

Parvathi Menon

List of Publications by Year in descending order

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Version: 2024-02-01

43
papers

1,976
citations

257450

24
h-index

276875

41
g-index

45
all docs

45
docs citations

45
times ranked

1906
citing authors

#	ARTICLE	IF	CITATIONS
1	Pathophysiological associations of transcallosal dysfunction in ALS. <i>European Journal of Neurology</i> , 2021, 28, 1172-1180.	3.3	12
2	Study protocol of RESCUE-ALS: A Phase 2, randomised, double-blind, placebo-controlled study in early symptomatic amyotrophic lateral sclerosis patients to assess bioenergetic catalysis with CNM-Au8 as a mechanism to slow disease progression. <i>BMJ Open</i> , 2021, 11, e041479.	1.9	33
3	Diagnostic Utility of Gold Coast Criteria in <scp>Amyotrophic Lateral Sclerosis</scp>. <i>Annals of Neurology</i> , 2021, 89, 979-986.	5.3	68
4	Association of Cortical Hyperexcitability and Cognitive Impairment in Patients With Amyotrophic Lateral Sclerosis. <i>Neurology</i> , 2021, 96, e2090-e2097.	1.1	12
5	Utility of Transcranial Magnetic Stimulation in Studying Upper Motor Neuron Dysfunction in Amyotrophic Lateral Sclerosis. <i>Brain Sciences</i> , 2021, 11, 906.	2.3	4
6	The Upper Motor Neuronâ€™Improved Knowledge from ALS and Related Clinical Disorders. <i>Brain Sciences</i> , 2021, 11, 958.	2.3	3
7	Split-hand index: A diagnostic and prognostic marker in amyotrophic lateral sclerosis across varying regions of onset. <i>Clinical Neurophysiology</i> , 2021, 132, 2130-2135.	1.5	7
8	ALS is a multistep process in South Korean, Japanese, and Australian patients. <i>Neurology</i> , 2020, 94, e1657-e1663.	1.1	39
9	Cortical hyperexcitability evolves with disease progression in ALS. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 733-741.	3.7	45
10	Regional motor cortex dysfunction in amyotrophic lateral sclerosis. <i>Annals of Clinical and Translational Neurology</i> , 2019, 6, 1373-1382.	3.7	19
11	Amyotrophic lateral sclerosis diagnostic index. <i>Neurology</i> , 2019, 92, e536-e547.	1.1	17
12	Association of Regulatory T-Cell Expansion With Progression of Amyotrophic Lateral Sclerosis. <i>JAMA Neurology</i> , 2018, 75, 681.	9.0	120
13	Utility of threshold tracking transcranial magnetic stimulation in ALS. <i>Clinical Neurophysiology Practice</i> , 2018, 3, 164-172.	1.4	51
14	Imbalance of cortical facilitatory and inhibitory circuits underlies hyperexcitability in ALS. <i>Neurology</i> , 2018, 91, e1669-e1676.	1.1	67
15	Cortical excitability varies across different muscles. <i>Journal of Neurophysiology</i> , 2018, 120, 1397-1403.	1.8	14
16	Physiological Processes Underlying Short Interval Intracortical Facilitation in the Human Motor Cortex. <i>Frontiers in Neuroscience</i> , 2018, 12, 240.	2.8	31
17	Physiological processes influencing motor-evoked potential duration with voluntary contraction. <i>Journal of Neurophysiology</i> , 2017, 117, 1156-1162.	1.8	23
18	The evolution of motor cortical dysfunction in amyotrophic lateral sclerosis. <i>Clinical Neurophysiology</i> , 2017, 128, 1075-1082.	1.5	34

