Tamas Revesz

List of Publications by Year in Descending Order

193 papers	19,263 citations	71 h-index	137 g-index
208 ext. papers	21,952 ext. citations	10.6 avg, IF	6.24 L-index

#	Paper	IF	Citations
193	Prion-like -synuclein pathology in the brain of infants with Krabbe disease.. *Brain*, **2022**,	11.2	2
192	Age-dependent formation of TMEM106B amyloid filaments in human brains.. *Nature*, **2022**,	50.4	6
191	Dissecting the Phenotype and Genotype of PLA2G6-Related Parkinsonism. *Movement Disorders*, **2021**,	7	8
190	A Clinicopathologic Study of Movement Disorders in Frontotemporal Lobar Degeneration. *Movement Disorders*, **2021**, 36, 632-641	7	2
189	Neuropathological and Biomarker Findings in Parkinson Disease and Alzheimer Disease: From Protein Aggregates to Synaptic Dysfunction. *Journal of Parkinsons Disease*, **2021**, 11, 107-121	5.3	12
188	Genetic determinants of survival in progressive supranuclear palsy: a genome-wide association study. *Lancet Neurology, The*, **2021**, 20, 107-116	24.1	23
187	Structure-based classification of tauopathies. *Nature*, **2021**, 598, 359-363	50.4	59
186	Association of clusterin with the BRI2-derived amyloid molecules ABri and ADan. *Neurobiology of Disease*, **2021**, 158, 105452	7.5	0
185	MOBP and HIP1 in multiple system atrophy: New -synuclein partners in glial cytoplasmic inclusions implicated in the disease pathogenesis. *Neuropathology and Applied Neurobiology*, **2021**, 47, 640-652	5.2	4
184	Fulminant corticobasal degeneration: a distinct variant with predominant neuronal tau aggregates. *Acta Neuropathologica*, **2020**, 139, 717-734	14.3	8
183	Analysis of neurodegenerative disease-causing genes in dementia with Lewy bodies. *Acta Neuropathologica Communications*, **2020**, 8, 5	7.3	15
182	Diagnosis Across the Spectrum of Progressive Supranuclear Palsy and Corticobasal Syndrome. *JAMA Neurology*, **2020**, 77, 377-387	17.2	44
181	Neuronal intranuclear inclusion disease is genetically heterogeneous. *Annals of Clinical and Translational Neurology*, **2020**, 7, 1716-1725	5.3	18
180	Fibrillation and molecular characteristics are coherent with clinical and pathological features of 4-repeat tauopathy caused by MAPT variant G273R. *Neurobiology of Disease*, **2020**, 146, 105079	7.5	0
179	White matter DNA methylation profiling reveals deregulation of HIP1, LMAN2, MOBP, and other loci in multiple system atrophy. *Acta Neuropathologica*, **2020**, 139, 135-156	14.3	21
178	Tau Isoform-Driven CBD Pathology Transmission in Oligodendrocytes in Humanized Tau Mice. *Frontiers in Neurology*, **2020**, 11, 589471	4.1	1
177	A novel mutation in a family with diverse frontotemporal dementia spectrum disorders. *Journal of Physical Education and Sports Management*, **2019**, 5,	2.8	15

176	Heritability and genetic variance of dementia with Lewy bodies. *Neurobiology of Disease*, **2019**, 127, 492-501	7.9	15
175	LATE to the PART-y. *Brain*, **2019**, 142, e47	11.2	25
174	The genetic and clinico-pathological profile of early-onset progressive supranuclear palsy. *Movement Disorders*, **2019**, 34, 1307-1314	7	8
173	Primum non nocere: a call for balance when reporting on CTE. *Lancet Neurology, The*, **2019**, 18, 231-233	24.1	34
172	A comprehensive screening of copy number variability in dementia with Lewy bodies. *Neurobiology of Aging*, **2019**, 75, 223.e1-223.e10	5.6	10
171	Neuroaxonal Dystrophy/Neurodegeneration with Brain Iron Accumulation **2018**, 455-468		
170	Variation at the TRIM11 locus modifies progressive supranuclear palsy phenotype. *Annals of Neurology*, **2018**, 84, 485-496	9.4	28
