Dongshen Fan

List of Publications by Year
in descending order

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147 papers	3,490 citations	236925 25 h-index	189892 50 g-index
187 all docs	187 docs citations	187 times ranked	4762 citing authors

#	Article	IF	Citations
1	Stroke in China: advances and challenges in epidemiology, prevention, and management. Lancet Neurology, The, 2019, 18, 394-405.	10.2	903
2	Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. Nature Genetics, 2021, 53, 1636-1648.	21.4	223
3	Improved precision of epigenetic clock estimates across tissues and its implication for biological ageing. Genome Medicine, 2019, 11, 54.	8.2	191
4	Global variation in prevalence and incidence of amyotrophic lateral sclerosis: a systematic review and meta-analysis. Journal of Neurology, 2020, 267, 944-953.	3.6	153
5	Natural history and clinical features of sporadic amyotrophic lateral sclerosis in China. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 1075-1081.	1.9	103
6	High Manganese, A Risk for Alzheimer's Disease: High Manganese Induces Amyloid-Î² Related Cognitive Impairment. Journal of Alzheimer's Disease, 2014, 42, 865-878.	2.6	99
7	A 36-week multicenter, randomized, double-blind, placebo-controlled, parallel-group, phase 3 clinical trial of sodium oligomannate for mild-to-moderate Alzheimerâ€™s dementia. Alzheimer's Research and Therapy, 2021, 13, 62.	6.2	99
8	Cross-ethnic meta-analysis identifies association of the GPX3-TNIP1 locus with amyotrophic lateral sclerosis. Nature Communications, 2017, 8, 611.	12.8	93
9	Diagnostic Accuracy of the Chinese Version of the Trailâ€Making Test for Screening Cognitive Impairment. Journal of the American Geriatrics Society, 2018, 66, 92-99.	2.6	84
10	Increased Interleukin-6 Levels in the Astrocyte-Derived Exosomes of Sporadic Amyotrophic Lateral Sclerosis Patients. Frontiers in Neuroscience, 2019, 13, 574.	2.8	61
11	C9orf72 hexanucleotide repeat expansions in Chinese sporadic amyotrophic lateral sclerosis. Neurobiology of Aging, 2015, 36, 2660.e1-2660.e8.	3.1	50
12	The Edinburgh Cognitive and Behavioural ALS Screen in a Chinese Amyotrophic Lateral Sclerosis Population. PLoS ONE, 2016, 11, e0155496.	2.5	50
13	Incidence and prevalence of amyotrophic lateral sclerosis in urban China: a national population-based study. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 520-525.	1.9	37
14	The epidemiology and genetics of Amyotrophic lateral sclerosis in China. Brain Research, 2018, 1693, 121-126.	2.2	36
15	Risk Score to Predict Hospital-Acquired Pneumonia After Spontaneous Intracerebral Hemorrhage. Stroke, 2014, 45, 2620-2628.	2.0	35
16	Amyotrophic lateral sclerosis in Beijing: Epidemiologic features and prognosis from 2010 to 2015. Brain and Behavior, 2018, 8, e01131.	2.2	35
17	Bloodâ€“brain barrier dysfunction in mice induced by lipopolysaccharide is attenuated by dapsone. Biochemical and Biophysical Research Communications, 2014, 453, 419-424.	2.1	34
18	Phospholipid transfer protein (PLTP) deficiency impaired bloodâ€“brain barrier integrity by increasing cerebrovascular oxidative stress. Biochemical and Biophysical Research Communications, 2014, 445, 352-356.	2.1	32

