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	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
2	Hypermetabolism associated with worse prognosis of amyotrophic lateral sclerosis. Journal of Neurology, 2022, 269, 1447-1455.	3.6	19
3	An identical DCTN1 mutation in two Chinese siblings manifest as dHMN and ALS respectively: a case report. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, 23, 149-153.	1.7	3
4	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
5	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
6	The Clinical Features of In-Hospital Recurrence in Acute Ischaemic Stroke Patients over Time: A Real-World Observation at a Single Center. Brain Sciences, 2022, 12, 123.	2.3	3
7	Association between type 2 diabetes and amyotrophic lateral sclerosis. Scientific Reports, 2022, 12, 2544.	3.3	11
8	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
9	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
10	Dietary-Derived Essential Nutrients and Amyotrophic Lateral Sclerosis: A Two-Sample Mendelian Randomization Study. Nutrients, 2022, 14, 920.	4.1	10
11	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
12	CT-Visible Convexity Subarachnoid Hemorrhage Predicts Early Recurrence of Lobar Hemorrhage. Frontiers in Neurology, 2022, 13, 843851.	2.4	1
13	Exercise Physiology Impairments of Patients With Amyotrophic Lateral Sclerosis: Cardiopulmonary Exercise Testing Findings. Frontiers in Physiology, 2022, 13, 792660.	2.8	3
14	Amyotrophic lateral sclerosis: new era, new challenges. Lancet Neurology, The, 2022, 21, 400-401.	10.2	6
15	Corneal confocal microscopy in the evaluation of immune-related motor neuron disease syndrome. BMC Neurology, 2022, 22, 138.	1.8	2
16	Analysis of ERBB4 Variants in Amyotrophic Lateral Sclerosis Within a Chinese Cohort. Frontiers in Neurology, 2022, 13, 865264.	2.4	1
17	SIRT1 Interacts with Prepro-Orexin in the Hypothalamus in SOD1G93A Mice. Brain Sciences, 2022, 12, 490.	2.3	1
18	Eye Movement Abnormalities in Amyotrophic Lateral Sclerosis. Brain Sciences, 2022, 12, 489.	2.3	2

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19	A two-sample Mendelian randomization analysis of modifiable risk factors and intracranial aneurysms. Scientific Reports, 2022, 12, 7659.	3.3	6
20	Neuroimmune Crosstalk Between the Peripheral and the Central Immune System in Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2022, 14, 890958.	3.4	10
21	Different electrophysiology patterns in GNE myopathy. Orphanet Journal of Rare Diseases, 2022, 17, 206.	2.7	3
22	Lipids, Apolipoproteins, Statins, and Intracerebral Hemorrhage: A Mendelian Randomization Study. Annals of Neurology, 2022, 92, 390-399.	5.3	16
23	TBK1 variants in Chinese patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2021, 97, 149.e9-149.e15.	3.1	8
24	The complement C7 variant rs3792646 is associated with amyotrophic lateral sclerosis in a Han Chinese population. Neurobiology of Aging, 2021, 99, 103.e1-103.e7.	3.1	3
25	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
26	Discontinuation Rate of Newly Prescribed Donepezil in Alzheimer's Disease Patients in Asia. Journal of		

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37	Characterization of genotypeâ€“phenotype correlation with MORC2 mutated Axonal Charcotâ€“Marieâ€“Tooth disease in a cohort of Chinese patients. Orphanet Journal of Rare Diseases, 2021, 16, 244.	2.7	3
38	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
39	GLT8D1 may not be significant in Chinese sporadic amyotrophic lateral sclerosis patients. Neurobiology of Aging, 2021, 102, 224.e1-224.e3.	3.1	3
40	Mutation analysis of the GLA gene in Chinese patients with intracerebral hemorrhage. Neurobiology of Aging, 2021, 102, 220.e1-220.e4.	3.1	2
41	A Mendelian randomization analysis of the relationship between cardioembolic risk factors and ischemic stroke. Scientific Reports, 2021, 11, 14583.	3.3	4
42	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
43	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
44	Prognosis of amyotrophic lateral sclerosis with cognitive and behavioural changes based on a sixty-month longitudinal follow-up. PLoS ONE, 2021, 16, e0253279.	2.5	3
45	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
46	Physical activity and amyotrophic lateral sclerosis: a Mendelian randomization study. Neurobiology of Aging, 2021, 105, 374.e1-374.e4.	3.1	6
47	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
48	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
49	Disease duration of progression is helpful in identifying isolated bulbar palsy of amyotrophic lateral sclerosis. BMC Neurology, 2021, 21, 405.	1.8	5
50	The Genotype and Phenotype Features in a Large Chinese MFN2 Mutation Cohort. Frontiers in Neurology, 2021, 12, 757518.	2.4	7
51	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
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	Selective and Inverse U-Shaped Curve Alteration of the Retinal Nerve in Amyotrophic Lateral Sclerosis: A Potential Mirror of the Disease. Frontiers in Aging Neuroscience, 2021, 13, 783431.	3.4	4
54	MRI Volumetric Analysis of the Thalamus and Hypothalamus in Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2021, 13, 610332.	3.4	3

