012, 135, 794-806.	3.7	355
461	Atypical Motor and Behavioral Presentations of Alzheimer Disease. <i>Neurologist</i> , 2012, 18, 266-272.	0.4	37
462	Age-specific and Sex-specific Prevalence and Incidence of Mild Cognitive Impairment, Dementia, and Alzheimer Dementia in Blacks and Whites. <i>Alzheimer Disease and Associated Disorders</i> , 2012, 26, 335-343.	0.6	297
463	Evidence for a role of the rare p.A152T variant in MAPT in increasing the risk for FTD-spectrum and Alzheimer's diseases. <i>Human Molecular Genetics</i> , 2012, 21, 3500-3512.	1.4	198
464	Characterization of frontotemporal dementia and/or amyotrophic lateral sclerosis associated with the GGGGCC repeat expansion in C9ORF72. <i>Brain</i> , 2012, 135, 765-783.	3.7	322
465	Mutations in the colony stimulating factor 1 receptor (CSF1R) gene cause hereditary diffuse leukoencephalopathy with spheroids. <i>Nature Genetics</i> , 2012, 44, 200-205.	9.4	428
466	An evaluation of the impact of <i>MAPT</i> , <i>SNCA</i> and <i>APOE</i> on the burden of Alzheimer's and Lewy body pathology. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2012, 83, 424-429.	0.9	50
467	Misregulation of human sortilin splicing leads to the generation of a nonfunctional progranulin receptor. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 21510-21515.	3.3	82
468	A Quantitative Postmortem MRI Design Sensitive to White Matter Hyperintensity Differences and Their Relationship With Underlying Pathology. <i>Journal of Neuropathology and Experimental Neurology</i> , 2012, 71, 1113-1122.	0.9	78

#	ARTICLE	IF	CITATIONS
469	<i>TMEM106B</i> risk variant is implicated in the pathologic presentation of Alzheimer disease. <i>Neurology</i> , 2012, 79, 717-718.	1.5	81
470	Neuropathologically defined subtypes of Alzheimer's disease differ significantly from neurofibrillary tangle-predominant dementia. <i>Acta Neuropathologica</i> , 2012, 124, 681-692.	3.9	103
471	National Institute on Aging's Alzheimer's Association guidelines for the neuropathologic assessment of Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2012, 8, 1-13.	0.4	1,968
472	Association and heterogeneity at the GAPDH locus in Alzheimer's disease. <i>Neurobiology of Aging</i> , 2012, 33, 203.e25-203.e33.	1.5	17
473	Multimodality imaging characteristics of dementia with Lewy bodies. <i>Neurobiology of Aging</i> , 2012, 33, 2091-2105.	1.5	162
474	Functional and genetic analysis of haplotypic sequence variation at the nicastrin genomic locus. <i>Neurobiology of Aging</i> , 2012, 33, 1848.e1-1848.e13.	1.5	5
475	Length of normal alleles of C9ORF72 GGGGCC repeat do not influence disease phenotype. <i>Neurobiology of Aging</i> , 2012, 33, 2950.e5-2950.e7.	1.5	83
476	Hippocampal-sparing Alzheimer's disease presenting as corticobasal syndrome. <i>Parkinsonism and Related Disorders</i> , 2012, 18, 683-685.	1.1	5
477	Characteristics of TBS-Extractable Hyperphosphorylated Tau Species: Aggregation Intermediates in rTg4510 Mouse Brain. <i>Journal of Alzheimer's Disease</i> , 2012, 33, 249-263.	1.2	81
478	Novel late-onset Alzheimer disease loci variants associate with brain gene expression. <i>Neurology</i> , 2012, 79, 221-228.	1.5	144
479	Right temporal variant frontotemporal dementia with motor neuron disease. <i>Journal of Clinical Neuroscience</i> , 2012, 19, 85-91.	0.8	20
480	Polymorphic genes of detoxification and mitochondrial enzymes and risk for progressive supranuclear palsy: a case control study. <i>BMC Medical Genetics</i> , 2012, 13, 16.	2.1	3
481	Neuroimaging correlates of pathologically defined subtypes of Alzheimer's disease: a case-control study. <i>Lancet Neurology</i> , The, 2012, 11, 868-877.	4.9	355
482	Hereditary diffuse leukoencephalopathy with axonal spheroids (HDLS): A misdiagnosed disease entity. <i>Journal of the Neurological Sciences</i> , 2012, 314, 130-137.	0.3	73
483	Parkinson's Disease and Parkinsonism: Neuropathology. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2012, 2, a009258-a009258.	2.9	593
484	Expression of Fused in sarcoma mutations in mice recapitulates the neuropathology of FUS proteinopathies and provides insight into disease pathogenesis. <i>Molecular Neurodegeneration</i> , 2012, 7, 53.	4.4	61
485	LRRK2 knockout mice have an intact dopaminergic system but display alterations in exploratory and motor co-ordination behaviors. <i>Molecular Neurodegeneration</i> , 2012, 7, 25.	4.4	165
486	Glutathione S-transferase omega genes in Alzheimer and Parkinson disease risk, age-at-diagnosis and brain gene expression: an association study with mechanistic implications. <i>Molecular Neurodegeneration</i> , 2012, 7, 13.	4.4	75

#	ARTICLE	IF	CITATIONS
487	Overlapping profiles of Aβ peptides in the Alzheimer's disease and pathological aging brains. <i>Alzheimer's Research and Therapy</i> , 2012, 4, 18.	3.0	92
488	Focal atrophy on MRI and neuropathologic classification of dementia with Lewy bodies. <i>Neurology</i> , 2012, 79, 553-560.	1.5	91
489	MRI characteristics and scoring in HDLS due to <i>CSF1R</i> gene mutations. <i>Neurology</i> , 2012, 79, 566-574.	1.5	153
490	Neuronal sensitivity to TDP-43 overexpression is dependent on timing of induction. <i>Acta Neuropathologica</i> , 2012, 123, 807-823.	3.9	46
491	Strikingly Different Clinicopathological Phenotypes Determined by Progranulin-Mutation Dosage. <i>American Journal of Human Genetics</i> , 2012, 90, 1102-1107.	2.6	414
492	Neuropathological analysis of brainstem cholinergic and catecholaminergic nuclei in relation to rapid eye movement (REM) sleep behaviour disorder. <i>Neuropathology and Applied Neurobiology</i> , 2012, 38, 142-152.	1.8	72
493	Ultrastructure of ubiquitin-positive, TDP-43-negative neuronal inclusions in cerebral cortex of C9ORF72-linked frontotemporal lobar degeneration/amyotrophic lateral sclerosis. <i>Neuropathology</i> , 2012, 32, 679-681.	0.7	5
494	Rapid eye movement sleep behavior disorder and subtypes in autopsy-confirmed dementia with Lewy bodies. <i>Movement Disorders</i> , 2012, 27, 72-78.	2.2	99
495	National Institute on Aging's Alzheimer's Association guidelines for the neuropathologic assessment of Alzheimer's disease: a practical approach. <i>Acta Neuropathologica</i> , 2012, 123, 1-11.	3.9	2,002
496	C9ORF72 repeat expansions and other FTD gene mutations in a clinical AD patient series from Mayo Clinic. <i>American Journal of Neurodegenerative Disease</i> , 2012, 1, 107-18.	0.1	32
497	Antemortem differential diagnosis of dementia pathology using structural MRI: Differential-STAND. <i>NeuroImage</i> , 2011, 55, 522-531.	2.1	90
498	Identification of common variants influencing risk of the tauopathy progressive supranuclear palsy. <i>Nature Genetics</i> , 2011, 43, 699-705.	9.4	502
499	Nuclear translocation of AMPK-1 potentiates striatal neurodegeneration in Huntington's disease. <i>Journal of Cell Biology</i> , 2011, 194, 209-227.	2.3	166
500	Disease specificity and pathologic progression of tau pathology in brainstem nuclei of Alzheimer's disease and progressive supranuclear palsy. <i>Neuroscience Letters</i> , 2011, 491, 122-126.	1.0	53
501	Glucocerebrosidase mutations in diffuse Lewy body disease. <i>Parkinsonism and Related Disorders</i> , 2011, 17, 55-57.	1.1	43
502	Cytokine expression and microglial activation in progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2011, 17, 683-688.	1.1	64
503	Incidental Lewy body disease: Do some cases represent a preclinical stage of dementia with Lewy bodies?. <i>Neurobiology of Aging</i> , 2011, 32, 857-863.	1.5	136
504	Temporoparietal atrophy: A marker of AD pathology independent of clinical diagnosis. <i>Neurobiology of Aging</i> , 2011, 32, 1531-1541.	1.5	105

