List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Distinct populations of highly potent TAU seed conformers in rapidly progressing Alzheimer's disease. Science Translational Medicine, 2022, 14, eabg0253.	5.8	26
2	Elevated 4Râ€ŧau in astrocytes from asymptomatic carriers of the <i>MAPT</i> 10+16 intronic mutation. Journal of Cellular and Molecular Medicine, 2022, 26, 1327-1331.	1.6	6
3	MIR-NATs repress MAPT translation and aid proteostasis in neurodegeneration. Nature, 2021, 594, 117-123.	13.7	29
4	A clinical, molecular genetics and pathological study of a FTDP-17 family with a heterozygous splicing variant c.823-10G>T at the intron 9/exon 10 of the MAPT gene. Neurobiology of Aging, 2021, 106, 343.e1-343.e8.	1.5	5
5	MOBP and HIP1 in multiple system atrophy: New αâ€synuclein partners in glial cytoplasmic inclusions implicated in the disease pathogenesis. Neuropathology and Applied Neurobiology, 2021, 47, 640-652.	1.8	11
6	Reply to â€~Impulse control disorders are associated with lower ventral striatum dopamine D3 receptor availability in Parkinson's disease: A [11C]-PHNO PET study.'. Parkinsonism and Related Disorders, 2021, 93, 31-32.	1.1	1
7	Fulminant corticobasal degeneration: a distinct variant with predominant neuronal tau aggregates. Acta Neuropathologica, 2020, 139, 717-734.	3.9	15
8	Pre-clinical characterisation of E2814, a high-affinity antibody targeting the microtubule-binding repeat domain of tau for passive immunotherapy in Alzheimer's disease. Acta Neuropathologica Communications, 2020, 8, 13.	2.4	61
9	Lower nucleus accumbens α-synuclein load and D3 receptor levels in Parkinson's disease with impulsive compulsive behaviours. Brain, 2019, 142, 3580-3591.	3.7	17
10	A walk through tau therapeutic strategies. Acta Neuropathologica Communications, 2019, 7, 22.	2.4	211
11	Association of <i>MAPT</i> haplotypeâ€tagging polymorphisms with cerebrospinal fluid biomarkers of Alzheimer's disease: A preliminary study in a Croatian cohort. Brain and Behavior, 2018, 8, e01128.	1.0	20
12	Variation at the <i>TRIM11</i> locus modifies progressive supranuclear palsy phenotype. Annals of Neurology, 2018, 84, 485-496.	2.8	37
13	Foamy Virus Vectors Transduce Visceral Organs and Hippocampal Structures following InÂVivo Delivery to Neonatal Mice. Molecular Therapy - Nucleic Acids, 2018, 12, 626-634.	2.3	7
14	Monoaminergic neuropathology in Alzheimer's disease. Progress in Neurobiology, 2017, 151, 101-138.	2.8	206
15	Tau Protein Hyperphosphorylation and Aggregation in Alzheimer's Disease and Other Tauopathies, and Possible Neuroprotective Strategies. Biomolecules, 2016, 6, 6.	1.8	503
16	The role of tau in the pathological process and clinical expression of Huntington's disease. Brain, 2015, 138, 1907-1918.	3.7	115
17	The analysis of C9orf72 repeat expansions in a large series of clinically and pathologically diagnosed cases with atypical parkinsonism. Neurobiology of Aging, 2015, 36, 1221.e1-1221.e6.	1.5	39
18	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.	3.0	74

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19	A cognitive chameleon: Lessons from a novel <i>MAPT</i> mutation case. Neurocase, 2014, 20, 684-694.	0.2	12
20	Reduced Vascular Endothelial Growth Factor and Capillary Density in the Occipital Cortex in Dementia with <scp>L</scp> ewy Bodies. Brain Pathology, 2014, 24, 334-343.	2.1	39
21	Genetic variability in the regulation of gene expression in ten regions of the human brain. Nature Neuroscience, 2014, 17, 1418-1428.	7.1	620
22	Assessment of common variability and expression quantitative trait loci for genome-wide associations for progressive supranuclear palsy. Neurobiology of Aging, 2014, 35, 1514.e1-1514.e12.	1.5	33
23	Widespread RNA metabolism impairment in sporadic inclusion body myositis TDP43-proteinopathy. Neurobiology of Aging, 2014, 35, 1491-1498.	1.5	36
24	Validation of next-generation sequencing technologies in genetic diagnosis of dementia. Neurobiology of Aging, 2014, 35, 261-265.	1.5	59
25	Variation in tau isoform expression in different brain regions and disease states. Neurobiology of Aging, 2013, 34, 1922.e7-1922.e12.	1.5	49
