Keith A Josephs

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

148 23,793 323 72 h-index g-index citations papers 6.58 28,236 6.4 338 L-index avg, IF ext. citations ext. papers

| # | Paper | IF | Citations |
|-----|--|------|-----------|
| 323 | Diffuse Lewy body disease presenting as Parkinsonß disease with progressive aphasia Neuropathology, 2022, | 2 | O |
| 322 | TDP-43-associated atrophy in brains with and without frontotemporal lobar degeneration <i>NeuroImage: Clinical</i> , 2022 , 34, 102954 | 5.3 | 0 |
| 321 | White matter damage due to vascular, tau, and TDP-43 pathologies and its relevance to cognition <i>Acta Neuropathologica Communications</i> , 2022 , 10, 16 | 7.3 | 1 |
| 320 | Tractography of supplementary motor area projections in progressive speech apraxia and aphasia <i>NeuroImage: Clinical</i> , 2022 , 34, 102999 | 5.3 | 1 |
| 319 | Does limited EMG denervation in early primary lateral sclerosis predict amyotrophic lateral sclerosis?. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2022 , 1-8 | 3.6 | 1 |
| 318 | Depression and Apathy across Different Variants of Progressive Supranuclear Palsy <i>Movement Disorders Clinical Practice</i> , 2022 , 9, 212-217 | 2.2 | 0 |
| 317 | Frontotemporal lobar degeneration with TAR DNA-binding protein 43 (TDP-43): its journey of more than 100 years <i>Journal of Neurology</i> , 2022 , 1 | 5.5 | O |
| 316 | Cross-Sectional and Longitudinal Assessment of Behavior in Primary Progressive Apraxia of Speech and Agrammatic Aphasia <i>Dementia and Geriatric Cognitive Disorders</i> , 2022 , 1-10 | 2.6 | |
| 315 | Frequency and distribution of TAR DNA-binding protein 43 (TDP-43) pathology increase linearly with age in a large cohort of older adults with and without dementia <i>Acta Neuropathologica</i> , 2022 , 1 | 14.3 | O |
| 314 | Diffusion tractography of Superior Cerebellar Peduncle and Dentatorubrothalamic Tracts in two Autopsy Confirmed Progressive Supranuclear Palsy Variants: Richardson syndrome and the speech-language variant. <i>NeuroImage: Clinical</i> , 2022 , 103030 | 5.3 | 0 |
| 313 | In Vivo Imaging and Autoradiography in a Case of Autopsy-Confirmed Pick Disease. <i>Neurology: Clinical Practice</i> , 2021 , 11, e11-e14 | 1.7 | 4 |
| 312 | Survival Analysis in Primary Progressive Apraxia of Speech and Agrammatic Aphasia. <i>Neurology: Clinical Practice</i> , 2021 , 11, 249-255 | 1.7 | 3 |
| 311 | Neuroimaging correlates of gait abnormalities in progressive supranuclear palsy. <i>NeuroImage: Clinical</i> , 2021 , 32, 102850 | 5.3 | 1 |
| 310 | Word Fluency Test Performance in Primary Progressive Aphasia and Primary Progressive Apraxia of Speech. <i>American Journal of Speech-Language Pathology</i> , 2021 , 30, 2635-2642 | 3.1 | 0 |
| 309 | Autopsy Validation of Progressive Supranuclear Palsy-Predominant Speech/Language Disorder Criteria. <i>Movement Disorders</i> , 2021 , | 7 | 2 |
| 308 | Diffusion tensor imaging analysis in three progressive supranuclear palsy variants. <i>Journal of Neurology</i> , 2021 , 268, 3409-3420 | 5.5 | 8 |
| 307 | TAR DNA-Binding Protein 43 Is Associated with Rate of Memory, Functional and Global Cognitive Decline in the Decade Prior to Death. <i>Journal of Alzheimerm Disease</i> , 2021 , 80, 683-693 | 4.3 | 2 |

(2021-2021)

| 306 | Long-read targeted sequencing uncovers clinicopathological associations for C9orf72-linked diseases. <i>Brain</i> , 2021 , 144, 1082-1088 | 11.2 | 2 | |
|-------------|---|------|----|--|
| 305 | A Cognitive Psychometric Investigation of Word Production and Phonological Error Rates in Logopenic Progressive Aphasia. <i>American Journal of Speech-Language Pathology</i> , 2021 , 30, 1194-1202 | 3.1 | | |
| 304 | Old age genetically confirmed frontotemporal lobar degeneration with TDP-43 has limbic predominant TDP-43 deposition. <i>Neuropathology and Applied Neurobiology</i> , 2021 , 47, 1050-1059 | 5.2 | 3 | |
