

Hui Fang Shang

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

250
papers

2,929
citations

29
h-index

39
g-index

286
ext. papers

4,098
ext. citations

4.6
avg, IF

5.27
L-index

#	Paper	IF	Citations
250	Magnetic Resonance Imaging Markers for Cognitive Impairment in Parkinson's Disease: Current View.. <i>Frontiers in Aging Neuroscience</i> , 2022 , 14, 788846	5.3	0
249	High neutrophil-to-lymphocyte ratio predicts short survival in multiple system atrophy.. <i>Npj Parkinson's Disease</i> , 2022 , 8, 11	9.7	2
248	Voxel-based meta-analysis of gray matter abnormalities in idiopathic dystonia.. <i>Journal of Neurology</i> , 2022 , 1	5.5	1
247	ANXA1 and the risk for early-onset Parkinson's disease.. <i>Neurobiology of Aging</i> , 2022 , 112, 212-214	5.6	0
246	Self-Stigma in Parkinson's Disease: A 3-Year Prospective Cohort Study.. <i>Frontiers in Aging Neuroscience</i> , 2022 , 14, 790897	5.3	0
245	Enrichment of rare variants of BIN1 but not APOE genes in Chinese patients with Parkinson's disease.. <i>Journal of Internal Medicine</i> , 2022 ,	10.8	
244	Neutrophil-to-lymphocyte ratio in sporadic amyotrophic lateral sclerosis. <i>Neural Regeneration Research</i> , 2022 , 17, 875-880	4.5	5
243	Genetic heterogeneity on sleep disorders in Parkinson's disease: a systematic review and meta-analysis.. <i>Translational Neurodegeneration</i> , 2022 , 11, 21	10.3	1
242	The expression discrepancy and characteristics of long non-coding RNAs in peripheral blood leukocytes from amyotrophic lateral sclerosis patients.. <i>Molecular Neurobiology</i> , 2022 , 1	6.2	1
241	De Novo Missense Mutation of in a Chinese Patient with Generalized Dystonia with Myoclonus.. <i>Movement Disorders Clinical Practice</i> , 2022 , 9, 551-552	2.2	
240	Shared genetic links between frontotemporal dementia and psychiatric disorders.. <i>BMC Medicine</i> , 2022 , 20, 131	11.4	0
239	Impaired Topographic Organization in Patients With Idiopathic Blepharospasm.. <i>Frontiers in Neurology</i> , 2021 , 12, 708634	4.1	0
238	Changes in Serum Cystatin C Levels and the Associations With Cognitive Function in Alzheimer's Disease Patients.. <i>Frontiers in Aging Neuroscience</i> , 2021 , 13, 790939	5.3	0
237	Rare Variants in Chinese Patients With Amyotrophic Lateral Sclerosis. <i>Frontiers in Genetics</i> , 2021 , 12, 740052	4.5	1
236	Association Analysis of , and Polymorphisms in Chinese Patients With Parkinson's Disease and Multiple System Atrophy. <i>Frontiers in Genetics</i> , 2021 , 12, 765833	4.5	0
235	Carnitine and COVID-19 Susceptibility and Severity: A Mendelian Randomization Study.. <i>Frontiers in Nutrition</i> , 2021 , 8, 780205	6.2	2
234	Different Associated Factors of Subjective Cognitive Complaints in Patients With Early- and Late-Onset Parkinson's Disease. <i>Frontiers in Neurology</i> , 2021 , 12, 749471	4.1	

