Janice Holton

List of Publications by Citations

306 papers	19,511 citations	75 h-index	131 g-index
324 ext. papers	22,307 ext. citations	7.6 avg, IF	6.25 L-index

#	Paper	IF	Citations
306	Lewy bodies in grafted neurons in subjects with Parkinson disease suggest host-to-graft disease propagation. *Nature Medicine*, **2008**, 14, 501-3	50.5	1293
305	Characteristics of two distinct clinical phenotypes in pathologically proven progressive supranuclear palsy: Richardson syndrome and PSP-parkinsonism. *Brain*, **2005**, 128, 1247-58	11.2	583
304	Glucocerebrosidase mutations in clinical and pathologically proven Parkinson disease. *Brain*, **2009**, 132, 1783-94	11.2	488
303	Lewy- and Alzheimer-type pathologies in Parkinson disease dementia: which is more important?. *Brain*, **2011**, 134, 1493-1505	11.2	399
302	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
301	A clinico-pathological study of subtypes in Parkinson disease. *Brain*, **2009**, 132, 2947-57	11.2	320
300	Does corticobasal degeneration exist? A clinicopathological re-evaluation. *Brain*, **2010**, 133, 2045-57	11.2	302
299	Synucleinopathy associated with G51D SNCA mutation: a link between Parkinson disease and multiple system atrophy?. *Acta Neuropathologica*, **2013**, 125, 753-69	14.3	298
298	Neuropathology underlying clinical variability in patients with synucleinopathies. *Acta Neuropathologica*, **2011**, 122, 187-204	14.3	292
297	Pathological tau burden and distribution distinguishes progressive supranuclear palsy-parkinsonism from Richardson syndrome. *Brain*, **2007**, 130, 1566-76	11.2	289
296	Clinical outcomes of progressive supranuclear palsy and multiple system atrophy. *Brain*, **2008**, 131, 1362-72	11.2	287
295	Mutations in the gene LRRK2 encoding dardarin (PARK8) cause familial Parkinson disease: clinical, pathological, olfactory and functional imaging and genetic data. *Brain*, **2005**, 128, 2786-96	11.2	283
294	A common LRRK2 mutation in idiopathic Parkinson disease. *Lancet, The*, **2005**, 365, 415-6	40	283
293	Relationships between age and late progression of Parkinson disease: a clinico-pathological study. *Brain*, **2010**, 133, 1755-62	11.2	280
292	PINK1 protein in normal human brain and Parkinson disease. *Brain*, **2006**, 129, 1720-31	11.2	267
291	A decamer duplication in the 3 region of the BRI gene originates an amyloid peptide that is associated with dementia in a Danish kindred. *Proceedings of the National Academy of Sciences of the United States of America*, **2000**, 97, 4920-5	11.5	255
290	Clinical and neuroanatomical signatures of tissue pathology in frontotemporal lobar degeneration. *Brain*, **2011**, 134, 2565-81	11.2	251

289	Research in motion: the enigma of Parkinson disease pathology spread. *Nature Reviews Neuroscience*, **2008**, 9, 741-5	13.5	251
288	SNCA variants are associated with increased risk for multiple system atrophy. *Annals of Neurology*, **2009**, 65, 610-4	9.4	232
287	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
286	Genetics and molecular pathogenesis of sporadic and hereditary cerebral amyloid angiopathies. *Acta Neuropathologica*, **2009**, 118, 115-30	14.3	212
285	A two-stage meta-analysis identifies several new loci for Parkinson disease. *PLoS Genetics*, **2011**, 7, e1002142	6	209
284	Cerebral amyloid angiopathies: a pathologic, biochemical, and genetic view. *Journal of Neuropathology and Experimental Neurology*, **2003**, 62, 885-98	3.1	207
283	FUS pathology defines the majority of tau- and TDP-43-negative frontotemporal lobar degeneration. *Acta Neuropathologica*, **2010**, 120, 33-41	14.3	198
282	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
281	Pure akinesia with gait freezing: a third clinical phenotype of progressive supranuclear palsy. *Movement Disorders*, **2007**, 22, 2235-41	7	188
280	The neuropathology, pathophysiology and genetics of multiple system atrophy. *Neuropathology and Applied Neurobiology*, **2012**, 38, 4-24	5.2	179
279	Cloning and sequence analysis of desmosomal glycoproteins 2 and 3 (desmocollins): cadherin-like desmosomal adhesion molecules with heterogeneous cytoplasmic domains. *Journal of Cell Biology*, **1991**, 113, 381-91	7.3	171
278	Pathological inclusion bodies in tauopathies contain distinct complements of tau with three or four microtubule-binding repeat domains as demonstrated by new specific monoclonal antibodies. *Neuropathology and Applied Neurobiology*, **2003**, 29, 288-302	5.2	167
277	C9orf72 expansions in frontotemporal dementia and amyotrophic lateral sclerosis. *Lancet Neurology, The*, **2015**, 14, 291-301	24.1	165
276	Neurofilament inclusion body disease: a new proteinopathy?. *Brain*, **2003**, 126, 2291-303	11.2	162
275	The novel Parkinson disease linked mutation G51D attenuates in vitro aggregation and membrane binding of -synuclein, and enhances its secretion and nuclear localization in cells. *Human Molecular Genetics*, **2014**, 23, 4491-509	5.6	153
274	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
273	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
272	Cognitive impairment in multiple system atrophy: a position statement by the Neuropsychology Task Force of the MDS Multiple System Atrophy (MODIMSA) study group. *Movement Disorders*, **2014**, 29, 857-67	7	148