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55	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
56	Omics feature learning for cross individual ALS disease identification with EMG signal. , 2021, , .		1
57	Wavelet-based Multi-branch Convolutional Neural Network for Cross-individual ALS Disease Identification with EMG Signal. , 2021, , .		0
58	Leukocyte telomere length and amyotrophic lateral sclerosis: a Mendelian randomization study. Orphanet Journal of Rare Diseases, 2021, 16, 508.	2.7	7
59	Application Value of the Motor Unit Number Index in Patients With Kennedy Disease. Frontiers in Neurology, 2021, 12, 705816.	2.4	1
60	Twelve-month duration as an appropriate criterion for flail arm syndrome. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 29-33.	1.7	5
61	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
62	Life Course Adiposity and Amyotrophic Lateral Sclerosis: A Mendelian Randomization Study. Annals of Neurology, 2020, 87, 434-441.	5.3	30
63	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
64	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
65	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
66	Screening for REEP1 Mutations in 31 Chinese Hereditary Spastic Paraplegia Families. Frontiers in Neurology, 2020, 11, 499.	2.4	5
67	Acute Myelitis, Recurrent Optic Neuritis, and Seizures Over 17 Years. Frontiers in Neurology, 2020, 11, 541146.	2.4	3
68	Cross-Sectional Study in a Large Cohort of Chinese Patients With GJB1 Gene Mutations. Frontiers in Neurology, 2020, 11, 690.	2.4	5
69	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
70	Education, intelligence, and amyotrophic lateral sclerosis: A Mendelian randomization study. Annals of Clinical and Translational Neurology, 2020, 7, 1642-1647.	3.7	12
71	Reply to â€œLife Course Adiposity and Amyotrophic Lateral Sclerosisâ€• Annals of Neurology, 2020, 88, 203-204.	5.3	1
72	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

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73	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
74	Molecular analysis and clinical diversity of distal hereditary motor neuropathy. European Journal of Neurology, 2020, 27, 1319-1326.	3.3	28
75	Multicentre, prospective registry study of amyotrophic lateral sclerosis in mainland China (CHALSR): study protocol. BMJ Open, 2020, 10, e042603.	1.9	5
76	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
77	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
78	In-hospital recurrence in a Chinese large cohort with acute ischemic stroke. Scientific Reports, 2019, 9, 14945.	3.3	9
79	Improved precision of epigenetic clock estimates across tissues and its implication for biological ageing. Genome Medicine, 2019, 11, 54.	8.2	191
80	Better survival in female SOD1-mutant patients with ALS: a study of SOD1-related natural history. Translational Neurodegeneration, 2019, 8, 2.	8.0	22
81	Increased Interleukin-6 Levels in the Astrocyte-Derived Exosomes of Sporadic Amyotrophic Lateral Sclerosis Patients. Frontiers in Neuroscience, 2019, 13, 574.	2.8	61
82	Stroke in China: advances and challenges in epidemiology, prevention, and management. Lancet Neurology, The, 2019, 18, 394-405.	10.2	903
83	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
84	Vestibular evoked myogenic potentials and their clinical utility in patients with amyotrophic lateral sclerosis. Clinical Neurophysiology, 2019, 130, 647-654.	1.5	11
85	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
86	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
87	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
88	The epidemiology and genetics of Amyotrophic lateral sclerosis in China. Brain Research, 2018, 1693, 121-126.	2.2	36
89	Serial Magnetic Resonance Imaging Changes in a Patient With Late-Onset Cobalamin C Disease With a Misdiagnosis of Metachromatic Leukodystrophy. JAMA Neurology, 2018, 75, 374.	9.0	3
90	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