233	Predictors of survival in patients with amyotrophic lateral sclerosis: A large meta-analysis. <i>EBioMedicine</i> , 2021 , 74, 103732	8.8	3
232	Prevalence and associated factors of malnutrition in patients with Parkinson's disease using CONUT and GNRI. <i>Parkinsonism and Related Disorders</i> , 2021 ,	3.6	2
231	Progression of Fatigue in Early Parkinson's Disease: A 3-Year Prospective Cohort Study. <i>Frontiers in Aging Neuroscience</i> , 2021 , 13, 701906	5.3	0
230	Vascular Risk Factors and Cognition in Multiple System Atrophy. <i>Frontiers in Neuroscience</i> , 2021 , 15, 749949	5.1	0
229	Comprehensive Analysis of LIN28A in Chinese Patients With Early Onset Parkinson's Disease. <i>Frontiers in Genetics</i> , 2021 , 12, 740096	4.5	1
228	Neurofilament Light Chain Predicts Disease Severity and Progression in Multiple System Atrophy. <i>Movement Disorders</i> , 2021 ,	7	1
227	Effect of diabetes control status on the progression of Parkinson's disease: A prospective study. <i>Annals of Clinical and Translational Neurology</i> , 2021 , 8, 887-897	5.3	2
226	Pathological laughter and crying in multiple system atrophy with different subtypes: Frequency and related factors. <i>Journal of Affective Disorders</i> , 2021 , 283, 60-65	6.6	1
225	Sarcopenia in Patients With Parkinson's Disease: A Systematic Review and Meta-Analysis. <i>Frontiers in Neurology</i> , 2021 , 12, 598035	4.1	5
224	Genetic Analysis of ZNF Protein Family Members for Early-Onset Parkinson's Disease in Chinese Population. <i>Molecular Neurobiology</i> , 2021 , 58, 3435-3442	6.2	6
223	Rheumatoid arthritis decreases risk for Parkinson's disease: a Mendelian randomization study. <i>Npj Parkinson's Disease</i> , 2021 , 7, 17	9.7	4
222	Mutation analysis of seven SLC family transporters for early-onset Parkinson's disease in Chinese population. <i>Neurobiology of Aging</i> , 2021 , 103, 152.e1-152.e6	5.6	1
221	Excessive daytime sleepiness in Parkinson's disease: A systematic review and meta-analysis. <i>Parkinsonism and Related Disorders</i> , 2021 , 85, 133-140	3.6	5
220	Genetic Modifiers of Age at Onset for Parkinson's Disease in Asians: A Genome-Wide Association Study. <i>Movement Disorders</i> , 2021 , 36, 2077-2084	7	1
219	Evidence for Peripheral Immune Activation in Parkinson's Disease. <i>Frontiers in Aging Neuroscience</i> , 2021 , 13, 617370	5.3	2
218	Camptocormia in patients with multiple system atrophy at different disease durations: frequency and related factors. <i>BMC Neurology</i> , 2021 , 21, 181	3.1	1
217	Prevalence and profile of nocturnal disturbances in Chinese patients with advanced-stage Parkinson's disease: a cross-sectional epidemiology study. <i>BMC Neurology</i> , 2021 , 21, 194	3.1	0
216	Mutation screening and burden analysis of GLT8D1 in Chinese patients with amyotrophic lateral sclerosis. <i>Neurobiology of Aging</i> , 2021 , 101, 298.e17-298.e21	5.6	2