271	Patterns of levodopa response in Parkinson's disease: a clinico-pathological study. *Brain*, **2007**, 130, 2123-8	11.2	145
270	Sporadic and familial cerebral amyloid angiopathies. *Brain Pathology*, **2002**, 12, 343-57	6	137
269	Globular glial tauopathies (GGT): consensus recommendations. *Acta Neuropathologica*, **2013**, 126, 537-544	14.3	136
268	Altered cleavage and localization of PINK1 to aggresomes in the presence of proteasomal stress. *Journal of Neurochemistry*, **2006**, 98, 156-69	6	136
267	Gene expression in Huntington's disease skeletal muscle: a potential biomarker. *Human Molecular Genetics*, **2005**, 14, 1863-76	5.6	134
266	Degeneration in different parkinsonian syndromes relates to astrocyte type and astrocyte protein expression. *Journal of Neuropathology and Experimental Neurology*, **2009**, 68, 1073-83	3.1	132
265	A multidisciplinary team approach to skull base chordomas. *Journal of Neurosurgery*, **2001**, 95, 175-83	3.2	131
264	Dysregulation of glucose metabolism is an early event in sporadic Parkinson's disease. *Neurobiology of Aging*, **2014**, 35, 1111-5	5.6	129
263	Mixed pathologies including chronic traumatic encephalopathy account for dementia in retired association football (soccer) players. *Acta Neuropathologica*, **2017**, 133, 337-352	14.3	128
262	Conventional magnetic resonance imaging in confirmed progressive supranuclear palsy and multiple system atrophy. *Movement Disorders*, **2012**, 27, 1754-62	7	126
261	Adenosine monophosphate-activated protein kinase disease mimicks hypertrophic cardiomyopathy and Wolff-Parkinson-White syndrome: natural history. *Journal of the American College of Cardiology* , **2005**, 45, 922-30	15.1	125
260	Alpha-synuclein mRNA expression in oligodendrocytes in MSA. *Glia*, **2014**, 62, 964-70	9	121
259	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
258	Frontotemporal lobar degeneration and ubiquitin immunohistochemistry. *Neuropathology and Applied Neurobiology*, **2004**, 30, 369-73	5.2	121
257	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
256	Magnetic resonance imaging signatures of tissue pathology in frontotemporal dementia. *Archives of Neurology*, **2005**, 62, 1402-8		120
255	Prognosis and Neuropathologic Correlation of Clinical Subtypes of Parkinson Disease. *JAMA Neurology*, **2019**, 76, 470-479	17.2	118
254	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