#	ARTICLE	IF	CITATIONS
505	Association of common KIBRA variants with episodic memory and AD risk. <i>Neurobiology of Aging</i> , 2011, 32, 557.e1-557.e9.	1.5	40
506	Corticobasal degeneration: a pathologically distinct 4R tauopathy. <i>Nature Reviews Neurology</i> , 2011, 7, 263-272.	4.9	270
507	Ataxin-2 repeat-length variation and neurodegeneration. <i>Human Molecular Genetics</i> , 2011, 20, 3207-3212.	1.4	147
508	Investigating Statistical Epistasis in Complex Disorders. <i>Journal of Alzheimer's Disease</i> , 2011, 25, 635-644.	1.2	8
509	Immunoelectron Microscopic and Biochemical Studies of Caspase-Cleaved Tau in a Mouse Model of Tauopathy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2011, 70, 779-787.	0.9	15
510	Clinical Characterization of a Kindred With a Novel 12-Octapeptide Repeat Insertion in the Prion Protein Gene. <i>Archives of Neurology</i> , 2011, 68, 1165.	4.9	25
511	Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. <i>Nature Genetics</i> , 2011, 43, 436-441.	9.4	1,676
512	Common variants at ABCA7, MS4A6A/MS4A4E, EPHA1, CD33 and CD2AP are associated with Alzheimer's disease. <i>Nature Genetics</i> , 2011, 43, 429-435.	9.4	1,708
513	Neuropathologically defined subtypes of Alzheimer's disease with distinct clinical characteristics: a retrospective study. <i>Lancet Neurology</i> , The, 2011, 10, 785-796.	4.9	733
514	Association of LRRK2 exonic variants with susceptibility to Parkinson's disease: a case-control study. <i>Lancet Neurology</i> , The, 2011, 10, 898-908.	4.9	294
515	Expanded GGGGCC Hexanucleotide Repeat in Noncoding Region of C9ORF72 Causes Chromosome 9p-Linked FTD and ALS. <i>Neuron</i> , 2011, 72, 245-256.	3.8	4,176
516	Translation Initiator EIF4G1 Mutations in Familial Parkinson Disease. <i>American Journal of Human Genetics</i> , 2011, 89, 398-406.	2.6	250
517	Investigation of 15 of the top candidate genes for late-onset Alzheimer's disease. <i>Human Genetics</i> , 2011, 129, 273-282.	1.8	57
518	AMPK is abnormally activated in tangle- and pre-tangle-bearing neurons in Alzheimer's disease and other tauopathies. <i>Acta Neuropathologica</i> , 2011, 121, 337-349.	3.9	247
519	Neuropathological background of phenotypical variability in frontotemporal dementia. <i>Acta Neuropathologica</i> , 2011, 122, 137-153.	3.9	375
520	Neuropathology underlying clinical variability in patients with synucleinopathies. <i>Acta Neuropathologica</i> , 2011, 122, 187-204.	3.9	357
521	A proteomic study identifies different levels of light chain ferritin in corticobasal degeneration and progressive supranuclear palsy. <i>Acta Neuropathologica</i> , 2011, 122, 727-736.	3.9	4
522	Clinical and neuropathologic heterogeneity of c9FTD/ALS associated with hexanucleotide repeat expansion in C9ORF72. <i>Acta Neuropathologica</i> , 2011, 122, 673-690.	3.9	277

#	ARTICLE	IF	CITATIONS
523	Imaging Signatures of Molecular Pathology in Behavioral Variant Frontotemporal Dementia. <i>Journal of Molecular Neuroscience</i> , 2011, 45, 372-8.	1.1	61
524	Neuropathology of Frontotemporal Lobar Degeneration-Tau (FTLD-Tau). <i>Journal of Molecular Neuroscience</i> , 2011, 45, 384-389.	1.1	295
525	Replication of EPHA1 and CD33 associations with late-onset Alzheimer's disease: a multi-centre case-control study. <i>Molecular Neurodegeneration</i> , 2011, 6, 54.	4.4	67
526	Expression of mutant TDP-43 induces neuronal dysfunction in transgenic mice. <i>Molecular Neurodegeneration</i> , 2011, 6, 73.	4.4	137
527	Altered microRNA expression in frontotemporal lobar degeneration with TDP-43 pathology caused by progranulin mutations. <i>BMC Genomics</i> , 2011, 12, 527.	1.2	48
528	Anatomy of disturbed sleep in pallidoâ€pontoâ€nigral degeneration. <i>Annals of Neurology</i> , 2011, 69, 1014-1025.	2.8	10
529	Mutations in <i>LRRK2</i> increase phosphorylation of peroxiredoxin 3 exacerbating oxidative stress-induced neuronal death. <i>Human Mutation</i> , 2011, 32, 1390-1397.	1.1	111
530	Genetic and Clinical Features of Progranulin-Associated Frontotemporal Lobar Degeneration. <i>Archives of Neurology</i> , 2011, 68, 488.	4.9	108
531	Hippocampal Sclerosis in the Elderly. <i>Alzheimer Disease and Associated Disorders</i> , 2011, 25, 364-368.	0.6	78
532	Association of Crossword Puzzle Participation with Memory Decline in Persons Who Develop Dementia. <i>Journal of the International Neuropsychological Society</i> , 2011, 17, 1006-1013.	1.2	112
533	Replication of BIN1 Association with Alzheimer's Disease and Evaluation of Genetic Interactions. <i>Journal of Alzheimer's Disease</i> , 2011, 24, 751-758.	1.2	61
534	Neuropathological features of corticobasal degeneration presenting as corticobasal syndrome or Richardson syndrome. <i>Brain</i> , 2011, 134, 3264-3275.	3.7	119
535	A yeast functional screen predicts new candidate ALS disease genes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 20881-20890.	3.3	365
536	TDP-43 in aging and Alzheimer's disease - a review. <i>International Journal of Clinical and Experimental Pathology</i> , 2011, 4, 147-55.	0.5	118
537	Neuropathology of variants of progressive supranuclear palsy. <i>Current Opinion in Neurology</i> , 2010, 23, 394-400.	1.8	312
538	Alzheimer Diseaseâ€like Phenotype Associated With the c.154delA Mutation in Progranulin. <i>Archives of Neurology</i> , 2010, 67, 171-7.	4.9	59
539	Functional Impact of White Matter Hyperintensities in Cognitively Normal Elderly Subjects. <i>Archives of Neurology</i> , 2010, 67, 1379-85.	4.9	146
540	Changes in the Expression of Genes Associated with Intraneuronal Amyloid-Î² and Tau in Alzheimer's Disease. <i>Journal of Alzheimer's Disease</i> , 2010, 19, 97-109.	1.2	6

#	ARTICLE	IF	CITATIONS
541	Nomenclature and nosology for neuropathologic subtypes of frontotemporal lobar degeneration: an update. <i>Acta Neuropathologica</i> , 2010, 119, 1-4.	3.9	854
542	FUS pathology defines the majority of tau- and TDP-43-negative frontotemporal lobar degeneration. <i>Acta Neuropathologica</i> , 2010, 120, 33-41.	3.9	222
543	Cell type specific sequestration of choline acetyltransferase and tyrosine hydroxylase within Lewy bodies. <i>Acta Neuropathologica</i> , 2010, 120, 633-639.	3.9	38
544	Heterodimerization of Lrrk1&Larrk2: Implications for LRRK2-associated Parkinson disease. <i>Mechanisms of Ageing and Development</i> , 2010, 131, 210-214.	2.2	18
545	Reply to: SNCA variants are associated with increased risk of multiple system atrophy. <i>Annals of Neurology</i> , 2010, 67, 414-415.	2.8	39
546	Evidence in favor of Braak staging of Parkinson's disease. <i>Movement Disorders</i> , 2010, 25, S78-82.	2.2	112
547	Anatomical differences between CBS&Larrcorticobasal degeneration and CBS&LAlzheimer's disease. <i>Movement Disorders</i> , 2010, 25, 1246-1252.	2.2	71
548	Expression and functional profiling of neprilysin, insulin&Ldegrading enzyme, and endothelin&Lconverting enzyme in prospectively studied elderly and Alzheimer&Ls brain. <i>Journal of Neurochemistry</i> , 2010, 115, 47-57.	2.1	144
549	Common variants at 7p21 are associated with frontotemporal lobar degeneration with TDP-43 inclusions. <i>Nature Genetics</i> , 2010, 42, 234-239.	9.4	479
550	Extensive FUS&Limmunoreactive Pathology in Juvenile Amyotrophic Lateral Sclerosis with Basophilic Inclusions. <i>Brain Pathology</i> , 2010, 20, 1069-1076.	2.1	116
551	Three Repeat Isoforms of Tau Inhibit Assembly of Four Repeat Tau Filaments. <i>PLoS ONE</i> , 2010, 5, e10810.	1.1	82
552	Genome-wide Analysis of Genetic Loci Associated With Alzheimer Disease. <i>JAMA - Journal of the American Medical Association</i> , 2010, 303, 1832.	3.8	1,064
553	Association of I&L±, I&L²-, and I&L³-Synuclein With Diffuse Lewy Body Disease. <i>Archives of Neurology</i> , 2010, 67, 970-5.	4.9	63
554	Wild-Type Human TDP-43 Expression Causes TDP-43 Phosphorylation, Mitochondrial Aggregation, Motor Deficits, and Early Mortality in Transgenic Mice. <i>Journal of Neuroscience</i> , 2010, 30, 10851-10859.	1.7	457
555	Concordant Association of Insulin Degrading Enzyme Gene (IDE) Variants with IDE mRNA, A&LŸ, and Alzheimer's Disease. <i>PLoS ONE</i> , 2010, 5, e8764.	1.1	48
556	Replication of CLU, CR1, and PICALM Associations With Alzheimer Disease. <i>Archives of Neurology</i> , 2010, 67, 961-4.	4.9	188
557	Leucine-Rich Repeat Kinase 2 Gene-Associated Disease: Redefining Genotype-Phenotype Correlation. <i>Neurodegenerative Diseases</i> , 2010, 7, 175-179.	0.8	127
558	Common Variant in &L&LGRN&L; Is a Genetic Risk Factor for Hippocampal Sclerosis in the Elderly. <i>Neurodegenerative Diseases</i> , 2010, 7, 170-174.	0.8	82