215	A case of juvenile-onset amyotrophic lateral sclerosis with a frameshift gene mutation presenting with bilateral abducens palsy. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2021 , 1-2	3.6	1
214	Early weight instability is associated with cognitive decline and poor survival in amyotrophic lateral sclerosis. <i>Brain Research Bulletin</i> , 2021 , 171, 10-15	3.9	2
213	Shared genetic links between amyotrophic lateral sclerosis and obesity-related traits: a genome-wide association study. <i>Neurobiology of Aging</i> , 2021 , 102, 211.e1-211.e9	5.6	1
212	Rare Variants Analysis of Lysosomal Related Genes in Early-Onset and Familial Parkinson's Disease in a Chinese Cohort. <i>Journal of Parkinson's Disease</i> , 2021 , 11, 1845-1855	5.3	0
211	Mutation analysis of LRP10 in a large Chinese familial Parkinson disease cohort. <i>Neurobiology of Aging</i> , 2021 , 99, 99.e1-99.e6	5.6	2
210	Mutation analysis of TMEM family members for early-onset Parkinson's disease in Chinese population. <i>Neurobiology of Aging</i> , 2021 , 101, 299.e1-299.e6	5.6	2
209	Genetic Analysis of Prosaposin, the Lysosomal Storage Disorder Gene in Parkinson's Disease. <i>Molecular Neurobiology</i> , 2021 , 58, 1583-1592	6.2	5
208	A novel nonsense mutation in the TYMP gene causing MNGIE with multiple intracranial hemorrhages on brain MRI. <i>Neurological Sciences</i> , 2021 , 42, 2119-2122	3.5	1
207	SQSTM1 variant that is linked to sporadic ALS exhibits impaired association with MAP1LC3 in cultured cells. <i>ENeurologicalSci</i> , 2021 , 22, 100301	2.1	3
206	Suicidal ideation in early-onset Parkinson's disease. <i>Journal of Neurology</i> , 2021 , 268, 1876-1884	5.5	2
205	Aberrations of biochemical indicators in amyotrophic lateral sclerosis: a systematic review and meta-analysis. <i>Translational Neurodegeneration</i> , 2021 , 10, 3	10.3	7
204	Creatine kinase in the diagnosis and prognostic prediction of amyotrophic lateral sclerosis: a retrospective case-control study. <i>Neural Regeneration Research</i> , 2021 , 16, 591-595	4.5	4
203	Genome-wide genetic links between amyotrophic lateral sclerosis and autoimmune diseases. <i>BMC Medicine</i> , 2021 , 19, 27	11.4	7
202	Facial tremor in patients with Parkinson's disease: prevalence, determinants and impacts on disease progression. <i>BMC Neurology</i> , 2021 , 21, 86	3.1	3
201	Risk factors for cognitive impairment in amyotrophic lateral sclerosis: a systematic review and meta-analysis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2021 , 92, 688-693	5.5	1
200	Brain Calcification in a Young Adult with Abnormal Copper Metabolism. <i>Movement Disorders Clinical Practice</i> , 2021 , 8, 476-479	2.2	
199	Longitudinal evolution of non-motor symptoms in early Parkinson's disease: a 3-year prospective cohort study. <i>Npj Parkinson's Disease</i> , 2021 , 7, 58	9.7	1
198	Multivariable clinical-genetic model for predicting dyskinesia in early-onset Parkinson's disease. <i>Translational Neurodegeneration</i> , 2021 , 10, 26	10.3	1

