

David M A Mann

List of Publications by Citations

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188
papers

29,760
citations

70
h-index

172
g-index

195
ext. papers

33,996
ext. citations

9.4
avg, IF

6.16
L-index

#	Paper	IF	Citations
188	A hexanucleotide repeat expansion in C9ORF72 is the cause of chromosome 9p21-linked ALS-FTD. <i>Neuron</i> , 2011 , 72, 257-68	13.9	3018
187	Association of missense and 5Qsplice-site mutations in tau with the inherited dementia FTDP-17. <i>Nature</i> , 1998 , 393, 702-5	50.4	2903
186	Genome-wide association study identifies variants at CLU and PICALM associated with AlzheimerQ disease. <i>Nature Genetics</i> , 2009 , 41, 1088-93	36.3	2018
185	TDP-43 is a component of ubiquitin-positive tau-negative inclusions in frontotemporal lobar degeneration and amyotrophic lateral sclerosis. <i>Biochemical and Biophysical Research Communications</i> , 2006 , 351, 602-11	3.4	1771
184	Mutations in progranulin cause tau-negative frontotemporal dementia linked to chromosome 17. <i>Nature</i> , 2006 , 442, 916-9	50.4	1549
183	Common variants at ABCA7, MS4A6A/MS4A4E, EPHA1, CD33 and CD2AP are associated with AlzheimerQ disease. <i>Nature Genetics</i> , 2011 , 43, 429-35	36.3	1421
182	Genetic meta-analysis of diagnosed AlzheimerQ disease identifies new risk loci and implicates A□ tau, immunity and lipid processing. <i>Nature Genetics</i> , 2019 , 51, 414-430	36.3	917
181	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	14.3	837
180	Nomenclature and nosology for neuropathologic subtypes of frontotemporal lobar degeneration: an update. <i>Acta Neuropathologica</i> , 2010 , 119, 1-4	14.3	711
179	A harmonized classification system for FTLTDP pathology. <i>Acta Neuropathologica</i> , 2011 , 122, 111-3	14.3	656
178	Prion-like spreading of pathological Bsynuclein in brain. <i>Brain</i> , 2013 , 136, 1128-38	11.2	551
177	Magnetite pollution nanoparticles in the human brain. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016 , 113, 10797-801	11.5	516
176	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in AlzheimerQ disease. <i>Nature Genetics</i> , 2017 , 49, 1373-1384	36.3	508
175	Frontotemporal dementia. <i>Lancet Neurology</i> , 2005 , 4, 771-80	24.1	434
174	Distinct clinical and pathological characteristics of frontotemporal dementia associated with C9ORF72 mutations. <i>Brain</i> , 2012 , 135, 693-708	11.2	420
173	Disease-specific patterns of locus coeruleus cell loss. <i>Annals of Neurology</i> , 1992 , 32, 667-76	9.4	402
172	Common variants at 7p21 are associated with frontotemporal lobar degeneration with TDP-43 inclusions. <i>Nature Genetics</i> , 2010 , 42, 234-9	36.3	361

