## Danielle D Seilhean

List of Publications by Year in descending order

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110 papers 9,011 citations

41344 49 h-index 91 g-index

113 all docs

113 docs citations

113 times ranked

13723 citing authors

#	Article	IF	CITATIONS
1	Neuroinvasion of SARS-CoV-2 in human and mouse brain. Journal of Experimental Medicine, 2021, 218, .	8.5	677
2	Common variants at 7p21 are associated with frontotemporal lobar degeneration with TDP-43 inclusions. Nature Genetics, 2010, 42, 234-239.	21.4	479
3	Aging-related tau astrogliopathy (ARTAG): harmonized evaluation strategy. Acta Neuropathologica, 2016, 131, 87-102.	7.7	380
4	Staging of Neurofibrillary Pathology in Alzheimer's Disease: A Study of the BrainNet Europe Consortium. Brain Pathology, 2008, 18, 484-496.	4.1	361
5	Remyelination in multiple sclerosis. Progress in Brain Research, 2009, 175, 453-464.	1.4	311
6	Activation of the subventricular zone in multiple sclerosis: Evidence for early glial progenitors. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 4694-4699.	7.1	299
7	Inhibition of the mTORC Pathway in the Antiphospholipid Syndrome. New England Journal of Medicine, 2014, 371, 303-312.	27.0	282
8	Perforin-Dependent Brain-Infiltrating Cytotoxic CD8+ T Lymphocytes Mediate Experimental Cerebral Malaria Pathogenesis. Journal of Immunology, 2003, 170, 2221-2228.	0.8	267
9	SOD1, ANG, VAPB, TARDBP, and FUS mutations in familial amyotrophic lateral sclerosis: genotype-phenotype correlations. Journal of Medical Genetics, 2010, 47, 554-560.	3.2	266
10	Re-expression of PSA-NCAM by demyelinated axons: an inhibitor of remyelination in multiple sclerosis?. Brain, 2002, 125, 1972-1979.	7.6	246
11	Mutations in SQSTM1 encoding p62 in amyotrophic lateral sclerosis: genetics and neuropathology. Acta Neuropathologica, 2013, 125, 511-522.	7.7	201
12	Critical role of IL-21 in modulating TH17 and regulatory TÂcells in Behçet disease. Journal of Allergy and Clinical Immunology, 2011, 128, 655-664.	2.9	196
13	Consensus classification of human prion disease histotypes allows reliable identification of molecular subtypes: an inter-rater study among surveillance centres in Europe and USA. Acta Neuropathologica, 2012, 124, 517-529.	7.7	184
14	A mutation of spastin is responsible for swellings and impairment of transport in a region of axon characterized by changes in microtubule composition. Human Molecular Genetics, 2006, 15, 3544-3558.	2.9	169
15	Phenotype difference between ALS patients with expanded repeats in <i>C9ORF72</i> and patients with mutations in other ALS-related genes. Journal of Medical Genetics, 2012, 49, 258-263.	3.2	157
16	Doxycycline in Creutzfeldt-Jakob disease: a phase 2, randomised, double-blind, placebo-controlled trial. Lancet Neurology, The, 2014, 13, 150-158.	10.2	157
17	In vivo transcranial brain surgery with an ultrasonic time reversal mirror. Journal of Neurosurgery, 2007, 106, 1061-1066.	1.6	155
18	Mixed Brain Pathologies in Dementia: The BrainNet Europe Consortium Experience. Dementia and Geriatric Cognitive Disorders, 2008, 26, 343-350.	1.5	148