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19	Optineurin mutations in patients with sporadic amyotrophic lateral sclerosis in China. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2015, 16, 485-489.	1.7	32
20	Full sequencing and haplotype analysis of MAPT in Parkinson's disease and rapid eye movement sleep behavior disorder. Movement Disorders, 2018, 33, 1016-1020.	3.9	31
21	Effects of diet on adenosine monophosphate-activated protein kinase activity and disease progression in an amyotrophic lateral sclerosis model. Journal of International Medical Research, 2015, 43, 67-79.	1.0	30
22	Life Course Adiposity and Amyotrophic Lateral Sclerosis: A Mendelian Randomization Study. Annals of Neurology, 2020, 87, 434-441.	5.3	30
23	Long-Term Use of Riluzole Could Improve the Prognosis of Sporadic Amyotrophic Lateral Sclerosis Patients: A Real-World Cohort Study in China. Frontiers in Aging Neuroscience, 2016, 8, 246.	3.4	29
24	Molecular analysis and clinical diversity of distal hereditary motor neuropathy. European Journal of Neurology, 2020, 27, 1319-1326.	3.3	28
25	MATR3 mutation analysis in a Chinese cohort with sporadic amyotrophic lateral sclerosis. Neurobiology of Aging, 2016, 38, 218.e3-218.e4.	3.1	27
26	Phospholipid transfer protein (PLTP) deficiency accelerates memory dysfunction through altering amyloid precursor protein (APP) processing in a mouse model of Alzheimer's disease. Human Molecular Genetics, 2015, 24, 5388-5403.	2.9	24
27	Amyotrophic Lateral Sclerosis Genetic Studies. Neuroscientist, 2015, 21, 599-615.	3.5	23
28	Six SQSTM1 mutations in a Chinese amyotrophic lateral sclerosis cohort. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2015, 16, 378-384.	1.7	23
29	Whole-exome sequencing in amyotrophic lateral sclerosis suggests NEK1 is a risk gene in Chinese. Genome Medicine, 2017, 9, 97.	8.2	23
30	DCTN1 gene analysis in Chinese patients with sporadic amyotrophic lateral sclerosis. PLoS ONE, 2017, 12, e0182572.	2.5	23
31	A Randomized, Doubleâ€blind, Placeboâ€controlled Trial to Evaluate the Efficacy and Safety of Pregabalin for Postherpetic Neuralgia in a Population of Chinese Patients. Pain Practice, 2017, 17, 62-69.	1.9	22
32	Better survival in female SOD1-mutant patients with ALS: a study of SOD1-related natural history. Translational Neurodegeneration, 2019, 8, 2.	8.0	22
33	Trends in stroke subtypes and vascular risk factors in a stroke center in China over 10 years. Scientific Reports, 2018, 8, 5037.	3.3	21
34	Efficacy and safety of pregabalin for painful diabetic peripheral neuropathy in a population of Chinese patients: A randomized placeboâ€controlled trial. Journal of Diabetes, 2018, 10, 256-265.	1.8	19
35	Hypermetabolism associated with worse prognosis of amyotrophic lateral sclerosis. Journal of Neurology, 2022, 269, 1447-1455.	3.6	19
36	Different post label delay cerebral blood flow measurements in patients with Alzheimer's disease using 3D arterial spin labeling. Magnetic Resonance Imaging, 2015, 33, 1019-1025.	1.8	17

