## Liying Cui

## List of Publications by Year in descending order

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		201385	1	89595	
159	3,580	27		50	
papers	citations	h-index		g-index	
187	187	187		4859	
all docs	docs citations	times ranked		citing authors	

#	Article	IF	CITATIONS
1	Autoimmune Cerebellar Ataxia: Etiology and Clinical Characteristics of a Case Series from China. Cerebellum, 2023, 22, 379-385.	1.4	3
2	Slow progression of amyotrophic lateral sclerosis in a Chinese patient carrying SOD1 p.S135T mutation. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, 23, 143-145.	1.1	0
3	Whole-exome sequencing identifies a novel de novo variant in DYNC1H in a patient with intractable epilepsy. Neurological Sciences, 2022, 43, 2853-2858.	0.9	2
4	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
5	Typical and atypical phenotype and neuroimaging of X-linked adrenoleukodystrophy in a Chinese cohort. Neurological Sciences, 2022, , $1.$	0.9	3
6	Nerve ultrasound may help predicting response to immune treatment in chronic inflammatory demyelinating polyradiculoneuropathy. Neurological Sciences, 2022, 43, 3929-3937.	0.9	3
7	Changes of Functional, Morphological, and Inflammatory Reactions in Spontaneous Peripheral Nerve Reinnervation after Thermal Injury. Oxidative Medicine and Cellular Longevity, 2022, 2022, 1-11.	1.9	4
8	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
9	Association Between Common Variants of APOE, ABCA7, A2M, BACE1, and Cerebrospinal Fluid Biomarkers in Alzheimer's Disease: Data from the PUMCH Dementia Cohort. Journal of Alzheimer's Disease, 2022, 85, 1511-1518.	1.2	3
10	Survival analysis of clinical and genetic factors in an amyotrophic lateral sclerosis cohort from China. Neurological Research, 2022, 44, 651-658.	0.6	1
11	Nerve Ultrasound Performances in Differentiating POEMS Syndrome from CIDP. Neurotherapeutics, 2022, 19, 455-463.	2.1	6
12	Genotype–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
13	Altered executive control network connectivity in antiâ€NMDA receptor encephalitis. Annals of Clinical and Translational Neurology, 2022, 9, 30-40.	1.7	10
14	Real-world outcomes of teriflunomide in relapsing–remitting multiple sclerosis: a prospective cohort study. Journal of Neurology, 2022, 269, 4808-4816.	1.8	11
15	PSEN2 Mutation Spectrum and Novel Functionally Validated Mutations in Alzheimer's Disease: Data from PUMCH Dementia Cohort. Journal of Alzheimer's Disease, 2022, , 1-8.	1.2	2
16	Study on sleep-wake disorders in patients with genetic and non-genetic amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 96-102.	0.9	8
17	Phenotype Heterogeneity and Genotype Correlation of MAPT Mutations in a Chinese PUMCH Cohort. Journal of Molecular Neuroscience, 2021, 71, 1015-1022.	1.1	3
18	Sensitivity and specificity of single and combined clouds analyses compared with quantitative motor unit potential analysis. Muscle and Nerve, 2021, 63, 225-230.	1.0	1