#	ARTICLE	IF	CITATIONS
559	Frontotemporal Dementia. Blue Books of Neurology, 2010, 34, 397-416.	0.1	0
560	Elucidating the genetics and pathology of Perry syndrome. Journal of the Neurological Sciences, 2010, 289, 149-154.	0.3	112
561	Symmetric corticobasal degeneration (S-CBD). Parkinsonism and Related Disorders, 2010, 16, 208-214.	1.1	56
562	Iron and reactive oxygen species activity in parkinsonian substantia nigra. Parkinsonism and Related Disorders, 2010, 16, 329-333.	1.1	97
563	Contribution of vascular pathology to the clinical expression of dementia. Neurobiology of Aging, 2010, 31, 1710-1720.	1.5	94
564	O1-07-01: Accelerated lipofuscinosis and ubiquitination in granulin knockout mice suggests a role for progranulin in successful aging. , 2010, 6, S83-S83.		0
565	Accelerated Lipofuscinosis and Ubiquitination in Granulin Knockout Mice Suggest a Role for Progranulin in Successful Aging. American Journal of Pathology, 2010, 177, 311-324.	1.9	262
566	Hereditary diffuse leukoencephalopathy with spheroids: ultrastructural and immunoelectron microscopic studies. International Journal of Clinical and Experimental Pathology, 2010, 3, 665-74.	0.5	16
567	Genetics of Vascular Dementia. Minerva Psichiatrica, 2010, 51, 9-25.	1.2	1
568	MRI Correlates of Protein Deposition and Disease Severity in Postmortem Frontotemporal Lobar Degeneration. Neurodegenerative Diseases, 2009, 6, 106-117.	0.8	47
569	Plasma progranulin levels predict progranulin mutation status in frontotemporal dementia patients and asymptomatic family members. Brain, 2009, 132, 583-591.	3.7	344
570	Mimicking Aspects of Frontotemporal Lobar Degeneration and Lou Gehrig's Disease in Rats via TDP-43 Overexpression. Molecular Therapy, 2009, 17, 607-613.	3.7	76
571	Acceleration and persistence of neurofibrillary pathology in a mouse model of tauopathy following anesthesia. FASEB Journal, 2009, 23, 2595-2604.	0.2	130
572	Neuropathology of Cockayne syndrome: Evidence for impaired development, premature aging, and neurodegeneration. Mechanisms of Ageing and Development, 2009, 130, 619-636.	2.2	125
573	Neuropathological assessment of Parkinson's disease: refining the diagnostic criteria. Lancet Neurology, The, 2009, 8, 1150-1157.	4.9	734
574	<i>ATP13A2</i> variability in Parkinson disease. Human Mutation, 2009, 30, 406-410.	1.1	37
575	Corticobasal syndrome with Alzheimer's disease pathology. Movement Disorders, 2009, 24, 152-153.	2.2	16
576	<i>FGF20</i> and Parkinson's disease: No evidence of association or pathogenicity via α -synuclein expression. Movement Disorders, 2009, 24, 455-459.	2.2	41

#	ARTICLE	IF	CITATIONS
577	Nomenclature for neuropathologic subtypes of frontotemporal lobar degeneration: consensus recommendations. <i>Acta Neuropathologica</i> , 2009, 117, 15-18.	3.9	377
578	Evaluation of subcortical pathology and clinical correlations in FTLD-U subtypes. <i>Acta Neuropathologica</i> , 2009, 118, 349-358.	3.9	114
579	Brainstem atrophy on routine MR study in pallidopontonigral degeneration. <i>Journal of Neurology</i> , 2009, 256, 827-829.	1.8	5
580	Familial idiopathic basal ganglia calcification: a challenging clinical pathologic correlation. <i>Journal of Neurology</i> , 2009, 256, 839-842.	1.8	38
581	DCTN1 mutations in Perry syndrome. <i>Nature Genetics</i> , 2009, 41, 163-165.	9.4	285
582	Genetic variation in PCDH11X is associated with susceptibility to late-onset Alzheimer's disease. <i>Nature Genetics</i> , 2009, 41, 192-198.	9.4	279
583	Interphase Cytogenetics for 1p19q and t(1;19)(q10;p10) may Distinguish Prognostically Relevant Subgroups in Extraventricular Neurocytoma. <i>Brain Pathology</i> , 2009, 19, 623-629.	2.1	58
584	Distinct anatomical subtypes of the behavioural variant of frontotemporal dementia: a cluster analysis study. <i>Brain</i> , 2009, 132, 2932-2946.	3.7	277
585	Prominent phenotypic variability associated with mutations in Progranulin. <i>Neurobiology of Aging</i> , 2009, 30, 739-751.	1.5	166
586	Neuropathology of nondemented aging: Presumptive evidence for preclinical Alzheimer disease. <i>Neurobiology of Aging</i> , 2009, 30, 1026-1036.	1.5	558
587	Pallidonigral TDP-43 pathology in Perry syndrome. <i>Parkinsonism and Related Disorders</i> , 2009, 15, 281-286.	1.1	89
588	Glucosidase-beta variations and Lewy body disorders. <i>Parkinsonism and Related Disorders</i> , 2009, 15, 414-416.	1.1	36
589	GCH1 expression in human cerebellum from healthy individuals is not gender dependant. <i>Neuroscience Letters</i> , 2009, 462, 73-75.	1.0	2
590	Aberrant cleavage of TDP-43 enhances aggregation and cellular toxicity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 7607-7612.	3.3	523
591	Overexpression of Wild-Type Murine Tau Results in Progressive Tauopathy and Neurodegeneration. <i>American Journal of Pathology</i> , 2009, 175, 1598-1609.	1.9	56
592	Neuropathology of non-motor features of Parkinson disease. <i>Parkinsonism and Related Disorders</i> , 2009, 15, S1-S5.	1.1	228
593	Comparison of Risk Factor Profiles in Incidental Lewy Body Disease and Parkinson Disease. <i>Archives of Neurology</i> , 2009, 66, 1114-9.	4.9	34
594	Transactivation Response DNA-Binding Protein 43 Microvasculopathy in Frontotemporal Degeneration and Familial Lewy Body Disease. <i>Journal of Neuropathology and Experimental Neurology</i> , 2009, 68, 1167-1176.	0.9	48