171	Amyloid beta protein (A beta) deposition: A beta 42(43) precedes A beta 40 in Down syndrome. <i>Annals of Neurology</i> , 1995 , 37, 294-9	9.4	351
170	Nomenclature for neuropathologic subtypes of frontotemporal lobar degeneration: consensus recommendations. <i>Acta Neuropathologica</i> , 2009 , 117, 15-8	14.3	325
169	Prion-like properties of pathological TDP-43 aggregates from diseased brains. <i>Cell Reports</i> , 2013 , 4, 124-34.6	14.6	322
168	A multicenter study of glucocerebrosidase mutations in dementia with Lewy bodies. <i>JAMA Neurology</i> , 2013 , 70, 727-35	17.2	285
167	Alzheimer@ presenile dementia, senile dementia of Alzheimer type and Down@ syndrome in middle age form an age related continuum of pathological changes. <i>Neuropathology and Applied Neurobiology</i> , 1984 , 10, 185-207	5.2	283
166	Heterogeneity of ubiquitin pathology in frontotemporal lobar degeneration: classification and relation to clinical phenotype. <i>Acta Neuropathologica</i> , 2006 , 112, 539-49	14.3	264
165	Ubiquitinated pathological lesions in frontotemporal lobar degeneration contain the TAR DNA-binding protein, TDP-43. <i>Acta Neuropathologica</i> , 2007 , 113, 521-33	14.3	252
164	Changes in the monoamine containing neurones of the human CNS in senile dementia. <i>British Journal of Psychiatry</i> , 1980 , 136, 533-41	5.4	248
163	Frontotemporal lobar degeneration: clinical and pathological relationships. <i>Acta Neuropathologica</i> , 2007 , 114, 31-8	14.3	244
162	Progressive language disorder due to lobar atrophy. <i>Annals of Neurology</i> , 1992 , 31, 174-83	9.4	231
161	Frontotemporal dementia and its subtypes: a genome-wide association study. <i>Lancet Neurology</i> , 2014 , 13, 686-99	24.1	207
160	Amyloid beta protein (Abeta) deposition in chromosome 14-linked Alzheimer@ disease: predominance of Abeta42(43). <i>Annals of Neurology</i> , 1996 , 40, 149-56	9.4	196
159	Presynaptic serotonergic dysfunction in patients with Alzheimer@ disease. <i>Journal of Neurochemistry</i> , 1987 , 48, 8-15	6	193
158	Alzheimer@ disease and Down@ syndrome. <i>Histopathology</i> , 1988 , 13, 125-37	7.3	187
157	The clinical diagnosis of early-onset dementias: diagnostic accuracy and clinicopathological relationships. <i>Brain</i> , 2011 , 134, 2478-92	11.2	178
156	Cognitive phenotypes in Alzheimer@ disease and genetic risk. <i>Cortex</i> , 2007 , 43, 835-45	3.8	170
155	Progranulin gene mutations associated with frontotemporal dementia and progressive non-fluent aphasia. <i>Brain</i> , 2006 , 129, 3091-102	11.2	166
154	Frequency and clinical characteristics of progranulin mutation carriers in the Manchester frontotemporal lobar degeneration cohort: comparison with patients with MAPT and no known mutations. <i>Brain</i> , 2008 , 131, 721-31	11.2	163

153	Genetic analysis implicates APOE, SNCA and suggests lysosomal dysfunction in the etiology of dementia with Lewy bodies. <i>Human Molecular Genetics</i> , 2014 , 23, 6139-46	5.6	152
152	Increased TDP-43 protein in cerebrospinal fluid of patients with amyotrophic lateral sclerosis. <i>Acta Neuropathologica</i> , 2009 , 117, 55-62	14.3	148
151	Vascular cognitive impairment neuropathology guidelines (VCING): the contribution of cerebrovascular pathology to cognitive impairment. <i>Brain</i> , 2016 , 139, 2957-2969	11.2	141
150	Pathological basis for neurotransmitter changes in Parkinson's disease. <i>Neuropathology and Applied Neurobiology</i> , 1983 , 9, 3-19	5.2	140
149	Lipoprotein pigments--their relationship to ageing in the human nervous system. I. The lipofuscin content of nerve cells. <i>Brain</i> , 1974 , 97, 481-8	11.2	139
148	Dipeptide repeat proteins are present in the p62 positive inclusions in patients with frontotemporal lobar degeneration and motor neurone disease associated with expansions in C9ORF72. <i>Acta Neuropathologica Communications</i> , 2013 , 1, 68	7.3	131
147	Sporadic Pick's disease: a tauopathy characterized by a spectrum of pathological tau isoforms in gray and white matter. <i>Annals of Neurology</i> , 2002 , 51, 730-9	9.4	130
146	The topography of plaques and tangles in Down's syndrome patients of different ages. <i>Neuropathology and Applied Neurobiology</i> , 1986 , 12, 447-57	5.2	126
145	The progression of the pathological changes of Alzheimer's disease in frontal and temporal neocortex examined both at biopsy and at autopsy. <i>Neuropathology and Applied Neurobiology</i> , 1988 , 14, 177-95	5.2	124
144	Biochemical classification of tauopathies by immunoblot, protein sequence and mass spectrometric analyses of sarkosyl-insoluble and trypsin-resistant tau. <i>Acta Neuropathologica</i> , 2016 , 131, 267-280	14.3	122
143	The prevalence of amyloid (A4) protein deposits within the cerebral and cerebellar cortex in Down's syndrome and Alzheimer's disease. <i>Acta Neuropathologica</i> , 1990 , 80, 318-27	14.3	122
142	The apolipoprotein E epsilon2 allele and the pathological features in cerebral amyloid angiopathy-related hemorrhage. <i>Journal of Neuropathology and Experimental Neurology</i> , 1999 , 58, 711-8 ^{3.1}		121
141	Investigating the genetic architecture of dementia with Lewy bodies: a two-stage genome-wide association study. <i>Lancet Neurology</i> , 2018 , 17, 64-74	24.1	121
140	Histopathological changes underlying frontotemporal lobar degeneration with clinicopathological correlation. <i>Acta Neuropathologica</i> , 2005 , 110, 501-12	14.3	117
139	Pick's disease is associated with mutations in the tau gene. <i>Annals of Neurology</i> , 2000 , 48, 859-867	9.4	116
138	TDP-43 protein in plasma may index TDP-43 brain pathology in Alzheimer's disease and frontotemporal lobar degeneration. <i>Acta Neuropathologica</i> , 2008 , 116, 141-6	14.3	115
137	The selective vulnerability of nerve cells in Huntington's disease. <i>Neuropathology and Applied Neurobiology</i> , 2001 , 27, 1-21	5.2	114
136	Inherited frontotemporal dementia in nine British families associated with intronic mutations in the tau gene. <i>Brain</i> , 2002 , 125, 732-51	11.2	110

