Suvankar Pal

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9634961/publications.pdf

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53	1,976	361413	254184
papers	citations	20	43
		h-index	g-index

57	57	57	2910
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical presentation and pre-mortem diagnosis of variant Creutzfeldt-Jakob disease associated with blood transfusion: a case report. Lancet, The, 2006, 368, 2061-2067.	13.7	374
2	Safety and efficacy of quinacrine in human prion disease (PRION-1 study): a patient-preference trial. Lancet Neurology, The, 2009, 8, 334-344.	10.2	226
3	ALS-specific cognitive and behavior changes associated with advancing disease stage in ALS. Neurology, 2018, 91, e1370-e1380.	1.1	170
4	Influence of Intracerebral Hemorrhage Location on Incidence, Characteristics, and Outcome. Stroke, 2015, 46, 361-368.	2.0	142
5	A systematic review and metaâ€analysis of retinal nerve fiber layer change in dementia, using optical coherence tomography. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2015, 1, 136-143.	2.4	133
6	Functional (Psychogenic) Cognitive Disorders: A Perspective from the Neurology Clinic. Journal of Alzheimer's Disease, 2015, 48, S5-S17.	2.6	83
7	The importance of ongoing international surveillance for Creutzfeldtâ€"Jakob disease. Nature Reviews Neurology, 2021, 17, 362-379.	10.1	69
8	TDP-43 as a potential biomarker for amyotrophic lateral sclerosis: a systematic review and meta-analysis. BMC Neurology, 2018, 18, 90.	1.8	63
9	A systematic review and metaâ€analysis of ¹⁸Fâ€labeled amyloid imaging in Alzheimer's disease. Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring, 2015, 1, 5-13.	2.4	59
10	Systematic review of the diagnostic utility of SPECT imaging in dementia. European Archives of Psychiatry and Clinical Neuroscience, 2013, 263, 539-552.	3.2	56
11	Executive, language and fluency dysfunction are markers of localised TDP-43 cerebral pathology in non-demented ALS. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 149-157.	1.9	54
12	The diagnostic utility of EEG in early-onset dementia: a systematic review of the literature with narrative analysis. Journal of Neural Transmission, 2014, 121, 59-69.	2.8	53
13	Improved PCR based methods for detecting C9orf72 hexanucleotide repeat expansions. Molecular and Cellular Probes, 2016, 30, 218-224.	2.1	42
14	Non-Motor Symptoms Profile and Burden in Drug NaÃ¯ve Versus Long-Term Parkinson's Disease Patients. Journal of Parkinson's Disease, 2014, 4, 541-547.	2.8	41
15	Changing epidemiology of motor neurone disease in Scotland. Journal of Neurology, 2019, 266, 817-825.	3.6	40
16	Genetic epidemiology of motor neuron disease-associated variants in the Scottish population. Neurobiology of Aging, 2017, 51, 178.e11-178.e20.	3.1	37
17	Clinical trials in amyotrophic lateral sclerosis: a systematic review and perspective. Brain Communications, 2021, 3, fcab242.	3.3	32
18	Validation of Revised International Creutzfeldt-Jakob Disease Surveillance Network Diagnostic Criteria for Sporadic Creutzfeldt-Jakob Disease. JAMA Network Open, 2022, 5, e2146319.	5.9	28