26	Fine-Mapping, Gene Expression and Splicing Analysis of the Disease Associated LRRK2 Locus. PLoS ONE, 2013, 8, e70724.	1.1	45
27	Development and assessment of sensitive immunoâ€ <scp>PCR</scp> assays for the quantification of cerebrospinal fluid threeâ€and fourâ€repeat tau isoforms in tauopathies. Journal of Neurochemistry, 2012, 123, 396-405.	2.1	64
28	MAPT expression and splicing is differentially regulated by brain region: relation to genotype and implication for tauopathies. Human Molecular Genetics, 2012, 21, 4094-4103.	1.4	191
29	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.	1.5	60
30	Identification of common variants influencing risk of the tauopathy progressive supranuclear palsy. Nature Genetics, 2011, 43, 699-705.	9.4	502
31	Tau-positive grains are constant in centenarians' hippocampus. Neurobiology of Aging, 2011, 32, 1296-1303.	1.5	20
32	Globular glial tauopathies (GGT) presenting with motor neuron disease or frontotemporal dementia: an emerging group of 4-repeat tauopathies. Acta Neuropathologica, 2011, 122, 415-428.	3.9	67
33	Clinical and neuroanatomical signatures of tissue pathology in frontotemporal lobar degeneration. Brain, 2011, 134, 2565-2581.	3.7	306
34	Lewy- and Alzheimer-type pathologies in Parkinson's disease dementia: which is more important?. Brain, 2011, 134, 1493-1505.	3.7	497
35	Novel L284R <i>MAPT </i> Mutation in a Family with an Autosomal Dominant Progressive Supranuclear Palsy Syndrome. Neurodegenerative Diseases, 2011, 8, 149-152.	0.8	47
36	Rational therapeutic approaches to progressive supranuclear palsy. Brain, 2010, 133, 1578-1590.	3.7	83

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37	Adapting the Sniffin' Sticks to diagnose Parkinson's disease in Sri Lanka. Movement Disorders, 2009, 24, 1229-1233.	2.2	38
38	Development of a sensitive ELISA for quantification of three- and four-repeat tau isoforms in tauopathies. Journal of Neuroscience Methods, 2009, 180, 34-42.	1.3	23
39	Association of MAPT haplotype-tagging SNPs with sporadic Parkinson's disease. Neurobiology of Aging, 2009, 30, 1477-1482.	1.5	48
40	Concomitant progressive supranuclear palsy and multiple system atrophy: More than a simple twist of fate?. Neuroscience Letters, 2009, 467, 208-211.	1.0	19
41	Aging Analysis Reveals Slowed Tau Turnover and Enhanced Stress Response in a Mouse Model of Tauopathy. American Journal of Pathology, 2009, 174, 228-238.	1.9	73
42	MAPT S305I mutation: implications for argyrophilic grain disease. Acta Neuropathologica, 2008, 116, 103-118.	3.9	52
43	Genetics of progressive supranuclear palsy. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2008, 89, 475-485.	1.0	0
44	Differential Incorporation of Tau Isoforms in Alzheimer's Disease. Journal of Alzheimer's Disease, 2008, 14, 1-16.	1.2	107
45	Pathological tau burden and distribution distinguishes progressive supranuclear palsy-parkinsonism from Richardson's syndrome. Brain, 2007, 130, 1566-1576.	3.7	364
46	Expression of embryonic tau protein isoforms persist during adult neurogenesis in the hippocampus. Hippocampus, 2007, 17, 98-102.	0.9	51
47	Genetic variation at the tau locus and clinical syndromes associated with progressive supranuclear palsy. Movement Disorders, 2007, 22, 895-897.	2.2	25
48	Pick's disease with Pick bodies: An unusual autopsy case showing degeneration of the pontine nucleus, dentate nucleus, Clarke's column, and lower motor neuron. Neuropathology, 2007, 27, 81-89.	0.7	3
49	The MAPT H1c risk haplotype is associated with increased expression of tau and especially of 4 repeat containing transcripts. Neurobiology of Disease, 2007, 25, 561-570.	2.1	231
50	DJ-1 (PARK7) is associated with 3R and 4R tau neuronal and glial inclusions in neurodegenerative disorders. Neurobiology of Disease, 2007, 28, 122-132.	2.1	32
51	Tangle Diseases and the Tau Haplotypes. Alzheimer Disease and Associated Disorders, 2006, 20, 60-62.	0.6	6
52	Microdeletion encompassing MAPT at chromosome 17q21.3 is associated with developmental delay and learning disability. Nature Genetics, 2006, 38, 1032-1037.	9.4	344