135	TDP-43 pathological changes in early onset familial and sporadic Alzheimer disease, late onset Alzheimer disease and Down syndrome: association with age, hippocampal sclerosis and clinical phenotype. <i>Acta Neuropathologica</i> , 2011 , 122, 703-13	14.3	106
134	The most common type of FTL-D-FUS (aFTLD-U) is associated with a distinct clinical form of frontotemporal dementia but is not related to mutations in the FUS gene. <i>Acta Neuropathologica</i> , 2011 , 122, 99-110	14.3	90
133	Apolipoprotein E epsilon 2 allele promotes longevity and protects patients with Down syndrome from dementia. <i>NeuroReport</i> , 1994 , 5, 2583-5	1.7	86
132	Early changes in extracellular matrix in Alzheimer disease. <i>Neuropathology and Applied Neurobiology</i> , 2017 , 43, 167-182	5.2	85
131	Mutations in progranulin explain atypical phenotypes with variants in MAPT. <i>Brain</i> , 2006 , 129, 3124-6	11.2	85
130	Generation and characterization of novel conformation-specific monoclonal antibodies for Bsynuclein pathology. <i>Neurobiology of Disease</i> , 2015 , 79, 81-99	7.5	83
129	An immunohistochemical study of cases of sporadic and inherited frontotemporal lobar degeneration using 3R- and 4R-specific tau monoclonal antibodies. <i>Acta Neuropathologica</i> , 2006 , 111, 329-40	14.3	81
128	The quantitative assessment of lipofuscin pigment, cytoplasmic RNA and nucleolar volume in senile dementia. <i>Neuropathology and Applied Neurobiology</i> , 1978 , 4, 129-35	5.2	80
127	Differential diagnosis of Alzheimer disease using spectrochemical analysis of blood. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, E7929-E7938	11.5	79
126	Phosphorylated TDP-43 pathology and hippocampal sclerosis in progressive supranuclear palsy. <i>Acta Neuropathologica</i> , 2010 , 120, 55-66	14.3	77
125	Frontotemporal lobar degeneration: Pathogenesis, pathology and pathways to phenotype. <i>Brain Pathology</i> , 2017 , 27, 723-736	6	76
124	Molecular analysis and biochemical classification of TDP-43 proteinopathy. <i>Brain</i> , 2012 , 135, 3380-91	11.2	75
123	Dementia lacking distinctive histology (DLDH) revisited. <i>Acta Neuropathologica</i> , 2006 , 112, 551-9	14.3	75
122	Accuracy of single-photon emission computed tomography in differentiating frontotemporal dementia from Alzheimer disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2007 , 78, 350-5	5.5	74
121	The topographic distribution of senile plaques and neurofibrillary tangles in the brains of non-demented persons of different ages. <i>Neuropathology and Applied Neurobiology</i> , 1987 , 13, 123-39	5.2	74
120	Prevalence of amyloid- β pathology in distinct variants of primary progressive aphasia. <i>Annals of Neurology</i> , 2018 , 84, 729-740	9.4	74
119	Frontotemporal dementia with Pick-type histology associated with Q336R mutation in the tau gene. <i>Brain</i> , 2004 , 127, 1415-26	11.2	73
118	Mechanisms of disease in frontotemporal lobar degeneration: gain of function versus loss of function effects. <i>Acta Neuropathologica</i> , 2012 , 124, 373-82	14.3	69

