

Andrew King

List of Publications by Year in descending order

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Version: 2024-02-01

60
papers

4,094
citations

236612

25
h-index

161609

54
g-index

61
all docs

61
docs citations

61
times ranked

6438
citing authors

#	ARTICLE	IF	CITATIONS
1	Genome-wide Analyses Identify KIF5A as a Novel ALS Gene. <i>Neuron</i> , 2018, 97, 1268-1283.e6.	3.8	517
2	Staging of Neurofibrillary Pathology in Alzheimer's Disease: A Study of the BrainNet Europe Consortium. <i>Brain Pathology</i> , 2008, 18, 484-496.	2.1	361
3	The genetics and neuropathology of amyotrophic lateral sclerosis. <i>Acta Neuropathologica</i> , 2012, 124, 339-352.	3.9	346
4	Exome-wide Rare Variant Analysis Identifies TUBA4A Mutations Associated with Familial ALS. <i>Neuron</i> , 2014, 84, 324-331.	3.8	308
5	Staging/typing of Lewy body related α -synuclein pathology: a study of the BrainNet Europe Consortium. <i>Acta Neuropathologica</i> , 2009, 117, 635-652.	3.9	249
6	Vascular cognitive impairment neuropathology guidelines (VCING): the contribution of cerebrovascular pathology to cognitive impairment. <i>Brain</i> , 2016, 139, 2957-2969.	3.7	220
7	NEK1 variants confer susceptibility to amyotrophic lateral sclerosis. <i>Nature Genetics</i> , 2016, 48, 1037-1042.	9.4	218
8	Plasma p-tau181 accurately predicts Alzheimer's disease pathology at least 8 years prior to post-mortem and improves the clinical characterisation of cognitive decline. <i>Acta Neuropathologica</i> , 2020, 140, 267-278.	3.9	209
9	Aluminium in brain tissue in familial Alzheimer's disease. <i>Journal of Trace Elements in Medicine and Biology</i> , 2017, 40, 30-36.	1.5	182
10	Mutations in the vesicular trafficking protein annexin A11 are associated with amyotrophic lateral sclerosis. <i>Science Translational Medicine</i> , 2017, 9, .	5.8	129
11	Aluminium in brain tissue in autism. <i>Journal of Trace Elements in Medicine and Biology</i> , 2018, 46, 76-82.	1.5	112
12	TDP-43 is consistently co-localized with ubiquitinated inclusions in sporadic and Guam amyotrophic lateral sclerosis but not in familial amyotrophic lateral sclerosis with and without SOD1 mutations. <i>Neuropathology</i> , 2009, 29, 672-683.	0.7	108
13	Abnormal TDP-43 expression is identified in the neocortex in cases of dementia pugilistica, but is mainly confined to the limbic system when identified in high and moderate stages of Alzheimer's disease. <i>Neuropathology</i> , 2010, 30, 408-419.	0.7	98
14	Genome-wide analyses as part of the international FTLTDP whole-genome sequencing consortium reveals novel disease risk factors and increases support for immune dysfunction in FTLTDP. <i>Acta Neuropathologica</i> , 2019, 137, 879-899.	3.9	90
15	Novel mutations support a role for Profilin 1 in the pathogenesis of ALS. <i>Neurobiology of Aging</i> , 2015, 36, 1602.e17-1602.e27.	1.5	87
16	Mixed tau, TDP-43 and p62 pathology in FTLTDP associated with a C9ORF72 repeat expansion and p.Ala239Thr MAPT (tau) variant. <i>Acta Neuropathologica</i> , 2013, 125, 303-310.	3.9	73
17	Ubiquitinated, p62 immunopositive cerebellar cortical neuronal inclusions are evident across the spectrum of TDP-43 proteinopathies but are only rarely additionally immunopositive for phosphorylation-dependent TDP-43. <i>Neuropathology</i> , 2011, 31, 239-249.	0.7	58
18	Two novel mutations in the gene for copper zinc superoxide dismutase in UK families with amyotrophic lateral sclerosis. <i>Human Molecular Genetics</i> , 1995, 4, 1239-1240.	1.4	54

