Dennis W Dickson

List of Publications by Year
in descending order

834	111,633	179 152	286 294
papers	citations	h-index	g-index

899	899	899	59675
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Expanded GGGGCC Hexanucleotide Repeat in Noncoding Region of C9ORF72 Causes Chromosome 9p-Linked FTD and ALS. Neuron, 2011, 72, 245-256.	3.8	4,176
2	Diagnosis and management of dementia with Lewy bodies. Neurology, 2017, 89, 88-100.	1.5	2,805
3	Mutations in LRRK2 Cause Autosomal-Dominant Parkinsonism with Pleomorphic Pathology. Neuron, 2004, 44, 601-607.	3.8	2,653
4	Clinical diagnostic criteria for dementia associated with Parkinson's disease. Movement Disorders, 2007, 22, 1689-1707.	2.2	2,497
5	National Institute on Agingâ€“Alzheimerâ€™s Association guidelines for the neuropathologic assessment of Alzheimerâ€™s disease: a practical approach. Acta Neuropathologica, 2012, 123, 1-11.	3.9	2,002
6	National Institute on Agingâ€“Alzheimer's Association guidelines for the neuropathologic assessment of Alzheimer's disease. Alzheimer's and Dementia, 2012, 8, 1-13.	0.4	1,968
7	Genetic meta-analysis of diagnosed Alzheimerâ€™s disease identifies new risk loci and implicates AÎ², tau, immunity and lipid processing. Nature Genetics, 2019, 51, 414-430.	9.4	1,962
8	Common variants at ABCA7, MS4A6A/MS4A4E, EPHA1, CD33 and CD2AP are associated with Alzheimer's disease. Nature Genetics, 2011, 43, 429-435.	9.4	1,708
9	Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. Nature Genetics, 2011, 43, 436-441.	9.4	1,676
10	The neuropathological diagnosis of Alzheimerâ€™s disease. Molecular Neurodegeneration, 2019, 14, 32.	4.4	1,497
11	Criteria for the diagnosis of corticobasal degeneration. Neurology, 2013, 80, 496-503.	1.5	1,445
12	Enhanced Neurofibrillary Degeneration in Transgenic Mice Expressing Mutant Tau and APP. Science, 2001, 293, 1487-1491.	6.0	1,409
13	Clinical diagnosis of progressive supranuclear palsy: The movement disorder society criteria. Movement Disorders, 2017, 32, 853-864.	2.2	1,402
14	Neurofibrillary tangles, amyotrophy and progressive motor disturbance in mice expressing mutant (P301L) tau protein. Nature Genetics, 2000, 25, 402-405.	9.4	1,254
15	Genome-wide Analysis of Genetic Loci Associated With Alzheimer Disease. JAMA - Journal of the American Medical Association, 2010, 303, 1832.	3.8	1,064
16	Primary age-related tauopathy (PART): a common pathology associated with human aging. Acta Neuropathologica, 2014, 128, 755-766.	3.9	1,060
17	Neuropathologic diagnostic and nosologic criteria for frontotemporal lobar degeneration: consensus of the Consortium for Frontotemporal Lobar Degeneration. Acta Neuropathologica, 2007, 114, 5-22.	3.9	978
18	Unconventional Translation of C9ORF72 GGGGCC Expansion Generates Insoluble Polypeptides Specific to c9FTD/ALS. Neuron, 2013, 77, 639-646.	3.8	962

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19	Limbic-predominant age-related TDP-43 encephalopathy (LATE): consensus working group report. Brain, 2019, 142, 1503-1527.	3.7	873
20	Pathology of Neurodegenerative Diseases. Cold Spring Harbor Perspectives in Biology, 2017, 9, a028035.	2.3	865
21	Nomenclature and nosology for neuropathologic subtypes of frontotemporal lobar degeneration: an update. Acta Neuropathologica, 2010, 119, 1-4.	3.9	854
22	Microglia and cytokines in neurological disease, with special reference to AIDS and Alzheimer's disease. Glia, 1993, 7, 75-83.	2.5	828
23	Molecular basis of phenotypic variability in sporadc creudeldt-jakob disease. Annals of Neurology, 1996, 39, 767-778.	2.8	819
24	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. Nature Genetics, 2017, 49, 1373-1384.	9.4	783
25	TDP-43 immunoreactivity in hippocampal sclerosis and Alzheimer's disease. Annals of Neurology, 2007, 61, 435-445.	2.8	753