117	An analysis of the morphology of senile plaques in DownQ syndrome patients of different ages using immunocytochemical and lectin histochemical techniques. <i>Neuropathology and Applied Neurobiology</i> , 1989 , 15, 317-29	5.2	66
116	Plasma phosphorylated-TDP-43 protein levels correlate with brain pathology in frontotemporal lobar degeneration. <i>Acta Neuropathologica</i> , 2009 , 118, 647-58	14.3	65
115	Pathological correlates of frontotemporal lobar degeneration in the elderly. <i>Acta Neuropathologica</i> , 2011 , 121, 365-71	14.3	64
114	Early senile plaques in DownQ syndrome brains show a close relationship with cell bodies of neurons. <i>Neuropathology and Applied Neurobiology</i> , 1989 , 15, 531-42	5.2	64
113	The neuropathology of frontotemporal lobar degeneration with respect to the cytological and biochemical characteristics of tau protein. <i>Neuropathology and Applied Neurobiology</i> , 2004 , 30, 1-18	5.2	62
112	Relationships between arteriosclerosis, cerebral amyloid angiopathy and myelin loss from cerebral cortical white matter in AlzheimerQ disease. <i>Neuropathology and Applied Neurobiology</i> , 2004 , 30, 46-56	5.2	62
111	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> , 2018 , 17, 548-558	24.1	60
110	Sensitivity and specificity of FTDC criteria for behavioral variant frontotemporal dementia. <i>Neurology</i> , 2013 , 80, 1881-7	6.5	60
109	The topographic distribution of brain atrophy in frontal lobe dementia. <i>Acta Neuropathologica</i> , 1993 , 85, 334-40	14.3	59
108	Accumulation of dipeptide repeat proteins predates that of TDP-43 in frontotemporal lobar degeneration associated with hexanucleotide repeat expansions in C9ORF72 gene. <i>Neuropathology and Applied Neurobiology</i> , 2015 , 41, 601-12	5.2	53
107	Amyloid (Abeta) deposition in chromosome 1-linked AlzheimerQ disease: the Volga German families. <i>Annals of Neurology</i> , 1997 , 41, 52-7	9.4	53
106	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	14.3	50
105	Patterns of microglial cell activation in frontotemporal lobar degeneration. <i>Neuropathology and Applied Neurobiology</i> , 2014 , 40, 686-96	5.2	50
104	Imbalance of a serotonergic system in frontotemporal dementia: implication for pharmacotherapy. <i>Psychopharmacology</i> , 2008 , 196, 603-10	4.7	50
103	Genome-wide analysis of genetic correlation in dementia with Lewy bodies, ParkinsonQ and AlzheimerQ diseases. <i>Neurobiology of Aging</i> , 2016 , 38, 214.e7-214.e10	5.6	49
102	Neurodegeneration in frontotemporal lobar degeneration and motor neurone disease associated with expansions in C9orf72 is linked to TDP-43 pathology and not associated with aggregated forms of dipeptide repeat proteins. <i>Neuropathology and Applied Neurobiology</i> , 2016 , 42, 242-54	5.2	48
101	The age of onset and evolution of Braak tangle stage and Thal amyloid pathology of AlzheimerQ disease in individuals with Down syndrome. <i>Acta Neuropathologica Communications</i> , 2018 , 6, 56	7.3	48
100	Microglial cells and amyloid beta protein (A beta) deposition; association with A beta 40-containing plaques. <i>Acta Neuropathologica</i> , 1995 , 90, 472-7	14.3	48