#	ARTICLE	IF	CITATIONS
595	Cortical Alzheimer type pathology does not influence tau pathology in progressive supranuclear palsy. <i>International Journal of Clinical and Experimental Pathology</i> , 2009, 2, 399-406.	0.5	11
596	Neuropathology of non-Alzheimer degenerative disorders. <i>International Journal of Clinical and Experimental Pathology</i> , 2009, 3, 1-23.	0.5	68
597	Neurodegeneration involving putative respiratory neurons in Perry syndrome. <i>Acta Neuropathologica</i> , 2008, 115, 263-268.	3.9	56
598	Evidence that incidental Lewy body disease is pre-symptomatic Parkinson's disease. <i>Acta Neuropathologica</i> , 2008, 115, 437-444.	3.9	329
599	Co-localization of tau and α -synuclein in the olfactory bulb in Alzheimer's disease with amygdala Lewy bodies. <i>Acta Neuropathologica</i> , 2008, 116, 17-24.	3.9	70
600	Frontotemporal lobar degeneration with ubiquitin-positive, but TDP-43-negative inclusions. <i>Acta Neuropathologica</i> , 2008, 116, 159-167.	3.9	50
601	Glial cytoplasmic inclusions in neurologically normal elderly: prodromal multiple system atrophy?. <i>Acta Neuropathologica</i> , 2008, 116, 269-275.	3.9	53
602	Temporal lobar predominance of TDP-43 neuronal cytoplasmic inclusions in Alzheimer disease. <i>Acta Neuropathologica</i> , 2008, 116, 215-220.	3.9	124
603	Ultrastructural localization of TDP-43 in filamentous neuronal inclusions in various neurodegenerative diseases. <i>Acta Neuropathologica</i> , 2008, 116, 205-213.	3.9	119
604	Evaluation of α -synuclein immunohistochemical methods used by invited experts. <i>Acta Neuropathologica</i> , 2008, 116, 277-288.	3.9	157
605	MR imaging of brainstem atrophy in progressive supranuclear palsy. <i>Journal of Neurology</i> , 2008, 255, 37-44.	1.8	46
606	Identification of proteins in human substantia nigra. <i>Proteomics - Clinical Applications</i> , 2008, 2, 776-782.	0.8	33
607	Cardiac sympathetic denervation correlates with clinical and pathologic stages of Parkinson's disease. <i>Movement Disorders</i> , 2008, 23, 1085-1092.	2.2	167
608	Progranulin gene mutation with an unusual clinical and neuropathologic presentation. <i>Movement Disorders</i> , 2008, 23, 1168-1173.	2.2	36
609	β -amyloid burden is not associated with rates of brain atrophy. <i>Annals of Neurology</i> , 2008, 63, 204-212.	2.8	187
610	A novel human disease with abnormal prion protein sensitive to protease. <i>Annals of Neurology</i> , 2008, 63, 697-708.	2.8	250
611	Neuropathology of Parkinson's Disease. , 2008, , 35-48.		6
612	Age and apoE associations with complex pathologic features in Alzheimer's disease. <i>Journal of the Neurological Sciences</i> , 2008, 273, 34-39.	0.3	30

#	ARTICLE	IF	CITATIONS
613	Voxel-based morphometry in autopsy proven PSP and CBD. <i>Neurobiology of Aging</i> , 2008, 29, 280-289.	1.5	221
614	Argyrophilic grains: A distinct disease or an additive pathology?. <i>Neurobiology of Aging</i> , 2008, 29, 566-573.	1.5	70
615	Neuropsychological findings in clinically atypical autopsy confirmed corticobasal degeneration and progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2008, 14, 376-378.	1.1	26
616	Rates of brain atrophy over time in autopsy-proven frontotemporal dementia and Alzheimer disease. <i>NeuroImage</i> , 2008, 39, 1034-1040.	2.1	52
617	Antemortem MRI based STructural Abnormality iNdex (STAND)-scores correlate with postmortem Braak neurofibrillary tangle stage. <i>NeuroImage</i> , 2008, 42, 559-567.	2.1	152
618	TDP-43 in neurodegenerative disorders. <i>Expert Opinion on Biological Therapy</i> , 2008, 8, 969-978.	1.4	39
619	Neuropathology of Hippocampal Sclerosis. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2008, 89, 569-572.	1.0	26
620	Leflunomide-Associated Progressive Multifocal Leukoencephalopathy. <i>Archives of Neurology</i> , 2008, 65, 1538.	4.9	58
621	Validation of the Neuropathologic Criteria of the Third Consortium for Dementia With Lewy Bodies for Prospectively Diagnosed Cases. <i>Journal of Neuropathology and Experimental Neurology</i> , 2008, 67, 649-656.	0.9	137
622	Clinical and neuropathologic features of progressive supranuclear palsy with severe pallido-nigro-luysial degeneration and axonal dystrophy. <i>Brain</i> , 2008, 131, 460-472.	3.7	94
623	Neuropathology of Progressive Supranuclear Palsy. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2008, 89, 487-491.	1.0	7
624	Novel Mutations in TARDBP (TDP-43) in Patients with Familial Amyotrophic Lateral Sclerosis. <i>PLoS Genetics</i> , 2008, 4, e1000193.	1.5	393
625	Incidental Lewy Body Disease and Preclinical Parkinson Disease. <i>Archives of Neurology</i> , 2008, 65, 1074-80.	4.9	166
626	Expanded-Polyglutamine Huntingtin Protein Suppresses the Secretion and Production of a Chemokine (CCL5/RANTES) by Astrocytes. <i>Journal of Neuroscience</i> , 2008, 28, 3277-3290.	1.7	100
627	Altered Expression of Zonula Occludens-2 Precedes Increased Blood-Brain Barrier Permeability in a Murine Model of Fulminant Hepatic Failure. <i>Journal of Investigative Surgery</i> , 2008, 21, 101-108.	0.6	48
628	Common variation in the miR-659 binding-site of GRN is a major risk factor for TDP43-positive frontotemporal dementia. <i>Human Molecular Genetics</i> , 2008, 17, 3631-3642.	1.4	271
629	Alzheimer Disease: Postmortem Neuropathologic Correlates of Antemortem ¹ H MR Spectroscopy Metabolite Measurements. <i>Radiology</i> , 2008, 248, 210-220.	3.6	147
630	IN DEMENTIA WITH LEWY BODIES, BRAAK STAGE DETERMINES PHENOTYPE, NOT LEWY BODY DISTRIBUTION. <i>Neurology</i> , 2008, 70, 2087-2089.	1.5	2

#	ARTICLE	IF	CITATIONS
631	Early Onset Familial Alzheimer Disease With Spastic Paraparesis, Dysarthria, and Seizures and N135S Mutation in PSEN1. <i>Alzheimer Disease and Associated Disorders</i> , 2008, 22, 299-307.	0.6	35
632	Frontotemporal Dementia Mimicking Dementia With Lewy Bodies. <i>Cognitive and Behavioral Neurology</i> , 2008, 21, 157-163.	0.5	50
633	Cognitive Performance Correlates with Cortical Isopeptide Immunoreactivity as Well as Alzheimer Type Pathology. <i>Journal of Alzheimer's Disease</i> , 2008, 13, 53-66.	1.2	20
634	Differential Incorporation of Tau Isoforms in Alzheimer's Disease. <i>Journal of Alzheimer's Disease</i> , 2008, 14, 1-16.	1.2	107
635	Coexistence of diffuse multisystem tauopathy and cerebral amyloid angiopathy in an elderly patient with dementia. <i>FASEB Journal</i> , 2008, 22, 707.10.	0.2	0
636	TDP43 Neuronal Cytoplasmic Inclusions in the Amygdala of Patients with Advanced Alzheimer Disease. <i>FASEB Journal</i> , 2008, 22, 58.6.	0.2	0
637	Tau Negative FTLD Without Abnormal TDP43 Immunoreactivity. <i>FASEB Journal</i> , 2008, 22, 707.13.	0.2	0
638	Immunoelectron microscopy of TDP43 in frontotemporal lobar degeneration, amyotrophic lateral sclerosis and Lewy body disease. <i>FASEB Journal</i> , 2008, 22, 58.12.	0.2	0
639	Limbic lobe microvacuolation is minimal in Alzheimer's disease in the absence of concurrent Lewy body disease. <i>International Journal of Clinical and Experimental Pathology</i> , 2008, 1, 369-75.	0.5	4
640	A β 40 Inhibits Amyloid Deposition In Vivo. <i>Journal of Neuroscience</i> , 2007, 27, 627-633.	1.7	327
641	Single-dose intracerebroventricular administration of galactocerebrosidase improves survival in a mouse model of globoid cell leukodystrophy. <i>FASEB Journal</i> , 2007, 21, 2520-2527.	0.2	85
642	Actin-binding Proteins Coronin-1a and IBA-1 are Effective Microglial Markers for Immunohistochemistry. <i>Journal of Histochemistry and Cytochemistry</i> , 2007, 55, 687-700.	1.3	214
643	A novel locus for dementia with Lewy bodies: a clinically and genetically heterogeneous disorder. <i>Brain</i> , 2007, 130, 2277-2291.	3.7	56
644	Clinical Features of Pathologic Subtypes of Behavioral-Variant Frontotemporal Dementia. <i>Archives of Neurology</i> , 2007, 64, 1611.	4.9	35
645	FRONTOTEMPORAL LOBAR DEGENERATION WITH UPPER MOTOR NEURON DISEASE/ PRIMARY LATERAL SCLEROSIS. <i>Neurology</i> , 2007, 69, 1800-1801.	1.5	24
646	Progranulin Mutations in Primary Progressive Aphasia. <i>Archives of Neurology</i> , 2007, 64, 43.	4.9	146
647	Sex-dependent association of a common low-density lipoprotein receptor polymorphism with RNA splicing efficiency in the brain and Alzheimer's disease. <i>Human Molecular Genetics</i> , 2007, 17, 929-935.	1.4	52
648	Voxel-Based Morphometry in Frontotemporal Lobar Degeneration With Ubiquitin-Positive Inclusions With and Without Progranulin Mutations. <i>Archives of Neurology</i> , 2007, 64, 371.	4.9	82