253	The midbrain to pons ratio: a simple and specific MRI sign of progressive supranuclear palsy. *Neurology*, **2013**, 80, 1856-61	6.5	114
252	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
251	Clinical phenotype and genetic associations in autosomal dominant familial Alzheimer disease: a case series. *Lancet Neurology, The*, **2016**, 15, 1326-1335	24.1	109
250	UCHL-1 is not a Parkinson disease susceptibility gene. *Annals of Neurology*, **2006**, 59, 627-33	9.4	107
249	Neuropathological criteria of anti-IgLON5-related tauopathy. *Acta Neuropathologica*, **2016**, 132, 531-43	14.3	107
248	Brain biopsy in dementia. *Brain*, **2005**, 128, 2016-25	11.2	105
247	A genome-wide association study in multiple system atrophy. *Neurology*, **2016**, 87, 1591-1598	6.5	104
246	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
245	Parkin disease: a clinicopathologic entity?. *JAMA Neurology*, **2013**, 70, 571-9	17.2	101
244	International consensus on a proposed score system for muscle biopsy evaluation in patients with juvenile dermatomyositis: a tool for potential use in clinical trials. *Arthritis and Rheumatism*, **2007**, 57, 1192-201		100
243	A pathway-based analysis provides additional support for an immune-related genetic susceptibility to Parkinson disease. *Human Molecular Genetics*, **2013**, 22, 1039-49	5.6	96
242	Regional differences in the severity of Lewy body pathology across the olfactory cortex. *Neuroscience Letters*, **2009**, 453, 77-80	3.3	96
241	Association of Autonomic Dysfunction With Disease Progression and Survival in Parkinson Disease. *JAMA Neurology*, **2017**, 74, 970-976	17.2	94
240	A novel prion disease associated with diarrhea and autonomic neuropathy. *New England Journal of Medicine*, **2013**, 369, 1904-14	59.2	93
239	A multidisciplinary team approach to skull base chondrosarcomas. *Journal of Neurosurgery*, **2001**, 95, 184-9	3.2	88
238	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
237	Astrogliopathy predominates the earliest stage of corticobasal degeneration pathology. *Brain*, **2016**, 139, 3237-3252	11.2	80
236	Development of phospho-specific Rab protein antibodies to monitor activity of the LRRK2 Parkinson disease kinase. *Biochemical Journal*, **2018**, 475, 1-22	3.8	79

235	Genetic dysfunction of MT-ATP6 causes axonal Charcot-Marie-Tooth disease. *Neurology*, **2012**, 79, 1145-54	6.5	77
234	Patients with a novel neurofilamentopathy: dementia with neurofilament inclusions. *Neuroscience Letters*, **2003**, 341, 177-80	3.3	76
233	Histological evidence of chronic traumatic encephalopathy in a large series of neurodegenerative diseases. *Acta Neuropathologica*, **2015**, 130, 891-3	14.3	75
232	Disentangling the relationship between lewy bodies and nigral neuronal loss in Parkinson's disease. *Journal of Parkinson's Disease*, **2011**, 1, 277-86	5.3	74
231	Rhabdomyolysis: a genetic perspective. *Orphanet Journal of Rare Diseases*, **2015**, 10, 51	4.2	73
230	Multiple system atrophy-parkinsonism with slow progression and prolonged survival: a diagnostic catch. *Movement Disorders*, **2012**, 27, 1186-90	7	72
229	Conspicuous involvement of desmin tail mutations in diverse cardiac and skeletal myopathies. *Human Mutation*, **2007**, 28, 374-86	4.7	72
228	The use of nerve and muscle biopsy in the diagnosis of vasculitis: a 5 year retrospective study. *Journal of Neurology, Neurosurgery and Psychiatry*, **2008**, 79, 1376-81	5.5	71
227	High resolution MR anatomy of the subthalamic nucleus: imaging at 9.4 T with histological validation. *NeuroImage*, **2012**, 59, 2035-44	7.9	70
226	Gangliocytomas of the pituitary: a heterogeneous group of lesions with differing histogenesis. *American Journal of Surgical Pathology*, **2000**, 24, 607-13	6.7	70
225	Targeting protein homeostasis in sporadic inclusion body myositis. *Science Translational Medicine*, **2016**, 8, 331ra41	17.5	69
224	Muscle Biopsy Findings in Combination With Myositis-Specific Autoantibodies Aid Prediction of Outcomes in Juvenile Dermatomyositis. *Arthritis and Rheumatology*, **2016**, 68, 2806-2816	9.5	67
223	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
222	The bovine desmocollin family: a new gene and expression patterns reflecting epithelial cell proliferation and differentiation. *Journal of Cell Biology*, **1994**, 126, 507-18	7.3	66
221	A comparative clinical, pathological, biochemical and genetic study of fused in sarcoma proteinopathies. *Brain*, **2011**, 134, 2548-64	11.2	65
220	A retrospective cohort study identifying the principal pathological features useful in the diagnosis of inclusion body myositis. *BMJ Open*, **2014**, 4, e004552	3	63
219	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
218	Distinct clinical and neuropathological features of G51D SNCA mutation cases compared with SNCA duplication and H50Q mutation. *Molecular Neurodegeneration*, **2015**, 10, 41	19	62

