James Burrell

List of Publications by Year in descending order

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304743 155660 4,580 60 22 55 h-index citations g-index papers 60 60 60 6082 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Longitudinal changes in behaviour, mood and functional capacity in the primary progressive aphasia variants. European Journal of Neuroscience, 2022, 56, 5601-5614.	2.6	7
2	Plasma Oxytocin Is Not Associated with Social Cognition or Behavior in Frontotemporal Dementia and Alzheimer's Disease Syndromes. Dementia and Geriatric Cognitive Disorders, 2022, 51, 241-248.	1.5	3
3	Utility of the Addenbrooke's Cognitive Examination III online calculator to differentiate the primary progressive aphasia variants. Brain Communications, 2022, 4, .	3. 3	6
4	Longitudinal cognitive and functional changes in primary progressive aphasia. Journal of Neurology, 2021, 268, 1951-1961.	3.6	16
5	Heterogeneity of behavioural and language deficits in FTD–MND. Journal of Neurology, 2021, 268, 2876-2889.	3 . 6	4
6	"More than words―– Longitudinal linguistic changes in the works of a writer diagnosed with semantic dementia. Neurocase, 2021, 27, 243-252.	0.6	6
7	The Box Task: A novel tool to differentiate the primary progressive aphasias. European Journal of Neurology, 2021, 28, 3945-3954.	3. 3	3
8	Verbal Short-Term Memory Disturbance in the Primary Progressive Aphasias: Challenges and Distinctions in a Clinical Setting. Brain Sciences, 2021, 11, 1060.	2.3	11
9	Amyotrophic lateral sclerosis features predict TDP-43 pathology in frontotemporal lobar degeneration. Neurobiology of Aging, 2021, 107, 11-20.	3.1	1
10	Visuospatial short-term and working memory disturbance in the primary progressive aphasias: Neuroanatomical and clinical implications. Cortex, 2020, 132, 223-237.	2.4	15
11	Sleep and orexin: A new paradigm for understanding behavioural-variant frontotemporal dementia?. Sleep Medicine Reviews, 2020, 54, 101361.	8.5	8
12	Using a second-person approach to identify disease-specific profiles of social behavior in frontotemporal dementia and Alzheimer's disease. Cortex, 2020, 133, 236-246.	2.4	2
13	What to make of equivocal amyloid imaging results. Neurocase, 2020, 26, 137-146.	0.6	2
14	Correlates of anomia in non-semantic variants of primary progressive aphasia converge over time. Cortex, 2019, 120, 201-211.	2.4	16
15	Clinical and neuroimaging investigations of language disturbance in frontotemporal dementia–motor neuron disease patients. Journal of Neurology, 2019, 266, 921-933.	3.6	14
16	Visuospatial dysfunction in Alzheimer's disease and behavioural variant frontotemporal dementia. Journal of the Neurological Sciences, 2019, 402, 74-80.	0.6	27
17	Sustained attention failures on a 3-min reaction time task is a sensitive marker of dementia. Journal of Neurology, 2019, 266, 1323-1331.	3.6	12
18	Can visuospatial measures improve the diagnosis of Alzheimer's disease?. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2018, 10, 66-74.	2.4	63

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19	Falls in frontotemporal dementia and related syndromes. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 159, 195-203.	1.8	1
20	Intrafamilial Phenotypic Variability in the C9orf72 Gene Expansion: 2 Case Studies. Frontiers in Psychology, 2018, 9, 1615.	2.1	9
21	Addenbrooke's Cognitive Examination III: Psychometric Characteristics and Relations to Functional Ability in Dementia. Journal of the International Neuropsychological Society, 2018, 24, 854-863.	1.8	66
22	Predicting Development of Amyotrophic Lateral Sclerosis in Frontotemporal Dementia. Journal of Alzheimer's Disease, 2017, 58, 163-170.	2.6	17
23	The midbrainâ€toâ€pons ratio distinguishes progressive supranuclear palsy from nonâ€fluent primary progressive aphasias. European Journal of Neurology, 2017, 24, 956-965.	3.3	11
24	Aphasia in Progressive Supranuclear Palsy: As Severe as Progressive Non-Fluent Aphasia. Journal of Alzheimer's Disease, 2017, 61, 705-715.	2.6	20
25	Dissociation of Structural and Functional Integrities of the Motor System in Amyotrophic Lateral		