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19	Assessment of $\hat{l}^2$ -amyloid deposits in human brain: a study of the BrainNet Europe Consortium. Acta Neuropathologica, 2009, 117, 309-320.	7.7	143
20	Causes of death in a postâ€mortem series of ALS patients. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2008, 9, 59-62.	2.1	140
21	Effects of Antemortem and Postmortem Variables on Human Brain mRNA Quality: A BrainNet Europe Study. Journal of Neuropathology and Experimental Neurology, 2010, 69, 70-81.	1.7	139
22	Oxidation of SQSTM1/p62 mediates the link between redox state and protein homeostasis. Nature Communications, 2018, 9, 256.	12.8	132
23	Retrospective Observational Study of Brain MRI Findings in Patients with Acute SARS-CoV-2 Infection and Neurologic Manifestations. Radiology, 2020, 297, E313-E323.	7.3	131
24	Human endogenous retrovirus (HERV)-W ENV and GAG proteins: Physiological expression in human brain and pathophysiological modulation in multiple sclerosis lesions. Journal of NeuroVirology, 2005, 11, 23-33.	2.1	128
25	Common mechanisms in neurodegeneration and neuroinflammation: a BrainNet Europe gene expression microarray study. Journal of Neural Transmission, 2015, 122, 1055-1068.	2.8	126
26	TMEM106B is a genetic modifier of frontotemporal lobar degeneration with C9orf72 hexanucleotide repeat expansions. Acta Neuropathologica, 2014, 127, 407-418.	7.7	123
27	Gain of Olig2 function in oligodendrocyte progenitors promotes remyelination. Brain, 2015, 138, 120-135.	7.6	119
28	Nogo expression in muscle correlates with amyotrophic lateral sclerosis severity. Annals of Neurology, 2005, 57, 553-556.	5.3	113
29	ApoE immunoreactivity and microglial cells in Alzheimer's disease brain. Neuroscience Letters, 1995, 195, 5-8.	2.1	112
30	Selection of novel reference genes for use in the human central nervous system: a BrainNet Europe Study. Acta Neuropathologica, 2012, 124, 893-903.	7.7	110
31	A stable proportion of Lewy body bearing neurons in the substantia nigra suggests a model in which the Lewy body causes neuronal death. Neurobiology of Aging, 2010, 31, 99-103.	3.1	107
32	Management of a twenty-first century brain bank: experience in the BrainNet Europe consortium. Acta Neuropathologica, 2008, 115, 497-507.	7.7	101
33	Interlaboratory Comparison of Assessments of Alzheimer Disease-Related Lesions: A Study of the BrainNet Europe Consortium. Journal of Neuropathology and Experimental Neurology, 2006, 65, 740-757.	1.7	95
34	C9ORF72 Repeat Expansions in the Frontotemporal Dementias Spectrum of Diseases: A Flow-chart for Genetic Testing. Journal of Alzheimer's Disease, 2013, 34, 485-499.	2.6	93
35	System xC $\hat{a}$ ' is a mediator of microglial function and its deletion slows symptoms in amyotrophic lateral sclerosis mice. Brain, 2015, 138, 53-68.	7.6	85
36	Alteration of Blood–Brain Barrier Integrity by Retroviral Infection. PLoS Pathogens, 2008, 4, e1000205.	4.7	84