197	Health-related quality of life in amyotrophic lateral sclerosis using EQ-5D-5L. <i>Health and Quality of Life Outcomes</i> , 2021 , 19, 181	3	0
196	Different resting-state network disruptions in newly diagnosed drug-naïve Parkinson's disease patients with mild cognitive impairment. <i>BMC Neurology</i> , 2021 , 21, 327	3.1	1
195	Excessive daytime sleepiness in idiopathic blepharospasm. <i>Parkinsonism and Related Disorders</i> , 2021 , 89, 134-138	3.6	
194	Sex related differences in nonmotor symptoms of patients with idiopathic blepharospasm. <i>Scientific Reports</i> , 2021 , 11, 17856	4.9	0
193	Disruption of the white matter structural network and its correlation with baseline progression rate in patients with sporadic amyotrophic lateral sclerosis. <i>Translational Neurodegeneration</i> , 2021 , 10, 35	10.3	2
192	Role of genetics in amyotrophic lateral sclerosis: a large cohort study in Chinese mainland population. <i>Journal of Medical Genetics</i> , 2021 ,	5.8	3
191	Genetic analysis of TRIM family genes for early-onset Parkinson's disease in Chinese population. <i>Parkinsonism and Related Disorders</i> , 2021 , 90, 105-113	3.6	0
190	Prevalence and Factors Related to Pathological Laughter and Crying in Patients With Amyotrophic Lateral Sclerosis. <i>Frontiers in Neurology</i> , 2021 , 12, 655674	4.1	
189	Abnormal eye movements in spinocerebellar ataxia type 3. <i>BMC Neurology</i> , 2021 , 21, 28	3.1	1
188	Overlapping Genetic Architecture Between Schizophrenia and Neurodegenerative Disorders.. <i>Frontiers in Cell and Developmental Biology</i> , 2021 , 9, 797072	5.7	0
187	Role of lipoic acid in multiple sclerosis.. <i>CNS Neuroscience and Therapeutics</i> , 2021 ,	6.8	3
186	The Cold Hand Sign in Multiple System Atrophy: Frequency-Associated Factors and Its Impact on Survival.. <i>Frontiers in Aging Neuroscience</i> , 2021 , 13, 767211	5.3	0
185	Association between positive history of essential tremor and disease progression in patients with Parkinson's disease. <i>Scientific Reports</i> , 2020 , 10, 21749	4.9	3
184	Analysis of Chinese patients with sporadic Creutzfeldt-Jakob disease. <i>Prion</i> , 2020 , 14, 137-142	2.3	3
183	Serum creatinine levels in patients with amyotrophic lateral sclerosis: a systematic review and meta-analysis. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2020 , 21, 502-508	3.6	2
182	Abnormal Serum Iron-Status Indicator Changes in Amyotrophic Lateral Sclerosis (ALS) Patients: A Meta-Analysis. <i>Frontiers in Neurology</i> , 2020 , 11, 380	4.1	5
181	Mutation screening and burden analysis of VPS13C in Chinese patients with early-onset Parkinson's disease. <i>Neurobiology of Aging</i> , 2020 , 94, 311.e1-311.e4	5.6	9
180	MicroRNA-183-5p is stress-inducible and protects neurons against cell death in amyotrophic lateral sclerosis. <i>Journal of Cellular and Molecular Medicine</i> , 2020 , 24, 8614-8622	5.6	8

179	Delivering patient-centered care in Parkinson's disease: Challenges and consensus from an international panel. <i>Parkinsonism and Related Disorders</i> , 2020 , 72, 82-87	3.6	13
178	Effects of Higher Serum Lipid Levels on the Risk of Parkinson's Disease: A Systematic Review and Meta-Analysis. <i>Frontiers in Neurology</i> , 2020 , 11, 597	4.1	5
177	The role of genetics in Parkinson's disease: a large cohort study in Chinese mainland population. <i>Brain</i> , 2020 , 143, 2220-2234	11.2	31
176	Reference values for the motor unit number index and the motor unit size index in five muscles. <i>Muscle and Nerve</i> , 2020 , 61, 657-661	3.4	2
175	Primary disruption of the default mode network subsystems in drug-naïve Parkinson's disease with mild cognitive impairments. <i>Neuroradiology</i> , 2020 , 62, 685-692	3.2	8
174	Impaired topographic organization in Parkinson's disease with mild cognitive impairment. <i>Journal of the Neurological Sciences</i> , 2020 , 414, 116861	3.2	8
173	Altered oligomeric states in pathogenic ALS2 variants associated with juvenile motor neuron diseases cause loss of ALS2-mediated endosomal function. <i>FASEB Journal</i> , 2020 , 34, 1-1	0.9	
172	Patterns of brain regional functional coherence in cognitive impaired ALS. <i>International Journal of Neuroscience</i> , 2020 , 130, 751-758	2	4
171	Chinese families with autosomal recessive hereditary spastic paraplegia caused by mutations in SPG11. <i>BMC Neurology</i> , 2020 , 20, 2	3.1	1
170	Grey matter abnormalities in Parkinson's disease: a voxel-wise meta-analysis. <i>European Journal of Neurology</i> , 2020 , 27, 653-659	6	11
169	Contribution of Five Functional Loci of Dopamine Metabolism-Related Genes to Parkinson's Disease and Multiple System Atrophy in a Chinese Population. <i>Frontiers in Neuroscience</i> , 2020 , 14, 889	5.1	0
168	ATP10B and the Risk for Early-Onset Parkinson's Disease. <i>Movement Disorders</i> , 2020 , 35, 2359-2360	7	3
167	Mutation Analysis of DNAJC Family for Early-Onset Parkinson's Disease in a Chinese Cohort. <i>Movement Disorders</i> , 2020 , 35, 2068-2076	7	8
166	Evaluating the Role of SNCA, LRRK2, and GBA in Chinese Patients With Early-Onset Parkinson's Disease. <i>Movement Disorders</i> , 2020 , 35, 2046-2055	7	13
165	Voxel-Based Meta-Analysis of Gray Matter Abnormalities in Multiple System Atrophy. <i>Frontiers in Aging Neuroscience</i> , 2020 , 12, 591666	5.3	4
164	Lipid Profile in Patients With Amyotrophic Lateral Sclerosis: A Systematic Review and Meta-Analysis. <i>Frontiers in Neurology</i> , 2020 , 11, 567753	4.1	3
163	Causal Association of Leukocytes Count and Amyotrophic Lateral Sclerosis: a Mendelian Randomization Study. <i>Molecular Neurobiology</i> , 2020 , 57, 4622-4627	6.2	5
162	Suicidal and death ideation in patients with progressive supranuclear palsy and corticobasal syndrome. <i>Journal of Affective Disorders</i> , 2020 , 276, 1061-1068	6.6	3