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19	Isolated nerve plasmacytoma in a patient previously in systemic myeloma remission. <i>Muscle and Nerve</i> , 2017, 55, E27-E28.	2.2	0
20	Hyperpolarization-activated cyclic-nucleotide-gated channels potentially modulate axonal excitability at different thresholds. <i>Journal of Neurophysiology</i> , 2017, 118, 3044-3050.	1.8	9
21	Cortical hyperexcitability may contribute to disease spread in als. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017, 88, e1.16-e1.	1.9	0
22	Imbalance in cortical inhibition-excitation networks underlies als. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017, 88, e1.14-e1.	1.9	1
23	Awaji criteria improves the diagnostic sensitivity in amyotrophic lateral sclerosis: A systematic review using individual patient data. <i>Clinical Neurophysiology</i> , 2016, 127, 2684-2691.	1.5	74
24	Pathophysiological and diagnostic implications of cortical dysfunction in ALS. <i>Nature Reviews Neurology</i> , 2016, 12, 651-661.	10.1	165
25	Diagnostic criteria in amyotrophic lateral sclerosis. <i>Neurology</i> , 2016, 87, 684-690.	1.1	46
26	Motor cortical function determines prognosis in sporadic ALS. <i>Neurology</i> , 2016, 87, 513-520.	1.1	76
27	Riluzole exerts transient modulating effects on cortical and axonal hyperexcitability in ALS. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2016, 17, 580-588.	1.7	58
28	Cortical contributions to the flail leg syndrome: Pathophysiological insights. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2016, 17, 389-396.	1.7	23
29	Potential structural and functional biomarkers of upper motor neuron dysfunction in ALS. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2016, 17, 85-92.	1.7	32
30	Rate of disease progression: a prognostic biomarker in ALS. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, 628-632.	1.9	123
31	Sensitivity and specificity of threshold tracking transcranial magnetic stimulation for diagnosis of amyotrophic lateral sclerosis: a prospective study. <i>Lancet Neurology</i> , The, 2015, 14, 478-484.	10.2	164
32	Cortical hyperexcitability precedes lower motor neuron dysfunction in ALS. <i>Clinical Neurophysiology</i> , 2015, 126, 803-809.	1.5	140
33	Cortical Dysfunction Underlies the Development of the Split-Hand in Amyotrophic Lateral Sclerosis. <i>PLoS ONE</i> , 2014, 9, e87124.	2.5	75
34	Cortical hyperexcitability and the split-hand plus phenomenon: Pathophysiological insights in ALS. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2014, 15, 250-256.	1.7	27
35	Cortical excitability differences in hand muscles follow a split-hand pattern in healthy controls. <i>Muscle and Nerve</i> , 2014, 49, 836-844.	2.2	22
36	ALS pathophysiology: Insights from the split-hand phenomenon. <i>Clinical Neurophysiology</i> , 2014, 125, 186-193.	1.5	44

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37	Utility of Dissociated Intrinsic Hand Muscle Atrophy in the Diagnosis of Amyotrophic Lateral Sclerosis. <i>Journal of Visualized Experiments</i> , 2014, , .	0.3	7
38	Motor cortical function and the precision grip. <i>Physiological Reports</i> , 2014, 2, e12120.	1.7	9
39	Split-hand plus sign in ALS: Differential involvement of the flexor pollicis longus and intrinsic hand muscles. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2013, 14, 315-318.	1.7	46
40	Split-hand index for the diagnosis of amyotrophic lateral sclerosis. <i>Clinical Neurophysiology</i> , 2013, 124, 410-416.	1.5	97
41	Riluzole exerts central and peripheral modulating effects in amyotrophic lateral sclerosis. <i>Brain</i> , 2013, 136, 1361-1370.	7.6	123
42	Abnormalities of neuromuscular transmission in patients with Millerâ€Fisher syndrome. <i>Journal of Clinical Neuroscience</i> , 2012, 19, 1599-1601.	1.5	3
43	Appearance, phenomenology and diagnostic utility of the split hand in amyotrophic lateral sclerosis. <i>Neurodegenerative Disease Management</i> , 2011, 1, 457-462.	2.2	12