Mingsheng Liu

List of Publications by Year in descending order

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59 papers	662 citations	14 h-index	752256 20 g-index
65	65	65	880
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	<i>ANXA11</i> mutations prevail in Chinese ALS patients with and without cognitive dementia. Neurology: Genetics, 2018, 4, e237.	0.9	40
2	Environmental risk factors and amyotrophic lateral sclerosis (ALS): A case-control study of ALS in China. Journal of Clinical Neuroscience, 2019, 66, 12-18.	0.8	33
3	Phenotypic differences of amyotrophic lateral sclerosis (ALS) in China and Germany. Journal of Neurology, 2018, 265, 774-782.	1.8	31
4	Comparison of the Upper Marginal Neurons of Cortical Layer 2 with Layer 2/3 Pyramidal Neurons in Mouse Temporal Cortex. Frontiers in Neuroanatomy, 2017, 11, 115.	0.9	28
5	Correlation of Creatine Kinase Levels with Clinical Features and Survival in Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 2017, 8, 322.	1.1	28
6	Cognitive Impairment in Chinese Patients with Sporadic Amyotrophic Lateral Sclerosis. PLoS ONE, 2015, 10, e0137921.	1.1	24
7	Excessive daytime sleepiness in Chinese patients with sporadic amyotrophic lateral sclerosis and its association with cognitive and behavioural impairments. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 1038-1043.	0.9	24
8	Crossâ€sectional area reference values for sonography of nerves in the upper extremities. Muscle and Nerve, 2020, 61, 338-346.	1.0	24
9	Multiple Sites Ultrasonography of Peripheral Nerves in Differentiating Charcot–Marie–Tooth Type 1A from Chronic Inflammatory Demyelinating Polyradiculoneuropathy. Frontiers in Neurology, 2017, 8, 181.	1.1	22
10	Amyotrophic Lateral Sclerosis and Myasthenia Gravis Overlap Syndrome: A Review of Two Cases and the Associated Literature. Frontiers in Neurology, 2017, 8, 218.	1.1	20
11	Motor nerve conduction study and muscle strength in newly diagnosed poems syndrome. Muscle and Nerve, 2015, 51, 19-23.	1.0	19
12	Diagnostic Performance of Neurofilaments in Chinese Patients With Amyotrophic Lateral Sclerosis: A Prospective Study. Frontiers in Neurology, 2018, 9, 726.	1.1	19
13	Brain Structural and Perfusion Signature of Amyotrophic Lateral Sclerosis With Varying Levels of Cognitive Deficit. Frontiers in Neurology, 2018, 9, 364.	1.1	17
14	Microneedle Electrode Array for Electrical Impedance Myography to Characterize Neurogenic Myopathy. Annals of Biomedical Engineering, 2016, 44, 1566-1575.	1.3	16
15	The Awaji criteria increases the diagnostic sensitivity of the revised El Escorial criteria for amyotrophic lateral sclerosis diagnosis in a Chinese population. PLoS ONE, 2017, 12, e0171522.	1.1	15
16	Clinical diagnosis and treatment recommendations for immune checkpoint inhibitorâ€related adverse reactions in the nervous system. Thoracic Cancer, 2020, 11, 481-487.	0.8	15
17	Differences in Dysfunction of Thenar and Hypothenar Motoneurons in Amyotrophic Lateral Sclerosis. Frontiers in Human Neuroscience, 2016, 10, 99.	1.0	14
18	Split-hand index in amyotrophic lateral sclerosis: an F-wave study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 562-567.	1.1	14

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19	Serial nerve ultrasound and motor nerve conduction studies in chronic inflammatory demyelinating polyradiculoneuropathy. Muscle and Nerve, 2019, 60, 254-262.	1.0	14
20	Neurophysiological Differences between Flail Arm Syndrome and Amyotrophic Lateral Sclerosis. PLoS ONE, 2015, 10, e0127601.	1.1	13
21	Vagus Nerve Ultrasound in Chronic Inflammatory Demyelinating Polyradiculoneuropathy and Charcotâ€Marieâ€Tooth Disease Type 1A. Journal of Neuroimaging, 2020, 30, 910-916.	1.0	13
22	Singleâ€fiber electromyography in amyotrophic lateral sclerosis and cervical spondylosis. Muscle and Nerve, 2013, 48, 137-139.	1.0	12
23	Mutation analysis of KIF5A in Chinese amyotrophic lateral sclerosis patients. Neurobiology of Aging, 2019, 73, 229.e1-229.e4.	1.5	12
24	A prospective study on split-hand index as a biomarker for the diagnosis of amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 574-583.	1.1	12
25	TheÂGold Coast criteria increases the diagnostic sensitivity for amyotrophic lateral sclerosis in a Chinese population. Translational Neurodegeneration, 2021, 10, 28.	3.6	12
26	Creatine kinase level and its relationship with quantitative electromyographic characteristics in amyotrophic lateral sclerosis. Clinical Neurophysiology, 2018, 129, 926-930.	0.7	11
27	Reassessment of Split-Leg Signs in Amyotrophic Lateral Sclerosis: Differential Involvement of the Extensor Digitorum Brevis and Abductor Hallucis Muscles. Frontiers in Neurology, 2019, 10, 565.	1.1	11
28	Fasciculation differences between ALS and non-ALS patients: an ultrasound study. BMC Neurology, 2021, 21, 441.	0.8	11
29	Afterdischarges following M waves in patients with voltage-gated potassium channels antibodies. Clinical Neurophysiology Practice, 2017, 2, 72-75.	0.6	9
30	Genetic analysis of TIA1 gene in Chinese patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2018, 67, 201.e9-201.e10.	1,5	9
31	Split-Hand Syndrome in Amyotrophic Lateral Sclerosis: Differences in Dysfunction of the FDI and ADM Spinal Motoneurons. Frontiers in Neuroscience, 2019, 13, 371.	1.4	8
32	Split hand in amyotrophic lateral sclerosis: A systematic review and meta-analysis. Journal of Clinical Neuroscience, 2021, 90, 293-301.	0.8	8
33	A Retrospective Study of the Characteristics and Clinical Significance of A-Waves in Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 2017, 8, 515.	1.1	7
34	Reference Values and Influencing Factors Analysis for Current Perception Threshold Testing Based on Study of 166 Healthy Chinese. Frontiers in Neuroscience, 2018, 12, 14.	1.4	7
35	Early onset but long survival and other prognostic factors in Chinese sporadic amyotrophic lateral sclerosis. Journal of Clinical Neuroscience, 2019, 69, 74-80.	0.8	7
36	Split phenomenon of antagonistic muscle groups in amyotrophic lateral sclerosis: relative preservation of flexor muscles. Neurological Research, 2021, 43, 372-380.	0.6	7