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19	Mutations of <i>DNAJC7</i> are rare in Chinese amyotrophic lateral sclerosis patients. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, 22, 312-315.	1.1	8
20	Increased Premature Cerebral Small Vessel Diseases in Dialysis Patients: A Retrospective Cross-Sectional Study. Nephron, 2021, 145, 330-341.	0.9	3
21	Differential Expression of miRNA in the Peripheral Blood Mononuclear Cells in Myasthenia Gravis with Muscle-Specific Receptor Tyrosine Kinase Antibodies. Critical Reviews in Eukaryotic Gene Expression, 2021, 31, 1-15.	0.4	4
22	Correlation between total homocysteine and cerebral small vessel disease: A Mendelian randomization study. European Journal of Neurology, 2021, 28, 1931-1938.	1.7	31
23	Peripheral nerve hyperexcitability syndrome: A clinical, electrophysiological, and immunological study. Muscle and Nerve, 2021, 63, 697-702.	1.0	2
24	The Impact of COVID-19 on Patients With Neuromyelitis Optica Spectrum Disorder Beyond Infection Risk. Frontiers in Neurology, 2021, 12, 657037.	1.1	9
25	Nerve ultrasound studies in POEMS syndrome. Muscle and Nerve, 2021, 63, 758-764.	1.0	4
26	White matter hyperintensities and patterns of atrophy in early onset Alzheimer's disease with causative gene mutations. Clinical Neurology and Neurosurgery, 2021, 203, 106552.	0.6	5
27	Reference values for lower limb nerve ultrasound and its diagnostic sensitivity. Journal of Clinical Neuroscience, 2021, 86, 276-283.	0.8	6
28	The Epidemiology of COVID-19 and MS-Related Characteristics in a National Sample of People With MS in China. Frontiers in Neurology, 2021, 12, 682729.	1.1	3
29	A longitudinal observation of brain structure between AD and FTLD. Clinical Neurology and Neurosurgery, 2021, 205, 106604.	0.6	5
30	The role of glymphatic system in the cerebral edema formation after ischemic stroke. Experimental Neurology, 2021, 340, 113685.	2.0	31
31	Differences in Multimodal Electroencephalogram and Clinical Correlations Between Early-Onset Alzheimer's Disease and Frontotemporal Dementia. Frontiers in Neuroscience, 2021, 15, 687053.	1.4	8
32	TheÂGold Coast criteria increases the diagnostic sensitivity for amyotrophic lateral sclerosis in a Chinese population. Translational Neurodegeneration, 2021, 10, 28.	3.6	12
33	Clinical Phenotype and Mutation Spectrum of Alzheimer's Disease with Causative Genetic Mutation in a Chinese Cohort. Current Alzheimer Research, 2021, 18, 265-272.	0.7	5
34	Abnormal Brain Activation During Verbal Memory Encoding in Postacute Anti-N-Methyl-d-Aspartate Receptor Encephalitis. Brain Connectivity, 2021, , .	0.8	1
35	Split phenomenon of antagonistic muscle groups in amyotrophic lateral sclerosis: relative preservation of flexor muscles. Neurological Research, 2021, 43, 372-380.	0.6	7
36	A pilot study of multiple time points and multidomain assessment in cerebrospinal fluid tap test for patients with idiopathic normal pressure hydrocephalus. Clinical Neurology and Neurosurgery, 2021, 210, 107012.	0.6	3

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37	Fasciculation differences between ALS and non-ALS patients: an ultrasound study. BMC Neurology, 2021, 21, 441.	0.8	11
38	Exonic rearrangements in <i>DMD</i> in Chinese Han individuals affected with Duchenne and Becker muscular dystrophies. Human Mutation, 2020, 41, 668-677.	1.1	29
39	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
40	Single-molecule optical mapping enables quantitative measurement of D4Z4 repeats in facioscapulohumeral muscular dystrophy (FSHD). Journal of Medical Genetics, 2020, 57, 109-120.	1.5	43
41	Standards of instrumentation of EMG. Clinical Neurophysiology, 2020, 131, 243-258.	0.7	109
42	Anti-NMDAR encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	3.1	106
43	Crossâ€sectional area reference values for sonography of nerves in the upper extremities. Muscle and Nerve, 2020, 61, 338-346.	1.0	24
44	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
45	Novel <scp><i>NEXMIF</i></scp> gene pathogenic variant in a female patient with refractory epilepsy and intellectual disability. American Journal of Medical Genetics, Part A, 2020, 182, 2765-2772.	0.7	7
46	Efficacy and safety of cinepazide maleate injection in patients with acute ischemic stroke: a multicenter, randomized, double-blind, placebo-controlled trial. BMC Neurology, 2020, 20, 282.	0.8	3
47	Lacune and Large Perivascular Space: Two Kinds of Cavities Are of Different Risk Factors and Stroke Risk. Cerebrovascular Diseases, 2020, 49, 522-530.	0.8	10
48	Elevated fasting blood glucose is predictive of the severity and poor outcome in nondiabetic patients with cerebral venous thrombosis. Journal of the Neurological Sciences, 2020, 417, 117017.	0.3	9
49	Neurological Manifestations in Critically Ill Patients With COVID-19: A Retrospective Study. Frontiers in Neurology, 2020, 11, 806.	1.1	61
50	Study of B Cell Repertoire in Patients With Anti-N-Methyl-D-Aspartate Receptor Encephalitis. Frontiers in Immunology, 2020, 11, 1539.	2.2	9
51	Disrupted white matter integrity and network connectivity are related to poor motor performance. Scientific Reports, 2020, 10, 18369.	1.6	16
52	Carotid artery stiffness in rural adult Chinese: a cross-sectional analysis of the community-based China stroke cohort study. BMJ Open, 2020, 10, e036398.	0.8	2
53	Prediction of long-term disability in Chinese patients with multiple sclerosis: A prospective cohort study. Multiple Sclerosis and Related Disorders, 2020, 46, 102461.	0.9	8
54	A pilot study on the positive rate of multiâ€time point and multiple domain evaluation in cerebrospinal fluid tap test of patients with normal pressure hydrocephalus. Alzheimer's and Dementia, 2020, 16, e037463.	0.4	0

