Suzee E Lee

List of Publications by Year
in descending order

39 papers	3,728 citations	279798 23 h-index	330143 37 g-index
40 all docs	40 docs citations	40 times ranked	5189 citing authors

#	Article	IF	Citations
1	Criteria for the diagnosis of corticobasal degeneration. Neurology, 2013, 80, 496-503.	1.1	1,445
2	Clinicopathological correlations in behavioural variant frontotemporal dementia. Brain, 2017, 140, 3329-3345.	7.6	226
3	Evidence for a role of the rare p.A152T variant in MAPT in increasing the risk for FTD-spectrum and Alzheimer's diseases. Human Molecular Genetics, 2012, 21, 3500-3512.	2.9	198
4	Frontotemporal dementia due to C9ORF72 mutations. Neurology, 2012, 79, 1002-1011.	1.1	183
5	Atypical, slowly progressive behavioural variant frontotemporal dementia associated withC9ORF72hexanucleotide expansion. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 358-364.	1.9	172
6	Altered network connectivity in frontotemporal dementia with C9orf72 hexanucleotide repeat expansion. Brain, 2014, 137, 3047-3060.	7.6	140
7	Suberoylanilide Hydroxamic Acid (Vorinostat) Up-regulates Progranulin Transcription. Journal of Biological Chemistry, 2011, 286, 16101-16108.	3.4	138
8	Network degeneration and dysfunction in presymptomatic C9ORF72 expansion carriers. NeuroImage: Clinical, 2017, 14, 286-297.	2.7	129
9	18F-flortaucipir (AV-1451) tau PET in frontotemporal dementia syndromes. Alzheimer's Research and Therapy, 2019, 11, 13.	6.2	121
10	Distinct Subtypes of Behavioral Variant Frontotemporal Dementia Based on Patterns of Network Degeneration. JAMA Neurology, 2016, 73, 1078.	9.0	115
11	Microglial NFÎºB-TNFÎ± hyperactivation induces obsessiveâ€“compulsive behavior in mouse models of progranulin-deficient frontotemporal dementia. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 5029-5034.	7.1	96
12	Patient-Tailored, Connectivity-Based Forecasts of Spreading Brain Atrophy. Neuron, 2019, 104, 856-868.e5.	8.1	85
13	A152T tau allele causes neurodegeneration that can be ameliorated in a zebrafish model by autophagy induction. Brain, 2017, 140, 1128-1146.	7.6	84
14	Cognition and neuropsychiatry in behavioral variant frontotemporal dementia by disease stage. Neurology, 2016, 86, 600-610.	1.1	73
15	A Comprehensive Resource for Induced Pluripotent Stem Cells from Patients with Primary Tauopathies. Stem Cell Reports, 2019, 13, 939-955.	4.8	62
16	Poly(GP), neurofilament and grey matter deficits in C9orf72 expansion carriers. Annals of Clinical and Translational Neurology, 2018, 5, 583-597.	3.7	48
17	Altered topology of the functional speech production network in non-fluent/agrammatic variant of PPA. Cortex, 2018, 108, 252-264.	2.4	41
18	Neurodegenerative Disease Phenotypes in Carriers of MAPT p.A152T, A Risk Factor for Frontotemporal Dementia Spectrum Disorders and Alzheimer Disease. Alzheimer Disease and Associated Disorders, 2013, 27, 302-309.	1.3	40

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19	MAPT H1 Haplotype is Associated with Late-Onset Alzheimerâ€™s Disease Risk in APOE É›4 Noncarriers: Results from the Dementia Genetics Spanish Consortium. Journal of Alzheimer's Disease, 2015, 49, 343-352.	2.6	32
20	Predicting amyloid status in corticobasal syndrome using modified clinical criteria, magnetic resonance imaging and fluorodeoxyglucose positron emission tomography. Alzheimer's Research and Therapy, 2015, 7, 8.	6.2	32
21	Thalamo-cortical network hyperconnectivity in preclinical progranulin mutation carriers. NeuroImage: Clinical, 2019, 22, 101751.	2.7	30
22	Rare variants in the neuronal ceroid lipofuscinosis gene MFSD8 are candidate risk factors for frontotemporal dementia. Acta Neuropathologica, 2019, 137, 71-88.	7.7	29
23	Advancing functional dysconnectivity and atrophy in progressive supranuclear palsy. NeuroImage: Clinical, 2017, 16, 564-574.	2.7	26
24	Gyrification abnormalities in presymptomatic c9orf72 expansion carriers. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 1005-1010.	1.9	24
25	Neuroimaging in genetic frontotemporal dementia and amyotrophic lateral sclerosis. Neurobiology of Disease, 2020, 145, 105063.	4.4	23
26	Brain volumetric deficits in MAPT mutation carriers: a multisite study. Annals of Clinical and Translational Neurology, 2021, 8, 95-110.	3.7	21
27	Rates of Brain Atrophy Across Disease Stages in Familial Frontotemporal Dementia Associated With MAPT, GRN, and C9orf72 Pathogenic Variants. JAMA Network Open, 2020, 3, e2022847.	5.9	19
28	Guam dementia syndrome revisited in 2011. Current Opinion in Neurology, 2011, 24, 517-524.	3.6	16
29	Frontotemporal dementia spectrum: first genetic screen in a Greek cohort. Neurobiology of Aging, 2019, 75, 224.e1-224.e8.	3.1	16
30	Amyloid in dementia associated with familial FTLD: not an innocent bystander. Neurocase, 2016, 22, 76-83.	0.6	12
31	A novel temporalâ€predominantÂneuroâ€astroglial tauopathyÂassociated with TMEM106B gene polymorphism in FTLD/ALSâ€TDP. Brain Pathology, 2021, 31, 267-282.	4.1	12
32	Lack of Association Between the CCR5-delta32 Polymorphism and Neurodegenerative Disorders. Alzheimer Disease and Associated Disorders, 2020, 34, 244-247.	1.3	11
33	Youngâ€onset frontotemporal dementia in a homozygous tau R406W mutation carrier. Annals of Clinical and Translational Neurology, 2015, 2, 1124-1128.	3.7	7
34	Frequency of frontotemporal dementia gene variants in C9ORF72, MAPT, and GRN in academic versus commercial laboratory cohorts. Advances in Genomics and Genetics, 2018, Volume 8, 23-33.	0.8	7
35	Frequency of the TREM2 R47H Variant in Various Neurodegenerative Disorders. Alzheimer Disease and Associated Disorders, 2019, 33, 327-330.	1.3	6
36	The unexpected co-occurrence of GRN and MAPT p.A152T in Basque families: Clinical and pathological characteristics. PLoS ONE, 2017, 12, e0178093.	2.5	5

#	Article	IF	Citations
37	Early-onset Alzheimerâ€™s disease versus frontotemporal dementia: resolution with genetic diagnoses?. Neurocase, 2016, 22, 161-167.	0.6	3
38	Patient-Tailored, Connectivity-Based Forecasts of Spreading Brain Atrophy. SSRN Electronic Journal, 0, , .	0.4	1
39	P1â€433: GRAY MATTER DEFICITS IN SYMPTOMATIC AND PRESYMPTOMATIC MAPT MUTATION CARRIERS. Alzheimer's and Dementia, 2018, 14, P475.	0.8	0