99	A morphological analysis of senile plaques in the brains of non-demented persons of different ages using silver, immunocytochemical and lectin histochemical staining techniques. <i>Neuropathology and Applied Neurobiology</i> , 1990 , 16, 17-25	5.2	48
98	Extensive deamidation at asparagine residue 279 accounts for weak immunoreactivity of tau with RD4 antibody in Alzheimer's disease brain. <i>Acta Neuropathologica Communications</i> , 2013 , 1, 54	7.3	47
97	Neurofibrillary pathology and protein synthetic capability in nerve cells in Alzheimer's disease. <i>Neuropathology and Applied Neurobiology</i> , 1981 , 7, 37-47	5.2	47
96	An ultrastructural analysis of the effects of accumulation of neurofibrillary tangle in pyramidal neurons of the cerebral cortex in Alzheimer's disease. <i>Neuropathology and Applied Neurobiology</i> , 1986 , 12, 305-19	5.2	45
95	Frontotemporal dementia with amyotrophic lateral sclerosis: a clinical comparison of patients with and without repeat expansions in C9orf72. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2013 , 14, 172-6	3.6	44
94	Relationships in Alzheimer's disease between the extent of Aβ deposition in cerebral blood vessel walls, as cerebral amyloid angiopathy, and the amount of cerebrovascular smooth muscle cells and collagen. <i>Neuropathology and Applied Neurobiology</i> , 2006 , 32, 332-40	5.2	43
93	Immunohistochemical staining of senile plaques. <i>Neuropathology and Applied Neurobiology</i> , 1982 , 8, 55-61	5.2	42
92	TDP-43 gene analysis in frontotemporal lobar degeneration. <i>Neuroscience Letters</i> , 2007 , 419, 1-4	3.3	40
91	Negative association between amyloid plaques and cerebral amyloid angiopathy in Alzheimer's disease. <i>Neuroscience Letters</i> , 2003 , 352, 137-40	3.3	39
90	Raman Spectroscopy to Diagnose Alzheimer's Disease and Dementia with Lewy Bodies in Blood. <i>ACS Chemical Neuroscience</i> , 2018 , 9, 2786-2794	5.7	38
89	Deposition of amyloid beta protein in non-Alzheimer dementias: evidence for a neuronal origin of parenchymal deposits of beta protein in neurodegenerative disease. <i>Acta Neuropathologica</i> , 1992 , 83, 415-9	14.3	37
88	Analysis of the hexanucleotide repeat in C9ORF72 in Alzheimer's disease. <i>Neurobiology of Aging</i> , 2012 , 33, 1846.e5-6	5.6	36
87	Histone deacetylase class II and acetylated core histone immunohistochemistry in human brains with Huntington's disease. <i>Brain Research</i> , 2013 , 1504, 16-24	3.7	35
86	Patterns and severity of vascular amyloid in Alzheimer's disease associated with duplications and missense mutations in APP gene, Down syndrome and sporadic Alzheimer's disease. <i>Acta Neuropathologica</i> , 2018 , 136, 569-587	14.3	35
85	Patterns of cerebral amyloid angiopathy define histopathological phenotypes in Alzheimer's disease. <i>Neuropathology and Applied Neurobiology</i> , 2014 , 40, 136-48	5.2	34
84	TREM2 analysis and increased risk of Alzheimer's disease. <i>Neurobiology of Aging</i> , 2015 , 36, 546.e9-13	5.6	33
83	Co-Occurrence of Language and Behavioural Change in Frontotemporal Lobar Degeneration. <i>Dementia and Geriatric Cognitive Disorders Extra</i> , 2016 , 6, 205-13	2.5	31
82	DJ-1 (PARK7) is associated with 3R and 4R tau neuronal and glial inclusions in neurodegenerative disorders. <i>Neurobiology of Disease</i> , 2007 , 28, 122-32	7.5	31