169	The presubiculum is preserved from neurodegenerative changes in Alzheimer's disease. *Acta Neuropathologica Communications*, **2018**, 6, 62	7.3	3
168	Pathological correlates of white matter hyperintensities in a case of progranulin mutation associated frontotemporal dementia. *Neurocase*, **2018**, 24, 166-174	0.8	24
167	Investigating the genetic architecture of dementia with Lewy bodies: a two-stage genome-wide association study. *Lancet Neurology, The*, **2018**, 17, 64-74	24.1	121
166	The aftermath of boxing revisited: identifying chronic traumatic encephalopathy pathology in the original Corsellis boxer series. *Acta Neuropathologica*, **2018**, 136, 973-974	14.3	19
165	The clinical, neuroanatomical, and neuropathologic phenotype of -associated frontotemporal dementia: A longitudinal case report. *Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring*, **2017**, 6, 75-81	5.2	23
164	Mixed pathologies including chronic traumatic encephalopathy account for dementia in retired association football (soccer) players. *Acta Neuropathologica*, **2017**, 133, 337-352	14.3	128
163	Clinical, pathological and functional characterization of riboflavin-responsive neuropathy. *Brain*, **2017**, 140, 2820-2837	11.2	40
162	1115 Chronic traumatic encephalopathy in retired footballers with dementia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2017**, 88, A1.1-A1	5.5	
161	[P2.041]: PATHOLOGICAL CORRELATES OF WHITE MATTER HYPERINTENSITIES ON CADAVERIC MRI IN PROGRANULIN-ASSOCIATED FRONTOTEMPORAL DEMENTIA **2017**, 13, P805-P805		
160	Association of Autonomic Dysfunction With Disease Progression and Survival in Parkinson Disease. *JAMA Neurology*, **2017**, 74, 970-976	17.2	94
159	Analysis of C9orf72 repeat expansions in a large international cohort of dementia with Lewy bodies. *Neurobiology of Aging*, **2017**, 49, 214.e13-214.e15	5.6	10

158	[P2158]: IS THE PRESUBICULUM PROTECTED FROM NEURODEGENERATIVE CHANGES? A PATHOLOGICAL AND BIOCHEMICAL INVESTIGATION **2017**, 13, P668-P668		
157	Parkinson disease without nigral degeneration: a pathological correlate of scans without evidence of dopaminergic deficit (SWEDD)?. *Journal of Neurology, Neurosurgery and Psychiatry*, **2016**, 87, 633-41	5.5	8
156	A genome-wide association study in multiple system atrophy. *Neurology*, **2016**, 87, 1591-1598	6.5	104
155	The presence of heterogeneous nuclear ribonucleoproteins in frontotemporal lobar degeneration with FUS-positive inclusions. *Neurobiology of Aging*, **2016**, 46, 192-203	5.6	16
154	Astrogliopathy predominates the earliest stage of corticobasal degeneration pathology. *Brain*, **2016**, 139, 3237-3252	11.2	80
153	Clinical phenotype and genetic associations in autosomal dominant familial Alzheimer disease: a case series. *Lancet Neurology, The*, **2016**, 15, 1326-1335	24.1	109
152	Apomorphine: A potential modifier of amyloid deposition in Parkinson disease?. *Movement Disorders*, **2016**, 31, 668-75	7	23
151	Genome-wide analysis of genetic correlation in dementia with Lewy bodies, Parkinson and Alzheimer diseases. *Neurobiology of Aging*, **2016**, 38, 214.e7-214.e10	5.6	49
150	Characterization of tau positron emission tomography tracer [F]AV-1451 binding to postmortem tissue in Alzheimer disease, primary tauopathies, and other dementias. *Alzheimer's and Dementia*, **2016**, 12, 1116-1124	1.2	139
149	Loss of VPS13C Function in Autosomal-Recessive Parkinsonism Causes Mitochondrial Dysfunction and Increases PINK1/Parkin-Dependent Mitophagy. *American Journal of Human Genetics*, **2016**, 98, 500-513	11	225
148	Neuropathological criteria of anti-IgLON5-related tauopathy. *Acta Neuropathologica*, **2016**, 132, 531-43	14.3	107