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19	Neuropathology of the hippocampus in FTLDâ€”Tau with Pick bodies: a study of the BrainNet Europe Consortium. <i>Neuropathology and Applied Neurobiology</i> , 2013, 39, 166-178.	1.8	54
20	The Identification of Aluminum in Human Brain Tissue Using Lumogallion and Fluorescence Microscopy. <i>Journal of Alzheimer's Disease</i> , 2016, 54, 1333-1338.	1.2	48
21	Association of Variants in the <i>SPTLC1</i> Gene With Juvenile Amyotrophic Lateral Sclerosis. <i>JAMA Neurology</i> , 2021, 78, 1236.	4.5	46
22	The Increased Densities, But Different Distributions, of Both C3 and S100A10 Immunopositive Astrocyte-Like Cells in Alzheimerâ€™s Disease Brains Suggest Possible Roles for Both A1 and A2 Astrocytes in the Disease Pathogenesis. <i>Brain Sciences</i> , 2020, 10, 503.	1.1	43
23	Mitochondrial abnormalities and low grade inflammation are present in the skeletal muscle of a minority of patients with amyotrophic lateral sclerosis; an observational myopathology study. <i>Acta Neuropathologica Communications</i> , 2014, 2, 165.	2.4	40
24	Multisite Assessment of Aging-Related Tau Astroglialopathy (ARTAG). <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 605-619.	0.9	38
25	Comparison of clinical and neuropathological diagnoses of neurodegenerative diseases in two centres from the Brains for Dementia Research (BDR) cohort. <i>Journal of Neural Transmission</i> , 2019, 126, 327-337.	1.4	33
26	Aluminium in Brain Tissue in Multiple Sclerosis. <i>International Journal of Environmental Research and Public Health</i> , 2018, 15, 1777.	1.2	31
27	Clusterin expression is upregulated following acute head injury and localizes to astrocytes in old head injury. <i>Neuropathology</i> , 2017, 37, 12-24.	0.7	24
28	Phenotypic variability in the brains of a family with a prion disease characterized by a 144-base pair insertion in the prion protein gene. <i>Neuropathology and Applied Neurobiology</i> , 2003, 29, 98-105.	1.8	23
29	<i>RBCK1</i> -related disease: A rare multisystem disorder with polyglucosan storage, autoinflammation, recurrent infections, skeletal, and cardiac myopathyâ€”Four additional patients and a review of the current literature. <i>Journal of Inherited Metabolic Disease</i> , 2020, 43, 1002-1013.	1.7	23
30	ALS-FUS pathology revisited: singleton FUS mutations and an unusual case with both a FUS and TARDBP mutation. <i>Acta Neuropathologica Communications</i> , 2015, 3, 62.	2.4	22
31	The Neuropathological Diagnosis of Alzheimerâ€™s Diseaseâ€”The Challenges of Pathological Mimics and Concomitant Pathology. <i>Brain Sciences</i> , 2020, 10, 479.	1.1	22
32	Frontotemporal lobar degeneration with ubiquitinated tau-negative inclusions and additional Î±-synuclein pathology but also unusual cerebellar ubiquitinated p62-positive, TDPâ€”43-negative inclusions. <i>Neuropathology</i> , 2009, 29, 466-471.	0.7	19
33	Extended post-mortem delay times should not be viewed as a deterrent to the scientific investigation of human brain tissue: a study from the Brains for Dementia Research Network Neuropathology Study Group, UK. <i>Acta Neuropathologica</i> , 2016, 132, 753-755.	3.9	18
34	Comparative gene expression profiling of ADAMs, MMPs, TIMPs, EMMPRIN, EGF-R and VEGFA in low grade meningioma. <i>International Journal of Oncology</i> , 2016, 49, 2309-2318.	1.4	17
35	Intracellular Aluminium in Inflammatory and Glial Cells in Cerebral Amyloid Angiopathy: A Case Report. <i>International Journal of Environmental Research and Public Health</i> , 2019, 16, 1459.	1.2	17
36	Assessment of the degree of asymmetry of pathological features in neurodegenerative diseases. What is the significance for brain banks?. <i>Journal of Neural Transmission</i> , 2015, 122, 1499-1508.	1.4	16