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37	A Novel Missense Mutation of the DDHD1 Gene Associated with Juvenile Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2016, 8, 291.	3.4	17
38	Corneal subâ€basal whorlâ€like nerve plexus: a landmark for early and followâ€up evaluation in transthyretin familial amyloid polyneuropathy. European Journal of Neurology, 2021, 28, 630-638.	3.3	17
39	Surgical stress induced depressive and anxiety like behavior are improved by dapsone via modulating NADPH oxidase level. Neuroscience Letters, 2015, 585, 103-108.	2.1	16
40	Lipids, Apolipoproteins, Statins, and Intracerebral Hemorrhage: A Mendelian Randomization Study. Annals of Neurology, 2022, 92, 390-399.	5.3	16
41	Changes in the concentrations of trimethylamine N-oxide (TMAO) and its precursors in patients with amyotrophic lateral sclerosis. Scientific Reports, 2020, 10, 15198.	3.3	15
42	Prognostic models for amyotrophic lateral sclerosis: a systematic review. Journal of Neurology, 2021, 268, 3361-3370.	3.6	15
43	Screening for TUBA4A mutations in a large Chinese cohort of patients with ALS: re-evaluating the pathogenesis of TUBA4A in ALS. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 1350-1352.	1.9	13
44	Increased extrasynaptic GluN2B expression is involved in cognitive impairment after isoflurane anesthesia. Experimental and Therapeutic Medicine, 2016, 12, 161-168.	1.8	12
45	CHCHD10 mutations in patients with amyotrophic lateral sclerosis in Mainland China. Neurobiology of Aging, 2017, 54, 214.e7-214.e10.	3.1	12
46	The Analysis of Two BDNF Polymorphisms G196A/C270T in Chinese Sporadic Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2017, 9, 135.	3.4	12
47	Two rare variants of the ANXA11 gene identified in Chinese patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2019, 74, 235.e9-235.e12.	3.1	12
48	Education, intelligence, and amyotrophic lateral sclerosis: A Mendelian randomization study. Annals of Clinical and Translational Neurology, 2020, 7, 1642-1647.	3.7	12
49	Treatment Adherence and Secondary Prevention of Ischemic Stroke Among Discharged Patients Using Mobile Phone- and WeChat-Based Improvement Services: Cohort Study. JMIR MHealth and UHealth, 2020, 8, e16496.	3.7	12
50	Adenosine monophosphate-activated protein kinase activation enhances embryonic neural stem cell apoptosis in a mouse model of amyotrophic lateral sclerosis. Neural Regeneration Research, 2014, 9, 1770.	3.0	12
51	Vestibular evoked myogenic potentials and their clinical utility in patients with amyotrophic lateral sclerosis. Clinical Neurophysiology, 2019, 130, 647-654.	1.5	11
52	Clinical and Genetic Features of Biallelic Mutations in SORD in a Series of Chinese Patients With Charcot-Marie-Tooth and Distal Hereditary Motor Neuropathy. Frontiers in Neurology, 2021, 12, 733926.	2.4	11
53	Association between type 2 diabetes and amyotrophic lateral sclerosis. Scientific Reports, 2022, 12, 2544.	3.3	11
54	Comparison of optical coherence tomography findings and visual field changes in patients with primary open-angle glaucoma and amyotrophic lateral sclerosis. Journal of Clinical Neuroscience, 2018, 48, 233-237.	1.5	10

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55	Sixteen-Week Interventional Study to Evaluate the Clinical Effects and Safety of Rivastigmine Capsules in Chinese Patients with Alzheimerâ€™s Disease. Journal of Alzheimer's Disease, 2019, 72, 1313-1322.	2.6	10
56	Urban prevalence of multiple sclerosis in China: A populationâ€based study in six provinces. European Journal of Neurology, 2021, 28, 1636-1644.	3.3	10
57	Human endogenous retrovirus K (HERV-K) env in neuronal extracellular vesicles: a new biomarker of motor neuron disease. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, 23, 100-107.	1.7	10
58	Assessment of bidirectional relationships between 98 genera of the human gut microbiota and amyotrophic lateral sclerosis: a 2-sample Mendelian randomization study. BMC Neurology, 2022, 22, 8.	1.8	10
59	Dietary-Derived Essential Nutrients and Amyotrophic Lateral Sclerosis: A Two-Sample Mendelian Randomization Study. Nutrients, 2022, 14, 920.	4.1	10
60	Neuroimmune Crosstalk Between the Peripheral and the Central Immune System in Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2022, 14, 890958.	3.4	10
61	TUBA4A may not be a significant genetic factor in Chinese ALS patients. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2016, 17, 148-150.	1.7	9
62	No Evidence for Pathogenic Role of UBQLN2 Mutations in Sporadic Amyotrophic Lateral Sclerosis in the Mainland Chinese Population. PLoS ONE, 2017, 12, e0170943.	2.5	9
63	Identification of an A4V SOD1 mutation in a Chinese patient with amyotrophic lateral sclerosis without the A4V founder effect common in North America. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2018, 19, 466-468.	1.7	9
64	In-hospital recurrence in a Chinese large cohort with acute ischemic stroke. Scientific Reports, 2019, 9, 14945.	3.3	9
65	Whole-exome sequencing identified novel KIF5A mutations in Chinese patients with amyotrophic lateral sclerosis and Charcot-Marie-Tooth type 2. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 326-328.	1.9	9
66	Is the C677T polymorphism in methylenetetrahydrofolate reductase gene or plasma homocysteine a risk factor for diabetic peripheral neuropathy in Chinese individuals?. Neural Regeneration Research, 2012, 7, 2384-91.	3.0	9
67	A natural history comparison of SOD1-mutant patients with amyotrophic lateral sclerosis between Chinese and German populations. Translational Neurodegeneration, 2021, 10, 42.	8.0	9
68	1.4 times increase in atrial fibrillation-related ischemic stroke and TIA over 12 years in a stroke center. Journal of the Neurological Sciences, 2017, 379, 1-6.	0.6	8
69	A novel mutation of BICD2 gene associated with juvenile amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2017, 18, 454-456.	1.7	8
70	Incidence and possible causes of nontraumatic convexal subarachnoid haemorrhage in Chinese patients: A retrospective review. Journal of International Medical Research, 2017, 45, 1870-1878.	1.0	8
71	The rs696880 Polymorphism in the Nogo-A Receptor Gene (RTN4R) Is Associated With Susceptibility to Sporadic Amyotrophic Lateral Sclerosis in the Chinese Population. Frontiers in Aging Neuroscience, 2018, 10, 108.	3.4	8
72	The protective role of pre-morbid type 2 diabetes in patients with amyotrophic lateral sclerosis: a center-based survey in China. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 209-215.	1.7	8