147	Alterations in global DNA methylation and hydroxymethylation are not detected in Alzheimer disease. *Neuropathology and Applied Neurobiology*, **2015**, 41, 497-506	5.2	61
146	Genetic determinants of white matter hyperintensities and amyloid angiopathy in familial Alzheimer disease. *Neurobiology of Aging*, **2015**, 36, 3140-3151	5.6	46
145	Histological evidence of chronic traumatic encephalopathy in a large series of neurodegenerative diseases. *Acta Neuropathologica*, **2015**, 130, 891-3	14.3	75
144	A 30-unit hexanucleotide repeat expansion in C9orf72 induces pathological lesions with dipeptide-repeat proteins and RNA foci, but not TDP-43 inclusions and clinical disease. *Acta Neuropathologica*, **2015**, 130, 599-601	14.3	27
143	Neuropathy target esterase impairments cause Oliver-McFarlane and Laurence-Moon syndromes. *Journal of Medical Genetics*, **2015**, 52, 85-94	5.8	77
142	Review: an update on clinical, genetic and pathological aspects of frontotemporal lobar degenerations. *Neuropathology and Applied Neurobiology*, **2015**, 41, 858-81	5.2	130
141	Distinct clinical and neuropathological features of G51D SNCA mutation cases compared with SNCA duplication and H50Q mutation. *Molecular Neurodegeneration*, **2015**, 10, 41	19	62

140	Serotonergic markers in Parkinson's disease and levodopa-induced dyskinesias. *Movement Disorders*, **2015**, 30, 796-804	7	24
139	Minimal change multiple system atrophy: an aggressive variant?. *Movement Disorders*, **2015**, 30, 960-7	7	30
138	Temporal Variant Frontotemporal Dementia is Associated with Globular Glial Tauopathy. *Cognitive and Behavioral Neurology*, **2015**, 28, 92-7	1.6	18
137	Concomitant fragile X-associated tremor ataxia syndrome and Parkinson's disease: a clinicopathological report of two cases. *Journal of Neurology, Neurosurgery and Psychiatry*, **2015**, 86, 934-6	5.5	16
136	Screening a UK amyotrophic lateral sclerosis cohort provides evidence of multiple origins of the C9orf72 expansion. *Neurobiology of Aging*, **2015**, 36, 546.e1-7	5.6	41
135	Analysis of C9orf72 repeat expansions in a large series of clinically and pathologically diagnosed cases with atypical parkinsonism. *Neurobiology of Aging*, **2015**, 36, 1221.e1-6	5.6	32
134	Spontaneous ARIA (amyloid-related imaging abnormalities) and cerebral amyloid angiopathy related inflammation in presenilin 1-associated familial Alzheimer's disease. *Journal of Alzheimer's Disease*, **2015**, 44, 1069-74	4.3	16
133	C9orf72 expansions in frontotemporal dementia and amyotrophic lateral sclerosis. *Lancet Neurology, The*, **2015**, 14, 291-301	24.1	165
132	Multiple system atrophy is not caused by C9orf72 hexanucleotide repeat expansions. *Neurobiology of Aging*, **2015**, 36, 1223.e1-2	5.6	19
131	Brain amyloid-beta fragment signatures in pathological ageing and Alzheimer's disease by hybrid immunoprecipitation mass spectrometry. *Neurodegenerative Diseases*, **2015**, 15, 50-7	2.3	33
130	Neuropathological assessments of the pathology in frontotemporal lobar degeneration with TDP43-positive inclusions: an inter-laboratory study by the BrainNet Europe consortium. *Journal of Neural Transmission*, **2015**, 122, 957-72	4.3	19
129	Evaluating the relationship between amyloid-β and α-synuclein phosphorylated at Ser129 in dementia with Lewy bodies and Parkinson's disease. *Alzheimer's Research and Therapy*, **2014**, 6, 77	9	60
128	Genetic analysis implicates APOE, SNCA and suggests lysosomal dysfunction in the etiology of dementia with Lewy bodies. *Human Molecular Genetics*, **2014**, 23, 6139-46	5.6	152
