Kevin F Bieniek

List of Publications by Year in Descending Order

40 papers	4,328 citations	25 h-index	41 g-index
41 ext. papers	5,076 ext. citations	10.1 avg, IF	4.68 L-index

#	Paper	IF	Citations
40	AuthorsSResponse. *Journal of Neuropathology and Experimental Neurology*, **2021**, 80, 1008-1010	3.1	1
39	Re: The Second NINDS/NIBIB Consensus Meeting to Define Neuropathological Criteria for the Diagnosis of Chronic Traumatic Encephalopathy. J Neuropathol Exp Neurol 2021;80(3):210-9. *Journal of Neuropathology and Experimental Neurology*, **2021**, 80, 1007-1008	3.1	1
38	COVID-19 Patients With CNS Complications and Neuropathologic Features of Acute Disseminated Encephalomyelitis and Acute Hemorrhagic Leukoencephalopathy. *Journal of Neuropathology and Experimental Neurology*, **2021**, 80, 628-631	3.1	7
37	The Second NINDS/NIBIB Consensus Meeting to Define Neuropathological Criteria for the Diagnosis of Chronic Traumatic Encephalopathy. *Journal of Neuropathology and Experimental Neurology*, **2021**, 80, 210-219	3.1	32
36	Practical Considerations in the Diagnosis of Mild Chronic Traumatic Encephalopathy and Distinction From Age-Related Tau Astrogliopathy. *Journal of Neuropathology and Experimental Neurology*, **2020**, 79, 921-924	3.1	6
35	Association between contact sports participation and chronic traumatic encephalopathy: a retrospective cohort study. *Brain Pathology*, **2020**, 30, 63-74	6	38
34	PET-detectable tau pathology correlates with long-term neuropsychiatric outcomes in patients with traumatic brain injury. *Brain*, **2019**, 142, 3265-3279	11.2	31
33	Ethnoracial differences in Alzheimer's disease from the FLorida Autopsied Multi-Ethnic (FLAME) cohort. *Alzheimer's and Dementia*, **2019**, 15, 635-643	1.2	17
32	Genome-wide analyses as part of the international FTLD-TDP whole-genome sequencing consortium reveals novel disease risk factors and increases support for immune dysfunction in FTLD. *Acta Neuropathologica*, **2019**, 137, 879-899	14.3	50
31	Microglia in frontotemporal lobar degeneration with progranulin or C9ORF72 mutations. *Annals of Clinical and Translational Neurology*, **2019**, 6, 1782-1796	5.3	11
30	A quantitative risk assessment for chronic traumatic encephalopathy (CTE) in football: How public health science evaluates evidence. *Human and Ecological Risk Assessment (HERA)*, **2019**, 25, 564-589	4.9	8
29	Potential genetic modifiers of disease risk and age at onset in patients with frontotemporal lobar degeneration and GRN mutations: a genome-wide association study. *Lancet Neurology, The*, **2018**, 17, 548-558	24.1	60
28	Poly-GR dipeptide repeat polymers correlate with neurodegeneration and Clinicopathological subtypes in C9ORF72-related brain disease. *Acta Neuropathologica Communications*, **2018**, 6, 63	7.3	51
27	In-depth clinico-pathological examination of RNA foci in a large cohort of C9ORF72 expansion carriers. *Acta Neuropathologica*, **2017**, 134, 255-269	14.3	57
26	Multisite Assessment of Aging-Related Tau Astrogliopathy (ARTAG). *Journal of Neuropathology and Experimental Neurology*, **2017**, 76, 605-619	3.1	28
25	Abnormal expression of homeobox genes and transthyretin in expansion carriers. *Neurology: Genetics*, **2017**, 3, e161	3.8	9
24	Chronic Traumatic Encephalopathy Pathology in Multiple System Atrophy. *Journal of Neuropathology and Experimental Neurology*, **2016**, 75, 963-970	3.1	45