#	ARTICLE	IF	CITATIONS
649	Chapter 7 Ubiquitinopathies. Blue Books of Neurology, 2007, , 165-185.	0.1	2
650	Clinical Features and Survival of 3R and 4R Tauopathies Presenting as Behavioral Variant Frontotemporal Dementia. Alzheimer Disease and Associated Disorders, 2007, 21, S39-S43.	0.6	23
651	Neuropathologic Features of Frontotemporal Lobar Degeneration With Ubiquitin-Positive Inclusions With Progranulin Gene (PGRN) Mutations. Journal of Neuropathology and Experimental Neurology, 2007, 66, 142-151.	0.9	184
652	The Etiopathogenesis of Parkinson Disease and Suggestions for Future Research. Part I. Journal of Neuropathology and Experimental Neurology, 2007, 66, 251-257.	0.9	104
653	The Etiopathogenesis of Parkinson Disease and Suggestions for Future Research. Part II. Journal of Neuropathology and Experimental Neurology, 2007, 66, 329-336.	0.9	41
654	Clinical-pathologic study of biomarkers in FTDP-17 (PPND family with N279K tau mutation). Parkinsonism and Related Disorders, 2007, 13, 230-239.	1.1	47
655	Quantitative PCR-based screening of α -synuclein multiplication in multiple system atrophy. Parkinsonism and Related Disorders, 2007, 13, 340-342.	1.1	35
656	Hippocampal sclerosis in tau-negative frontotemporal lobar degeneration. Neurobiology of Aging, 2007, 28, 1718-1722.	1.5	47
657	Identification of a Novel Risk Locus for Progressive Supranuclear Palsy by a Pooled Genomewide Scan of 500,288 Single-Nucleotide Polymorphisms. American Journal of Human Genetics, 2007, 80, 769-778.	2.6	68
658	Linking Selective Vulnerability to Cell Death Mechanisms in Parkinson's Disease. American Journal of Pathology, 2007, 170, 16-19.	1.9	32
659	Neuropathology of Parkinson's disease dementia and dementia with Lewy bodies with reference to striatal pathology. Parkinsonism and Related Disorders, 2007, 13, S221-S224.	1.1	105
660	Progranulin in frontotemporal lobar degeneration and neuroinflammation. Journal of Neuroinflammation, 2007, 4, 7.	3.1	194
661	TDP-43 immunoreactivity in hippocampal sclerosis and Alzheimer's disease. Annals of Neurology, 2007, 61, 435-445.	2.8	753
662	The ups and downs of α -synuclein mRNA expression. Movement Disorders, 2007, 22, 293-295.	2.2	47
663	Clinical diagnostic criteria for dementia associated with Parkinson's disease. Movement Disorders, 2007, 22, 1689-1707.	2.2	2,497
664	Progressive Supranuclear Palsy: Pathology and Genetics. Brain Pathology, 2007, 17, 74-82.	2.1	249
665	A presenilin 1 mutation (L420R) in a family with early onset Alzheimer disease, seizures and cotton wool plaques, but not spastic paraparesis. Neuropathology, 2007, 27, 228-232.	0.7	38
666	Lrrk2 G2019S substitution in frontotemporal lobar degeneration with ubiquitin-immunoreactive neuronal inclusions. Acta Neuropathologica, 2007, 113, 601-606.	3.9	55

#	ARTICLE	IF	CITATIONS
667	Hippocampal sclerosis dementia differs from hippocampal sclerosis in frontal lobe degeneration. <i>Acta Neuropathologica</i> , 2007, 113, 245-252.	3.9	87
668	TDP-43 in differential diagnosis of motor neuron disorders. <i>Acta Neuropathologica</i> , 2007, 114, 71-79.	3.9	131
669	Neuropathologic diagnostic and nosologic criteria for frontotemporal lobar degeneration: consensus of the Consortium for Frontotemporal Lobar Degeneration. <i>Acta Neuropathologica</i> , 2007, 114, 5-22.	3.9	978
670	TDP-43 immunoreactivity in neurodegenerative disorders: disease versus mechanism specificity. <i>Acta Neuropathologica</i> , 2007, 115, 147-149.	3.9	22
671	The high-affinity HSP90-CHIP complex recognizes and selectively degrades phosphorylated tau client proteins. <i>Journal of Clinical Investigation</i> , 2007, 117, 648-658.	3.9	545
672	Coronin-1a: A novel microglial marker for use in paraffin embedded tissue. <i>FASEB Journal</i> , 2007, 21, A20.	0.2	0
673	Dual pathologies: Utility of TAR DNA-Binding Protein 43 (TDP-43) Staining in Patients with Frontal and Temporal Lobe Abnormalities and Alzheimer disease. <i>FASEB Journal</i> , 2007, 21, .	0.2	0
674	Frontotemporal lobar degeneration with upper motor neuron disease/primary lateral sclerosis. <i>FASEB Journal</i> , 2007, 21, A21.	0.2	2
675	Detection of TDP-43 in Alzheimer's disease and hippocampal sclerosis. <i>FASEB Journal</i> , 2007, 21, A25.	0.2	0
676	Progranulin is located in secretory granules and vesicles of neutrophils and macrophages by immunogold electron microscopy. <i>FASEB Journal</i> , 2007, 21, A22.	0.2	1
677	Clinicopathological and imaging correlates of progressive aphasia and apraxia of speech. <i>Brain</i> , 2006, 129, 1385-1398.	3.7	624
678	Neuropsychological Differentiation of Dementia with Lewy Bodies from Normal Aging and Alzheimer's Disease. <i>Clinical Neuropsychologist</i> , 2006, 20, 623-636.	1.5	170
679	Effect of MAPT and APOE on prognosis of progressive supranuclear palsy. <i>Neuroscience Letters</i> , 2006, 405, 116-119.	1.0	10
680	Neurofibrillary tangle-related synaptic alterations of spinal motor neurons of P301L tau transgenic mice. <i>Neuroscience Letters</i> , 2006, 409, 95-99.	1.0	26
681	The relationship between histopathological features of progressive supranuclear palsy and disease duration. <i>Parkinsonism and Related Disorders</i> , 2006, 12, 109-112.	1.1	21
682	Absence of Rapid Eye Movement Sleep Behavior Disorder in 11 Members of the Pallidopontonigral Degeneration Kindred. <i>Archives of Neurology</i> , 2006, 63, 268.	4.9	27
683	Frontotemporal Lobar Degeneration Without Lobar Atrophy. <i>Archives of Neurology</i> , 2006, 63, 1632.	4.9	52
684	Alzheimer Disease With Amygdala Lewy Bodies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2006, 65, 685-697.	0.9	279