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55	The changing of cognitive function and the influence of learning effect on the results before and after the CSF tap test in patients with normal pressure hydrocephalus. Alzheimer's and Dementia, 2020, 16, e037467.	0.4	0
56	Factors related to CSF biomarkers examination in Alzheimer's disease. Alzheimer's and Dementia, 2020, 16, e041636.	0.4	0
57	Cognitionâ€related structure in the progression of Alzheimer's disease. Alzheimer's and Dementia, 2020, 16, e041640.	0.4	0
58	Pseudorabies virus encephalitis in humans: a case series study. Journal of NeuroVirology, 2020, 26, 556-564.	1.0	35
59	Age-related characteristics and normative values of F waves in healthy infants. Clinical Neurophysiology, 2020, 131, 1068-1074.	0.7	1
60	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
61	Spinal Intrathecal Actinomycosis Causes Multisegmental Root Failure: A Case Report. Frontiers in Neurology, 2020, 11, 621.	1.1	0
62	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
63	Overexpression of MicroRNA-9a-5p Ameliorates NLRP1 Inflammasome-mediated Ischemic Injury in Rats Following Ischemic Stroke. Neuroscience, 2020, 444, 106-117.	1.1	38
64	Association between lipoprotein(a) concentration and the risk of stroke in the Chinese Han population: a retrospective case-control study. Annals of Translational Medicine, 2020, 8, 212-212.	0.7	14
65	Long-term efficacy of mycophenolate mofetil in myelin oligodendrocyte glycoprotein antibody-associated disorders. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	3.1	46
66	Strategy for screening cognitive impairment in Chinese patients with amyotrophic lateral sclerosis. Journal of Clinical Neuroscience, 2020, 81, 105-110.	0.8	3
67	Construction of a long non‑coding RNA‑mediated transcription factor and gene regulatory triplet network reveals global patterns and biomarkers for ischemic stroke. International Journal of Molecular Medicine, 2020, 45, 333-342.	1.8	4
68	Mutation analysis of KIF5A in Chinese amyotrophic lateral sclerosis patients. Neurobiology of Aging, 2019, 73, 229.e1-229.e4.	1.5	12
69	Split-hand index in amyotrophic lateral sclerosis: an F-wave study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 562-567.	1.1	14
70	Early secondâ€line therapy is associated with improved episodic memory in antiâ€NMDA receptor encephalitis. Annals of Clinical and Translational Neurology, 2019, 6, 1202-1213.	1.7	14
71	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
72	Magnetic resonance fingerprinting of temporal lobe white matter in mesial temporal lobe epilepsy. Annals of Clinical and Translational Neurology, 2019, 6, 1639-1646.	1.7	18