| 303 | Progressive apraxia of speech: delays to diagnosis and rates of alternative diagnoses. <i>Journal of Neurology</i> , 2021 , 268, 4752-4758 | 5.5 | 2 | |
| 302 | A molecular pathology, neurobiology, biochemical, genetic and neuroimaging study of progressive apraxia of speech. <i>Nature Communications</i> , 2021 , 12, 3452 | 17.4 | 10 | |
| 301 | Neurodegeneration of the visual word form area in a patient with word form alexia. <i>Neurology and Clinical Neuroscience</i> , 2021 , 9, 359-360 | 0.3 | O | |
| 300 | Motor Speech Disorders and Communication Limitations in Progressive Supranuclear Palsy. <i>American Journal of Speech-Language Pathology</i> , 2021 , 30, 1361-1372 | 3.1 | 5 | |
| 299 | Neuropsychological Profiles of Patients with Progressive Apraxia of Speech and Aphasia. <i>Journal of the International Neuropsychological Society</i> , 2021 , 1-11 | 3.1 | | |
| 298 | Tau and Amyloid Relationships with Resting-state Functional Connectivity in Atypical Alzheimerß Disease. <i>Cerebral Cortex</i> , 2021 , 31, 1693-1706 | 5.1 | 13 | |
| 297 | Timeline of Rapid Eye Movement Sleep Behavior Disorder in Overt Alpha-Synucleinopathies. <i>Annals of Neurology</i> , 2021 , 89, 293-303 | 9.4 | 3 | |
| 296 | Primary Progressive Apraxia of Speech: From Recognition to Diagnosis and Care. <i>Aphasiology</i> , 2021 , 35, 560-591 | 1.6 | 15 | |
| 295 | Lewy Body Disease is a Contributor to Logopenic Progressive Aphasia Phenotype. <i>Annals of Neurology</i> , 2021 , 89, 520-533 | 9.4 | 6 | |
| 294 | Association of amyloid angiopathy with microbleeds in logopenic progressive aphasia: an imaging-pathology study. <i>European Journal of Neurology</i> , 2021 , 28, 670-675 | 6 | 2 | |
| 293 | Neurobehavioral Characteristics of FDG-PET Defined Right-Dominant Semantic Dementia: A Longitudinal Study. <i>Dementia and Geriatric Cognitive Disorders</i> , 2021 , 50, 17-28 | 2.6 | Ο | |
| 292 | Phonological Errors in Posterior Cortical Atrophy. <i>Dementia and Geriatric Cognitive Disorders</i> , 2021 , 50, 195-203 | 2.6 | 1 | |
| 291 | Progressive Supranuclear Palsy and Corticobasal Degeneration. <i>Advances in Experimental Medicine and Biology</i> , 2021 , 1281, 151-176 | 3.6 | 2 | |
| 29 0 | A Longitudinal Evaluation of Speech Rate in Primary Progressive Apraxia of Speech. <i>Journal of Speech, Language, and Hearing Research</i> , 2021 , 64, 392-404 | 2.8 | 5 | |
| 289 | Natural History of "Pure" Primary Lateral Sclerosis. <i>Neurology</i> , 2021 , 96, e2231-e2238 | 6.5 | 4 | |
| | | | | |

| 288 | Underlying pathology identified after 20 years of disease course in two cases of slowly progressive frontotemporal dementia syndromes. <i>Neurocase</i> , 2021 , 27, 212-222 | 0.8 | 1 |
|-----|---|---------------|----|
| 287 | Gray and White Matter Correlates of Dysphagia in Progressive Supranuclear Palsy. <i>Movement Disorders</i> , 2021 , 36, 2669-2675 | 7 | 2 |
| 286 | Posterior cortical atrophy phenotypic heterogeneity revealed by decoding F-FDG-PET. <i>Brain Communications</i> , 2021 , 3, fcab182 | 4.5 | 3 |
| 285 | Progressive Auditory Verbal Agnosia Secondary to Alzheimer Disease. <i>Neurology</i> , 2021 , 97, 908-909 | 6.5 | 3 |
| 284 | Selecting software pipelines for change in flortaucipir SUVR: Balancing repeatability and group separation. <i>NeuroImage</i> , 2021 , 238, 118259 | 7.9 | 4 |
| 283 | Assessing Change in Communication Limitations in Primary Progressive Apraxia of Speech and Aphasia: A 1-Year Follow-Up Study. <i>American Journal of Speech-Language Pathology</i> , 2021 , 30, 2368-237 | ' 8³.1 | |
| 282 | Sleep disturbances in the speech-language variant of progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2021 , 91, 9-12 | 3.6 | 0 |
| 281 | Relationship of APOE, age at onset, amyloid and clinical phenotype in Alzheimer disease. <i>Neurobiology of Aging</i> , 2021 , 108, 90-98 | 5.6 | 1 |