161	Elevated Percentage of CD3 T-Cells and CD4/CD8 Ratios in Multiple System Atrophy Patients. <i>Frontiers in Neurology</i> , 2020 , 11, 658	4.1	3
160	Clinical Reasoning: A 24-year-old man with head tremor and decreased ceruloplasmin level. <i>Neurology</i> , 2020 , 95, e1906-e1910	6.5	
159	Anti-N-methyl-D-Aspartate Receptor Encephalitis Mimicking Sporadic Creutzfeldt-Jakob Disease. <i>Frontiers in Neurology</i> , 2020 , 11, 593680	4.1	0
158	LanCL1 promotes motor neuron survival and extends the lifespan of amyotrophic lateral sclerosis mice. <i>Cell Death and Differentiation</i> , 2020 , 27, 1369-1382	12.7	14
157	Identification of Risk Loci for Parkinson Disease in Asians and Comparison of Risk Between Asians and Europeans: A Genome-Wide Association Study. <i>JAMA Neurology</i> , 2020 , 77, 746-754	17.2	84
156	Evolution of Apathy in Early Parkinson Disease: A 4-Years Prospective Cohort Study. <i>Frontiers in Aging Neuroscience</i> , 2020 , 12, 620762	5.3	4
155	Establish a Nomogram to Predict Falls in Spinocerebellar Ataxia Type 3. <i>Frontiers in Neurology</i> , 2020 , 11, 602003	4.1	1
154	Decreased Glycogenolysis by Promotes Regional Glycogen Accumulation Within the Spinal Cord of Amyotrophic Lateral Sclerosis Mice. <i>Frontiers in Molecular Neuroscience</i> , 2019 , 12, 114	6.1	13
153	Parkinson disease in the Western Pacific Region. <i>Lancet Neurology</i> , 2019 , 18, 865-879	24.1	55
152	Neurophysiological index is associated with the survival of patients with amyotrophic lateral sclerosis. <i>Clinical Neurophysiology</i> , 2019 , 130, 1730-1733	4.3	4
151	Extra-Cerebellar Signs and Non-motor Features in Chinese Patients With Spinocerebellar Ataxia Type 3. <i>Frontiers in Neurology</i> , 2019 , 10, 110	4.1	11
150	Unique characteristics of the genetics epidemiology of amyotrophic lateral sclerosis in China. <i>Science China Life Sciences</i> , 2019 , 62, 517-525	8.5	12
149	Prevalence of and factors associated with postural deformities in Chinese patients with multiple system atrophy. <i>Parkinsonism and Related Disorders</i> , 2019 , 64, 324-327	3.6	2
148	Frontal lobe function, behavioral changes and quality of life in patients with multiple system atrophy. <i>Restorative Neurology and Neuroscience</i> , 2019 , 37, 11-19	2.8	2
147	Mutation screening of the gene in Chinese patients with amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019 , 90, 245-246	5.5	4
146	Pisa Syndrome in Chinese Patients With Parkinson Disease. <i>Frontiers in Neurology</i> , 2019 , 10, 651	4.1	3
145	Predictors of Pisa syndrome in Chinese patients with Parkinson disease: A prospective study. <i>Parkinsonism and Related Disorders</i> , 2019 , 69, 1-6	3.6	1
144	RNM-01 Weight stability is associated with longer survival in amyotrophic lateral sclerosis. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2019 , 20, 309-326	3.6	0