81	Comparison of extent of tau pathology in patients with frontotemporal dementia with Parkinsonism linked to chromosome 17 (FTDP-17), frontotemporal lobar degeneration with Pick bodies and early onset Alzheimer's disease. <i>Neuropathology and Applied Neurobiology</i> , 2006 , 32, 374-87	5.2	31
80	Cases of Alzheimer's disease due to deletion of exon 9 of the presenilin-1 gene show an unusual but characteristic beta-amyloid pathology known as cotton wool plaques. <i>Neuropathology and Applied Neurobiology</i> , 2001 , 27, 189-96	5.2	31
79	Amyloid or tau: the chicken or the egg?. <i>Acta Neuropathologica</i> , 2013 , 126, 609-13	14.3	30
78	Ubiquitin associated protein 1 is a risk factor for frontotemporal lobar degeneration. <i>Neurobiology of Aging</i> , 2009 , 30, 656-65	5.6	29
77	The DNA content of Purkinje cells in mammals. <i>Journal of Comparative Neurology</i> , 1978 , 180, 345-7	3.4	29
76	ADAM30 Downregulates APP-Linked Defects Through Cathepsin D Activation in Alzheimer's Disease. <i>EBioMedicine</i> , 2016 , 9, 278-292	8.8	28
75	Dysregulation of C-X-C motif ligand 10 during aging and association with cognitive performance. <i>Neurobiology of Aging</i> , 2018 , 63, 54-64	5.6	28
74	Association study and meta-analysis of low-density lipoprotein receptor related protein in Alzheimer's disease. <i>Neuroscience Letters</i> , 2005 , 382, 221-6	3.3	27
73	Pathological tau deposition in Motor Neurone Disease and frontotemporal lobar degeneration associated with TDP-43 proteinopathy. <i>Acta Neuropathologica Communications</i> , 2016 , 4, 33	7.3	26
72	Polygenic risk score in postmortem diagnosed sporadic early-onset Alzheimer's disease. <i>Neurobiology of Aging</i> , 2018 , 62, 244.e1-244.e8	5.6	25
71	TDP-43 in ubiquitinated inclusions in the inferior olives in frontotemporal lobar degeneration and in other neurodegenerative diseases: a degenerative process distinct from normal ageing. <i>Acta Neuropathologica</i> , 2009 , 118, 359-69	14.3	25
70	Atypical amyloid (A beta) deposition in the cerebellum in Alzheimer's disease: an immunohistochemical study using end-specific A beta monoclonal antibodies. <i>Acta Neuropathologica</i> , 1996 , 91, 647-53	14.3	25
69	Effect of topographical distribution of Synuclein pathology on TDP-43 accumulation in Lewy body disease. <i>Acta Neuropathologica</i> , 2010 , 120, 789-801	14.3	24
68	Plasma levels of progranulin and interleukin-6 in frontotemporal lobar degeneration. <i>Neurobiology of Aging</i> , 2015 , 36, 1603.e1-4	5.6	22
67	Association between apolipoprotein E e4 allele and arteriosclerosis, cerebral amyloid angiopathy, and cerebral white matter damage in Alzheimer's disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2004 , 75, 696-9	5.5	22
66	Semantic dementia, progressive non-fluent aphasia and their association with amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017 , 88, 711-712	5.5	20
65	Symmetric dimethylation of poly-GR correlates with disease duration in C9orf72 FTLN and ALS and reduces poly-GR phase separation and toxicity. <i>Acta Neuropathologica</i> , 2020 , 139, 407-410	14.3	20
64	Pathological evidence for neurotransmitter deficits in Down's syndrome of middle age. <i>Journal of Intellectual Disability Research</i> , 1985 , 29 (Pt 2), 125-35	3.2	20