#	Article	IF	Citations
19	Phenotypic diversity of genetic Creutzfeldtâ€“Jakob disease: a histo-molecular-based classification. Acta Neuropathologica, 2021, 142, 707-728.	7.7	24
20	The brief Dimensional Apathy Scale: A short clinical assessment of apathy. Clinical Neuropsychologist, 2020, 34, 423-435.	2.3	22
21	A systematic review of neuropsychiatric and cognitive assessments used in clinical trials for amyotrophic lateral sclerosis. Journal of Neurology, 2021, 268, 4510-4521.	3.6	22
22	A systematic review of non-motor symptom evaluation in clinical trials for amyotrophic lateral sclerosis. Journal of Neurology, 2022, 269, 411-426.	3.6	21
23	Clinical audit research and evaluation of motor neuron disease (CARE-MND): a national electronic platform for prospective, longitudinal monitoring of MND in Scotland. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 242-250.	1.7	19
24	Investigating Domain-Specific Cognitive Impairment Among Patients With Multiple Sclerosis Using Touchscreen Cognitive Testing in Routine Clinical Care. Frontiers in Neurology, 2018, 9, 331.	2.4	17
25	Relationship between neuropsychiatric disorders and cognitive and behavioural change in MND. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 245-253.	1.9	15
26	Reliability and validity of the brief dimensional apathy scale. Archives of Clinical Neuropsychology, 2020, 35, 539-544.	0.5	13
27	Cerebrospinal fluid cytotoxicity in amyotrophic lateral sclerosis: a systematic review of in vitro studies. Brain Communications, 2020, 2, fcaa121.	3.3	10
28	40 Years of CSF Toxicity Studies in ALS: What Have We Learnt About ALS Pathophysiology?. Frontiers in Molecular Neuroscience, 2021, 14, 647895.	2.9	10
29	Motor Neuron Disease Systematic Multi-Arm Adaptive Randomised Trial (MND-SMART): a multi-arm, multi-stage, adaptive, platform, phase III randomised, double-blind, placebo-controlled trial of repurposed drugs in motor neuron disease. BMJ Open, 2022, 12, e064173.	1.9	10
30	Trials for neurodegenerative diseases: time to innovate. Lancet Neurology, The, 2021, 20, 984.	10.2	9
31	Incidental Findings Identified on Head MRI for Investigation of Cognitive Impairment: A Retrospective Review. Dementia and Geriatric Cognitive Disorders, 2019, 48, 123-130.	1.5	8
32	Bilateral lower limb weakness in acute severe ulcerative colitis. Lancet, The, 2016, 388, 101-102.	13.7	7
33	Surveillance for variant CJD: should more children with neurodegenerative diseases have autopsies?. Archives of Disease in Childhood, 2019, 104, 360-365.	1.9	7
34	Application of telehealth for comprehensive Creutzfeldt-Jakob disease surveillance in the United Kingdom. Journal of the Neurological Sciences, 2021, 420, 117221.	0.6	7
35	An epidemiological profile of dysarthria incidence and assistive technology use in the living population of people with MND in Scotland. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 116-122.	1.7	6
36	The immediate impact of the COVID-19 pandemic on motor neuron disease services and mortality in Scotland. Journal of Neurology, 2021, 268, 2038-2040.	3.6	6

#	Article	IF	Citations
37	Prospective observational cohort study of factors influencing trial participation in people with motor neuron disease (FIT-participation-MND): a protocol. BMJ Open, 2021, 11, e044996.	1.9	6
38	Validation of The Edinburgh cognitive and behavioural ALS screen (ECAS) in behavioural variant frontotemporal dementia and Alzheimerâ€™s disease. International Journal of Geriatric Psychiatry, 2021, 36, 1576-1587.	2.7	5
39	The Prevalence and Management of Saliva Problems in Motor Neuron Disease: A 4-Year Analysis of the Scottish Motor Neuron Disease Register. Neurodegenerative Diseases, 2020, 20, 147-152.	1.4	5
40	Progressive unsteadiness in a 68-year-old man with longstanding abdominal pain and altered bowel habit. Practical Neurology, 2009, 9, 210-220.	1.1	4
41	Excellent reliability of the ALSFRS-R administered via videoconferencing: A study of people with motor neuron disease in Scotland. Journal of the Neurological Sciences, 2020, 416, 116991.	0.6	4
42	Riluzole prescribing, uptake and treatment discontinuation in people with amyotrophic lateral sclerosis in Scotland. Journal of Neurology, 2020, 267, 2459-2461.	3.6	4
43	Impact of the COVIDâ€19 pandemic on Creutzfeldtâ€“Jakob disease surveillance and patient care in the United Kingdom. European Journal of Neurology, 2021, , .	3.3	4
44	Screening for cognitive impairment among patients with neuromyelitis optica using touchscreen cognitive testing in routine clinical care. Journal of Neurology, 2019, 266, 2571-2572.	3.6	2
45	The prevalence of multimorbidity and its impact on survival in people with motor neuron disease. European Journal of Neurology, 2021, 28, 2756-2765.	3.3	2
46	31st Advanced Clinical Neurology Course, Edinburgh 2009: progressive cognitive impairment, behavioural change and upper motor neuron signs in a 57-year-old woman. Practical Neurology, 2011, 11, 71-80.	1.1	1
47	070â€... A pilot study assessing choroidal thickness in frontotemporal dementia and motor neuron disease. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, A122.2-A122.	1.9	1
48	A challenging case of double vision and altered facial sensation. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, 539-540.	1.9	0
49	P2-190: Detecting profiles of cognitive performance in dementia syndromes and parkinson's disease using brief computerized cognitive tests. , 2015, 11, P565-P565.		0
50	Clinicopathological case: progressive somnolence and dementia in an accountant: when the shine rubs off the gold standard. Practical Neurology, 2018, 18, 505-512.	1.1	0
51	Understanding the risk of incidental findings: A qualitative study of people with cognitive symptoms. Journal of the Neurological Sciences, 2020, 419, 117203.	0.6	0
52	Non-white cases of sporadic Creutzfeldt-Jakob disease: A 28Âyear review of United Kingdom National Surveillance Data. Journal of the Neurological Sciences, 2021, 424, 117416.	0.6	0
53	The assessment of visually guided reaching in prodromal Alzheimerâ€™s disease. Journal of Vision, 2020, 20, 1059.	0.3	0