26	Neuropathological assessment of Parkinson's disease: refining the diagnostic criteria. Lancet Neurology, The, 2009, 8, 1150-1157.	4.9	734
27	Neuropathologically defined subtypes of Alzheimer's disease with distinct clinical characteristics: a retrospective study. Lancet Neurology, The, 2011, 10, 785-796.	4.9	733
28	ALS/FTD Mutation-Induced Phase Transition of FUS Liquid Droplets and Reversible Hydrogels into Irreversible Hydrogels Impairs RNP Granule Function. Neuron, 2015, 88, 678-690.	3.8	716
29	The first NINDS/NIBIB consensus meeting to define neuropathological criteria for the diagnosis of chronic traumatic encephalopathy. Acta Neuropathologica, 2016, 131, 75-86.	3.9	708
30	Clinicopathological and imaging correlates of progressive aphasia and apraxia of speech. Brain, 2006, 129, 1385-1398.	3.7	624
31	The Pathogenesis of Senile Plaques. Journal of Neuropathology and Experimental Neurology, 1997, 56, 321-339.	0.9	619
32	Parkinson's Disease and Parkinsonism: Neuropathology. Cold Spring Harbor Perspectives in Medicine, 2012, 2, a009258-a009258.	2.9	593
33	Identification of normal and pathological aging in prospectively studied nondemented elderly humans. Neurobiology of Aging, 1992, 13, 179-189.	1.5	580
34	Neuropathologic Features of Amnestic Mild Cognitive Impairment. Archives of Neurology, 2006, 63, 665.	4.9	562
35	Neuropathology of nondemented aging: Presumptive evidence for preclinical Alzheimer disease. Neurobiology of Aging, 2009, 30, 1026-1036.	1.5	558
36	Large-scale proteomic analysis of Alzheimerâ€™s disease brain and cerebrospinal fluid reveals early changes in energy metabolism associated with microglia and astrocyte activation. Nature Medicine, 2020, 26, 769-780.	15.2	547

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37	The high-affinity HSP90-CHIP complex recognizes and selectively degrades phosphorylated tau client proteins. Journal of Clinical Investigation, 2007, 117, 648-658.	3.9	545
38	AÎ²42 Is Essential for Parenchymal and Vascular Amyloid Deposition in Mice. Neuron, 2005, 47, 191-199.	3.8	524
39	Aberrant cleavage of TDP-43 enhances aggregation and cellular toxicity. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 7607-7612.	3.3	523
40	Antisense transcripts of the expanded C9ORF72 hexanucleotide repeat form nuclear RNA foci and undergo repeat-associated non-ATG translation in c9FTD/ALS. Acta Neuropathologica, 2013, 126, 829-844.	3.9	506
41	Identification of common variants influencing risk of the tauopathy progressive supranuclear palsy. Nature Genetics, 2011, 43, 699-705.	9.4	502
42	TIA1 Mutations in Amyotrophic Lateral Sclerosis and Frontotemporal Dementia Promote Phase Separation and Alter Stress Granule Dynamics. Neuron, 2017, 95, 808-816.e9.	3.8	493
43	The Levels of Soluble versus Insoluble Brain AÎ² Distinguish Alzheimer's Disease from Normal and Pathologic Aging. Experimental Neurology, 1999, 158, 328-337.	2.0	490
44	Common variants at 7p21 are associated with frontotemporal lobar degeneration with TDP-43 inclusions. Nature Genetics, 2010, 42, 234-239.	9.4	479
45	ERâ€“mitochondria associations are regulated by the VAPBâ€“PTPIP51 interaction and are disrupted by ALS/FTD-associated TDP-43. Nature Communications, 2014, 5, 3996.	5.8	463
46	Wild-Type Human TDP-43 Expression Causes TDP-43 Phosphorylation, Mitochondrial Aggregation, Motor Deficits, and Early Mortality in Transgenic Mice. Journal of Neuroscience, 2010, 30, 10851-10859.	1.7	457
47	Genetic evidence for the involvement of ? in progressive supranuclear palsy. Annals of Neurology, 1997, 41, 277-281.	2.8	433
48	Mutations in the colony stimulating factor 1 receptor (CSF1R) gene cause hereditary diffuse leukoencephalopathy with spheroids. Nature Genetics, 2012, 44, 200-205.	9.4	428