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73	Increased substantia nigra echogenicity correlated with visual hallucinations in Parkinsonâ€™s disease: a Chinese population-based study. Neurological Sciences, 2020, 41, 661-667.	1.9	8
74	TBK1 variants in Chinese patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2021, 97, 149.e9-149.e15.	3.1	8
75	Early Axonal Dysfunction of the Peripheral Nervous System Influences Disease Progression of ALS: Evidence From Clinical Neuroelectrophysiology. Frontiers in Neurology, 2021, 12, 574919.	2.4	8
76	Painful Diabetic Peripheral Neuropathy Study of Chinese Outpatients (PDNSCOPE): A Multicentre Cross-Sectional Registry Study of Clinical Characteristics and Treatment in Mainland China. Pain and Therapy, 2021, 10, 1355-1373.	3.2	8
77	Small fiber neuropathy for assessment of disease severity in amyotrophic lateral sclerosis: corneal confocal microscopy findings. Orphanet Journal of Rare Diseases, 2022, 17, 7.	2.7	8
78	Impaired synaptic vesicle recycling contributes to presynaptic dysfunction in lipoprotein lipase-deficient mice. Neuroscience, 2014, 280, 275-281.	2.3	7
79	Cognitive and behavioral impairments in German and Chinese ALS populations â€“ a post-hoc comparison of national study data. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 28-36.	1.7	7
80	Daytime sleepiness might increase the risk of ALS: a 2-sample Mendelian randomization study. Journal of Neurology, 2021, 268, 4332-4339.	3.6	7
81	Loss of appetite in patients with amyotrophic lateral sclerosis is associated with weight loss and anxiety/depression. Scientific Reports, 2021, 11, 9119.	3.3	7
82	Trends in the clinical features of amyotrophic lateral sclerosis: A 14â€year Chinese cohort study. European Journal of Neurology, 2021, 28, 2893-2900.	3.3	7
83	A twoâ€sample Mendelian randomization analysis of heart rate variability and cerebral small vessel disease. Journal of Clinical Hypertension, 2021, 23, 1608-1614.	2.0	7
84	The Genotype and Phenotype Features in a Large Chinese MFN2 Mutation Cohort. Frontiers in Neurology, 2021, 12, 757518.	2.4	7
85	Glucocerebrosidase Mutations Cause Mitochondrial and Lysosomal Dysfunction in Parkinsonâ€™s Disease: Pathogenesis and Therapeutic Implications. Frontiers in Aging Neuroscience, 2022, 14, 851135.	3.4	7
86	Leukocyte telomere length and amyotrophic lateral sclerosis: a Mendelian randomization study. Orphanet Journal of Rare Diseases, 2021, 16, 508.	2.7	7
87	A Novel Asp121Asn Mutation of Myelin Protein Zero Is Associated with Late-Onset Axonal Charcot-Marie-Tooth Disease, Hearing Loss and Pupil Abnormalities. Frontiers in Aging Neuroscience, 2016, 8, 222.	3.4	6
88	Screening for CCNF Mutations in a Chinese Amyotrophic Lateral Sclerosis Cohort. Frontiers in Aging Neuroscience, 2018, 10, 185.	3.4	6
89	Painful Diabetic Peripheral Neuropathy Study of Chinese OutPatiEnts (PDN-SCOPE): protocol for a multicentre cross-sectional registry study of clinical characteristics and treatment in China. BMJ Open, 2019, 9, e025722.	1.9	6
90	Clinical and Genetic Diversity of PMP22 Mutations in a Large Cohort of Chinese Patients With Charcot-Marie-Tooth Disease. Frontiers in Neurology, 2020, 11, 630.	2.4	6