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37	Mechanism hypotheses for the electrophysiological manifestations of two cases of endplate acetylcholinesterase deficiency related congenital myasthenic syndrome. Journal of Clinical Neuroscience, 2018, 48, 229-232.	0.8	6
38	Restless Legs Syndrome in Chinese Patients With Sporadic Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 2018, 9, 735.	1.1	6
39	Motor Nerve Conduction Block Predicting Outcome of Guillain-Barre Syndrome. Frontiers in Neurology, 2018, 9, 399.	1.1	6
40	Reference values for lower limb nerve ultrasound and its diagnostic sensitivity. Journal of Clinical Neuroscience, 2021, 86, 276-283.	0.8	6
41	Nerve Ultrasound Performances in Differentiating POEMS Syndrome from CIDP. Neurotherapeutics, 2022, 19, 455-463.	2.1	6
42	Differences in F-Wave Characteristics between Spinobulbar Muscular Atrophy and Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2016, 8, 50.	1.7	5
43	Conduction Block and Nerve Cross-Sectional Area in Multifocal Motor Neuropathy. Frontiers in Neurology, 2019, 10, 1055.	1.1	4
44	Nerve ultrasound studies in POEMS syndrome. Muscle and Nerve, 2021, 63, 758-764.	1.0	4
45	Amyotrophic lateral sclerosis with frontotemporal dementia presented with prominent psychosis. Chinese Medical Journal, 2014, 127, 3996-8.	0.9	4
46	Genotypeâ€"phenotype association of TARDBP mutations in Chinese patients with amyotrophic lateral sclerosis: a single-center study and systematic review of published literature. Journal of Neurology, 2022, 269, 4204-4212.	1.8	4
47	Motor conduction block and conduction velocity in Lewis-Sumner syndrome and multifocal motor neuropathy. Journal of Clinical Neuroscience, 2019, 67, 10-13.	0.8	3
48	GJB1 Mutation-A Disease Spectrum: Report of Case Series. Frontiers in Neurology, 2019, 10, 1406.	1.1	3
49	Strategy for screening cognitive impairment in Chinese patients with amyotrophic lateral sclerosis. Journal of Clinical Neuroscience, 2020, 81, 105-110.	0.8	3
50	Neuropsychological Investigation in Chinese Patients with Progressive Muscular Atrophy. PLoS ONE, 2015, 10, e0128883.	1.1	3
51	Nerve ultrasound may help predicting response to immune treatment in chronic inflammatory demyelinating polyradiculoneuropathy. Neurological Sciences, 2022, 43, 3929-3937.	0.9	3
52	Single-fiber EMG with concentric electrodes in lambert-eaton myasthenia. Muscle and Nerve, 2017, 56, 253-257.	1.0	2
53	Plateaus and reversals evaluated by different methods in patients with limb-onset amyotrophic lateral sclerosis. Journal of Clinical Neuroscience, 2022, 97, 93-98.	0.8	2
54	Abduction range: A potential parameter for the long exercise test in hypokalemic periodic paralysis during interâ€attack periods. Muscle and Nerve, 2020, 61, 104-107.	1.0	1

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55	The frequency of ALSFRS-R reversals and plateaus in patients with limb-onset amyotrophic lateral sclerosis: a cohort study. Acta Neurologica Belgica, 2022, 122, 1567-1573.	0.5	1
56	Reference value of long-time exercise test in the diagnosis of primary periodic paralysis. Chinese Medical Journal, 2014, 127, 3219-23.	0.9	1
57	Survival analysis of clinical and genetic factors in an amyotrophic lateral sclerosis cohort from China. Neurological Research, 2022, 44, 651-658.	0.6	1
58	Chronic inflammatory demyelinating polyradiculoneuropathy concomitant with nephropathy. Neurological Sciences, 0, , .	0.9	1
59	A rare cause of fever of unknown origin - cervical spinal cord lesion. Chinese Medical Journal, 2014, 127, 3517-8.	0.9	0