| 280 | Brainstem Biomarkers of Clinical Variant and Pathology in Progressive Supranuclear Palsy <i>Movement Disorders</i> , 2021 , | 7 | 2 |
| 279 | Protein contributions to brain atrophy acceleration in Alzheimer® disease and primary age-related tauopathy. <i>Brain</i> , 2020 , 143, 3463-3476 | 11.2 | 13 |
| 278 | Cerebrovascular pathology and misdiagnosis of multiple system atrophy: An autopsy study. <i>Parkinsonism and Related Disorders</i> , 2020 , 75, 34-40 | 3.6 | 5 |
| 277 | Ioflupane 123I (DAT scan) SPECT identifies dopamine receptor dysfunction early in the disease course in progressive apraxia of speech. <i>Journal of Neurology</i> , 2020 , 267, 2603-2611 | 5.5 | 6 |
| 276 | Progressive dysexecutive syndrome due to Alzheimer disease: a description of 55 cases and comparison to other phenotypes. <i>Brain Communications</i> , 2020 , 2, fcaa068 | 4.5 | 36 |
| 275 | Utility of FDG-PET in diagnosis of Alzheimer-related TDP-43 proteinopathy. <i>Neurology</i> , 2020 , 95, e23-e3 | 34 6.5 | 11 |
| 274 | Longitudinal neuroimaging biomarkers differ across Alzheimerß disease phenotypes. <i>Brain</i> , 2020 , 143, 2281-2294 | 11.2 | 23 |
| 273 | Clinical and pathologic features of cognitive-predominant corticobasal degeneration. <i>Neurology</i> , 2020 , 95, e35-e45 | 6.5 | 3 |
| 272 | PSP-like syndrome after aortic surgery in adults (Mokri syndrome). <i>Neurology: Clinical Practice</i> , 2020 , 10, 245-254 | 1.7 | 2 |
| 271 | Longitudinal flortaucipir ([F]AV-1451) PET uptake in semantic dementia. <i>Neurobiology of Aging</i> , 2020 , 92, 135-140 | 5.6 | 2 |

(2020-2020)

| Brain volume and flortaucipir analysis of progressive supranuclear palsy clinical variants. NeuroImage: Clinical, 2020 , 25, 102152 | 5.3 | 20 |
|--|---|--|
| Effect Modifiers of TDP-43-Associated Hippocampal Atrophy Rates in Patients with Alzheimerß Disease Neuropathological Changes. <i>Journal of Alzheimerm Disease</i> , 2020 , 73, 1511-1523 | 4.3 | 5 |
| TDP-43 is associated with a reduced likelihood of rendering a clinical diagnosis of dementia with Lewy bodies in autopsy-confirmed cases of transitional/diffuse Lewy body disease. <i>Journal of Neurology</i> , 2020 , 267, 1444-1453 | 5.5 | 1 |
| MRI and flortaucipir relationships in Alzheimerß phenotypes are heterogeneous. <i>Annals of Clinical and Translational Neurology</i> , 2020 , 7, 707-721 | 5.3 | 6 |
| Truncated stathmin-2 is a marker of TDP-43 pathology in frontotemporal dementia. <i>Journal of Clinical Investigation</i> , 2020 , 130, 6080-6092 | 15.9 | 34 |
| Longitudinal anatomic, functional, and molecular characterization of Pick disease phenotypes. <i>Neurology</i> , 2020 , 95, e3190-e3202 | 6.5 | 4 |
| Pick® disease: clinicopathologic characterization of 21 cases. <i>Journal of Neurology</i> , 2020 , 267, 2697-270 | 4 5.5 | 8 |
| Western Aphasia Battery-Revised Profiles in Primary Progressive Aphasia and Primary Progressive Apraxia of Speech. <i>American Journal of Speech-Language Pathology</i> , 2020 , 29, 498-510 | 3.1 | 11 |
| Communication Limitations in Patients With Progressive Apraxia of Speech and Aphasia. <i>American Journal of Speech-Language Pathology</i> , 2020 , 29, 1976-1986 | 3.1 | 9 |
| Dysphagia in Progressive Supranuclear Palsy. <i>Dysphagia</i> , 2020 , 35, 667-676 | 3.7 | 12 |
| Incidence of frontotemporal disorders in Olmsted County: A population-based study. <i>Alzheimern</i> and Dementia, 2020 , 16, 482-490 | 1.2 | 6 |
| Longitudinal flortaucipir ([F]AV-1451) PET imaging in primary progressive apraxia of speech. <i>Cortex</i> , 2020 , 124, 33-43 | 3.8 | 3 |
| The evolution of parkinsonism in primary progressive apraxia of speech: A 6-year longitudinal study. <i>Parkinsonism and Related Disorders</i> , 2020 , 81, 34-40 | 3.6 | 9 |