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37	Striking phenotypic variation in a family with the P506S UBQLN2 mutation including amyotrophic lateral sclerosis, spastic paraplegia, and frontotemporal dementia. <i>Neurobiology of Aging</i> , 2019, 73, 229.e5-229.e9.	1.5	16
38	SIL1-related Marinescoâ€”Sjogren syndrome (MSS) with associated motor neuronopathy and bradykinetic movement disorder. <i>Neuromuscular Disorders</i> , 2015, 25, 585-588.	0.3	14
39	Heterogeneous Nuclear Ribonucleoprotein E2 (hnRNP E2) Is a Component of TDP-43 Aggregates Specifically in the A and C Pathological Subtypes of Frontotemporal Lobar Degeneration. <i>Frontiers in Neuroscience</i> , 2019, 13, 551.	1.4	13
40	<i>APOE</i> ϵ 4 is also required in <i>TREM2</i> R47H variant carriers for Alzheimer's disease to develop. <i>Neuropathology and Applied Neurobiology</i> , 2019, 45, 183-186.	1.8	12
41	Delayed presentation of late-onset cerebrospinal fluid rhinorrhoea following dopamine agonist therapy for giant prolactinoma. <i>Endocrinology, Diabetes and Metabolism Case Reports</i> , 2014, 2014, 140020.	0.2	12
42	Simulated surgical-type cerebral biopsies from post-mortem brains allows accurate neuropathological diagnoses in the majority of neurodegenerative disease groups. <i>Acta Neuropathologica Communications</i> , 2013, 1, 53.	2.4	10
43	Unusual neuropathological features and increased brain aluminium in a resident of Camelford, UK. <i>Neuropathology and Applied Neurobiology</i> , 2017, 43, 537-541.	1.8	8
44	Adult Parasellar Capillary Hemangioma with Intrasellar Extension. <i>World Neurosurgery</i> , 2019, 124, 184-191.	0.7	8
45	Primary intracerebral INI1-deficient rhabdoid tumor with CD34 immunopositivity in a young adult. , 2018, 9, 45.		8
46	Expression of the chondroitin sulphate proteoglycan, NG2, in paediatric brain tumors. <i>Anticancer Research</i> , 2014, 34, 6919-24.	0.5	7
47	â€œBrain on fireâ€: A new imaging sign. <i>Journal of Clinical Neuroscience</i> , 2014, 21, 2015-2017.	0.8	6
48	Spontaneous epidural spinal haematoma in children caused by vascular malformations. <i>European Spine Journal</i> , 2016, 25, 614-618.	1.0	6
49	IgG4-related hypophysitis in adolescence. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2021, 34, 395-399.	0.4	4
50	Expression of the candidate invasion suppressor gene, nm23, in human brain tumors. <i>Neuropathology</i> , 1998, 18, 315-320.	0.7	3
51	A case of Miller-Dieker syndrome in a family with neurofibromatosis type I. <i>Acta Neuropathologica</i> , 2000, 99, 425-427.	3.9	3
52	Lack of Correlation Between Immunohistochemical Expression of SPARC and Invasion in Different Grades of Meningiomas. <i>Anticancer Research</i> , 2020, 40, 3081-3089.	0.5	3
53	A pathologically confirmed case of combined amyotrophic lateral sclerosis with <i>C9orf72</i> mutation and multiple system atrophy. <i>Neuropathology</i> , 0, , .	0.7	3
54	GPI deep brain stimulation for palliation of hemidystonia and hemibody jerking in a patient with suspected adult onset neuronal ceroid lipofuscinosis. <i>Journal of the Neurological Sciences</i> , 2016, 362, 228-229.	0.3	2

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55	Reply: Atherosclerosis and vascular cognitive impairment neuropathological guideline. Brain, 2017, 140, e13-e13.	3.7	2
56	ATRX immunohistochemistry can help refine "not elsewhere classified"™ categorisation for grade II/III gliomas. British Journal of Neurosurgery, 2019, 33, 536-540.	0.4	1
57	Leptomeningeal melanoma and Creutzfeldt-Jakob disease in a patient with chronic lymphocytic leukaemia. Neuropathology and Applied Neurobiology, 1999, 25, 345-348.	1.8	0
58	P43...Parasellar capillary haemangioma with intrasellar extension. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, e35.4-e36.	0.9	0
59	An unusual case of primary leptomeningeal marginal zone B-cell lymphoma. , 1998, 17, 326-9.		0
60	A novel case of paravertebral glomangiomyomatosis. British Journal of Neurosurgery, 2022, , 1-6.	0.4	0