217	MHC Class I overexpression on muscles in early juvenile dermatomyositis. *Journal of Rheumatology*, **2004**, 31, 605-9	4.1	61
216	Improving diagnostic accuracy of multiple system atrophy: a clinicopathological study. *Brain*, **2019**, 142, 2813-2827	11.2	60
215	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
214	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
213	The effects of the tremorgenic mycotoxin penitrem A on the rat cerebellum. *Veterinary Pathology*, **1998**, 35, 53-63	2.8	59
212	Characteristics of progressive supranuclear palsy presenting with corticobasal syndrome: a cortical variant. *Neuropathology and Applied Neurobiology*, **2014**, 40, 149-63	5.2	58
211	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
210	Systemic amyloid deposits in familial British dementia. *Journal of Biological Chemistry*, **2001**, 276, 43909-14	5.4	57
209	Neuropathological findings in benign tremulous parkinsonism. *Movement Disorders*, **2013**, 28, 145-52	7	56
208	Down-regulation of the dopamine receptor D2 in mice lacking ataxin 1. *Human Molecular Genetics*, **2007**, 16, 2122-34	5.6	53
207	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
206	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
205	Cytoskeletal pathology in familial cerebral amyloid angiopathy (British type) with non-neuritic amyloid plaque formation. *Acta Neuropathologica*, **1999**, 97, 170-6	14.3	52
204	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
203	Transportin1: a marker of FTLD-FUS. *Acta Neuropathologica*, **2011**, 122, 591-600	14.3	50
202	Brain biopsy in dementia: clinical indications and diagnostic approach. *Acta Neuropathologica*, **2010**, 120, 327-41	14.3	49
201	Skull base chordomas: correlation of tumour doubling time with age, mitosis and Ki67 proliferation index. *Neuropathology and Applied Neurobiology*, **2000**, 26, 497-503	5.2	49
200	Parietal lobe deficits in frontotemporal lobar degeneration caused by a mutation in the progranulin gene. *Archives of Neurology*, **2008**, 65, 506-13		48

199	Complement activation in chromosome 13 dementias. Similarities with Alzheimer's disease. *Journal of Biological Chemistry*, **2002**, 277, 49782-90	5.4	47
198	Neuropathy target esterase: immunolocalization to neuronal cell bodies and axons. *Neuroscience*, **1998**, 83, 295-302	3.9	46
197	Frontotemporal lobar degeneration with ubiquitin-only-immunoreactive neuronal changes: broadening the clinical picture to include progressive supranuclear palsy. *Brain*, **2004**, 127, 2441-51	11.2	46
196	Does levodopa accelerate the pathologic process in Parkinson disease brain?. *Neurology*, **2011**, 77, 1420-6	6.5	45
195	Diagnosis Across the Spectrum of Progressive Supranuclear Palsy and Corticobasal Syndrome. *JAMA Neurology*, **2020**, 77, 377-387	17.2	44
194	Cerebral amyloidosis: amyloid subunits, mutants and phenotypes. *Cellular and Molecular Life Sciences*, **2010**, 67, 581-600	10.3	43
193	The phenomenon of disproportionate antecollis in Parkinson's disease and multiple system atrophy. *Movement Disorders*, **2007**, 22, 2325-31	7	43
192	Clinical features of the myasthenic syndrome arising from mutations in GMPPB. *Journal of Neurology, Neurosurgery and Psychiatry*, **2016**, 87, 802-9	5.5	43
191	Somatic copy number gains of α-synuclein (SNCA) in Parkinson's disease and multiple system atrophy brains. *Brain*, **2018**, 141, 2419-2431	11.2	41
190	Kearns-Sayre syndrome caused by defective R1/p53R2 assembly. *Journal of Medical Genetics*, **2011**, 48, 610-7	5.8	41
189	Neuronal intranuclear inclusion disease: report on a case originally diagnosed as dopa-responsive dystonia with Lewy bodies. *Movement Disorders*, **2005**, 20, 1345-9	7	40
188	Validation of a score tool for measurement of histological severity in juvenile dermatomyositis and association with clinical severity of disease. *Annals of the Rheumatic Diseases*, **2015**, 74, 204-10	2.4	39
187	Proteomics of rimmed vacuoles define new risk allele in inclusion body myositis. *Annals of Neurology*, **2017**, 81, 227-239	9.4	38
186	Hyposmia in progressive supranuclear palsy. *Movement Disorders*, **2010**, 25, 570-7	7	38
185	Neuropathology of primary adult-onset dystonia. *Neurology*, **2008**, 70, 695-9	6.5	38
184	LRRK2 and parkin immunoreactivity in multiple system atrophy inclusions. *Acta Neuropathologica*, **2008**, 116, 639-46	14.3	38
183	The alpha-synuclein gene in multiple system atrophy. *Journal of Neurology, Neurosurgery and Psychiatry*, **2006**, 77, 464-7	5.5	38
182	Genetic alterations of the BRI2 gene: familial British and Danish dementias. *Brain Pathology*, **2006**, 16, 71-9	6	38