49	Strikingly Different Clinicopathological Phenotypes Determined by Progranulin-Mutation Dosage. American Journal of Human Genetics, 2012, 90, 1102-1107.	2.6	414
50	TDP-43 pathology disrupts nuclear pore complexes and nucleocytoplasmic transport in ALS/FTD. Nature Neuroscience, 2018, 21, 228-239.	7.1	404
51	Novel Mutations in TARDBP (TDP-43) in Patients with Familial Amyotrophic Lateral Sclerosis. PLoS Genetics, 2008, 4, e1000193.	1.5	393
52	Correlations of synaptic and pathological markers with cognition of the elderly. Neurobiology of Aging, 1995, 16, 285-298.	1.5	391
53	An autoradiographic evaluation of AV-1451 Tau PET in dementia. Acta Neuropathologica Communications, 2016, 4, 58.	2.4	388
54	Aging-related tau astrogliopathy (ARTAG): harmonized evaluation strategy. Acta Neuropathologica, 2016, 131, 87-102.	3.9	380

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55	Nomenclature for neuropathologic subtypes of frontotemporal lobar degeneration: consensus recommendations. Acta Neuropathologica, 2009, 117, 15-18.	3.9	377
56	Neuropathological background of phenotypical variability in frontotemporal dementia. Acta Neuropathologica, 2011, 122, 137-153.	3.9	375
57	Parkinson Disease Neuropathology. Archives of Neurology, 2002, 59, 102.	4.9	366
58	A yeast functional screen predicts new candidate ALS disease genes. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 20881-20890.	3.3	365
59	Neuropathology of Parkinson disease. Parkinsonism and Related Disorders, 2018, 46, S30-S33.	1.1	363
60	Human whole genome genotype and transcriptome data for Alzheimerâ€™s and other neurodegenerative diseases. Scientific Data, 2016, 3, 160089.	2.4	361
61	Neuropathology underlying clinical variability in patients with synucleinopathies. Acta Neuropathologica, 2011, 122, 187-204.	3.9	357
62	Pathology and Biology of the Lewy Body. Journal of Neuropathology and Experimental Neurology, 1993, 52, 183-191.	0.9	356
63	Neuroimaging signatures of frontotemporal dementia genetics: C9ORF72, tau, progranulin and sporadics. Brain, 2012, 135, 794-806.	3.7	355
64	Neuroimaging correlates of pathologically defined subtypes of Alzheimer's disease: a case-control study. Lancet Neurology, The, 2012, 11, 868-877.	4.9	355
65	Plasma progranulin levels predict progranulin mutation status in frontotemporal dementia patients and asymptomatic family members. Brain, 2009, 132, 583-591.	3.7	344
66	CNS small vessel disease. Neurology, 2019, 92, 1146-1156.	1.5	343
67	TDP-43 is a key player in the clinical features associated with Alzheimerâ€™s disease. Acta Neuropathologica, 2014, 127, 811-824.	3.9	336
68	Dimeric Amyloid Â Protein Rapidly Accumulates in Lipid Rafts followed by Apolipoprotein E and Phosphorylated Tau Accumulation in the Tg2576 Mouse Model of Alzheimer's Disease. Journal of Neuroscience, 2004, 24, 3801-3809.	1.7	334
69	C9ORF72 repeat expansions in mice cause TDP-43 pathology, neuronal loss, and behavioral deficits. Science, 2015, 348, 1151-1154.	6.0	332
70	Distinct brain transcriptome profiles in C9orf72-associated and sporadic ALS. Nature Neuroscience, 2015, 18, 1175-1182.	7.1	330
71	Evidence that incidental Lewy body disease is pre-symptomatic Parkinsonâ€™s disease. Acta Neuropathologica, 2008, 115, 437-444.	3.9	329
72	AÂ40 Inhibits Amyloid Deposition In Vivo. Journal of Neuroscience, 2007, 27, 627-633.	1.7	327

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73	TREM2 in neurodegeneration: evidence for association of the p.R47H variant with frontotemporal dementia and Parkinsonâ€™s disease. Molecular Neurodegeneration, 2013, 8, 19.	4.4	323
74	Characterization of frontotemporal dementia and/or amyotrophic lateral sclerosis associated with the GGGGCC repeat expansion in C9ORF72. Brain, 2012, 135, 765-783.	3.7	322