23	Aging-related tau astrogliopathy (ARTAG): harmonized evaluation strategy. *Acta Neuropathologica*, **2016**, 131, 87-102	14.3	272
22	The first NINDS/NIBIB consensus meeting to define neuropathological criteria for the diagnosis of chronic traumatic encephalopathy. *Acta Neuropathologica*, **2016**, 131, 75-86	14.3	524
21	Chapter 14 Deep Brain Stimulation for Memory Dysfunction **2016**, 257-274		
20	A truncating SOD1 mutation, p.Gly141X, is associated with clinical and pathologic heterogeneity, including frontotemporal lobar degeneration. *Acta Neuropathologica*, **2015**, 130, 145-57	14.3	17
19	Distinct brain transcriptome profiles in C9orf72-associated and sporadic ALS. *Nature Neuroscience*, **2015**, 18, 1175-82	25.5	235
18	Neurodegeneration. C9ORF72 repeat expansions in mice cause TDP-43 pathology, neuronal loss, and behavioral deficits. *Science*, **2015**, 348, 1151-4	33.3	279
17	Whole-genome sequencing reveals important role for TBK1 and OPTN mutations in frontotemporal lobar degeneration without motor neuron disease. *Acta Neuropathologica*, **2015**, 130, 77-92	14.3	222
16	Concurrent neurodegenerative pathologies in periventricular nodular heterotopia. *Acta Neuropathologica*, **2015**, 130, 895-7	14.3	4
15	Cerebellar c9RAN proteins associate with clinical and neuropathological characteristics of C9ORF72 repeat expansion carriers. *Acta Neuropathologica*, **2015**, 130, 559-73	14.3	72
14	Chronic traumatic encephalopathy pathology in a neurodegenerative disorders brain bank. *Acta Neuropathologica*, **2015**, 130, 877-89	14.3	176
13	TMEM106B protects C9ORF72 expansion carriers against frontotemporal dementia. *Acta Neuropathologica*, **2014**, 127, 397-406	14.3	108
12	Concurrent variably protease-sensitive prionopathy and amyotrophic lateral sclerosis. *Acta Neuropathologica*, **2014**, 128, 313-315	14.3	8
11	Aggregation-prone c9FTD/ALS poly(GA) RAN-translated proteins cause neurotoxicity by inducing ER stress. *Acta Neuropathologica*, **2014**, 128, 505-24	14.3	227
10	Expanded C9ORF72 hexanucleotide repeat in depressive pseudodementia. *JAMA Neurology*, **2014**, 71, 775-81	17.2	24
9	Tau pathology in frontotemporal lobar degeneration with C9ORF72 hexanucleotide repeat expansion. *Acta Neuropathologica*, **2013**, 125, 289-302	14.3	75
8	Unconventional translation of C9ORF72 GGGGCC expansion generates insoluble polypeptides specific to c9FTD/ALS. *Neuron*, **2013**, 77, 639-46	13.9	783
7	Validation of the Mayo Sleep Questionnaire to screen for REM sleep behavior disorder in a community-based sample. *Journal of Clinical Sleep Medicine*, **2013**, 9, 475-80	3.1	98
6	Antisense transcripts of the expanded C9ORF72 hexanucleotide repeat form nuclear RNA foci and undergo repeat-associated non-ATG translation in c9FTD/ALS. *Acta Neuropathologica*, **2013**, 126, 829-44	14.3	392

5	Progressive amnestic dementia, hippocampal sclerosis, and mutation in C9ORF72. *Acta Neuropathologica*, **2013**, 126, 545-54	14.3	22
4	C9ORF72 repeat expansions in cases with previously identified pathogenic mutations. *Neurology*, **2013**, 81, 1332-41	6.5	75
3	Profilin-1 mutations are rare in patients with amyotrophic lateral sclerosis and frontotemporal dementia. *Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration*, **2013**, 14, 463-9	3.6	23
2	Subtle gait changes in patients with REM sleep behavior disorder. *Movement Disorders*, **2013**, 28, 1847-53	7	42
1	Validation of the Mayo Sleep Questionnaire to screen for REM sleep behavior disorder in an aging and dementia cohort. *Sleep Medicine*, **2011**, 12, 445-53	4.6	198