#	ARTICLE	IF	CITATIONS
685	Clinically Undetected Motor Neuron Disease in Pathologically Proven Frontotemporal Lobar Degeneration With Motor Neuron Disease. <i>Archives of Neurology</i> , 2006, 63, 506.	4.9	66
686	Atypical Progressive Supranuclear Palsy With Corticospinal Tract Degeneration. <i>Journal of Neuropathology and Experimental Neurology</i> , 2006, 65, 396-405.	0.9	129
687	Argyrophilic Grain Disease in Demented Subjects Presenting Initially With Amnesic Mild Cognitive Impairment. <i>Journal of Neuropathology and Experimental Neurology</i> , 2006, 65, 602-609.	0.9	48
688	Identification of G-Protein Coupled Receptor Kinase 2 in Paired Helical Filaments and Neurofibrillary Tangles. <i>Journal of Neuropathology and Experimental Neurology</i> , 2006, 65, 1157-1169.	0.9	19
689	Heterogeneous inclusions in neurofilament inclusion disease. <i>Neuropathology</i> , 2006, 26, 417-421.	0.7	21
690	Coexistence of PSP and MSA: a case report and review of the literature. <i>Acta Neuropathologica</i> , 2006, 111, 186-192.	3.9	36
691	Hereditary diffuse leukoencephalopathy with spheroids: clinical, pathologic and genetic studies of a new kindred. <i>Acta Neuropathologica</i> , 2006, 111, 300-311.	3.9	84
692	Dopamine β -hydroxylase deficiency involves the central autonomic network. <i>Acta Neuropathologica</i> , 2006, 112, 227-229.	3.9	9
693	Suppression of galactosylceramidase (GALC) expression in the twitcher mouse model of globoid cell leukodystrophy (GLD) is caused by nonsense-mediated mRNA decay (NMD). <i>Neurobiology of Disease</i> , 2006, 23, 273-280.	2.1	27
694	Rates of cerebral atrophy in autopsy-confirmed progressive supranuclear palsy. <i>Annals of Neurology</i> , 2006, 59, 200-203.	2.8	30
695	Lrrk2 and Lewy body disease. <i>Annals of Neurology</i> , 2006, 59, 388-393.	2.8	259
696	Frontotemporal dementia and parkinsonism associated with the IVS1+1G->A mutation in progranulin: a clinicopathologic study. <i>Brain</i> , 2006, 129, 3103-3114.	3.7	105
697	Lewy Bodies in Progressive Supranuclear Palsy Represent an Independent Disease Process. <i>Journal of Neuropathology and Experimental Neurology</i> , 2006, 65, 387-395.	0.9	53
698	Alpha1-antichymotrypsin, an inflammatory protein overexpressed in Alzheimer's disease brain, induces tau phosphorylation in neurons. <i>Brain</i> , 2006, 129, 3020-3034.	3.7	101
699	Rates of cerebral atrophy differ in different degenerative pathologies. <i>Brain</i> , 2006, 130, 1148-1158.	3.7	146
700	β -Amyloid Degradation and Alzheimer's Disease. <i>Journal of Biomedicine and Biotechnology</i> , 2006, 2006, 1-12.	3.0	151
701	Ageing is Neuroprotective During Global Ischemia but Leads to Increased Caspase-3 and Apoptotic Activity in Hippocampal Neurons. <i>Current Neurovascular Research</i> , 2006, 3, 181-186.	0.4	17
702	Neuropathologic Features of Amnesic Mild Cognitive Impairment. <i>Archives of Neurology</i> , 2006, 63, 665.	4.9	562

#	ARTICLE	IF	CITATIONS
703	Deletion of the Ubiquitin Ligase CHIP Leads to the Accumulation, But Not the Aggregation, of Both Endogenous Phospho- and Caspase-3-Cleaved Tau Species. <i>Journal of Neuroscience</i> , 2006, 26, 6985-6996.	1.7	234
704	Cockayne Syndrome in Adults: Review With Clinical and Pathologic Study of a New Case. <i>Journal of Child Neurology</i> , 2006, 21, 991-1006.	0.7	113
705	An inhibitor of tau hyperphosphorylation prevents severe motor impairments in tau transgenic mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006, 103, 9673-9678.	3.3	206
706	Increased Frequency of Argyrophilic Grain Disease in Alzheimer Disease with 4R Tau-Specific Immunohistochemistry. <i>Journal of Neuropathology and Experimental Neurology</i> , 2005, 64, 209-214.	0.9	51
707	Neuropathologic, Biochemical, and Molecular Characterization of the Frontotemporal Dementias. <i>Journal of Neuropathology and Experimental Neurology</i> , 2005, 64, 420-428.	0.9	92
708	Hippocampal progenitor cells express nestin following cerebral ischemia in rats. <i>NeuroReport</i> , 2005, 16, 1541-1544.	0.6	13
709	Ubiquitin Immunohistochemistry of Frontotemporal Lobar Degeneration Differentiates Cases With and Without Motor Neuron Disease. <i>Alzheimer Disease and Associated Disorders</i> , 2005, 19, S37-S43.	0.6	35
710	Progressive white matter pathology in the spinal cord of transgenic mice expressing mutant (P301L) human tau. <i>Journal of Neurocytology</i> , 2005, 34, 397-410.	1.6	36
711	Antemortem diagnosis of frontotemporal lobar degeneration. <i>Annals of Neurology</i> , 2005, 57, 480-488.	2.8	181
712	Tau kinases and Parkinson's disease: Guilt by association?. <i>Annals of Neurology</i> , 2005, 58, 819-820.	2.8	2
713	Increased tau burden in the cortices of progressive supranuclear palsy presenting with corticobasal syndrome. <i>Movement Disorders</i> , 2005, 20, 982-988.	2.2	111
714	Alpha-synuclein immunohistochemistry in two cases of co-occurring idiopathic Parkinson's disease and motor neuron disease. <i>Movement Disorders</i> , 2005, 20, 1515-1520.	2.2	18
715	Required techniques and useful molecular markers in the neuropathologic diagnosis of neurodegenerative diseases. <i>Acta Neuropathologica</i> , 2005, 109, 14-24.	3.9	71
716	Extending the clinicopathological spectrum of neurofilament inclusion disease. <i>Acta Neuropathologica</i> , 2005, 109, 427-432.	3.9	28
717	Decreased Nprilysin Immunoreactivity in Alzheimer Disease, but Not in Pathological Aging. <i>Journal of Neuropathology and Experimental Neurology</i> , 2005, 64, 378-385.	0.9	72
718	Neuropathology of Parkinson's disease. , 2005, , 575-585.		0
719	Screening for neurofilament inclusion disease using \hat{A} -internexin immunohistochemistry. <i>Neurology</i> , 2005, 64, 1658-1659.	1.5	22
720	Aging Blunts Ischemic-Preconditioning-Induced Neuroprotection Following Transient Global Ischemia in Rats. <i>Current Neurovascular Research</i> , 2005, 2, 365-374.	0.4	51

#	ARTICLE	IF	CITATIONS
721	Enzyme replacement therapy results in substantial improvements in early clinical phenotype in a mouse model of globoid cell leukodystrophy. <i>FASEB Journal</i> , 2005, 19, 1549-1551.	0.2	79
722	A Tribute to a Neuropathologist, Robert D. Terry. , 2005, 1, 74-76.		1
723	Tau gene transfer, but not alpha-synuclein, induces both progressive dopamine neuron degeneration and rotational behavior in the rat. <i>Neurobiology of Disease</i> , 2005, 20, 64-73.	2.1	36
724	Dementia with Lewy bodies and Parkinson's disease with dementia: Are they different?. <i>Parkinsonism and Related Disorders</i> , 2005, 11, S47-S51.	1.1	135
725	The Effect of tau genotype on clinical features in FTDP-17. <i>Parkinsonism and Related Disorders</i> , 2005, 11, 205-208.	1.1	31
726	A β 242 Is Essential for Parenchymal and Vascular Amyloid Deposition in Mice. <i>Neuron</i> , 2005, 47, 191-199.	3.8	524
727	Sporadic tauopathies: Pick's disease, corticobasal degeneration, progressive supranuclear palsy and argyrophilic grain disease. , 2004, , 227-256.		18
728	Hippocampal Sclerosis and Ubiquitin-Positive Inclusions in Dementia Lacking Distinctive Histopathology. <i>Dementia and Geriatric Cognitive Disorders</i> , 2004, 17, 342-345.	0.7	44
729	Apolipoprotein E ϵ 4 Is a Determinant for Alzheimer-Type Pathologic Features in Tauopathies, Synucleinopathies, and Frontotemporal Degeneration. <i>Archives of Neurology</i> , 2004, 61, 1579.	4.9	64
730	Dimeric Amyloid β Protein Rapidly Accumulates in Lipid Rafts followed by Apolipoprotein E and Phosphorylated Tau Accumulation in the Tg2576 Mouse Model of Alzheimer's Disease. <i>Journal of Neuroscience</i> , 2004, 24, 3801-3809.	1.7	334
731	Hippocampal sclerosis dementia. <i>Neurology</i> , 2004, 63, 414-415.	1.5	26
732	Nonvasculitic autoimmune inflammatory meningoencephalitis. <i>Neuropathology</i> , 2004, 24, 149-152.	0.7	39
733	Ballooned neurones in the limbic lobe are associated with Alzheimer type pathology and lack diagnostic specificity. <i>Neuropathology and Applied Neurobiology</i> , 2004, 30, 676-682.	1.8	26
734	Inferior olivary hypertrophy is uncommon in progressive supranuclear palsy. <i>Acta Neuropathologica</i> , 2004, 108, 143-6.	3.9	16
735	Neuropathology of primary restless leg syndrome: Absence of specific α - and β -synuclein pathology. <i>Movement Disorders</i> , 2004, 19, 695-699.	2.2	78
736	Is the neuropathological "gold standard" diagnosis dead? Implications of clinicopathological findings in an autosomal dominant neurodegenerative disorder. <i>Parkinsonism and Related Disorders</i> , 2004, 10, 461-463.	1.1	19
737	Mutations in LRRK2 Cause Autosomal-Dominant Parkinsonism with Pleomorphic Pathology. <i>Neuron</i> , 2004, 44, 601-607.	3.8	2,653
738	Building a More Perfect Beast. <i>American Journal of Pathology</i> , 2004, 164, 1143-1146.	1.9	23