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91	2020 expert consensus statement on neuro-protection after cardiac arrest in China. Annals of Translational Medicine, 2021, 9, 175-175.	1.7	6
92	Using corneal confocal microscopy to compare Mecobalamin intramuscular injections vs oral tablets in treating diabetic peripheral neuropathy: a RCT. Scientific Reports, 2021, 11, 14697.	3.3	6
93	Physical activity and amyotrophic lateral sclerosis: a Mendelian randomization study. Neurobiology of Aging, 2021, 105, 374.e1-374.e4.	3.1	6
94	Nuclear TAR DNA-binding protein 43: A new target for amyotrophic lateral sclerosis treatment. Neural Regeneration Research, 2013, 8, 3284-95.	3.0	6
95	Angiogenin gene polymorphism: A risk factor for diabetic peripheral neuropathy in the northern Chinese Han population. Neural Regeneration Research, 2013, 8, 3434-40.	3.0	6
96	Amyotrophic lateral sclerosis: new era, new challenges. Lancet Neurology, The, 2022, 21, 400-401.	10.2	6
97	A two-sample Mendelian randomization analysis of modifiable risk factors and intracranial aneurysms. Scientific Reports, 2022, 12, 7659.	3.3	6
98	Lipoprotein lipase deficiency leads to Î±-synuclein aggregation and ubiquitin C-terminal hydrolase L1 reduction. Neuroscience, 2015, 290, 1-10.	2.3	5
99	Twelve-month duration as an appropriate criterion for flail arm syndrome. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 29-33.	1.7	5
100	Screening for REEP1 Mutations in 31 Chinese Hereditary Spastic Paraplegia Families. Frontiers in Neurology, 2020, 11, 499.	2.4	5
101	Cross-Sectional Study in a Large Cohort of Chinese Patients With GJB1 Gene Mutations. Frontiers in Neurology, 2020, 11, 690.	2.4	5
102	Characteristics of Clinical and Electrophysiological Pattern in a Large Cohort of Chinese Patients With Charcot-Marie-Tooth 4C. Frontiers in Neurology, 2021, 12, 598168.	2.4	5
103	Chinese validation of the Raschâ€Built Overall Amyotrophic Lateral Sclerosis Disability Scale. European Journal of Neurology, 2021, 28, 1876-1883.	3.3	5
104	Effect of racial background on motor cortical function as measured by threshold tracking transcranial magnetic stimulation. Journal of Neurophysiology, 2021, 126, 840-844.	1.8	5
105	Validation of the pathogenic role of rare DNAJC7 variants in Chinese patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2021, 106, 314.e1-314.e6.	3.1	5
106	Multicentre, prospective registry study of amyotrophic lateral sclerosis in mainland China (CHALSR): study protocol. BMJ Open, 2020, 10, e042603.	1.9	5
107	Disease duration of progression is helpful in identifying isolated bulbar palsy of amyotrophic lateral sclerosis. BMC Neurology, 2021, 21, 405.	1.8	5
108	Madras pattern of motor neuron disease: improvement of symptoms with intravenous immunoglobulin. The National Medical Journal of India, 2004, 17, 141-2.	0.3	5