181	LRRK2 expression in idiopathic and G2019S positive Parkinson's disease subjects: a morphological and quantitative study. *Neuropathology and Applied Neurobiology*, **2011**, 37, 777-90	5.2	37
180	Normal dopamine transporter single photon-emission CT scan in corticobasal degeneration. *Movement Disorders*, **2008**, 23, 2424-6	7	37
179	Cryptic Amyloidogenic Elements in the 3'UTRs of Neurofilament Genes Trigger Axonal Neuropathy. *American Journal of Human Genetics*, **2016**, 98, 597-614	11	37
178	Variant Alzheimer's disease with spastic paraparesis and cotton wool plaques is caused by PS-1 mutations that lead to exceptionally high amyloid-beta concentrations. *Annals of Neurology*, **2000**, 48, 806-8	9.4	37
177	Pantothenate kinase-associated neurodegeneration is not a synucleinopathy. *Neuropathology and Applied Neurobiology*, **2013**, 39, 121-31	5.2	35
176	LRRK2 exonic variants and risk of multiple system atrophy. *Neurology*, **2014**, 83, 2256-61	6.5	34
175	Microdysgenesis with abnormal cortical myelinated fibres in temporal lobe epilepsy: a histopathological study with calbindin D-28-K immunohistochemistry. *Neuropathology and Applied Neurobiology*, **2000**, 26, 251-7	5.2	34
174	Primum non nocere: a call for balance when reporting on CTE. *Lancet Neurology, The*, **2019**, 18, 231-233	24.1	34
173	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
172	Is it really myositis? A consideration of the differential diagnosis. *Current Opinion in Rheumatology*, **2004**, 16, 684-91	5.3	33
171	A Clinical, Neuropathological and Genetic Study of Homozygous A467T POLG-Related Mitochondrial Disease. *PLoS ONE*, **2016**, 11, e0145500	3.7	33
170	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
169	Selective damage to the cerebellar vermis in chronic alcoholism: a contribution from neurotoxicology to an old problem of selective vulnerability. *Neuropathology and Applied Neurobiology*, **1997**, 23, 355-363	5.2	32
168	Neuropathological features of multiple system atrophy with cognitive impairment. *Movement Disorders*, **2014**, 29, 884-8	7	31
167	Characterisation of a novel NR4A2 mutation in Parkinson's disease brain. *Neuroscience Letters*, **2009**, 457, 75-9	3.3	31
166	Clinical features of congenital myasthenic syndrome due to mutations in DPAGT1. *Journal of Neurology, Neurosurgery and Psychiatry*, **2013**, 84, 1119-25	5.5	30
165	Minimal change multiple system atrophy: an aggressive variant?. *Movement Disorders*, **2015**, 30, 960-7	7	30
164	Review: genetics and neuropathology of primary pure dystonia. *Neuropathology and Applied Neurobiology*, **2012**, 38, 520-34	5.2	30