75	Neuropathology of variants of progressive supranuclear palsy. Current Opinion in Neurology, 2010, 23, 394-400.	1.8	312
76	Colocalization of Tau and Alpha-Synuclein Epitopes in Lewy Bodies. Journal of Neuropathology and Experimental Neurology, 2003, 62, 389-397.	0.9	306
77	Genome-Wide Association Meta-analysis of Neuropathologic Features of Alzheimer's Disease and Related Dementias. PLoS Genetics, 2014, 10, e1004606.	1.5	305
78	Frontotemporal dementia and its subtypes: a genome-wide association study. Lancet Neurology, The, 2014, 13, 686-699.	4.9	302
79	Neurodegenerative disorders with extensive tau pathology: A comparative study and review. Annals of Neurology, 1996, 40, 139-148.	2.8	301
80	Posttranslational Modifications Mediate the Structural Diversity of Tauopathy Strains. Cell, 2020, 180, 633-644.e12.	13.5	300
81	Age-specific and Sex-specific Prevalence and Incidence of Mild Cognitive Impairment, Dementia, and Alzheimer Dementia in Blacks and Whites. Alzheimer Disease and Associated Disorders, 2012, 26, 335-343.	0.6	297
82	Neuropathology of Frontotemporal Lobar Degeneration-Tau (FTLD-Tau). Journal of Molecular Neuroscience, 2011, 45, 384-389.	1.1	295
83	Association of LRRK2 exonic variants with susceptibility to Parkinson's disease: a caseâ€“control study. Lancet Neurology, The, 2011, 10, 898-908.	4.9	294
84	DCTN1 mutations in Perry syndrome. Nature Genetics, 2009, 41, 163-165.	9.4	285
85	Aggregation-prone c9FTD/ALS poly(GA) RAN-translated proteins cause neurotoxicity by inducing ER stress. Acta Neuropathologica, 2014, 128, 505-524.	3.9	284
86	Alzheimer Disease With Amygdala Lewy Bodies. Journal of Neuropathology and Experimental Neurology, 2006, 65, 685-697.	0.9	279
87	Genetic variation in PCDH11X is associated with susceptibility to late-onset Alzheimer's disease. Nature Genetics, 2009, 41, 192-198.	9.4	279
88	Staging TDP-43 pathology in Alzheimerâ€™s disease. Acta Neuropathologica, 2014, 127, 441-450.	3.9	278
89	Distinct anatomical subtypes of the behavioural variant of frontotemporal dementia: a cluster analysis study. Brain, 2009, 132, 2932-2946.	3.7	277
90	Clinical and neuropathologic heterogeneity of c9FTD/ALS associated with hexanucleotide repeat expansion in C9ORF72. Acta Neuropathologica, 2011, 122, 673-690.	3.9	277

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91	Clinical Correlations With Lewy Body Pathology inLRRK2-Related Parkinson Disease. JAMA Neurology, 2015, 72, 100.	4.5	272
92	When DLB, PD, and PSP masquerade as MSA. Neurology, 2015, 85, 404-412.	1.5	272
93	Common variation in the miR-659 binding-site of GRN is a major risk factor for TDP43-positive frontotemporal dementia. Human Molecular Genetics, 2008, 17, 3631-3642.	1.4	271
94	Corticobasal degeneration: a pathologically distinct 4R tauopathy. Nature Reviews Neurology, 2011, 7, 263-272.	4.9	270
95	Clinicopathologic and¹¹C-Pittsburgh compound B implications of Thal amyloid phase across the Alzheimerâ€™s disease spectrum. Brain, 2015, 138, 1370-1381.	3.7	270
96	C9ORF72 poly(GA) aggregates sequester and impair HR23 and nucleocytoplasmic transport proteins. Nature Neuroscience, 2016, 19, 668-677.	7.1	268
97	Reduced C9orf72 gene expression in c9FTD/ALS is caused by histone trimethylation, an epigenetic event detectable in blood. Acta Neuropathologica, 2013, 126, 895-905.	3.9	263
98	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
99	Tangential Flow Filtration for Highly Efficient Concentration of Extracellular Vesicles from Large Volumes of Fluid. Cells, 2018, 7, 273.	1.8	262
100	Lrrk2 and Lewy body disease. Annals of Neurology, 2006, 59, 388-393.	2.8	259