127	TDP-43 pathology is present in most post-encephalitic parkinsonism brains. *Neuropathology and Applied Neurobiology*, **2014**, 40, 654-7	5.2	8
126	Alpha-synuclein mRNA expression in oligodendrocytes in MSA. *Glia*, **2014**, 62, 964-70	9	121
125	A pathogenic progranulin mutation and C9orf72 repeat expansion in a family with frontotemporal dementia. *Neuropathology and Applied Neurobiology*, **2014**, 40, 502-13	5.2	31
124	A 6.4 Mb duplication of the α-synuclein locus causing frontotemporal dementia and Parkinsonism: phenotype-genotype correlations. *JAMA Neurology*, **2014**, 71, 1162-71	17.2	51
123	Concomitant progressive supranuclear palsy and chronic traumatic encephalopathy in a boxer. *Acta Neuropathologica Communications*, **2014**, 2, 24	7.3	16

122	The significance of -synuclein, amyloid- and tau pathologies in Parkinson's disease progression and related dementia. *Neurodegenerative Diseases*, **2014**, 13, 154-6	2.3	66
121	MM2 subtype of sporadic Creutzfeldt-Jakob disease may underlie the clinical presentation of progressive supranuclear palsy. *Journal of Neurology*, **2013**, 260, 1031-6	5.5	17
120	Homozygosity for the C9orf72 GGGGCC repeat expansion in frontotemporal dementia. *Acta Neuropathologica*, **2013**, 126, 401-9	14.3	119
119	A novel prion disease associated with diarrhea and autonomic neuropathy. *New England Journal of Medicine*, **2013**, 369, 1904-14	59.2	93
118	Globular glial tauopathies (GGT): consensus recommendations. *Acta Neuropathologica*, **2013**, 126, 537-544	14.3	136
117	Neuropathological findings in benign tremulous parkinsonism. *Movement Disorders*, **2013**, 28, 145-52	7	56
116	The midbrain to pons ratio: a simple and specific MRI sign of progressive supranuclear palsy. *Neurology*, **2013**, 80, 1856-61	6.5	114
115	TDP-43 pathology in a patient carrying G2019S LRRK2 mutation and a novel p.Q124E MAPT. *Neurobiology of Aging*, **2013**, 34, 2889.e5-9	5.6	33
114	Abundant pyroglutamate-modified ABri and ADan peptides in extracellular and vascular amyloid deposits in familial British and Danish dementias. *Neurobiology of Aging*, **2013**, 34, 1416-25	5.6	12
113	Parkin disease: a clinicopathologic entity?. *JAMA Neurology*, **2013**, 70, 571-9	17.2	101
112	Synucleinopathy associated with G51D SNCA mutation: a link between Parkinson's disease and multiple system atrophy?. *Acta Neuropathologica*, **2013**, 125, 753-69	14.3	298
111	Genetic analysis of inherited leukodystrophies: genotype-phenotype correlations in the CSF1R gene. *JAMA Neurology*, **2013**, 70, 875-882	17.2	58
110	Exome sequencing reveals a novel partial deletion in the progranulin gene causing primary progressive aphasia. *Journal of Neurology, Neurosurgery and Psychiatry*, **2013**, 84, 1411-2	5.5	8
109	A pathway-based analysis provides additional support for an immune-related genetic susceptibility to Parkinson's disease. *Human Molecular Genetics*, **2013**, 22, 1039-49	5.6	96
108	C9ORF72 expansions, parkinsonism, and Parkinson disease: a clinicopathologic study. *Neurology*, **2013**, 81, 808-11	6.5	49
107	A NOVEL TAUOPATHY PRESENTING WITH A MOTOR NEURONE DISEASE PHENOTYPE. *Journal of Neurology, Neurosurgery and Psychiatry*, **2013**, 84, e2.123-e2	5.5	
106	Identification and quantification of oligodendrocyte precursor cells in multiple system atrophy, progressive supranuclear palsy and Parkinson's disease. *Brain Pathology*, **2013**, 23, 263-73	6	52
105	Reply to letter: Multiple system atrophy-parkinsonism with slow progression and prolonged survival: a diagnostic catch. *Movement Disorders*, **2013**, 28, 408	7	