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37	Apraxia and Motor Dysfunction in Corticobasal Syndrome. PLoS ONE, 2014, 9, e92944.	2.5	26
38	Job variation in advanced training in adult neurology in Australia and New Zealand: a follow-up study. Internal Medicine Journal, 2014, 44, 554-561.	0.8	0
39	C9 <scp>ORF</scp> 72 familial motor neuron disease â^ frontotemporal dementia associated with lung adenocarcinoma and antiâ€Ma2/Ta antibodies: a chance association?. European Journal of Neurology, 2014, 21, e31-3.	3.3	5
40	Degradation of emotion processing ability in corticobasal syndrome and Alzheimer's disease. Brain, 2014, 137, 3061-3072.	7.6	88
41	Heritability in frontotemporal dementia: more missing pieces?. Journal of Neurology, 2014, 261, 2170-2177.	3.6	27
42	Measuring disease progression in corticobasal syndrome. Journal of Neurology, 2014, 261, 1598-1605.	3.6	5
43	Cognition in corticobasal syndrome and progressive supranuclear palsy: A review. Movement Disorders, 2014, 29, 684-693.	3.9	137
44	Frontotemporal Dementia Associated With the <i>C9ORF72</i> Mutation. JAMA Neurology, 2014, 71, 331.	9.0	144
45	Disability in atypical parkinsonian syndromes is more dependent on memory dysfunction than motor symptoms. Parkinsonism and Related Disorders, 2013, 19, 436-440.	2.2	7
46	Early saccades in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2013, 14, 294-301.	1.7	9
47	Could immunological mechanisms trigger neurodegeneration in frontotemporal dementia?. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 946-946.	1.9	2
48	Clinical Profile of PiB-Positive Corticobasal Syndrome. PLoS ONE, 2013, 8, e61025.	2.5	48
49	Response to Karam et al Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2012, 13, 159-160.	2.1	0
50	The Neural Basis of Logopenic Progressive Aphasia. Journal of Alzheimer's Disease, 2012, 32, 1051-1059.	2.6	53
51	Saccadic abnormalities in frontotemporal dementia. Neurology, 2012, 78, 1816-1823.	1.1	41
52	Grey and White Matter Changes across the Amyotrophic Lateral Sclerosis-Frontotemporal Dementia Continuum. PLoS ONE, 2012, 7, e43993.	2.5	168
53	Isolated bulbar phenotype of amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2011, 12, 283-289.	2.1	52
54	Amyotrophic lateral sclerosis. Lancet, The, 2011, 377, 942-955.	13.7	2,182

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55	Subtypes of progressive aphasia: application of the international consensus criteria and validation using \hat{l}^2 -amyloid imaging. Brain, 2011, 134, 3030-3043.	7.6	294
56	Motor Neuron dysfunction in frontotemporal dementia. Brain, 2011, 134, 2582-2594.	7.6	271
57	Predicting a Positive Response to Intravenous Immunoglobulin in Isolated Lower Motor Neuron Syndromes. PLoS ONE, 2011, 6, e27041.	2.5	13
58	From FUS to Fibs: What's New in Frontotemporal Dementia?. Journal of Alzheimer's Disease, 2010, 21, 349-360.	2.6	13
59	Coma and seizures due to gas emboli following extubation. Journal of Clinical Neuroscience, 2009, 16, 344-345.	1.5	0
60	Right trochlear and left oculomotor palsies. Journal of Clinical Neuroscience, 2009, 16, 1464.	1.5	O