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109	Assessing the role of blood pressure in amyotrophic lateral sclerosis: a Mendelian randomization study. Orphanet Journal of Rare Diseases, 2022, 17, 56.	2.7	5
110	The single-nucleotide polymorphism rs6690993 in FGGY is not associated with amyotrophic lateral sclerosisin a large Chinese cohort. Neurobiology of Aging, 2014, 35, 1512.e3-1512.e4.	3.1	4
111	Kennedyâ€™s disease 1234 scale: Preliminary design and test. Journal of Clinical Neuroscience, 2017, 40, 185-189.	1.5	4
112	Autologous Bone Marrow-Derived Stem Cells for Treating Diabetic Neuropathy in Metabolic Syndrome. BioMed Research International, 2017, 2017, 1-6.	1.9	4
113	Discontinuation Rate of Newly Prescribed Donepezil in Alzheimer's Disease Patients in Asia. Journal of		

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127	Exercise Physiology Impairments of Patients With Amyotrophic Lateral Sclerosis: Cardiopulmonary Exercise Testing Findings. Frontiers in Physiology, 2022, 13, 792660.	2.8	3
128	Different electrophysiology patterns in GNE myopathy. Orphanet Journal of Rare Diseases, 2022, 17, 206.	2.7	3
129	Mutation analysis of the GLA gene in Chinese patients with intracerebral hemorrhage. Neurobiology of Aging, 2021, 102, 220.e1-220.e4.	3.1	2
130	hTBK1-c.978T>A mutation promotes the ferroptosis in NSC-34 cells via mediation of KEAP1/NRF2/p62 signaling. American Journal of Translational Research (discontinued), 2020, 12, 7386-7394.	0.0	2
131	Serum Neurofilament Light Chain Levels May Be a Marker of Lower Motor Neuron Damage in Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 2022, 13, 833507.	2.4	2
132	Corneal confocal microscopy in the evaluation of immune-related motor neuron disease syndrome. BMC Neurology, 2022, 22, 138.	1.8	2
133	Eye Movement Abnormalities in Amyotrophic Lateral Sclerosis. Brain Sciences, 2022, 12, 489.	2.3	2
134	Comparison of Slow and Forced Vital Capacity on Ability to Evaluate Respiratory Function in Bulbar-Involved Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 0, 13, .	2.4	2
135	The presence of spontaneous EMG activity in sternocleidomastoid is associated with ventilatory dysfunction in ALS. Neurophysiologie Clinique, 2016, 46, 145-148.	2.2	1
136	Metachromatic Leukodystrophy: Too Frequent (Mis)Diagnosis?â€”Reply. JAMA Neurology, 2018, 75, 1027.	9.0	1
137	Reply to â€œLife Course Adiposity and Amyotrophic Lateral Sclerosisâ€• Annals of Neurology, 2020, 88, 203-204.	5.3	1
138	Upper Motor Neuron Signs in the Cervical Region of Patients With Flail Arm Syndrome. Frontiers in Neurology, 2021, 12, 610786.	2.4	1
139	CT-Visible Convexity Subarachnoid Hemorrhage Predicts Early Recurrence of Lobar Hemorrhage. Frontiers in Neurology, 2022, 13, 843851.	2.4	1
140	Omics feature learning for cross individual ALS disease identification with EMG signal. , 2021, , .		1
141	Application Value of the Motor Unit Number Index in Patients With Kennedy Disease. Frontiers in Neurology, 2021, 12, 705816.	2.4	1
142	Analysis of ERBB4 Variants in Amyotrophic Lateral Sclerosis Within a Chinese Cohort. Frontiers in Neurology, 2022, 13, 865264.	2.4	1
143	SIRT1 Interacts with Prepro-Orexin in the Hypothalamus in SOD1G93A Mice. Brain Sciences, 2022, 12, 490.	2.3	1
144	Stratifying disease stages with different progression rates determined by electrophysiological tests in patients with amyotrophic lateral sclerosis (reply). Muscle and Nerve, 2009, 40, 319-319.	2.2	0

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145	The Important Role of Lipids in Cognitive Impairment. , 0, , 268-272.		0
146	The Important Role of Lipids in Cognitive Impairment. , 0, , 206-211.		0
147	Wavelet-based Multi-branch Convolutional Neural Network for Cross-individual ALS Disease Identification with EMG Signal. , 2021, , .		0