| Video-tutorial for the Movement Disorder Society criteria for progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2020 , 78, 200-203 | 3.6 | 4 |
| Predicting future rates of tau accumulation on PET. <i>Brain</i> , 2020 , 143, 3136-3150 | 11.2 | 25 |
| Dementia with Lewy bodies presenting as Logopenic variant primary progressive Aphasia. <i>Neurocase</i> , 2020 , 26, 259-263 | 0.8 | 4 |
| Longitudinal Amyloid-IPET in Atypical Alzheimerß Disease and Frontotemporal Lobar Degeneration. <i>Journal of Alzheimer</i> Disease, 2020 , 74, 377-389 | 4.3 | 5 |
| Neuronal intranuclear inclusion disease is genetically heterogeneous. <i>Annals of Clinical and Translational Neurology</i> , 2020 , 7, 1716-1725 | 5.3 | 18 |
| | Effect Modifiers of TDP-43-Associated Hippocampal Atrophy Rates in Patients with Alzheimer® Disease Neuropathological Changes. <i>Journal of Alzheimer® Disease</i> , 2020, 73, 1511-1523 TDP-43 is associated with a reduced likelihood of rendering a clinical diagnosis of dementia with Lewy bodies in autopsy-confirmed cases of transitional/diffuse Lewy body disease. <i>Journal of Neurology</i> , 2020, 267, 1444-1453 MRI and flortaucipir relationships in Alzheimer® phenotypes are heterogeneous. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 707-721 Truncated stathmin-2 is a marker of TDP-43 pathology in frontotemporal dementia. <i>Journal of Clinical Investigation</i> , 2020, 130, 6080-6092 Longitudinal anatomic, functional, and molecular characterization of Pick disease phenotypes. <i>Neurology</i> , 2020, 95, e3190-e3202 Pick® disease: clinicopathologic characterization of 21 cases. <i>Journal of Neurology</i> , 2020, 267, 2697-270 Western Aphasia Battery-Revised Profiles in Primary Progressive Aphasia and Primary Progressive Apraxia of Speech. <i>American Journal of Speech-Language Pathology</i> , 2020, 29, 498-510 Communication Limitations in Patients With Progressive Apraxia of Speech and Aphasia. <i>American Journal of Speech-Language Pathology</i> , 2020, 29, 1976-1986 Dysphagia in Progressive Supranuclear Palsy. <i>Dysphagia</i> , 2020, 35, 667-676 Incidence of frontotemporal disorders in Olmsted County: A population-based study. <i>Alzheimerm and Dementia</i> , 2020, 16, 482-490 Longitudinal flortaucipir ([F]AV-1451) PET imaging in primary progressive apraxia of speech: A 6-year longitudinal study. <i>Parkinsonism and Related Disorders</i> , 2020, 78, 200-203 Predicting future rates of tau accumulation on PET. <i>Brain</i> , 2020, 143, 3136-3150 Dementia with Lewy bodies presenting as Logopenic variant primary progressive Aphasia. <i>Neurocase</i> , 2020, 26, 259-263 Longitudinal Amyloid-PET in Atypical Alzheimer® Disease and Frontotemporal Lobar Degeneration. <i>Journal of Alzheimer® Disease</i> , 2020, 74, 377-389 | Effect Modifiers of TDP-43-Associated Hippocampal Atrophy Rates in Patients with Alzheimeriß Disease Neuropathological Changes. <i>Journal of AlzheimermDisease</i> , 2020, 73, 1511-1523 TDP-43 is associated with a reduced likelihood of rendering a clinical diagnosis of dementia with Lewy bodies in autopsy-confirmed cases of transitional/diffuse Lewy body disease. <i>Journal of Neurology</i> , 2020, 267, 1444-1453 MRI and Riortaucipir relationships in Alzheimeriß phenotypes are heterogeneous. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 707-721 Truncated stathmin-2 is a marker of TDP-43 pathology in frontotemporal dementia. <i>Journal of Clinical Investigation</i> , 2020, 130, 6080-6092 Longitudinal anatomic, functional, and molecular characterization of Pick disease phenotypes. <i>Neurology</i> , 2020, 95, e3190-e3202 Pickß disease: clinicopathologic characterization of 21 cases. <i>Journal of Neurology</i> , 2020, 267, 2697-270-4;5 Western Aphasia Battery-Revised Profiles in Primary Progressive Aphasia and Primary Progressive Apraxia of Speech. <i>American