63	The nucleus basalis of Meynert in multi-infarct (vascular) dementia. <i>Acta Neuropathologica</i> , 1986 , 71, 332-7	14.3	20
62	Granulovacuolar degeneration in pyramidal cells of the hippocampus. <i>Acta Neuropathologica</i> , 1978 , 42, 149-51	14.3	20
61	Pathological Correlates of Cognitive Impairment in The University of Manchester Longitudinal Study of Cognition in Normal Healthy Old Age. <i>Journal of Alzheimers Disease</i> , 2018 , 64, 483-496	4.3	19
60	Comparison of Common and Disease-Specific Post-translational Modifications of Pathological Tau Associated With a Wide Range of Tauopathies. <i>Frontiers in Neuroscience</i> , 2020 , 14, 581936	5.1	18
59	Nuclear carrier and RNA-binding proteins in frontotemporal lobar degeneration associated with fused in sarcoma (FUS) pathological changes. <i>Neuropathology and Applied Neurobiology</i> , 2013 , 39, 157-65	5.2	18
58	Do NIA-AA criteria distinguish Alzheimer's disease from frontotemporal dementia?. <i>Alzheimers and Dementia</i> , 2015 , 11, 207-15	1.2	18
57	Granular expression of prolyl-peptidyl isomerase PIN1 is a constant and specific feature of Alzheimer's disease pathology and is independent of tau, A β and TDP-43 pathology. <i>Acta Neuropathologica</i> , 2011 , 121, 635-49	14.3	18
56	Evidence of a founder effect in families with frontotemporal dementia that harbor the tau +16 splice mutation. <i>American Journal of Medical Genetics Part A</i> , 2004 , 125B, 79-82		18
55	Small deletion in C9orf72 hides a proportion of expansion carriers in FTL. <i>Neurobiology of Aging</i> , 2015 , 36, 1601.e1-5	5.6	17
54	No interaction between tau and TDP-43 pathologies in either frontotemporal lobar degeneration or motor neurone disease. <i>Neuropathology and Applied Neurobiology</i> , 2014 , 40, 844-54	5.2	17
53	Epitope mapping of antibodies against TDP-43 and detection of protease-resistant fragments of pathological TDP-43 in amyotrophic lateral sclerosis and frontotemporal lobar degeneration. <i>Biochemical and Biophysical Research Communications</i> , 2012 , 417, 116-21	3.4	17
52	A 3QTR polymorphism in the oxidized LDL receptor 1 gene increases A β 40 load as cerebral amyloid angiopathy in Alzheimer's disease. <i>Acta Neuropathologica</i> , 2006 , 111, 15-20	14.3	17
51	A quantitative study of the ultrastructure of pyramidal neurons of the cerebral cortex in Alzheimer's disease in relationship to the degree of dementia. <i>Neuropathology and Applied Neurobiology</i> , 1986 , 12, 321-9	5.2	16
50	Nuclear inclusions in Alzheimer's disease. <i>Neuropathology and Applied Neurobiology</i> , 1980 , 6, 245-53	5.2	16
49	Heritability and genetic variance of dementia with Lewy bodies. <i>Neurobiology of Disease</i> , 2019 , 127, 492-501	5.1	15
48	Autopsy proven sporadic frontotemporal dementia due to microvacuolar-type histology, with onset at 21 years of age. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2004 , 75, 1337-9	5.5	15
47	Analysis of neurodegenerative disease-causing genes in dementia with Lewy bodies. <i>Acta Neuropathologica Communications</i> , 2020 , 8, 5	7.3	15
46	Extended post-mortem delay times should not be viewed as a deterrent to the scientific investigation of human brain tissue: a study from the Brains for Dementia Research Network Neuropathology Study Group, UK. <i>Acta Neuropathologica</i> , 2016 , 132, 753-755	14.3	13

45	Heterogeneous ribonuclear protein A3 (hnRNP A3) is present in dipeptide repeat protein containing inclusions in Frontotemporal Lobar Degeneration and Motor Neurone disease associated with expansions in C9orf72 gene. <i>Acta Neuropathologica Communications</i> , 2017 , 5, 31	7.3	13
44	No association between polymorphisms in the lectin-like oxidised low density lipoprotein receptor (ORL1) gene on chromosome 12 and Alzheimer's disease in a UK cohort. <i>Neuroscience Letters</i> , 2004 , 366, 126-9	3.3	13
43	UBQLN2 variant of unknown significance in frontotemporal lobar degeneration. <i>Neurobiology of Aging</i> , 2015 , 36, 546.e15-6	5.6	12
42	The role of lysosomes and autophagosomes in frontotemporal lobar degeneration. <i>Neuropathology and Applied Neurobiology</i> , 2019 , 45, 244-261	5.2	12
41	What's in a name? Neuronal intermediate filament inclusion disease (NIFID), frontotemporal lobar degeneration-intermediate filament (FTLD-IF) or frontotemporal lobar degeneration-fused in sarcoma (FTLD-FUS)? <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2011 , 82, 1412-4	5.5	12
40	Progressive anomia revisited: focal degeneration associated with progranulin gene mutation. <i>Neurocase</i> , 2007 , 13, 366-77	0.8	12
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