# 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 | Brain volumetric deficits in MAPT mutation carriers: a multisite study. Annals of Clinical and Translational Neurology, 2021, 8, 95-110. | 3.7 | 21 |
| 2 | A novel temporalâ€predominantÂneuroâ€astroglial tauopathyÂassociated with TMEM106B gene polymorphism in FTLD/ALSâ€TDP. Brain Pathology, 2021, 31, 267-282. | 4.1 | 12 |
| 3 | Neuroimaging in genetic frontotemporal dementia and amyotrophic lateral sclerosis. Neurobiology of Disease, 2020, 145, 105063. | 4.4 | 23 |
| 4 | Lack of Association Between the CCR5-delta32 Polymorphism and Neurodegenerative Disorders. Alzheimer Disease and Associated Disorders, 2020, 34, 244-247. | 1.3 | 11 |
| 5 | 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 |
| 6 | Patient-Tailored, Connectivity-Based Forecasts of Spreading Brain Atrophy. Neuron, 2019, 104, 856-868.e5. | 8.1 | 85 |
| 7 | A Comprehensive Resource for Induced Pluripotent Stem Cells from Patients with Primary Tauopathies. Stem Cell Reports, 2019, 13, 939-955. | 4.8 | 62 |
| 8 | 18F-flortaucipir (AV-1451) tau PET in frontotemporal dementia syndromes. Alzheimer's Research and Therapy, 2019, 11, 13. | 6.2 | 121 |
| 9 | Gyrification abnormalities in presymptomatic c9orf72 expansion carriers. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 1005-1010. | 1.9 | 24 |
| 10 | Thalamo-cortical network hyperconnectivity in preclinical progranulin mutation carriers. NeuroImage: Clinical, 2019, 22, 101751. | 2.7 | 30 |
| 11 | Frequency of the TREM2 R47H Variant in Various Neurodegenerative Disorders. Alzheimer Disease and Associated Disorders, 2019, 33, 327-330. | 1.3 | 6 |
| 12 | Frontotemporal dementia spectrum: first genetic screen in a Greek cohort. Neurobiology of Aging, 2019, 75, 224.e1-224.e8. | 3.1 | 16 |
| 13 | 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 |
| 14 | Poly(GP), neurofilament and grey matter deficits in C9orf72 expansion carriers. Annals of Clinical and Translational Neurology, 2018, 5, 583-597. | 3.7 | 48 |
| 15 | P1â€433: GRAY MATTER DEFICITS IN SYMPTOMATIC AND PRESYMPTOMATIC MAPT MUTATION CARRIERS. Alzheimer's and Dementia, 2018, 14, P475. | 0.8 | 0 |
| 16 | Frequency of frontotemporal dementia gene variants in <em>C9ORF72</em>, <em>MAPT</em>, and <em>GRN</em> in academic versus commercial laboratory cohorts. Advances in Genomics and Genetics, 2018, Volume 8, 23-33. | 0.8 | 7 |
| 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 | 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 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Network degeneration and dysfunction in presymptomatic C9ORF72 expansion carriers. NeuroImage: Clinical, 2017, 14, 286-297. | 2.7 | 129 |
| 20 | Clinicopathological correlations in behavioural variant frontotemporal dementia. Brain, 2017, 140, 3329-3345. | 7.6 | 226 |
| 21 | Advancing functional dysconnectivity and atrophy in progressive supranuclear palsy. NeuroImage: Clinical, 2017, 16, 564-574. | 2.7 | 26 |
| 22 | A152T tau allele causes neurodegeneration that can be ameliorated in a zebrafish model by autophagy induction. Brain, 2017, 140, 1128-1146. | 7.6 | 84 |
| 23 | 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 |
| 24 | Distinct Subtypes of Behavioral Variant Frontotemporal Dementia Based on Patterns of Network Degeneration. JAMA Neurology, 2016, 73, 1078. | 9.0 | 115 |
| 25 | Cognition and neuropsychiatry in behavioral variant frontotemporal dementia by disease stage. Neurology, 2016, 86, 600-610. | 1.1 | 73 |
| 26 | Early-onset Alzheimerâ€™s disease versus frontotemporal dementia: resolution with genetic diagnoses?. Neurocase, 2016, 22, 161-167. | 0.6 | 3 |
| 27 | Amyloid in dementia associated with familial FTLD: not an innocent bystander. Neurocase, 2016, 22, 76-83. | 0.6 | 12 |
| 28 | Youngâ€onset frontotemporal dementia in a homozygous tau R406W mutation carrier. Annals of Clinical and Translational Neurology, 2015, 2, 1124-1128. | 3.7 | 7 |
| 29 | 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 |
| 30 | 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 |
| 31 | Altered network connectivity in frontotemporal dementia with C9orf72 hexanucleotide repeat expansion. Brain, 2014, 137, 3047-3060. | 7.6 | 140 |
| 32 | Criteria for the diagnosis of corticobasal degeneration. Neurology, 2013, 80, 496-503. | 1.1 | 1,445 |
| 33 | 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 |
| 34 | Atypical, slowly progressive behavioural variant frontotemporal dementia associated withC9ORF72hexanucleotide expansion. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 358-364. | 1.9 | 172 |
| 35 | 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 |
| 36 | Frontotemporal dementia due to C9ORF72 mutations. Neurology, 2012, 79, 1002-1011. | 1.1 | 183 |

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|---|---|---|---|
| 37 | Guam dementia syndrome revisited in 2011. Current Opinion in Neurology, 2011, 24, 517-524. | 3.6 | 16 |
| 38 | Suberoylanilide Hydroxamic Acid (Vorinostat) Up-regulates Progranulin Transcription. Journal of Biological Chemistry, 2011, 286, 16101-16108. | 3.4 | 138 |
| 39 | Patient-Tailored, Connectivity-Based Forecasts of Spreading Brain Atrophy. SSRN Electronic Journal, 0, , . | 0.4 | 1 |