Journal of Speech-Language Pathology</i> , 2020, 29, 498-510 Communication Limitations in Patients With Progressive Apraxia of Speech and Aphasia. <i>American Journal of Speech-Language Pathology</i> , 2020, 29, 1976-1986 Dysphagia in Progressive Supranuclear Palsy. <i>Dysphagia</i> , 2020, 35, 667-676 37 Incidence of frontotemporal disorders in Olmsted County: A population-based study. <i>Alzheimerm and Dementia</i> , 2020, 16, 482-490 Longitudinal flortaucipir ([F]AV-1451) PET imaging in primary progressive apraxia of speech. <i>Cortex</i> , 2020, 124, 33-43 The evolution of parkinsonism in primary progressive apraxia of speech: A 6-year longitudinal study. <i>Parkinsonism and Related Disorders</i> , 2020, 81, 34-40 Predicting future rates of tau accumulation on PET. <i>Brain</i> , 2020, 143, 3136-3150 11.2 Dementia with Levy bodies presenting as Logopenic variant primary progressive Aphasia. <i>Neurocase</i> , 2020, 26, 259-263 Longitudinal Amyloid-IPET in Atypical Alzheimeris Disease and Frontotempora |

| 252 | Automated Hippocampal Subfield Volumetric Analyses in Atypical Alzheimer Disease. <i>Journal of Alzheimer Disease</i> , 2020 , 78, 927-937 | 4.3 | 2 |
|-----|--|--------|--------------|
| 251 | Sensitivity-Specificity of Tau and Amyloid IPositron Emission Tomography in Frontotemporal Lobar Degeneration. <i>Annals of Neurology</i> , 2020 , 88, 1009-1022 | 9.4 | 9 |
| 250 | Validation of the movement disorder society criteria for the diagnosis of 4-repeat tauopathies. <i>Movement Disorders</i> , 2020 , 35, 171-176 | 7 | 23 |
| 249 | Neuroanatomical correlates of phonologic errors in logopenic progressive aphasia. <i>Brain and Language</i> , 2020 , 204, 104773 | 2.9 | 7 |
| 248 | Association between transactive response DNA-binding protein of A3 kDa type and cognitive resilience to Alzheimer disease: a control study. <i>Neurobiology of Aging</i> , 2020 , 92, 92-97 | 5.6 | 4 |
| 247 | Neuropathologic basis of frontotemporal dementia in progressive supranuclear palsy. <i>Movement Disorders</i> , 2019 , 34, 1655-1662 | 7 | 8 |
| 246 | Progressive supranuclear palsy is not associated with neurogenic orthostatic hypotension. <i>Neurology</i> , 2019 , 93, e1339-e1347 | 6.5 | 7 |
| 245 | Antemortem volume loss mirrors TDP-43 staging in older adults with non-frontotemporal lobar degeneration. <i>Brain</i> , 2019 , 142, 3621-3635 | 11.2 | 22 |
| 244 | Transient Epileptic Amnesia: A Treatable Cause of Spells Associated With Persistent Cognitive Symptoms. <i>Frontiers in Neurology</i> , 2019 , 10, 939 | 4.1 | 9 |
| 243 | The influence of tau, amyloid, alpha-synuclein, TDP-43, and vascular pathology in clinically normal elderly individuals. <i>Neurobiology of Aging</i> , 2019 , 77, 26-36 | 5.6 | 32 |
| 242 | Progressive agrammatic aphasia without apraxia of speech as a distinct syndrome. <i>Brain</i> , 2019 , 142, 24 | 66-248 | 32 18 |
| 241 | An Evaluation of the Progressive Supranuclear Palsy Speech/Language Variant. <i>Movement Disorders Clinical Practice</i> , 2019 , 6, 452-461 | 2.2 | 20 |
| 240 | Cross-sectional associations of tau-PET signal with cognition in cognitively unimpaired adults. <i>Neurology</i> , 2019 , 93, e29-e39 | 6.5 | 36 |
| 239 | Longitudinal tau-PET uptake and atrophy in atypical Alzheimerß disease. <i>NeuroImage: Clinical</i> , 2019 , 23, 101823 | 5.3 | 27 |
| 238 | Brain atrophy in primary age-related tauopathy is linked to transactive response DNA-binding protein of 43 kDa. <i>Alzheimermand Dementia</i> , 2019 , 15, 799-806 | 1.2 | 11 |
| 237 | How to apply the movement disorder society criteria for diagnosis of progressive supranuclear palsy. <i>Movement Disorders</i> , 2019 , 34, 1228-1232 | 7 | 56 |
| 236 | The role of age on tau PET uptake and gray matter atrophy in atypical Alzheimerß disease. <i>Alzheimermand Dementia</i> , 2019 , 15, 675-685 | 1.2 | 18 |