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37	Human Blood-Brain Barrier Disruption by Retroviral-Infected Lymphocytes: Role of Myosin Light Chain Kinase in Endothelial Tight-Junction Disorganization. Journal of Immunology, 2007, 179, 2576-2583.	0.8	82
38	Tumor necrosis factor-α, microglia and astrocytes in AIDS dementia complex. Acta Neuropathologica, 1997, 93, 508-517.	7.7	78
39	Association of Small-Vessel Disease With Dilatative Arteriopathy of the Brain. Stroke, 2007, 38, 1197-1202.	2.0	78
40	Assessment of $\hat{l}_{\pm}$ -Synuclein Pathology: A Study of the BrainNet Europe Consortium. Journal of Neuropathology and Experimental Neurology, 2008, 67, 125-143.	1.7	73
41	Accumulation of TDP-43 and $\hat{l}\pm$ -actin in an amyotrophic lateral sclerosis patient with the K17I ANG mutation. Acta Neuropathologica, 2009, 118, 561-573.	7.7	71
42	Neuropathology of iatrogenic Creutzfeldt–Jakob disease and immunoassay of French cadaver-sourced growth hormone batches suggest possible transmission of tauopathy and long incubation periods for the transmission of Abeta pathology. Acta Neuropathologica, 2018, 135, 201-212.	7.7	71
43	Contribution of <i>ATXN2</i> intermediary polyQ expansions in a spectrum of neurodegenerative disorders. Neurology, 2014, 83, 990-995.	1.1	70
44	Modifying macrophages at the periphery has the capacity to change microglial reactivity and to extend ALS survival. Nature Neuroscience, 2020, 23, 1339-1351.	14.8	69
45	Screening of OPTN in French familial amyotrophic lateral sclerosis. Neurobiology of Aging, 2011, 32, 557.e11-557.e13.	3.1	68
46	Dissociation of Alzheimer type pathology in a disconnected piece of cortex. Acta Neuropathologica, 1997, 93, 501-507.	7.7	60
47	The need to unify neuropathological assessments of vascular alterations in the ageing brain. Experimental Gerontology, 2012, 47, 825-833.	2.8	57
48	The autophagy/lysosome pathway is impaired in SCA7 patients and SCA7 knock-in mice. Acta Neuropathologica, 2014, 128, 705-722.	7.7	56
49	Bilateral Adrenal Infiltration in Erdheim-Chester Disease. Report of Seven Cases and Literature Review. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 2007-2012.	3.6	53
50	Novel UBQLN2 mutations linked to amyotrophic lateral sclerosis and atypical hereditary spastic paraplegia phenotype through defective HSP70-mediated proteolysis. Neurobiology of Aging, 2017, 58, 239.e11-239.e20.	3.1	50
51	Amyotrophic lateral sclerosis with neuronal intranuclear protein inclusions. Acta Neuropathologica, 2004, 108, 81-87.	7.7	49
52	Loss of paraplegin drives spasticity rather than ataxia in a cohort of 241 patients with <i>SPG7</i> Neurology, 2019, 92, e2679-e2690.	1.1	49
53	BrainNet Europe's Code of Conduct for brain banking. Journal of Neural Transmission, 2015, 122, 937-940.	2.8	46
54	TDP-43 Pathology Progression Along the Olfactory Pathway as a Possible Substrate for Olfactory Impairment in Amyotrophic Lateral Sclerosis. Journal of Neuropathology and Experimental Neurology, 2015, 74, 547-556.	1.7	41

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55	Paradoxical intracranial cryptococcoma in a human immunodeficiency virus–infected man being treated with combination antiretroviral therapy. American Journal of Medicine, 2002, 113, 155-157.	1.5	40
56	Neuropathology of Sleep Disorders: A Review. Journal of Neuropathology and Experimental Neurology, 2011, 70, 243-252.	1.7	39
57	Multisite Assessment of Aging-Related Tau Astrogliopathy (ARTAG). Journal of Neuropathology and Experimental Neurology, 2017, 76, 605-619.	1.7	38
58	Association of Clinical, Biological, and Brain Magnetic Resonance Imaging Findings With Electroencephalographic Findings for Patients With COVID-19. JAMA Network Open, 2021, 4, e211489.	5.9	38
59	Mutations in UBQLN2 are rare in French amyotrophic lateral sclerosis. Neurobiology of Aging, 2012, 33, 839.e1-839.e3.	3.1	34
60	Human immunodeficiency virus type 1 DNA and RNA load in brains of demented and nondemented patients with acquired immunodeficiency syndrome. Journal of NeuroVirology, 1997, 3, 299-303.	2.1	31
61	Long-standing Prion Dementia Manifesting as Posterior Cortical Atrophy. Alzheimer Disease and Associated Disorders, 2012, 26, 289-292.	1.3	29
62	Human T-Lymphotropic Virus Type 1-Induced Overexpression of Activated Leukocyte Cell Adhesion Molecule (ALCAM) Facilitates Trafficking of Infected Lymphocytes through the Blood-Brain Barrier. Journal of Virology, 2016, 90, 7303-7312.	3.4	29
63	Non-invasive ultrasonic surgery of the brain in non-human primates. Journal of the Acoustical Society of America, 2013, 134, 1632-1639.	1.1	26
64	Abnormal glycogen in astrocytes is sufficient to cause adult polyglucosan body disease. Gene, 2013, 515, 376-379.	2.2	26
65	Neuropathological assessments of the pathology in frontotemporal lobar degeneration with TDP43-positive inclusions: an inter-laboratory study by the BrainNet Europe consortium. Journal of Neural Transmission, 2015, 122, 957-972.	2.8	25
66	Inconstant apolipoprotein E (ApoE)-like immunoreactivity in amyloid $\hat{l}^2$ protein deposits: relationship with APOE genotype in aging brain and Alzheimer's disease. Acta Neuropathologica, 1996, 92, 180-185.	7.7	24
67	Laminar spongiosis of the dentate gyrus: a sign of disconnection, present in cases of severe Alzheimer's disease. Acta Neuropathologica, 1998, 95, 413-420.	7.7	24
68	Coronary and Basilar Artery Ectasia Are Associated. Stroke, 2016, 47, 224-227.	2.0	24
69	PrP immunohistochemistry: Different protocols, including a procedure for long formalin fixation, and a proposed schematic classification for deposits in sporadic Creutzfeldt-Jakob disease. Microscopy Research and Technique, 2000, 50, 26-31.	2.2	23
70	Severe Demyelinating Myelopathy with Low Human T Cell Lymphotropic Virus Type 1 Expression after Transfusion in an Immunosuppressed Patient. Clinical Infectious Diseases, 2002, 34, 855-860.	5.8	23
71	Abnormal TDP-43 and FUS proteins in muscles of sporadic IBM: similarities in a TARDBP-linked ALS patient. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 1414-1416.	1.9	22
72	Interrupted CAG expansions in ATXN2 gene expand the genetic spectrum of frontotemporal dementias. Acta Neuropathologica Communications, 2018, 6, 41.	5.2	21