143	Clinical disease stage related changes of serological factors in amyotrophic lateral sclerosis. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2019 , 20, 53-60	3.6	12
142	Identification of and as susceptibility genes for amyotrophic lateral sclerosis. <i>Neurology: Genetics</i> , 2019 , 5, e375	3.8	5
141	Rapid eye movement behavior disorder in drug-naïve patients with Parkinson's disease. <i>Journal of Clinical Neuroscience</i> , 2019 , 59, 254-258	2.2	14
140	Executive dysfunctions and behavioral changes in early drug-naïve patients with Parkinson's disease. <i>Journal of Affective Disorders</i> , 2019 , 243, 525-530	6.6	2
139	Predictors of camptocormia in patients with Parkinson's disease: A prospective study from southwest China. <i>Parkinsonism and Related Disorders</i> , 2018 , 52, 69-75	3.6	7
138	No association between 5 new GWAS-linked loci in Parkinson's disease and multiple system atrophy in a Chinese population. <i>Neurobiology of Aging</i> , 2018 , 67, 202.e7-202.e8	5.6	7
137	Altered white matter microarchitecture in amyotrophic lateral sclerosis: A voxel-based meta-analysis of diffusion tensor imaging. <i>NeuroImage: Clinical</i> , 2018 , 19, 122-129	5.3	24
136	Impact of sleep-related breathing disorder on motor and non-motor symptoms in multiple system atrophy. <i>Sleep and Breathing</i> , 2018 , 22, 981-987	3.1	1
135	Vascular risk factors and depression in Parkinson's disease. <i>European Journal of Neurology</i> , 2018 , 25, 637-643	6	7
134	An association study between SCFD1 rs10139154 variant and amyotrophic lateral sclerosis in a Chinese cohort. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2018 , 19, 413-418	3.6	3
133	Predictors of freezing of gait in Chinese patients with Parkinson's disease. <i>Brain and Behavior</i> , 2018 , 8, e00931	3.4	21
132	Clinical characteristics and quality of life in Chinese patients with Parkinson's disease beyond 20 years. <i>Neurological Research</i> , 2018 , 40, 312-317	2.7	2
131	Altered intrinsic brain functional connectivity in drug-naïve Parkinson's disease patients with LRRK2 mutations. <i>Neuroscience Letters</i> , 2018 , 675, 145-151	3.3	8
130	Determining the Effect of the HNMT, STK39, and NMD3 Polymorphisms on the Incidence of Parkinson's Disease, Amyotrophic Lateral Sclerosis, and Multiple System Atrophy in Chinese Populations. <i>Journal of Molecular Neuroscience</i> , 2018 , 64, 574-580	3.3	5
129	TMEM230 Mutations Are Rare in Han Chinese Patients with Autosomal Dominant Parkinson's Disease. <i>Molecular Neurobiology</i> , 2018 , 55, 2851-2855	6.2	7
128	Systemic overexpression of SQSTM1/p62 accelerates disease onset in a SOD1-expressing ALS mouse model. <i>Molecular Brain</i> , 2018 , 11, 30	4.5	23
127	Functional Variant rs3135500 in Increases the Risk of Multiple System Atrophy in a Chinese Population. <i>Frontiers in Aging Neuroscience</i> , 2018 , 10, 150	5.3	4
126	Clinical Staging of Amyotrophic Lateral Sclerosis in Chinese Patients. <i>Frontiers in Neurology</i> , 2018 , 9, 4424.1		8