#	ARTICLE	IF	CITATIONS
739	Apoptosis in oligodendrocytes is associated with axonal degeneration in P301L tau mice. <i>Neurobiology of Disease</i> , 2004, 15, 553-562.	2.1	43
740	Î±-Synuclein immunoreactivity in neuronal nuclear inclusions and neurites in multiple system atrophy. <i>Neuroscience Letters</i> , 2004, 354, 99-102.	1.0	57
741	Decreases in soluble Î±-synuclein in frontal cortex correlate with cognitive decline in the elderly. <i>Neuroscience Letters</i> , 2004, 359, 104-108.	1.0	25
742	Contribution of changes in ubiquitin and myelin basic protein to age-related cognitive decline. <i>Neuroscience Research</i> , 2004, 48, 93-100.	1.0	61
743	Biochemical characterization of torsinB. <i>Molecular Brain Research</i> , 2004, 127, 1-9.	2.5	12
744	Correlation Between Antemortem Magnetic Resonance Imaging Findings and Pathologically Confirmed Corticobasal Degeneration. <i>Archives of Neurology</i> , 2004, 61, 1881-4.	4.9	67
745	Apoptotic mechanisms in Alzheimer neurofibrillary degeneration: cause or effect?. <i>Journal of Clinical Investigation</i> , 2004, 114, 23-27.	3.9	163
746	Tau protein expression in adult bovine oligodendrocytes: functional and pathological significance. <i>Neurochemical Research</i> , 2003, 28, 1385-1392.	1.6	3
747	Ultrastructural neuronal pathology in transgenic mice expressing mutant (P301L) human tau. <i>Journal of Neurocytology</i> , 2003, 32, 1091-1105.	1.6	115
748	Diagnostic accuracy of progressive supranuclear palsy in the Society for Progressive Supranuclear Palsy Brain Bank. <i>Movement Disorders</i> , 2003, 18, 1018-1026.	2.2	155
749	The neuropathology and biochemistry of frontotemporal dementia. <i>Annals of Neurology</i> , 2003, 54, S24-S28.	2.8	83
750	ANTEMORTEM MEMORY IMPAIRMENT SCREEN PERFORMANCE IS CORRELATED WITH POSTMORTEM ALZHEIMER PATHOLOGY. <i>Journal of the American Geriatrics Society</i> , 2003, 51, 1043-1045.	1.3	9
751	Co-Localization of Glycogen Synthase Kinase-3 with Neurofibrillary Tangles and Granulovacuolar Degeneration in Transgenic Mice. <i>American Journal of Pathology</i> , 2003, 163, 1057-1067.	1.9	87
752	Filamentous Tau in Oligodendrocytes and Astrocytes of Transgenic Mice Expressing the Human Tau Isoform with the P301L Mutation. <i>American Journal of Pathology</i> , 2003, 162, 213-218.	1.9	95
753	Oxidized neprilysin in aging and Alzheimer's disease brains. <i>Biochemical and Biophysical Research Communications</i> , 2003, 310, 236-241.	1.0	132
754	In situ hybridization for detection of nocardial 16S rRNA: reactivity within intracellular inclusions in experimentally infected cynomolgus monkeys and in Lewy body-containing human brain specimens. <i>Experimental Neurology</i> , 2003, 184, 715-725.	2.0	32
755	Caught in the Act. <i>Neuron</i> , 2003, 40, 453-456.	3.8	184
756	Pin1 colocalization with phosphorylated tau in Alzheimer's disease and other tauopathies. <i>Neurobiology of Disease</i> , 2003, 14, 251-264.	2.1	78

#	ARTICLE	IF	CITATIONS
757	<i>APOE</i> E4 is a determinant for Alzheimer type pathology in progressive supranuclear palsy. <i>Neurology</i> , 2003, 60, 240-245.	1.5	43
758	Failure to Wean from a Ventilator Caused by ANNA-1 Seropositive Paraneoplastic Syndrome. <i>European Neurology</i> , 2003, 50, 112-114.	0.6	0
759	Colocalization of Tau and Alpha-Synuclein Epitopes in Lewy Bodies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2003, 62, 389-397.	0.9	306
760	Neurofilament inclusion body disease: a new proteinopathy?. <i>Brain</i> , 2003, 126, 2291-2303.	3.7	176
761	Familial Primary Progressive Aphasia. <i>Alzheimer Disease and Associated Disorders</i> , 2003, 17, 106-112.	0.6	24
762	Dementia with Lewy Bodies: Neuropathology. <i>Journal of Geriatric Psychiatry and Neurology</i> , 2002, 15, 210-216.	1.2	56
763	Selective Neurofibrillary Degeneration of the Hippocampal CA2 Sector Is Associated with Four-Repeat Tauopathies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2002, 61, 1040-1047.	0.9	61
764	Argyrophilic Grain Disease Is a Sporadic 4-Repeat Tauopathy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2002, 61, 547-556.	0.9	232
765	A Clinicopathological Study of Vascular Progressive Supranuclear Palsy. <i>Archives of Neurology</i> , 2002, 59, 1597.	4.9	64
766	Parkinson Disease Neuropathology. <i>Archives of Neurology</i> , 2002, 59, 102.	4.9	366
767	Dementia with Lewy bodies may present as dementia and REM sleep behavior disorder without parkinsonism or hallucinations. <i>Journal of the International Neuropsychological Society</i> , 2002, 8, 907-914.	1.2	124
768	The subthalamic nucleus has neurofibrillary tangles in argyrophilic grain disease and advanced Alzheimer's disease. <i>Neuroscience Letters</i> , 2002, 320, 81-85.	1.0	24
769	Clinical correlates of the pathology underlying parkinsonism: A population perspective. <i>Movement Disorders</i> , 2002, 17, 910-916.	2.2	72
770	Neuropathology of two members of a German-American kindred (Family C) with late onset parkinsonism. <i>Acta Neuropathologica</i> , 2002, 103, 344-350.	3.9	17
771	Ballooned neurons in progressive supranuclear palsy are usually due to concurrent argyrophilic grain disease. <i>Acta Neuropathologica</i> , 2002, 104, 53-56.	3.9	64
772	Tau accumulation in astrocytes in progressive supranuclear palsy is a degenerative rather than a reactive process. <i>Acta Neuropathologica</i> , 2002, 104, 398-402.	3.9	90
773	Contrasting genotypes of the tau gene in two phenotypically distinct patients with P301L mutation of frontotemporal dementia and parkinsonism linked to chromosome 17. <i>Journal of Neurology</i> , 2002, 249, 669-675.	1.8	19
774	Assembly of tau in transgenic animals expressing P301L tau: alteration of phosphorylation and solubility. <i>Journal of Neurochemistry</i> , 2002, 83, 1498-1508.	2.1	122

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775	The Distribution and Biochemical Properties of a Cdc2-Related Kinase, KKIALLRE, in Normal and Alzheimer Brains. <i>Journal of Neurochemistry</i> , 2002, 65, 2577-2584.	2.1	31
776	Argyrophilic Grain Disease: Neuropathology, Frequency in a Dementia Brain Bank and Lack of Relationship with Apolipoprotein E. <i>Brain Pathology</i> , 2002, 12, 45-52.	2.1	88
777	Misfolded, protease-resistant proteins in animal models and human neurodegenerative disease. <i>Journal of Clinical Investigation</i> , 2002, 110, 1403-1405.	3.9	6
778	Enhanced Neurofibrillary Degeneration in Transgenic Mice Expressing Mutant Tau and APP. <i>Science</i> , 2001, 293, 1487-1491.	6.0	1,409
779	Transfected synphilin-1 forms cytoplasmic inclusions in HEK293 cells. <i>Molecular Brain Research</i> , 2001, 97, 94-102.	2.5	57
780	Neuropathology of Alzheimer's disease and other dementias. <i>Clinics in Geriatric Medicine</i> , 2001, 17, 209-228.	1.0	97
781	Î±-Synuclein and the Lewy body disorders. <i>Current Opinion in Neurology</i> , 2001, 14, 423-432.	1.8	153
782	Frontal Lobe Dementia With Novel Tauopathy: Sporadic Multiple System Tauopathy With Dementia. <i>Journal of Neuropathology and Experimental Neurology</i> , 2001, 60, 328-341.	0.9	83
783	Relationship of the extended tau haplotype to tau biochemistry and neuropathology in progressive supranuclear palsy. <i>Annals of Neurology</i> , 2001, 50, 494-502.	2.8	73
784	Cotton Wool Plaques in Non-Familial Late-Onset Alzheimer Disease. <i>Journal of Neuropathology and Experimental Neurology</i> , 2001, 60, 1051-1061.	0.9	57
785	Microglial Activation parallels System Degeneration in progressive Supranuclear palsy and Corticobasal Degeneration. <i>Journal of Neuropathology and Experimental Neurology</i> , 2001, 60, 647-657.	0.9	176
786	Progressive Supranuclear Palsy and Corticobasal Degeneration. , 2001, , 155-171.		7
787	A Qualitative and Quantitative Study of Grumose Degeneration in Progressive Supranuclear Palsy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2000, 59, 513-524.	0.9	50
788	Induction of Alzheimer-specific tau epitope AT100 in apoptotic human fetal astrocytes. <i>Cytoskeleton</i> , 2000, 47, 236-252.	4.4	20
789	Research goals in progressive supranuclear palsy. <i>Movement Disorders</i> , 2000, 15, 446-458.	2.2	29
790	Neurofibrillary tangles, amyotrophy and progressive motor disturbance in mice expressing mutant (P301L) tau protein. <i>Nature Genetics</i> , 2000, 25, 402-405.	9.4	1,254
791	Tau and Synuclein and Their Role in Neuropathology. <i>Brain Pathology</i> , 1999, 9, 657-661.	2.1	75
792	Multiple System Atrophy: A Sporadic Synucleinopathy. <i>Brain Pathology</i> , 1999, 9, 721-732.	2.1	176