104	Difference in MSA phenotype distribution between populations: genetics or environment?. *Journal of Parkinsonıs Disease*, **2012**, 2, 7-18	5.3	27
103	Widespread Lewy body and tau accumulation in childhood and adult onset dystonia-parkinsonism cases with PLA2G6 mutations. *Neurobiology of Aging*, **2012**, 33, 814-23	5.6	151
102	Duplication of amyloid precursor protein (APP), but not prion protein (PRNP) gene is a significant cause of early onset dementia in a large UK series. *Neurobiology of Aging*, **2012**, 33, 426.e13-21	5.6	55
101	Tau acts as an independent genetic risk factor in pathologically proven PD. *Neurobiology of Aging*, **2012**, 33, 838.e7-11	5.6	20
100	The MAPT p.A152T variant is a risk factor associated with tauopathies with atypical clinical and neuropathological features. *Neurobiology of Aging*, **2012**, 33, 2231.e7-2231.e14	5.6	57
99	The spread of neurodegenerative disease. *New England Journal of Medicine*, **2012**, 366, 2126-8	59.2	67
98	Conventional magnetic resonance imaging in confirmed progressive supranuclear palsy and multiple system atrophy. *Movement Disorders*, **2012**, 27, 1754-62	7	126
97	Multiple system atrophy-parkinsonism with slow progression and prolonged survival: a diagnostic catch. *Movement Disorders*, **2012**, 27, 1186-90	7	72
96	Frontotemporal dementia with the C9ORF72 hexanucleotide repeat expansion: clinical, neuroanatomical and neuropathological features. *Brain*, **2012**, 135, 736-50	11.2	340
95	Identification of common variants influencing risk of the tauopathy progressive supranuclear palsy. *Nature Genetics*, **2011**, 43, 699-705	36.3	386
94	Glucocerebrosidase mutations do not cause increased Lewy body pathology in Parkinsonֿ disease. *Molecular Genetics and Metabolism*, **2011**, 103, 410-2	3.7	30
93	Mutational analysis of parkin and PINK1 in multiple system atrophy. *Neurobiology of Aging*, **2011**, 32, 548.e5-7	5.6	10
92	Disentangling the relationship between lewy bodies and nigral neuronal loss in Parkinsonֿ disease. *Journal of Parkinsonıs Disease*, **2011**, 1, 277-86	5.3	74
91	TDP-43 pathology may occur in the BRI2 gene-related dementias. *Acta Neuropathologica*, **2011**, 121, 559-60	14.3	3
90	Neuropathology underlying clinical variability in patients with synucleinopathies. *Acta Neuropathologica*, **2011**, 122, 187-204	14.3	292
89	Globular glial tauopathies (GGT) presenting with motor neuron disease or frontotemporal dementia: an emerging group of 4-repeat tauopathies. *Acta Neuropathologica*, **2011**, 122, 415-28	14.3	60
88	Transportin1: a marker of FTLD-FUS. *Acta Neuropathologica*, **2011**, 122, 591-600	14.3	50
87	Postural instability, frontotemporal dementia, and ophthalmoplegia: clinicopathological case. *Movement Disorders*, **2011**, 26, 1808-13	7	2

86	A comparative clinical, pathological, biochemical and genetic study of fused in sarcoma proteinopathies. *Brain*, **2011**, 134, 2548-64	11.2	65
85	Clinical and neuroanatomical signatures of tissue pathology in frontotemporal lobar degeneration. *Brain*, **2011**, 134, 2565-81	11.2	251
84	Lewy- and Alzheimer-type pathologies in Parkinson's disease dementia: which is more important?. *Brain*, **2011**, 134, 1493-1505	11.2	399
83	The clinical and neuroanatomical phenotype of FUS associated frontotemporal lobar degeneration. *Journal of Neurology, Neurosurgery and Psychiatry*, **2011**, 82, 1405-7	5.5	30
82	Testing an aetiological model of visual hallucinations in Parkinson's disease. *Brain*, **2011**, 134, 3299-309	11.2	106
81	Genetic variability at the PARK16 locus. *European Journal of Human Genetics*, **2010**, 18, 1356-9	5.3	69