101	The ALS disease-associated mutant TDP-43 impairs mitochondrial dynamics and function in motor neurons. Human Molecular Genetics, 2013, 22, 4706-4719.	1.4	251
102	A novel human disease with abnormal prion protein sensitive to protease. Annals of Neurology, 2008, 63, 697-708.	2.8	250
103	Translation Initiator EIF4G1 Mutations in Familial Parkinson Disease. American Journal of Human Genetics, 2011, 89, 398-406.	2.6	250
104	Progressive Supranuclear Palsy: Pathology and Genetics. Brain Pathology, 2007, 17, 74-82.	2.1	249
105	C9orf72 BAC Transgenic Mice Display Typical Pathologic Features of ALS/FTD. Neuron, 2015, 88, 892-901.	3.8	249
106	A large-scale comparison of cortical thickness and volume methods for measuring Alzheimer's disease severity. NeuroImage: Clinical, 2016, 11, 802-812.	1.4	249
107	Neuropathologic Overlap of Progressive Supranuclear Palsy, PickÊ¼s Disease and Corticobasal Degeneration. Journal of Neuropathology and Experimental Neurology, 1996, 55, 53-67.	0.9	248
108	AMPK is abnormally activated in tangle- and pre-tangle-bearing neurons in Alzheimerâ€™s disease and other tauopathies. Acta Neuropathologica, 2011, 121, 337-349.	3.9	247

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109	Exceptionally low likelihood of Alzheimerâ€™s dementia in APOE2 homozygotes from a 5,000-person neuropathological study. Nature Communications, 2020, 11, 667.	5.8	246
110	Updated TDP-43 in Alzheimerâ€™s disease staging scheme. Acta Neuropathologica, 2016, 131, 571-585.	3.9	244
111	Poly(GR) impairs protein translation and stress granule dynamics in C9orf72-associated frontotemporal dementia and amyotrophic lateral sclerosis. Nature Medicine, 2018, 24, 1136-1142.	15.2	241
112	Ribosomal Protein s15 Phosphorylation Mediates LRRK2 Neurodegeneration in Parkinsonâ€™s Disease. Cell, 2014, 157, 472-485.	13.5	239
113	Chronic traumatic encephalopathy pathology in a neurodegenerative disorders brain bank. Acta Neuropathologica, 2015, 130, 877-889.	3.9	235
114	Deletion of the Ubiquitin Ligase CHIP Leads to the Accumulation, But Not the Aggregation, of Both Endogenous Phospho- and Caspase-3-Cleaved Tau Species. Journal of Neuroscience, 2006, 26, 6985-6996.	1.7	234
115	Argyrophilic Grain Disease Is a Sporadic 4-Repeat Tauopathy. Journal of Neuropathology and Experimental Neurology, 2002, 61, 547-556.	0.9	232
116	Association between repeat sizes and clinical and pathological characteristics in carriers of C9ORF72 repeat expansions (Xpansize-72): a cross-sectional cohort study. Lancet Neurology, The, 2013, 12, 978-988.	4.9	232
117	Neuropathology of non-motor features of Parkinson disease. Parkinsonism and Related Disorders, 2009, 15, S1-S5.	1.1	228
118	Brain Expression Genome-Wide Association Study (eGWAS) Identifies Human Disease-Associated Variants. PLoS Genetics, 2012, 8, e1002707.	1.5	225
119	FUS pathology defines the majority of tau- and TDP-43-negative frontotemporal lobar degeneration. Acta Neuropathologica, 2010, 120, 33-41.	3.9	222
120	Voxel-based morphometry in autopsy proven PSP and CBD. Neurobiology of Aging, 2008, 29, 280-289.	1.5	221
121	Actin-binding Proteins Coronin-1a and IBA-1 are Effective Microglial Markers for Immunohistochemistry. Journal of Histochemistry and Cytochemistry, 2007, 55, 687-700.	1.3	214
122	An inhibitor of tau hyperphosphorylation prevents severe motor impairments in tau transgenic mice. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 9673-9678.	3.3	206
123	Pathology of cryptococcal meningoencephalitis: Analysis of 27 patients with pathogenetic implications. Human Pathology, 1996, 27, 839-847.	1.1	201
124	Evidence for a role of the rare p.A152T variant in MAPT in increasing the risk for FTD-spectrum and Alzheimer's diseases. Human Molecular Genetics, 2012, 21, 3500-3512.	1.4	198