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73	Fatal HHV-6 associated encephalitis in an HIV-1 infected patient treated with cidofovir. Journal of Infection, 2006, 52, 237-242.	3.3	20
74	Genetic screening of ANXA11 revealed novel mutations linked to amyotrophic lateral sclerosis. Neurobiology of Aging, 2021, 99, 102.e11-102.e20.	3.1	20
75	Accuracy of diagnosis criteria in patients with suspected diagnosis of sporadic Creutzfeldt-Jakob disease and detection of 14-3-3 protein, France, 1992 to 2009. Eurosurveillance, 2017, 22, .	7.0	20
76	Miliary Brain Metastases in Lung Cancer. Journal of Clinical Oncology, 2010, 28, e714-e716.	1.6	19
77	Increased prevalence of granulovacuolar degeneration in C9orf72 mutation. Acta Neuropathologica, 2019, 138, 783-793.	7.7	19
78	Transmission of amyloid-beta and tau pathologies is associated with cognitive impairments in a primate. Acta Neuropathologica Communications, 2021, 9, 165.	5.2	18
79	JC Virus Variant Associated with Cerebellar Atrophy in a Patient with AIDS. Journal of Clinical Microbiology, 2011, 49, 2196-2199.	3.9	17
80	Axonal expression of sodium channels and neuropathology of the plaques in multiple sclerosis. Neuropathology and Applied Neurobiology, 2014, 40, 579-590.	3.2	16
81	The wide spectrum of COVID-19 neuropsychiatric complications within a multidisciplinary centre. Brain Communications, 2021, 3, fcab135.	3.3	16
82	Pathway from TDP-43-Related Pathology to Neuronal Dysfunction in Amyotrophic Lateral Sclerosis and Frontotemporal Lobar Degeneration. International Journal of Molecular Sciences, 2021, 22, 3843.	4.1	16
83	Amygdala TDP-43 Pathology in Frontotemporal Lobar Degeneration and Motor Neuron Disease. Journal of Neuropathology and Experimental Neurology, 2017, 76, 800-812.	1.7	14
84	Neuropathogenesis of acute coronavirus disease 2019. Current Opinion in Neurology, 2021, 34, 417-422.	3.6	14
85	Exclusive induction of tau2 epitope in microglia/macrophages in inflammatory lesions?tautwopathy distinct from degenerative tauopathies. Acta Neuropathologica, 2005, 109, 159-164.	7.7	12
86	Genetic analysis of CHCHD10 in French familial amyotrophic lateral sclerosis patients. Neurobiology of Aging, 2016, 42, 218.e1-218.e3.	3.1	12
87	Detection and partial discrimination of atypical and classical bovine spongiform encephalopathies in cattle and primates using real-time quaking-induced conversion assay. PLoS ONE, 2017, 12, e0172428.	2.5	12
88	Progressive multifocal leukoencephalopathy mimicking cerebral vasculitis in systemic granulomatosis. Journal of Infection, 2007, 54, e133-e135.	3.3	10
89	Aspergillus flavus Brain Abscesses Associated with Hepatic Amebiasis in a Non-neutropenic Man in Senegal. American Journal of Tropical Medicine and Hygiene, 2009, 81, 583-586.	1.4	10
90	Fronto-temporal lobar degeneration: neuropathology in 60 cases. Journal of Neural Transmission, 2011, 118, 753-764.	2.8	10