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73	Conduction Block and Nerve Cross-Sectional Area in Multifocal Motor Neuropathy. Frontiers in Neurology, 2019, 10, 1055.	1.1	4
74	Non-invasive brain stimulation for fatigue in multiple sclerosis patients: A systematic review and meta-analysis. Multiple Sclerosis and Related Disorders, 2019, 36, 101375.	0.9	32
75	Surgical outcomes in patients with anti-N-methyl D-aspartate receptor encephalitis with ovarian teratoma. American Journal of Obstetrics and Gynecology, 2019, 221, 485.e1-485.e10.	0.7	35
76	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
77	Serial nerve ultrasound and motor nerve conduction studies in chronic inflammatory demyelinating polyradiculoneuropathy. Muscle and Nerve, 2019, 60, 254-262.	1.0	14
78	Treatable cause of hereditary spastic paraplegia: eight cases of combined homocysteinaemia with methylmalonic aciduria. Journal of Neurology, 2019, 266, 2434-2439.	1.8	9
79	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
80	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
81	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
82	Guidelines for single fiber EMG. Clinical Neurophysiology, 2019, 130, 1417-1439.	0.7	63
83	Consecutive Slides on Axial View Is More Effective Than Transversal Diameter to Differentiate Mechanisms of Single Subcortical Infarctions in the Lenticulostriate Artery Territory. Frontiers in Neurology, 2019, 10, 336.	1.1	2
84	Stroke in China: advances and challenges in epidemiology, prevention, and management. Lancet Neurology, The, 2019, 18, 394-405.	4.9	903
85	l-Arginine prevents stroke-like episodes but not brain atrophy: a 20-year follow-up of a MELAS patient. Neurological Sciences, 2019, 40, 209-211.	0.9	7
86	Acute epileptic seizures in myelin oligodendrocyte glycoprotein encephalomyelitis and neuromyelitis optica spectrum disorder: A comparative cohort study. Multiple Sclerosis and Related Disorders, 2019, 27, 281-288.	0.9	16
87	Whole exome sequencing identified a novel <i><scp>DAG</scp>1</i> mutation in a patient with rare, mild and late age of onset muscular dystrophyâ€dystroglycanopathy. Journal of Cellular and Molecular Medicine, 2019, 23, 811-818.	1.6	56
88	Systemic autoimmune diseases complicated with hydrocephalus: pathogenesis and management. Neurosurgical Review, 2019, 42, 255-261.	1.2	7
89	A therapeutic regimen for 3-hydroxyisobutyryl-CoA hydrolase deficiency with exercise-induced dystonia. European Journal of Paediatric Neurology, 2019, 23, 755-759.	0.7	13
90	GJB1 Mutation-A Disease Spectrum: Report of Case Series. Frontiers in Neurology, 2019, 10, 1406.	1.1	3

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91	Creatine kinase level and its relationship with quantitative electromyographic characteristics in amyotrophic lateral sclerosis. Clinical Neurophysiology, 2018, 129, 926-930.	0.7	11
92	Autoimmune GFAP astrocytopathy after viral encephalitis: A case report. Multiple Sclerosis and Related Disorders, 2018, 21, 84-87.	0.9	42
93	Phenotypic differences of amyotrophic lateral sclerosis (ALS) in China and Germany. Journal of Neurology, 2018, 265, 774-782.	1.8	31
94	Comparison of myelin oligodendrocyte glycoprotein (MOG)-antibody disease and AQP4-lgG-positive neuromyelitis optica spectrum disorder (NMOSD) when they co-exist with anti-NMDA (N-methyl-D-aspartate) receptor encephalitis. Multiple Sclerosis and Related Disorders, 2018, 20, 144-152.	0.9	89
95	Genetic analysis of TIA1 gene in Chinese patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2018, 67, 201.e9-201.e10.	1.5	9
96	Volumetric Changes in Hippocampal Subregions and Memory Performance in Mesial Temporal Lobe Epilepsy with Hippocampal Sclerosis. Neuroscience Bulletin, 2018, 34, 389-396.	1.5	12
97	Next-generation sequencing of the cerebrospinal fluid in the diagnosis of neurobrucellosis. International Journal of Infectious Diseases, 2018, 67, 20-24.	1.5	71
98	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
99	Resection of melanocytic nevi as a potential treatment of anti-NMDAR encephalitis patients without tumor: report of three cases. Neurological Sciences, 2018, 39, 165-167.	0.9	5
100	P2â€351: REVERSIBLE WHITE MATTER LESIONS IN NPH: TWO CASE REPORTS. Alzheimer's and Dementia, 2018, 14, P823.	0.4	0
101	Atherosclerosis Might Be Responsible for Branch Artery Disease: Evidence From White Matter Hyperintensity Burden in Acute Isolated Pontine Infarction. Frontiers in Neurology, 2018, 9, 840.	1.1	17
102	Diagnostic Performance of Neurofilaments in Chinese Patients With Amyotrophic Lateral Sclerosis: A Prospective Study. Frontiers in Neurology, 2018, 9, 726.	1.1	19
103	Restless Legs Syndrome in Chinese Patients With Sporadic Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 2018, 9, 735.	1.1	6
104	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
105	Brain Structural and Perfusion Signature of Amyotrophic Lateral Sclerosis With Varying Levels of Cognitive Deficit. Frontiers in Neurology, 2018, 9, 364.	1.1	17
106	Motor Nerve Conduction Block Predicting Outcome of Guillain-Barre Syndrome. Frontiers in Neurology, 2018, 9, 399.	1.1	6
107	Next-Generation Sequencing of Cerebrospinal Fluid for the Diagnosis of Neurocysticercosis. Frontiers in Neurology, 2018, 9, 471.	1.1	35
108	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