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91	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
92	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
93	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
94	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
95	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
96	Amyotrophic lateral sclerosis in Beijing: Epidemiologic features and prognosis from 2010 to 2015. Brain and Behavior, 2018, 8, e01131.	2.2	35
97	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
98	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
99	Screening for CCNF Mutations in a Chinese Amyotrophic Lateral Sclerosis Cohort. Frontiers in Aging Neuroscience, 2018, 10, 185.	3.4	6
100	Metachromatic Leukodystrophy: Too Frequent (Mis)Diagnosis?â€”Reply. JAMA Neurology, 2018, 75, 1027.	9.0	1
101	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
102	CHCHD10 mutations in patients with amyotrophic lateral sclerosis in Mainland China. Neurobiology of Aging, 2017, 54, 214.e7-214.e10.	3.1	12
103	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
104	Kennedyâ€™s disease 1234 scale: Preliminary design and test. Journal of Clinical Neuroscience, 2017, 40, 185-189.	1.5	4
105	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
106	Cross-ethnic meta-analysis identifies association of the GPX3-TNIP1 locus with amyotrophic lateral sclerosis. Nature Communications, 2017, 8, 611.	12.8	93
107	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
108	The Analysis of Two BDNF Polymorphisms G196A/C270T in Chinese Sporadic Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2017, 9, 135.	3.4	12

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109	Autologous Bone Marrow-Derived Stem Cells for Treating Diabetic Neuropathy in Metabolic Syndrome. BioMed Research International, 2017, 2017, 1-6.	1.9	4
110	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
111	Whole-exome sequencing in amyotrophic lateral sclerosis suggests NEK1 is a risk gene in Chinese. Genome Medicine, 2017, 9, 97.	8.2	23
112	DCTN1 gene analysis in Chinese patients with sporadic amyotrophic lateral sclerosis. PLoS ONE, 2017, 12, e0182572.	2.5	23
113	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
114	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
115	A Novel Missense Mutation of the DDHD1 Gene Associated with Juvenile Amyotrophic Lateral Sclerosis. Frontiers in Aging Neuroscience, 2016, 8, 291.	3.4	17
116	Enrichment of SNPs in Functional Categories Reveals Genes Affecting Complex Traits. Human Mutation, 2016, 37, 820-826.	2.5	3
117	Increased extrasynaptic GluN2B expression is involved in cognitive impairment after isoflurane anesthesia. Experimental and Therapeutic Medicine, 2016, 12, 161-168.	1.8	12
118	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
119	The presence of spontaneous EMG activity in sternocleidomastoid is associated with ventilatory dysfunction in ALS. Neurophysiologie Clinique, 2016, 46, 145-148.	2.2	1
120	MATR3 mutation analysis in a Chinese cohort with sporadic amyotrophic lateral sclerosis. Neurobiology of Aging, 2016, 38, 218.e3-218.e4.	3.1	27
121	The Edinburgh Cognitive and Behavioural ALS Screen in a Chinese Amyotrophic Lateral Sclerosis Population. PLoS ONE, 2016, 11, e0155496.	2.5	50
122	Lipoprotein lipase deficiency leads to Î±-synuclein aggregation and ubiquitin C-terminal hydrolase L1 reduction. Neuroscience, 2015, 290, 1-10.	2.3	5
123	Amyotrophic Lateral Sclerosis Genetic Studies. Neuroscientist, 2015, 21, 599-615.	3.5	23
124	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
125	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
126	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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127	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
128	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
129	Six SQSTM1 mutations in a Chinese amyotrophic lateral sclerosis cohort. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2015, 16, 378-384.	1.7	23
130	Optineurin mutations in patients with sporadic amyotrophic lateral sclerosis in China. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2015, 16, 485-489.	1.7	32
131	C9orf72 hexanucleotide repeat expansions in Chinese sporadic amyotrophic lateral sclerosis. Neurobiology of Aging, 2015, 36, 2660.e1-2660.e8.	3.1	50
132	Impaired synaptic vesicle recycling contributes to presynaptic dysfunction in lipoprotein lipase-deficient mice. Neuroscience, 2014, 280, 275-281.	2.3	7
133	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
134	Risk Score to Predict Hospital-Acquired Pneumonia After Spontaneous Intracerebral Hemorrhage. Stroke, 2014, 45, 2620-2628.	2.0	35
135	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
136	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
137	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
138	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
139	Nuclear TAR DNA-binding protein 43: A new target for amyotrophic lateral sclerosis treatment. Neural Regeneration Research, 2013, 8, 3284-95.	3.0	6
140	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
141	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
142	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
143	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
144	Adeno-associated virus vector-mediated triple gene transfer of dopamine synthetic enzymes. Chinese Medical Journal, 2001, 114, 1276-9.	2.3	3

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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	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