| 235 | Rare Tauopathies. Seminars in Neurology, 2019 , 39, 264-273 | 3.2 | 2 |

| 234 | Prominent auditory deficits in primary progressive aphasia: A case study. <i>Cortex</i> , 2019 , 117, 396-406 | 3.8 | 7 |
|-----|--|------|----|
| 233 | Sensitivity and Specificity of Diagnostic Criteria for Progressive Supranuclear Palsy. <i>Movement Disorders</i> , 2019 , 34, 1144-1153 | 7 | 56 |
| 232 | 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. <i>Acta Neuropathologica</i> , 2019 , 137, 879-899 | 14.3 | 50 |
| 231 | Clinical and neuroimaging characteristics of clinically unclassifiable primary progressive aphasia. <i>Brain and Language</i> , 2019 , 197, 104676 | 2.9 | 21 |
| 230 | Clinicopathologic subtype of Alzheimerß disease presenting as corticobasal syndrome. <i>Alzheimern</i> and Dementia, 2019 , 15, 1218-1228 | 1.2 | 20 |
| 229 | LATE to the PART-y. <i>Brain</i> , 2019 , 142, e47 | 11.2 | 25 |
| 228 | Multimodal neuroimaging relationships in progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2019 , 66, 56-61 | 3.6 | 10 |
| 227 | C-terminal and full length TDP-43 specie differ according to FTLD-TDP lesion type but not genetic mutation. <i>Acta Neuropathologica Communications</i> , 2019 , 7, 100 | 7.3 | 9 |
| 226 | Submentalis Rapid Eye Movement Sleep Muscle Activity: A Potential Biomarker for Synucleinopathy. <i>Annals of Neurology</i> , 2019 , 86, 969-974 | 9.4 | 7 |
| 225 | Extensive transcriptomic study emphasizes importance of vesicular transport in C9orf72 expansion carriers. <i>Acta Neuropathologica Communications</i> , 2019 , 7, 150 | 7.3 | 18 |
| 224 | Pathological, imaging and genetic characteristics support the existence of distinct TDP-43 types in non-FTLD brains. <i>Acta Neuropathologica</i> , 2019 , 137, 227-238 | 14.3 | 32 |
| 223 | MRI Outperforms [18F]AV-1451 PET as a Longitudinal Biomarker in Progressive Supranuclear Palsy. <i>Movement Disorders</i> , 2019 , 34, 105-113 | 7 | 21 |
| 222 | F-AV-1451 uptake differs between dementia with lewy bodies and posterior cortical atrophy. <i>Movement Disorders</i> , 2019 , 34, 344-352 | 7 | 18 |
| 221 | The influence of Eamyloid on [F]AV-1451 in semantic variant of primary progressive aphasia. <i>Neurology</i> , 2019 , 92, e710-e722 | 6.5 | 8 |
| 220 | Electroencephalography in Primary Progressive Aphasia and Apraxia of Speech. <i>Aphasiology</i> , 2019 , 33, 1410-1417 | 1.6 | 4 |
| 219 | Regional multimodal relationships between tau, hypometabolism, atrophy, and fractional anisotropy in atypical Alzheimerß disease. <i>Human Brain Mapping</i> , 2019 , 40, 1618-1631 | 5.9 | 26 |
| 218 | [F] AV-1451 uptake in corticobasal syndrome: the influence of beta-amyloid and clinical presentation. <i>Journal of Neurology</i> , 2018 , 265, 1079-1088 | 5.5 | 18 |
| 217 | Regional Distribution, Asymmetry, and Clinical Correlates of Tau Uptake on [18F]AV-1451 PET in Atypical Alzheimerß Disease. <i>Journal of Alzheimer</i> ß Disease, 2018 , 62, 1713-1724 | 4.3 | 32 |

| 216 | Elevated medial temporal lobe and pervasive brain tau-PET signal in normal participants. <i>Alzheimerm and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2018 , 10, 210-216 | 5.2 | 11 |
|-----|---|------|-----|
| 215 | [F]AV-1451 tau-PET and primary progressive aphasia. <i>Annals of Neurology</i> , 2018 , 83, 599-611 | 9.4 | 46 |
| 214 | Rest in peace FTDP-17. <i>Brain</i> , 2018 , 141, 324-331 | 11.2 | 5 |
| 213 | Tau-PET imaging with [18F]AV-1451 in primary progressive apraxia of speech. <i>Cortex</i> , 2018 , 99, 358-374 | 3.8 | 31 |
| 212 | Pittsburgh Compound B and AV-1451 positron emission tomography assessment of molecular pathologies of Alzheimerß disease in progressive supranuclear palsy. <i>Parkinsonism and Related Disorders</i> , 2018 , 48, 3-9 | 3.6 | 22 |