125	Genome sequencing analysis identifies new loci associated with Lewy body dementia and provides insights into its genetic architecture. Nature Genetics, 2021, 53, 294-303.	9.4	198
126	GM-CSF promotes proliferation of human fetal and adult microglia in primary cultures. Glia, 1994, 12, 309-318.	2.5	197

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127	Investigating the genetic architecture of dementia with Lewy bodies: a two-stage genome-wide association study. Lancet Neurology, The, 2018, 17, 64-74.	4.9	195
128	Progranulin in frontotemporal lobar degeneration and neuroinflammation. Journal of Neuroinflammation, 2007, 4, 7.	3.1	194
129	TDP-43 represses cryptic exon inclusion in the FTDâ€“ALS gene UNC13A. Nature, 2022, 603, 124-130.	13.7	193
130	Nonamnestic mild cognitive impairment progresses to dementia with Lewy bodies. Neurology, 2013, 81, 2032-2038.	1.5	191
131	Replication of CLU, CR1, and PICALM Associations With Alzheimer Disease. Archives of Neurology, 2010, 67, 961-4.	4.9	188
132	Î²â€amyloid burden is not associated with rates of brain atrophy. Annals of Neurology, 2008, 63, 204-212.	2.8	187
133	Association ofGBAMutations and the E326K Polymorphism With Motor and Cognitive Progression in Parkinson Disease. JAMA Neurology, 2016, 73, 1217.	4.5	185
134	Caught in the Act. Neuron, 2003, 40, 453-456.	3.8	184
135	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
136	Antemortem diagnosis of frontotemporal lobar degeneration. Annals of Neurology, 2005, 57, 480-488.	2.8	181
137	Heterochromatin anomalies and double-stranded RNA accumulation underlie C9orf72 poly(PR) toxicity. Science, 2019, 363, .	6.0	181
138	Poly(GP) proteins are a useful pharmacodynamic marker for C9ORF72 -associated amyotrophic lateral sclerosis. Science Translational Medicine, 2017, 9, .	5.8	179
139	Genetic analysis implicates APOE, SNCA and suggests lysosomal dysfunction in the etiology of dementia with Lewy bodies. Human Molecular Genetics, 2014, 23, 6139-6146.	1.4	178
140	Multiple System Atrophy: A Sporadic Synucleinopathy. Brain Pathology, 1999, 9, 721-732.	2.1	176
141	Microglial Activation parallels System Degeneration in progressive Supranuclear palsy and Corticobasal Degeneration. Journal of Neuropathology and Experimental Neurology, 2001, 60, 647-657.	0.9	176
142	Neurofilament inclusion body disease: a new proteinopathy?. Brain, 2003, 126, 2291-2303.	3.7	176
143	Regional synaptic pathology in Alzheimer's disease. Neurobiology of Aging, 1992, 13, 375-382.	1.5	175
144	CCNF mutations in amyotrophic lateral sclerosis and frontotemporal dementia. Nature Communications, 2016, 7, 11253.	5.8	174

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145	Neuropsychological Differentiation of Dementia with Lewy Bodies from Normal Aging and Alzheimer's Disease. Clinical Neuropsychologist, 2006, 20, 623-636.	1.5	170
146	Genome-wide association study of corticobasal degeneration identifies risk variants shared with progressive supranuclear palsy. Nature Communications, 2015, 6, 7247.	5.8	170
147	Globular glial tauopathies (GGT): consensus recommendations. Acta Neuropathologica, 2013, 126, 537-544.	3.9	168
148	Cardiac sympathetic denervation correlates with clinical and pathologic stages of Parkinson's disease. Movement Disorders, 2008, 23, 1085-1092.	2.2	167
149	Pathological markers associated with normal aging and dementia in the elderly. Annals of Neurology, 1993, 34, 566-573.	2.8	166
150	Incidental Lewy Body Disease and Preclinical Parkinson Disease. Archives of Neurology, 2008, 65, 1074-80.	4.9	166
151	Prominent phenotypic variability associated with mutations in Progranulin. Neurobiology of Aging, 2009, 30, 739-751.	1.5	166
152	Nuclear translocation of AMPK-Î±1 potentiates striatal neurodegeneration in Huntingtonâ€™s disease. Journal of Cell Biology, 2011, 194, 209-227.	2.3	166