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91	Human subiculo-fornico-mamillary system in Alzheimer's disease: Tau seeding by the pillar of the fornix. Acta Neuropathologica, 2020, 139, 443-461.	7.7	10
92	The Amyotrophic Lateral Sclerosis M114T PFN1 Mutation Deregulates Alternative Autophagy Pathways and Mitochondrial Homeostasis. International Journal of Molecular Sciences, 2022, 23, 5694.	4.1	10
93	The nosology and neuropathology of human conditions related to unconventional infectious agents or prions. European Journal of Neurology, 1996, 3, 487-499.	3.3	9
94	Ensemencement et propagation des lésions dans les maladies neurodégénératives: un nouveau paradigme. Bulletin De L'Academie Nationale De Medecine, 2015, 199, 809-819.	0.0	8
95	Impact of a frequent nearsplice <i>SOD1</i> variant in amyotrophic lateral sclerosis: optimising <i>SOD1</i> genetic screening for gene therapy opportunities. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 942-949.	1.9	7
96	Region-specific protein misfolding cyclic amplification reproduces brain tropism of prion strains. Journal of Biological Chemistry, 2017, 292, 16688-16696.	3.4	6
97	Frontotemporal lobar degeneration: Diversity of FTLD lesions. Revue Neurologique, 2013, 169, 786-792.	1.5	5
98	Progressive Supranuclear Palsy Syndrome and Semantic Dementia in Neuropathologically Proven Lewy Body Disease: A Report of Two Cases. Journal of Alzheimer's Disease, 2015, 47, 95-101.	2.6	5
99	Autopsie et religions. Bulletin De L'Academie Nationale De Medecine, 2001, 185, 877-889.	0.0	5
100	Neuropathology of Central Nervous System Arterial Syndromes. Part I. Journal of Neuropathology and Experimental Neurology, 2009, 68, 113-124.	1.7	4
101	The Dejerine Foundation. , 2018, 37, 193-196.		4
102	First European case of Creutzfeldt-Jakob disease with a PRNP G114V mutation. Cortex, 2019, 117, 407-413.	2.4	3
103	Neuropathology in Pitiéâ€Salpêtrière hospital: Past, present and prospect. Neuropathology, 2020, 40, 3-13.	1.2	3
104	High densities of tumor necrosis factor-? in the cerebral cortex and basal ganglia in human immunodeficiency virus-1-associated cognitive/motor complex: A quantitative regional analysis study. Neuropathology, 1997, 17, 168-173.	1.2	1
105	The Human Tissue Bill. Lancet Neurology, The, 2004, 3, 685-690.	10.2	1
106	Association between a Primitive Brain Tumor and Cerebral Aspergillosis. Case Reports in Medicine, 2012, 2012, 1-5.	0.7	1
107	Activated leukocyte cell adhesion molecule (ALCAM) facilitates trafficking of HTLV-1 infected lymphocytes through the blood brain barrier. Retrovirology, 2015, 12, .	2.0	1
108	Diseases. , 2013, , 63-74.		0

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109	Genetic Creutzfeldt–Jakob disease with an 8-year disease course. Acta Neurologica Belgica, 2018, 118, 313-314.	1.1	0
110	Neuronal loss in HIV. Neurology, 1994, 44, 1365.	1.1	0