#	ARTICLE	IF	CITATIONS
793	Distinguishing primary angiitis of the central nervous system from cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy: The importance of family history. <i>Arthritis and Rheumatism</i> , 1999, 42, 2243-2248.	6.7	21
794	The Levels of Soluble versus Insoluble Brain A β Distinguish Alzheimer's Disease from Normal and Pathologic Aging. <i>Experimental Neurology</i> , 1999, 158, 328-337.	2.0	490
795	Inducible nitric oxide synthase expression is selectively induced in astrocytes isolated from adult human brain. <i>Brain Research</i> , 1998, 813, 402-405.	1.1	70
796	Polyglutamine-containing aggregates in neuronal intranuclear inclusion disease. <i>Lancet, The</i> , 1998, 351, 884.	6.3	54
797	Pick's Disease: A Modern Approach. <i>Brain Pathology</i> , 1998, 8, 339-354.	2.1	145
798	The Pathogenesis of Senile Plaques. <i>Journal of Neuropathology and Experimental Neurology</i> , 1997, 56, 321-339.	0.9	619
799	Chapter 3 Structural Changes in the Aged Brain. <i>Advances in Cell Aging and Gerontology</i> , 1997, , 51-76.	0.1	0
800	Enhanced binding of advanced glycation endproducts (AGE) by the ApoE4 isoform links the mechanism of plaque deposition in Alzheimer's disease. <i>Neuroscience Letters</i> , 1997, 226, 155-158.	1.0	58
801	Paired helical filaments in corticobasal degeneration: the fine fibrillary structure with NanoVan. <i>Brain Research</i> , 1997, 773, 33-44.	1.1	23
802	Genetic evidence for the involvement of ? in progressive supranuclear palsy. <i>Annals of Neurology</i> , 1997, 41, 277-281.	2.8	433
803	Neurodegenerative diseases with cytoskeletal pathology: A biochemical classification. <i>Annals of Neurology</i> , 1997, 42, 541-544.	2.8	81
804	Monoclonal antibodies to purified cortical lewy bodies recognize the mid-size neurofilament subunit. <i>Annals of Neurology</i> , 1997, 42, 595-603.	2.8	48
805	Glycation and microglial reaction in lesions of Alzheimer's disease. <i>Neurobiology of Aging</i> , 1996, 17, 733-743.	1.5	79
806	Senile cerebral amyloidosis (pathological aging) and cognitive status predictions: A neuropathology perspective. <i>Neurobiology of Aging</i> , 1996, 17, 936-937.	1.5	7
807	Pathology of cryptococcal meningoencephalitis: Analysis of 27 patients with pathogenetic implications. <i>Human Pathology</i> , 1996, 27, 839-847.	1.1	201
808	Neuropathologic Overlap of Progressive Supranuclear Palsy, Pick's Disease and Corticobasal Degeneration. <i>Journal of Neuropathology and Experimental Neurology</i> , 1996, 55, 53-67.	0.9	248
809	Molecular basis of phenotypic variability in sporadic Creutzfeldt-Jakob disease. <i>Annals of Neurology</i> , 1996, 39, 767-778.	2.8	819
810	Neurodegenerative disorders with extensive tau pathology: A comparative study and review. <i>Annals of Neurology</i> , 1996, 40, 139-148.	2.8	301

#	ARTICLE	IF	CITATIONS
811	Multicystic Encephalopathy. <i>Journal of Neuropathology and Experimental Neurology</i> , 1995, 54, 268-275.	0.9	20
812	The Role of Microglia and Astrocytes in Amyloid Deposition in Alzheimer's Disease. , 1995, , 108-127.		0
813	Correlations of synaptic and pathological markers with cognition of the elderly. <i>Neurobiology of Aging</i> , 1995, 16, 285-298.	1.5	391
814	Mismatch between plaques and tangles in staging Alzheimer pathology. <i>Neurobiology of Aging</i> , 1995, 16, 283-284.	1.5	2
815	Authors' response to commentaries. <i>Neurobiology of Aging</i> , 1995, 16, 302-304.	1.5	0
816	In Human Fetal Astrocytes Exposure to Interleukin-1 ² Stimulates Acquisition of the GD3 ⁺ Phenotype and Inhibits Cell Division. <i>Journal of Neurochemistry</i> , 1995, 64, 1800-1807.	2.1	15
817	NONHEREDITARY DIFFUSE LEUKOENCEPHALOPATHY WITH SPHEROIDS PRESENTING AS EARLY-ONSET, RAPIDLY-PROGRESSIVE DEMENTIA. <i>Journal of Neuropathology and Experimental Neurology</i> , 1995, 54, 471.	0.9	13
818	Microglia in HIV-Related CNS Neuropathology:. <i>Journal of Neuro-AIDS</i> , 1995, 1, 57-83.	0.2	15
819	GM-CSF promotes proliferation of human fetal and adult microglia in primary cultures. <i>Glia</i> , 1994, 12, 309-318.	2.5	197
820	Tau immunoreactivity and SDS solubility of two populations of paired helical filaments that differ in morphology. <i>Brain Research</i> , 1994, 649, 185-196.	1.1	30
821	Amino Acid Residues 226-240 of τ , Which Encompass the First Lysine-Pro Site of τ , Are Partially Phosphorylated in Alzheimer Paired Helical Filaments. <i>Journal of Neurochemistry</i> , 1994, 62, 1055-1061.	2.1	13
822	Pathological markers associated with normal aging and dementia in the elderly. <i>Annals of Neurology</i> , 1993, 34, 566-573.	2.8	166
823	Microglia and cytokines in neurological disease, with special reference to AIDS and Alzheimer's disease. <i>Glia</i> , 1993, 7, 75-83.	2.5	828
824	Neuroaxonal dystrophy in HTLV-1-associated myelopathy/tropical spastic paraparesis: neuropathologic and neuroimmunologic correlations. <i>Acta Neuropathologica</i> , 1993, 86, 224-235.	3.9	58
825	Central Nervous System Pathology in Pediatric AIDS. <i>Annals of the New York Academy of Sciences</i> , 1993, 693, 93-106.	1.8	33
826	Productive Infection of Human Fetal Microglia in Vitro by HIV-1. <i>Annals of the New York Academy of Sciences</i> , 1993, 693, 314-316.	1.8	25
827	Pathology and Biology of the Lewy Body. <i>Journal of Neuropathology and Experimental Neurology</i> , 1993, 52, 183-191.	0.9	356
828	Regional synaptic pathology in Alzheimer's disease. <i>Neurobiology of Aging</i> , 1992, 13, 375-382.	1.5	175

#	ARTICLE	IF	CITATIONS
829	Identification of normal and pathological aging in prospectively studied nondemented elderly humans. <i>Neurobiology of Aging</i> , 1992, 13, 179-189.	1.5	580
830	Ubiquitin immunoreactivity in kuru plaques in Creutzfeldt-Jakob disease. <i>Annals of Neurology</i> , 1990, 28, 174-177.	2.8	19
831	Primary central nervous system lymphoma in a pediatric patient with acquired immune deficiency syndrome: Treatment with radiation therapy. <i>Cancer</i> , 1990, 66, 2503-2508.	2.0	25
832	Immunohistochemical Localization of an Hiv Epitope in Cerebral Aneurysmal Arteriopathy in Pediatric Acquired Immunodeficiency Syndrome (AIDS). <i>Pediatric Pathology</i> , 1989, 9, 655-667.	0.5	83
833	Cerebral Granular Cell Tumor. <i>Journal of Neuropathology and Experimental Neurology</i> , 1986, 45, 304-316.	0.9	47
834	Postencephalitic Parkinsonism. , 0, , 179-187.		2