| 211 | [F]AV-1451 clustering of entorhinal and cortical uptake in Alzheimerß disease. <i>Annals of Neurology</i> , 2018 , 83, 248-257 | 9.4 | 42 |
| 210 | Longitudinal structural and molecular neuroimaging in agrammatic primary progressive aphasia. <i>Brain</i> , 2018 , 141, 302-317 | 11.2 | 23 |
| 209 | Widespread brain tau and its association with ageing, Braak stage and Alzheimer dementia. <i>Brain</i> , 2018 , 141, 271-287 | 11.2 | 139 |
| 208 | Potential genetic modifiers of disease risk and age at onset in patients with frontotemporal lobar degeneration and GRN mutations: a genome-wide association study. <i>Lancet Neurology, The</i> , 2018 , 17, 548-558 | 24.1 | 60 |
| 207 | Imaging correlations of tau, amyloid, metabolism, and atrophy in typical and atypical Alzheimerß disease. <i>Alzheimermand Dementia</i> , 2018 , 14, 1005-1014 | 1.2 | 47 |
| 206 | Disrupted functional connectivity in primary progressive apraxia of speech. <i>NeuroImage: Clinical</i> , 2018 , 18, 617-629 | 5.3 | 19 |
| 205 | Molecular neuroimaging in primary progressive aphasia with predominant agraphia. <i>Neurocase</i> , 2018 , 24, 121-123 | 0.8 | O |
| 204 | FDG-PET in tau-negative amnestic dementia resembles that of autopsy-proven hippocampal sclerosis. <i>Brain</i> , 2018 , 141, 1201-1217 | 11.2 | 46 |
| 203 | A Neuropsychiatric Analysis of the Cotard Delusion. <i>Journal of Neuropsychiatry and Clinical Neurosciences</i> , 2018 , 30, 58-65 | 2.7 | 17 |
| 202 | Corticobasal degeneration: key emerging issues. <i>Journal of Neurology</i> , 2018 , 265, 439-445 | 5.5 | 20 |
| 201 | Fitting TDP-43 into the APOE 4 and neurodegeneration story. <i>Lancet Neurology, The</i> , 2018 , 17, 735-737 | 24.1 | 2 |
| 200 | Prosodic and phonetic subtypes of primary progressive apraxia of speech. <i>Brain and Language</i> , 2018 , 184, 54-65 | 2.9 | 62 |
| 199 | TDP-43 and Alzheimerß Disease Pathologic Subtype in Non-Amnestic Alzheimerß Disease Dementia. <i>Journal of Alzheimerm Disease</i> , 2018 , 64, 1227-1233 | 4.3 | 16 |

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| 198 | Non-right handed primary progressive apraxia of speech. <i>Journal of the Neurological Sciences</i> , 2018 , 390, 246-254 | 3.2 | 4 |
|-----|--|------|-----|
| 197 | Quantitative Analysis of Agrammatism in Agrammatic Primary Progressive Aphasia and Dominant Apraxia of Speech. <i>Journal of Speech, Language, and Hearing Research</i> , 2018 , 61, 2337-2346 | 2.8 | 10 |
| 196 | Diffuse Lewy body disease manifesting as corticobasal syndrome: A rare form of Lewy body disease. <i>Neurology</i> , 2018 , 91, e268-e279 | 6.5 | 30 |
| 195 | The diagnosis of progressive supranuclear palsy: current opinions and challenges. <i>Expert Review of Neurotherapeutics</i> , 2018 , 18, 603-616 | 4.3 | 12 |
| 194 | Clinical and imaging progression over 10 years in a patient with primary progressive apraxia of speech and autopsy-confirmed corticobasal degeneration. <i>Neurocase</i> , 2018 , 24, 111-120 | 0.8 | 18 |
| 193 | Corticobasal degeneration with TDP-43 pathology presenting with progressive supranuclear palsy syndrome: a distinct clinicopathologic subtype. <i>Acta Neuropathologica</i> , 2018 , 136, 389-404 | 14.3 | 27 |
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| 35 | Imaging correlates of posterior cortical atrophy. <i>Neurobiology of Aging</i> , 2007 , 28, 1051-61 | 5.6 | 137 |
| 34 | Hippocampal sclerosis in tau-negative frontotemporal lobar degeneration. <i>Neurobiology of Aging</i> , 2007 , 28, 1718-22 | 5.6 | 41 |
| 33 | Rates of cerebral atrophy in autopsy-confirmed progressive supranuclear palsy. <i>Annals of Neurology</i> , 2006 , 59, 200-3 | 9.4 | 30 |
| 32 | Benign tremulous parkinsonism. <i>Archives of Neurology</i> , 2006 , 63, 354-7 | | 109 |
| 31 | Clinicopathological and imaging correlates of progressive aphasia and apraxia of speech. <i>Brain</i> , 2006 , 129, 1385-98 | 11.2 | 529 |