80	Relationships between age and late progression of Parkinson's disease: a clinico-pathological study. *Brain*, **2010**, 133, 1755-62	11.2	280
79	Does corticobasal degeneration exist? A clinicopathological re-evaluation. *Brain*, **2010**, 133, 2045-57	11.2	302
78	Nomenclature and nosology for neuropathologic subtypes of frontotemporal lobar degeneration: an update. *Acta Neuropathologica*, **2010**, 119, 1-4	14.3	711
77	FUS pathology defines the majority of tau- and TDP-43-negative frontotemporal lobar degeneration. *Acta Neuropathologica*, **2010**, 120, 33-41	14.3	198
76	Brain biopsy in dementia: clinical indications and diagnostic approach. *Acta Neuropathologica*, **2010**, 120, 327-41	14.3	49
75	Hyposmia in progressive supranuclear palsy. *Movement Disorders*, **2010**, 25, 570-7	7	38
74	Impulsive-compulsive spectrum behaviors in pathologically confirmed progressive supranuclear palsy. *Movement Disorders*, **2010**, 25, 638-42	7	30
73	CEREBRAL AMYLOID ANGIOPATHY AND ALZHEIMER'S DISEASE **2010**, 61, S111-S124		15
72	PYROGLUTAMATE FORMATION AT THE N-TERMINI OF ABRI MOLECULES IN FAMILIAL BRITISH DEMENTIA IS NOT RESTRICTED TO THE CENTRAL NERVOUS SYSTEM **2010**, 61, S262-S269		1
71	Glucocerebrosidase mutations in clinical and pathologically proven Parkinson's disease. *Brain*, **2009**, 132, 1783-94	11.2	488
70	SNCA variants are associated with increased risk for multiple system atrophy. *Annals of Neurology*, **2009**, 65, 610-4	9.4	232
69	Nomenclature for neuropathologic subtypes of frontotemporal lobar degeneration: consensus recommendations. *Acta Neuropathologica*, **2009**, 117, 15-8	14.3	325

68	Can olfactory bulb biopsy be justified for the diagnosis of Parkinson's disease? Comments on "olfactory bulb alpha-synucleinopathy has high specificity and sensitivity for Lewy body disorders". *Acta Neuropathologica*, **2009**, 117, 213-4; author reply 217-8	14.3	19
67	Assessment of beta-amyloid deposits in human brain: a study of the BrainNet Europe Consortium. *Acta Neuropathologica*, **2009**, 117, 309-20	14.3	119
66	Genetics and molecular pathogenesis of sporadic and hereditary cerebral amyloid angiopathies. *Acta Neuropathologica*, **2009**, 118, 115-30	14.3	212
65	The genetics of Parkinson's syndromes: a critical review. *Current Opinion in Genetics and Development*, **2009**, 19, 254-65	4.9	168
64	Clinical and pathological features of an Alzheimer's disease patient with the MAPT Delta K280 mutation. *Neurobiology of Aging*, **2009**, 30, 388-93	5.6	45
63	Regional differences in the severity of Lewy body pathology across the olfactory cortex. *Neuroscience Letters*, **2009**, 453, 77-80	3.3	96
62	Concomitant progressive supranuclear palsy and multiple system atrophy: more than a simple twist of fate?. *Neuroscience Letters*, **2009**, 467, 208-11	3.3	17
61	Parkinson's disease. *Lancet, The*, **2009**, 373, 2055-66	40	1541
60	Research in motion: the enigma of Parkinson's disease pathology spread. *Nature Reviews Neuroscience*, **2008**, 9, 741-5	13.5	251
59	Lewy bodies in grafted neurons in subjects with Parkinson's disease suggest host-to-graft disease propagation. *Nature Medicine*, **2008**, 14, 501-3	50.5	1293
58	A distinct clinical, neuropsychological and radiological phenotype is associated with progranulin gene mutations in a large UK series. *Brain*, **2008**, 131, 706-20	11.2	198
57	Parietal lobe deficits in frontotemporal lobar degeneration caused by a mutation in the progranulin gene. *Archives of Neurology*, **2008**, 65, 506-13		48
56	MAPT S305I mutation: implications for argyrophilic grain disease. *Acta Neuropathologica*, **2008**, 116, 103-18	14.3	44