163	Glucocerebrosidase mutations do not cause increased Lewy body pathology in Parkinson's disease. *Molecular Genetics and Metabolism*, **2011**, 103, 410-2	3.7	30
162	The clinical and neuroanatomical phenotype of FUS associated frontotemporal lobar degeneration. *Journal of Neurology, Neurosurgery and Psychiatry*, **2011**, 82, 1405-7	5.5	30
161	Impulsive-compulsive spectrum behaviors in pathologically confirmed progressive supranuclear palsy. *Movement Disorders*, **2010**, 25, 638-42	7	30
160	Coenzyme Q10 Levels Are Decreased in the Cerebellum of Multiple-System Atrophy Patients. *PLoS ONE*, **2016**, 11, e0149557	3.7	30
159	The effect of drug treatment on neurogenesis in Parkinson's disease. *Movement Disorders*, **2011**, 26, 45-50	7	29
158	Rare variants in SQSTM1 and VCP genes and risk of sporadic inclusion body myositis. *Neurobiology of Aging*, **2016**, 47, 218.e1-218.e9	5.6	29
157	Variation at the TRIM11 locus modifies progressive supranuclear palsy phenotype. *Annals of Neurology*, **2018**, 84, 485-496	9.4	28
156	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
155	Atypical periodic paralysis and myalgia: A novel phenotype. *Neurology*, **2018**, 90, e412-e418	6.5	27
154	The novel MAPT mutation K298E: mechanisms of mutant tau toxicity, brain pathology and tau expression in induced fibroblast-derived neurons. *Acta Neuropathologica*, **2014**, 127, 283-95	14.3	27
153	Difference in MSA phenotype distribution between populations: genetics or environment?. *Journal of Parkinson's Disease*, **2012**, 2, 7-18	5.3	27
152	CAV3 mutations causing exercise intolerance, myalgia and rhabdomyolysis: Expanding the phenotypic spectrum of caveolinopathies. *Neuromuscular Disorders*, **2016**, 26, 504-10	2.9	27
151	LRRK2 levels and phosphorylation in Parkinson's disease brain and cases with restricted Lewy bodies. *Movement Disorders*, **2017**, 32, 423-432	7	26
150	Reduced LRRK2 in association with retromer dysfunction in post-mortem brain tissue from LRRK2 mutation carriers. *Brain*, **2018**, 141, 486-495	11.2	26
149	Targeting of the pedunculopontine nucleus by an MRI-guided approach: a cadaver study. *Journal of Neural Transmission*, **2011**, 118, 1487-95	4.3	26
148	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
147	LATE to the PART-y. *Brain*, **2019**, 142, e47	11.2	25
146	Widespread RNA metabolism impairment in sporadic inclusion body myositis TDP43-proteinopathy. *Neurobiology of Aging*, **2014**, 35, 1491-8	5.6	25

145	Sporadic and familial dementia with ubiquitin-positive tau-negative inclusions: clinical features of one histopathological abnormality underlying frontotemporal lobar degeneration. *Archives of Neurology*, **2005**, 62, 1097-101		25
144	Cerebral mitochondrial electron transport chain dysfunction in multiple system atrophy and Parkinson disease. *Scientific Reports*, **2019**, 9, 6559	4.9	24
143	Neuropathology of Beta-propeller protein associated neurodegeneration (BPAN): a new tauopathy. *Acta Neuropathologica Communications*, **2015**, 3, 39	7.3	24
142	Pathological correlates of white matter hyperintensities in a case of progranulin mutation associated frontotemporal dementia. *Neurocase*, **2018**, 24, 166-174	0.8	24
141	COX10 mutations resulting in complex multisystem mitochondrial disease that remains stable into adulthood. *JAMA Neurology*, **2013**, 70, 1556-61	17.2	24
140	Preferential association of serum amyloid P component with fibrillar deposits in familial British and Danish dementias: similarities with Alzheimer disease. *Journal of the Neurological Sciences*, **2007**, 257, 88-96	3.2	24
139	Molecular chaperons, amyloid and preamyloid lesions in the BRI2 gene-related dementias: a morphological study. *Neuropathology and Applied Neurobiology*, **2006**, 32, 492-504	5.2	24
138	Evidence for pathological involvement of the spinal cord in motor neuron disease-inclusion dementia. *Acta Neuropathologica*, **2002**, 103, 221-7	14.3	24
137	Histological heterogeneity in a large clinical cohort of juvenile idiopathic inflammatory myopathy: analysis by myositis autoantibody and pathological features. *Neuropathology and Applied Neurobiology*, **2019**, 45, 495-512	5.2	24
136	Expression of BRI2 mRNA and protein in normal human brain and familial British dementia: its relevance to the pathogenesis of disease. *Neuropathology and Applied Neurobiology*, **2008**, 34, 492-505	5.2	23
135	Investigation of somatic CNVs in brains of synucleinopathy cases using targeted SNCA analysis and single cell sequencing. *Acta Neuropathologica Communications*, **2019**, 7, 219	7.3	22
134	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
133	Disease-related patterns of in vivo pathology in Corticobasal syndrome. *European Journal of Nuclear Medicine and Molecular Imaging*, **2018**, 45, 2413-2425	8.8	21
132	Tau acts as an independent genetic risk factor in pathologically proven PD. *Neurobiology of Aging*, **2012**, 33, 838.e7-11	5.6	20
131	Silver staining (Campbell-Switzer) of neuronal -synuclein assemblies induced by multiple system atrophy and Parkinson disease brain extracts in transgenic mice. *Acta Neuropathologica Communications*, **2019**, 7, 148	7.3	19
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List of Publications

1 Antibody-Mediated Muscle Disease? **2017**, 203-206