53	An immunohistochemical study of cases of sporadic and inherited frontotemporal lobar degeneration using 3R- and 4R-specific tau monoclonal antibodies. Acta Neuropathologica, 2006, 111, 329-340.	3.9	91
54	No alteration in tau exon 10 alternative splicing in tangle-bearing neurons of the Alzheimer's disease brain. Acta Neuropathologica, 2006, 112, 439-449.	3.9	41

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55	Amyotrophic lateral sclerosis with dementia: an autopsy case showing many Bunina bodies, tau-positive neuronal and astrocytic plaque-like pathologies, and pallido-nigral degeneration. Acta Neuropathologica, 2006, 112, 633-645.	3.9	28
56	Untangling the tau gene association with neurodegenerative disorders. Human Molecular Genetics, 2006, 15, R188-R195.	1.4	102
57	Sporadic four-repeat tauopathy with frontotemporal degeneration, parkinsonism and motor neuron disease. Acta Neuropathologica, 2005, 110, 600-609.	3.9	30
58	Characteristics of two distinct clinical phenotypes in pathologically proven progressive supranuclear palsy: Richardson's syndrome and PSP-parkinsonism. Brain, 2005, 128, 1247-1258.	3.7	743
59	The architecture of the tau haplotype block in different ethnicities. Neuroscience Letters, 2005, 377, 81-84.	1.0	13
60	Development, characterisation and epitope mapping of novel monoclonal antibodies for DJ-1 (PARK7) protein. Neuroscience Letters, 2005, 383, 225-230.	1.0	11
61	The structure of the tau haplotype in controls and in progressive supranuclear palsy. Human Molecular Genetics, 2004, 13, 1267-1274.	1.4	119
62	The expression of DJ-1 (PARK7) in normal human CNS and idiopathic Parkinson's disease. Brain, 2004, 127, 420-430.	3.7	404
63	Detecting tau isoforms in archival cases. Acta Neuropathologica, 2004, 107, 181-182.	3.9	3
64	Immunohistochemical study of tau accumulation in early stages of Alzheimer-type neurofibrillary lesions. Acta Neuropathologica, 2004, 107, 504-508.	3.9	37
65	Alzheimer's associated variant ubiquitin causes inhibition of the 26S proteasome and chaperone expression. Journal of Neurochemistry, 2004, 86, 394-404.	2.1	78
66	Cloning of the Gene Containing Mutations that Cause PARK8-Linked Parkinson's Disease. Neuron, 2004, 44, 595-600.	3.8	2,183
67	Failure in heat-shock protein expression in response to UBB+1 protein in progressive supranuclear palsy in humans. Neuroscience Letters, 2004, 359, 94-98.	1.0	5
68	The tau H2 haplotype is almost exclusively Caucasian in origin. Neuroscience Letters, 2004, 369, 183-185.	1.0	102
69	Dementia with Lewy bodies from the perspective of tauopathy. Acta Neuropathologica, 2003, 105, 265-270.	3.9	67
70	4-repeat tauopathy sharing pathological and biochemical features of corticobasal degeneration and progressive supranuclear palsy. Acta Neuropathologica, 2003, 106, 251-260.	3.9	45
71	The L266V tau mutation is associated with frontotemporal dementia and Pick-like 3R and 4R tauopathy. Acta Neuropathologica, 2003, 106, 323-336.	3.9	84
72	Hyperphosphorylation and aggregation of tau in mice expressing normal human tau isoforms. Journal of Neurochemistry, 2003, 86, 582-590.	2.1	662

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73	Argyrophilic Grain Disease Is a Sporadic 4-Repeat Tauopathy. Journal of Neuropathology and Experimental Neurology, 2002, 61, 547-556.	0.9	232
74	Tau neurotoxicity without the lesions: a fly challenges a tangled web. Trends in Neurosciences, 2002, 25, 327-329.	4.2	22
75	The tau locus is not significantly associated with pathologically confirmed sporadic Parkinson's disease. Neuroscience Letters, 2002, 330, 201-203.	1.0	39
76	The Slow Axonal Transport of the Microtubule-Associated Protein Tau and the Transport Rates of Different Isoforms and Mutants in Cultured Neurons. Journal of Neuroscience, 2002, 22, 6394-6400.	1.7	69
77	No pathogenic mutations in the synphilin-1 gene in Parkinson's disease. Neuroscience Letters, 2001, 307, 125-127.	1.0	18
78	Strong association of a novel Tau promoter haplotype in progressive supranuclear palsy. Neuroscience Letters, 2001, 311, 145-148.	1.0	49