55	Cortical alpha-synuclein load is associated with amyloid-beta plaque burden in a subset of Parkinson's disease patients. *Acta Neuropathologica*, **2008**, 115, 417-25	14.3	121
54	LRRK2 and parkin immunoreactivity in multiple system atrophy inclusions. *Acta Neuropathologica*, **2008**, 116, 639-46	14.3	38
53	Diseases of movement and system degenerations **2008**, 889-1030		5
52	Preferential association of serum amyloid P component with fibrillar deposits in familial British and Danish dementias: similarities with Alzheimer's disease. *Journal of the Neurological Sciences*, **2007**, 257, 88-96	3.2	24
51	Genetic variation at the tau locus and clinical syndromes associated with progressive supranuclear palsy. *Movement Disorders*, **2007**, 22, 895-7	7	20

50	Pure akinesia with gait freezing: a third clinical phenotype of progressive supranuclear palsy. *Movement Disorders*, **2007**, 22, 2235-41	7	188
49	Mutations in TTBK2, encoding a kinase implicated in tau phosphorylation, segregate with spinocerebellar ataxia type 11. *Nature Genetics*, **2007**, 39, 1434-6	36.3	152
48	The phagocytic capacity of neurones. *European Journal of Neuroscience*, **2007**, 25, 2947-55	3.5	30
47	DJ-1 (PARK7) is associated with 3R and 4R tau neuronal and glial inclusions in neurodegenerative disorders. *Neurobiology of Disease*, **2007**, 28, 122-32	7.5	31
46	Adult-onset neurodegeneration with brain iron accumulation and cortical alpha-synuclein and tau pathology: a distinct clinicopathological entity. *Archives of Neurology*, **2007**, 64, 280-2		34
45	Pathological tau burden and distribution distinguishes progressive supranuclear palsy-parkinsonism from Richardson's syndrome. *Brain*, **2007**, 130, 1566-76	11.2	289
44	NR4A2 genetic variation in sporadic Parkinson's disease: a genewide approach. *Movement Disorders* , **2006**, 21, 1960-3	7	14
43	UCHL-1 is not a Parkinson's disease susceptibility gene. *Annals of Neurology*, **2006**, 59, 627-33	9.4	107
42	An immunohistochemical study of cases of sporadic and inherited frontotemporal lobar degeneration using 3R- and 4R-specific tau monoclonal antibodies. *Acta Neuropathologica*, **2006**, 111, 329-40	14.3	81
41	Mutations in the gene LRRK2 encoding dardarin (PARK8) cause familial Parkinson's disease: clinical, pathological, olfactory and functional imaging and genetic data. *Brain*, **2005**, 128, 2786-96	11.2	283
40	A common LRRK2 mutation in idiopathic Parkinson's disease. *Lancet, The*, **2005**, 365, 415-6	40	283
39	Characteristics of two distinct clinical phenotypes in pathologically proven progressive supranuclear palsy: Richardson's syndrome and PSP-parkinsonism. *Brain*, **2005**, 128, 1247-58	11.2	583
38	UCHL-1 gene in multiple system atrophy: a haplotype tagging approach. *Movement Disorders*, **2005**, 20, 1338-43	7	15
37	Sporadic four-repeat tauopathy with frontotemporal degeneration, parkinsonism and motor neuron disease. *Acta Neuropathologica*, **2005**, 110, 600-9	14.3	28
36	Familial Danish dementia: co-existence of Danish and Alzheimer amyloid subunits (ADan AND A{beta}) in the absence of compact plaques. *Journal of Biological Chemistry*, **2005**, 280, 36883-94	5.4	53
35	The expression of DJ-1 (PARK7) in normal human CNS and idiopathic Parkinson's disease. *Brain*, **2004**, 127, 420-30	11.2	341
34	The spectrum of pathological involvement of the striatonigral and olivopontocerebellar systems in multiple system atrophy: clinicopathological correlations. *Brain*, **2004**, 127, 2657-71	11.2	359
33	Detecting tau isoforms in archival cases. *Acta Neuropathologica*, **2004**, 107, 181-2	14.3	3

