

Naomi Kouri

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

6,669
citations

304743

22
h-index

526287

27
g-index

30
all docs

30
docs citations

30
times ranked

10364
citing authors

#	ARTICLE	IF	CITATIONS
1	Latent trait modeling of tau neuropathology in progressive supranuclear palsy. <i>Acta Neuropathologica</i> , 2021, 141, 667-680.	7.7	5
2	MAPT subhaplotypes in corticobasal degeneration: assessing associations with disease risk, severity of tau pathology, and clinical features. <i>Acta Neuropathologica Communications</i> , 2020, 8, 218.	5.2	8
3	Selective Genetic Overlap Between Amyotrophic Lateral Sclerosis and Diseases of the Frontotemporal Dementia Spectrum. <i>JAMA Neurology</i> , 2018, 75, 860.	9.0	79
4	Replication of progressive supranuclear palsy genome-wide association study identifies <i>SLCO1A2</i> and <i>DUSP10</i> as new susceptibility loci. <i>Molecular Neurodegeneration</i> , 2018, 13, 37.	10.8	54
5	Divergent brain gene expression patterns associate with distinct cell-specific tau neuropathology traits in progressive supranuclear palsy. <i>Acta Neuropathologica</i> , 2018, 136, 709-727.	7.7	47
6	Immune-related genetic enrichment in frontotemporal dementia: An analysis of genome-wide association studies. <i>PLoS Medicine</i> , 2018, 15, e1002487.	8.4	111
7	Corticobasal degeneration with TDP-43 pathology presenting with progressive supranuclear palsy syndrome: a distinct clinicopathologic subtype. <i>Acta Neuropathologica</i> , 2018, 136, 389-404.	7.7	59
8	Shared genetic risk between corticobasal degeneration, progressive supranuclear palsy, and frontotemporal dementia. <i>Acta Neuropathologica</i> , 2017, 133, 825-837.	7.7	90
9	FTDP-17 with Pick body-like inclusions associated with a novel tau mutation, p.E372G. <i>Brain Pathology</i> , 2017, 27, 612-626.	4.1	11
10	Gene expression, methylation and neuropathology correlations at progressive supranuclear palsy risk loci. <i>Acta Neuropathologica</i> , 2016, 132, 197-211.	7.7	49
11	A novel tau mutation, p.K317N, causes globular glial tauopathy. <i>Acta Neuropathologica</i> , 2015, 130, 199-214.	7.7	38
12	Genome-wide association study of corticobasal degeneration identifies risk variants shared with progressive supranuclear palsy. <i>Nature Communications</i> , 2015, 6, 7247.	12.8	170
13	Novel mutation in MAPT exon 13 (p.N410H) causes corticobasal degeneration. <i>Acta Neuropathologica</i> , 2014, 127, 271-282.	7.7	66
14	Clinicopathologic assessment and imaging of tauopathies in neurodegenerative dementias. <i>Alzheimer's Research and Therapy</i> , 2014, 6, 1.	6.2	156
15	Corticobasal degeneration with olivopontocerebellar atrophy and TDP-43 pathology: an unusual clinicopathologic variant of CBD. <i>Acta Neuropathologica</i> , 2013, 125, 741-752.	7.7	40
16	Progressive amnesic dementia, hippocampal sclerosis, and mutation in <i>C9ORF72</i> . <i>Acta Neuropathologica</i> , 2013, 126, 545-554.	7.7	30
17	<i>LRRTM3</i> Interacts with APP and BACE1 and Has Variants Associating with Late-Onset Alzheimer's Disease (LOAD). <i>PLoS ONE</i> , 2013, 8, e64164.	2.5	12
18	Brain Expression Genome-Wide Association Study (eGWAS) Identifies Human Disease-Associated Variants. <i>PLoS Genetics</i> , 2012, 8, e1002707.	3.5	225

#	ARTICLE	IF	CITATIONS
19	Mutations in the colony stimulating factor 1 receptor (CSF1R) gene cause hereditary diffuse leukoencephalopathy with spheroids. <i>Nature Genetics</i> , 2012, 44, 200-205.	21.4	428
20	Corticobasal degeneration: a pathologically distinct 4R tauopathy. <i>Nature Reviews Neurology</i> , 2011, 7, 263-272.	10.1	270
21	Expanded GGGGCC Hexanucleotide Repeat in Noncoding Region of C9ORF72 Causes Chromosome 9p-Linked FTD and ALS. <i>Neuron</i> , 2011, 72, 245-256.	8.1	4,176
22	Neuropathology of Frontotemporal Lobar Degeneration-Tau (FTLD-Tau). <i>Journal of Molecular Neuroscience</i> , 2011, 45, 384-389.	2.3	295
23	Altered microRNA expression in frontotemporal lobar degeneration with TDP-43 pathology caused by progranulin mutations. <i>BMC Genomics</i> , 2011, 12, 527.	2.8	48
24	Neuropathological features of corticobasal degeneration presenting as corticobasal syndrome or Richardson syndrome. <i>Brain</i> , 2011, 134, 3264-3275.	7.6	119
25	Gene expression levels as endophenotypes in genome-wide association studies of Alzheimer disease. <i>Neurology</i> , 2010, 74, 480-486.	1.1	30
26	Spectrophotometric determination of aqueous cyanide using a revised phenolphthalin method. <i>Analytica Chimica Acta</i> , 2007, 589, 137-141.	5.4	45