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| 29 | The relationship between histopathological features of progressive supranuclear palsy and disease duration. <i>Parkinsonism and Related Disorders</i> , 2006 , 12, 109-12 | 3.6 | 13 |
| 28 | Visual hallucinations in posterior cortical atrophy. <i>Archives of Neurology</i> , 2006 , 63, 1427-32 | | 52 |
| 27 | Frontotemporal lobar degeneration without lobar atrophy. <i>Archives of Neurology</i> , 2006 , 63, 1632-8 | | 48 |
| 26 | Clinically undetected motor neuron disease in pathologically proven frontotemporal lobar degeneration with motor neuron disease. <i>Archives of Neurology</i> , 2006 , 63, 506-12 | | 62 |
| 25 | Atypical progressive supranuclear palsy with corticospinal tract degeneration. <i>Journal of Neuropathology and Experimental Neurology</i> , 2006 , 65, 396-405 | 3.1 | 110 |
| 24 | MRI correlates of alien leg-like phenomenon in corticobasal degeneration. <i>Movement Disorders</i> , 2005 , 20, 870-3 | 7 | 17 |
| 23 | Increased tau burden in the cortices of progressive supranuclear palsy presenting with corticobasal syndrome. <i>Movement Disorders</i> , 2005 , 20, 982-8 | 7 | 100 |
| 22 | Alpha-synuclein immunohistochemistry in two cases of co-occurring idiopathic Parkinson disease and motor neuron disease. <i>Movement Disorders</i> , 2005 , 20, 1515-20 | 7 | 17 |
| 21 | Extending the clinicopathological spectrum of neurofilament inclusion disease. <i>Acta Neuropathologica</i> , 2005 , 109, 427-32 | 14.3 | 26 |
| 20 | Mild cognitive impairment 2005 , 409-415 | | |
| 19 | Atypical progressive supranuclear palsy underlying progressive apraxia of speech and nonfluent aphasia. <i>Neurocase</i> , 2005 , 11, 283-96 | 0.8 | 148 |

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| 17 | Survival in two variants of tau-negative frontotemporal lobar degeneration: FTLD-U vs FTLD-MND. <i>Neurology</i> , 2005 , 65, 645-7 | 6.5 | 69 |
| 16 | Hippocampal sclerosis and ubiquitin-positive inclusions in dementia lacking distinctive histopathology. <i>Dementia and Geriatric Cognitive Disorders</i> , 2004 , 17, 342-5 | 2.6 | 33 |
| 15 | Apolipoprotein E epsilon 4 is a determinant for Alzheimer-type pathologic features in tauopathies, synucleinopathies, and frontotemporal degeneration. <i>Archives of Neurology</i> , 2004 , 61, 1579-84 | | 58 |
| 14 | Heterozygous Niemann-Pick disease type C presenting with tremor. <i>Neurology</i> , 2004 , 63, 2189-90 | 6.5 | 35 |
| 13 | Nonvasculitic autoimmune inflammatory meningoencephalitis. <i>Neuropathology</i> , 2004 , 24, 149-52 | 2 | 28 |
| 12 | Frontotemporal lobar degeneration and ubiquitin immunohistochemistry. <i>Neuropathology and Applied Neurobiology</i> , 2004 , 30, 369-73 | 5.2 | 121 |
| 11 | The Alien Limb. <i>Practical Neurology</i> , 2004 , 4, 44-45 | 2.4 | 12 |
| 10 | Creutzfeldt-Jakob disease presenting as progressive supranuclear palsy. <i>European Journal of Neurology</i> , 2004 , 11, 343-6 | 6 | 37 |
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| 8 | Correlation between antemortem magnetic resonance imaging findings and pathologically confirmed corticobasal degeneration. <i>Archives of Neurology</i> , 2004 , 61, 1881-4 | | 60 |
| 7 | Pathologically confirmed corticobasal degeneration presenting with visuospatial dysfunction. <i>Neurology</i> , 2003 , 61, 1134-5 | 6.5 | 92 |
| 6 | Neurofilament inclusion body disease: a new proteinopathy?. <i>Brain</i> , 2003 , 126, 2291-303 | 11.2 | 162 |
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| 4 | Atrophy of superior cerebellar peduncle in progressive supranuclear palsy. <i>Neurology</i> , 2003 , 60, 1766-9 | 6.5 | 104 |
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| 1 | Alpha-synuclein studies are negative in postencephalic parkinsonism of von Economo. <i>Neurology</i> , 2002 , 59, 645-6 | 6.5 | 22 |