32	Clinicopathological investigation of vascular parkinsonism, including clinical criteria for diagnosis. *Movement Disorders*, **2004**, 19, 630-40	7	273
31	Somatic and germline mosaicism in sporadic early-onset Alzheimer disease. *Human Molecular Genetics*, **2004**, 13, 1219-24	5.6	76
30	Familial and sporadic cerebral amyloid angiopathies associated with dementia and the BRI dementias **2004**, 330-352		6
29	Cerebral amyloid angiopathies: a pathologic, biochemical, and genetic view. *Journal of Neuropathology and Experimental Neurology*, **2003**, 62, 885-98	3.1	207
28	Neurofilament inclusion body disease: a new proteinopathy?. *Brain*, **2003**, 126, 2291-303	11.2	162
27	Anatamopathological spectrum of tauopathies. *Movement Disorders*, **2003**, 18 Suppl 6, S13-20	7	18
26	Sporadic and familial cerebral amyloid angiopathies. *Brain Pathology*, **2002**, 12, 343-57	6	137
25	Complement activation in chromosome 13 dementias. Similarities with Alzheimer disease. *Journal of Biological Chemistry*, **2002**, 277, 49782-90	5.4	47
24	Familial Danish dementia: a novel form of cerebral amyloidosis associated with deposition of both amyloid-Dan and amyloid-beta. *Journal of Neuropathology and Experimental Neurology*, **2002**, 61, 254-67	3.1	102
23	A multidisciplinary team approach to skull base chondrosarcomas. *Journal of Neurosurgery*, **2001**, 95, 184-9	3.2	88
22	Systemic amyloid deposits in familial British dementia. *Journal of Biological Chemistry*, **2001**, 276, 43909-14	5.4	57
21	A pathogenic presenilin-1 deletion causes abberrant Abeta 42 production in the absence of congophilic amyloid plaques. *Journal of Biological Chemistry*, **2001**, 276, 7233-9	5.4	63
20	A multidisciplinary team approach to skull base chordomas. *Journal of Neurosurgery*, **2001**, 95, 175-83	3.2	131
19	Regional distribution of amyloid-Bri deposition and its association with neurofibrillary degeneration in familial British dementia. *American Journal of Pathology*, **2001**, 158, 515-26	5.8	115
18	Sequence, genomic structure and tissue expression of Human BRI3, a member of the BRI gene family. *Gene*, **2001**, 266, 95-102	3.8	48
17	A novel presenilin mutation (M233V) causing very early onset Alzheimer disease with Lewy bodies. *Neuroscience Letters*, **2001**, 313, 93-5	3.3	47
16	Chromosome 13 dementia syndromes as models of neurodegeneration. *Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis*, **2001**, 8, 277-84	2.7	26
15	Variant Alzheimer disease with spastic paraparesis and cotton wool plaques is caused by PS-1 mutations that lead to exceptionally high amyloid- concentrations. *Annals of Neurology*, **2000**, 48, 806-808	8.4	113

14	Familial British dementia with amyloid angiopathy: early clinical, neuropsychological and imaging findings. *Brain*, **2000**, 123 (Pt 5), 975-91	11.2	86
13	Axonal lesions in multiple sclerosis: an old story revisited. *Brain*, **2000**, 123 (Pt 2), 203-4	11.2	6
12	Variant Alzheimer disease with spastic paraparesis and cotton wool plaques is caused by PS-1 mutations that lead to exceptionally high amyloid- concentrations **2000**, 48, 806		3
11	A stop-codon mutation in the BRI gene associated with familial British dementia. *Nature*, **1999**, 399, 776-81	30.4	423
10	A case of sporadic Pick disease with onset at 27 years. *Archives of Neurology*, **1999**, 56, 1289-91		11
9	Central benzodiazepine receptor autoradiography in hippocampal sclerosis. *British Journal of Pharmacology*, **1997**, 122, 358-64	8.6	40
8	Skull base chordomas: a review of 38 patients, 1958-88. *British Journal of Neurosurgery*, **1993**, 7, 241-8	1	93
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