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109	Mutation screening of NEK1 in Chinese ALS patients. Neurobiology of Aging, 2018, 71, 267.e1-267.e4.	1.5	13
110	Mitochondrial DNA mutations in late-onset Leigh syndrome. Journal of Neurology, 2018, 265, 2388-2395.	1.8	31
111	Reorganization of anterior and posterior hippocampal networks associated with memory performance in mesial temporal lobe epilepsy. Clinical Neurophysiology, 2017, 128, 830-838.	0.7	24
112	Re-evaluate the Efficacy and Safety of Human Urinary Kallidinogenase (RESK): Protocol for an Open-Label, Single-Arm, Multicenter Phase IV Trial for the Treatment of Acute Ischemic Stroke in Chinese Patients. Translational Stroke Research, 2017, 8, 341-346.	2.3	12
113	Statistical analysis plan for the Head Position in Stroke Trial (HeadPoST): An international cluster cross-over randomized trial. International Journal of Stroke, 2017, 12, 667-670.	2.9	2
114	Depolarized GABAergic Signaling in Subicular Microcircuits Mediates Generalized Seizure in Temporal Lobe Epilepsy. Neuron, 2017, 95, 92-105.e5.	3.8	97
115	Afterdischarges following M waves in patients with voltage-gated potassium channels antibodies. Clinical Neurophysiology Practice, 2017, 2, 72-75.	0.6	9
116	Single-fiber EMG with concentric electrodes in lambert-eaton myasthenia. Muscle and Nerve, 2017, 56, 253-257.	1.0	2
117	[P4–078]: APOLIPOROTEIN E POLYMORPHISM IN CHINESE POPULATION WITH VARIOUS TYPES OF COGNITIVE DISORDERS. Alzheimer's and Dementia, 2017, 13, P1287.	0.4	O
118	[P1â€"300]: CORRELATION BETWEEN THE CLINICAL, NEUROIMAGING CHARACTERS AND THE CEREBROSPINAL FLUID TAP TEST RESPONSE IN CHINESE IDIOPATHIC NORMAL PRESSURE HYDROCEPHALUS PATIENTS. Alzheimer's and Dementia, 2017, 13, P367.	0.4	0
119	[P1–319]: EFFECTS OF APOLIPOPROTEIN E POLYMORPHISM ON NEUROPSYCHOLOGICAL DOMAINS IN CHINES POPULATION WITH ALZHEIMER's DISEASE. Alzheimer's and Dementia, 2017, 13, P376.	SE 0.4	0
120	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
121	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
122	Correlation of Creatine Kinase Levels with Clinical Features and Survival in Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 2017, 8, 322.	1.1	28
123	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
124	An LMNB1 Duplication Caused Adult-Onset Autosomal Dominant Leukodystrophy in Chinese Family: Clinical Manifestations, Neuroradiology and Genetic Diagnosis. Frontiers in Molecular Neuroscience, 2017, 10, 215.	1.4	11
125	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
126	Malnutrition-inflammation is a risk factor for cerebral small vessel diseases and cognitive decline in peritoneal dialysis patients: a cross-sectional observational study. BMC Nephrology, 2017, 18, 366.	0.8	21