153	Effects of Multiple Genetic Loci on Age at Onset in Late-Onset Alzheimer Disease. JAMA Neurology, 2014, 71, 1394.	4.5	166
154	Transethnic genomeâ€wide scan identifies novel Alzheimer's disease loci. Alzheimer's and Dementia, 2017, 13, 727-738.	0.4	166
155	LRRK2 knockout mice have an intact dopaminergic system but display alterations in exploratory and motor co-ordination behaviors. Molecular Neurodegeneration, 2012, 7, 25.	4.4	165
156	Alterations in microRNA-124 and AMPA receptors contribute to social behavioral deficits in frontotemporal dementia. Nature Medicine, 2014, 20, 1444-1451.	15.2	165
157	Apoptotic mechanisms in Alzheimer neurofibrillary degeneration: cause or effect?. Journal of Clinical Investigation, 2004, 114, 23-27.	3.9	163
158	Multimodality imaging characteristics of dementia with Lewy bodies. Neurobiology of Aging, 2012, 33, 2091-2105.	1.5	162
159	Rates of hippocampal atrophy and presence of post-mortem TDP-43 in patients with Alzheimer's disease: a longitudinal retrospective study. Lancet Neurology, The, 2017, 16, 917-924.	4.9	159
160	White-matter integrity on DTI and the pathologic staging of Alzheimer's disease. Neurobiology of Aging, 2017, 56, 172-179.	1.5	158
161	Evaluation of Î±-synuclein immunohistochemical methods used by invited experts. Acta Neuropathologica, 2008, 116, 277-288.	3.9	157
162	Clinicopathologic assessment and imaging of tauopathies in neurodegenerative dementias. Alzheimer's Research and Therapy, 2014, 6, 1.	3.0	156

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163	Diagnostic accuracy of progressive supranuclear palsy in the Society for Progressive Supranuclear Palsy Brain Bank. Movement Disorders, 2003, 18, 1018-1026.	2.2	155
164	Î±-Synuclein and the Lewy body disorders. Current Opinion in Neurology, 2001, 14, 423-432.	1.8	153
165	MRI characteristics and scoring in HDLS due to CSF1R gene mutations. Neurology, 2012, 79, 566-574.	1.5	153
166	Distinct binding of PET ligands PBB3 and AV-1451 to tau fibril strains in neurodegenerative tauopathies. Brain, 2017, 140, aww339.	3.7	153
167	Antemortem MRI based STructural Abnormality iNDex (STAND)-scores correlate with postmortem Braak neurofibrillary tangle stage. NeuroImage, 2008, 42, 559-567.	2.1	152
168	Î²-Amyloid Degradation and Alzheimer's Disease. Journal of Biomedicine and Biotechnology, 2006, 2006, 1-12.	3.0	151
169	Alzheimer Disease: Postmortem Neuropathologic Correlates of Antemortem¹H MR Spectroscopy Metabolite Measurements¹. Radiology, 2008, 248, 210-220.	3.6	147
170	Ataxin-2 repeat-length variation and neurodegeneration. Human Molecular Genetics, 2011, 20, 3207-3212.	1.4	147
171	(Pathoâ€)physiological relevance of <scp>PINK</scp> 1â€dependent ubiquitin phosphorylation. EMBO Reports, 2015, 16, 1114-1130.	2.0	147
172	Propagation of tau pathology: hypotheses, discoveries, and yet unresolved questions from experimental and human brain studies. Acta Neuropathologica, 2016, 131, 27-48.	3.9	147
173	Rates of cerebral atrophy differ in different degenerative pathologies. Brain, 2006, 130, 1148-1158.	3.7	146
174	Progranulin Mutations in Primary Progressive Aphasia. Archives of Neurology, 2007, 64, 43.	4.9	146
175	Functional Impact of White Matter Hyperintensities in Cognitively Normal Elderly Subjects. Archives of Neurology, 2010, 67, 1379-85.	4.9	146
176	Pick's Disease: A Modern Approach. Brain Pathology, 1998, 8, 339-354.	2.1	145
177	Expression and functional profiling of neprilysin, insulinâ€degrading enzyme, and endothelinâ€converting enzyme in prospectively studied elderly and Alzheimerâ€™s brain. Journal of Neurochemistry, 2010, 115, 47-57.	2.1	144
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