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127	Military service and the risk of amyotrophic lateral sclerosis: A meta-analysis. Journal of Clinical Neuroscience, 2017, 45, 337-342.	0.8	17
128	Differences in F-Wave Characteristics between Spinobulbar Muscular Atrophy and Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2016, 8, 50.	1.7	5
129	Voxel-Wise Meta-Analysis of Gray Matter Changes in Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2016, 8, 64.	1.7	52
130	Neurofilaments in CSF As Diagnostic Biomarkers in Motor Neuron Disease: A Meta-Analysis. Frontiers in Aging Neuroscience, 2016, 8, 290.	1.7	25
131	Differences in Dysfunction of Thenar and Hypothenar Motoneurons in Amyotrophic Lateral Sclerosis. Frontiers in Human Neuroscience, 2016, 10, 99.	1.0	14
132	Elevated Fasting Blood Glucose Is Predictive of Poor Outcome in Non-Diabetic Stroke Patients: A Sub-Group Analysis of SMART. PLoS ONE, 2016, 11, e0160674.	1.1	47
133	Detection of <i>Listeria monocytogenes</i> ii>in CSF from Three Patients with Meningoencephalitis by		

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145	A Systematic Review and Meta-Analysis of the Functional MRI Investigation of Motor Neuron Disease. Frontiers in Neurology, 2015, 6, 246.	1.1	10
146	Neurophysiological Differences between Flail Arm Syndrome and Amyotrophic Lateral Sclerosis. PLoS ONE, 2015, 10, e0127601.	1.1	13
147	Cognitive Impairment in Chinese Patients with Sporadic Amyotrophic Lateral Sclerosis. PLoS ONE, 2015, 10, e0137921.	1.1	24
148	A comprehensive genetic diagnosis of Chinese muscular dystrophy and congenital myopathy patients by targeted next-generation sequencing. Neuromuscular Disorders, 2015, 25, 617-624.	0.3	29
149	Head Position in Stroke Trial (HeadPoST) $\hat{a}\in$ sitting-up vs lying-flat positioning of patients with acute stroke: study protocol for a cluster randomised controlled trial. Trials, 2015, 16, 256.	0.7	27
150	Neuropsychological Investigation in Chinese Patients with Progressive Muscular Atrophy. PLoS ONE, 2015, 10, e0128883.	1.1	3
151	Application of <sup>68</sup> Ga-PRGD2 PET/CT for α <sub>v</sub> β <sub>3</sub> -integrin Imaging of Myocardial Infarction and Stroke. Theranostics, 2014, 4, 778-786.	4.6	50
152	Symmetric Thalamic Lesions in a Patient With a Myoclonic Epilepsy with Ragged Red Fibers-Leigh Spectrum Phenotype due to the m.A8344G Mutation. Pediatric Neurology, 2014, 51, e19-e20.	1.0	3
153	A Novel Dystrophin Deletion Mutation in a Becker Muscular Dystrophy Patient With Early-Onset Dilated Cardiomyopathy. Canadian Journal of Cardiology, 2014, 30, 956.e1-956.e3.	0.8	5
154	Safflower yellow for acute ischemic stroke: A systematic review of randomized controlled trials. Complementary Therapies in Medicine, 2014, 22, 354-361.	1.3	35
155	Targeted next-generation sequencing as a comprehensive test for patients with and female carriers of DMD/BMD: a multi-population diagnostic study. European Journal of Human Genetics, 2014, 22, 110-118.	1.4	66
156	Specific Changes of Serum Proteins in Parkinson's Disease Patients. PLoS ONE, 2014, 9, e95684.	1.1	17
157	A rare case of limbic encephalitis with anti leucine-rich glioma inactivated-1 (LGI1) antibodies. Neuroendocrinology Letters, 2014, 35, 95-7.	0.2	7
158	Reference value of long-time exercise test in the diagnosis of primary periodic paralysis. Chinese Medical Journal, 2014, 127, 3219-23.	0.9	1
159	Amyotrophic lateral sclerosis with frontotemporal dementia presented with prominent psychosis. Chinese Medical Journal, 2014, 127, 3996-8.	0.9	4