125	A Voxel-Wise Meta-Analysis of Gray Matter Abnormalities in Essential Tremor. <i>Frontiers in Neurology</i> , 2018 , 9, 495	4.1	14
124	Survival analysis and prognostic nomogram model for multiple system atrophy. <i>Parkinsonism and Related Disorders</i> , 2018 , 54, 68-73	3.6	7
123	The relationship between four GWAS-identified loci in Alzheimer's disease and the risk of Parkinson's disease, amyotrophic lateral sclerosis, and multiple system atrophy. <i>Neuroscience Letters</i> , 2018 , 686, 205-210	3.3	4
122	Analysis of GWAS-linked variants in multiple system atrophy. <i>Neurobiology of Aging</i> , 2018 , 67, 201.e1-201.e4	5.64	10
121	Association analysis of SNP rs11868035 in SREBF1 with sporadic Parkinson's disease, sporadic amyotrophic lateral sclerosis and multiple system atrophy in a Chinese population. <i>Neuroscience Letters</i> , 2018 , 664, 128-132	3.3	5
120	Management Recommendations on Sleep Disturbance of Patients with Parkinson's Disease. <i>Chinese Medical Journal</i> , 2018 , 131, 2976-2985	2.9	13
119	Gender and onset age related-differences of non-motor symptoms and quality of life in drug-naïve Parkinson's disease. <i>Clinical Neurology and Neurosurgery</i> , 2018 , 175, 124-129	2	13
118	Association Between Serum Vitamin D Levels and Parkinson's Disease: A Systematic Review and Meta-Analysis. <i>Frontiers in Neurology</i> , 2018 , 9, 909	4.1	35
117	Prognostic Nomogram Associated with Longer Survival in Amyotrophic Lateral Sclerosis Patients 2018 , 9, 965-975		6
116	Resting-state network connectivity in cognitively unimpaired drug-naïve patients with rigidity-dominant Parkinson's disease. <i>Journal of the Neurological Sciences</i> , 2018 , 395, 147-152	3.2	10
115	Patterns of striatal and cerebellar functional connectivity in early-stage drug-naïve patients with Parkinson's disease subtypes. <i>Neuroradiology</i> , 2018 , 60, 1323-1333	3.2	11
114	Causes of Death in Chinese Patients with Multiple System Atrophy 2018 , 9, 102-108		16
113	Altered oligomeric states in pathogenic ALS2 variants associated with juvenile motor neuron diseases cause loss of ALS2-mediated endosomal function. <i>Journal of Biological Chemistry</i> , 2018 , 293, 17135-17153	5.4	14
112	Coding mutations in contribute to Parkinson's disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018 , 115, 11567-11572	11.5	50
111	Clinical and prognostic features of ALS/MND in different phenotypes-data from a hospital-based registry. <i>Brain Research Bulletin</i> , 2018 , 142, 403-408	3.9	5
110	Isaacs syndrome associated with GABA and AChR antibodies in sarcomatoid carcinoma. <i>Neurology</i> , 2018 , 91, 663-665	6.5	3
109	Voxel-based meta-analysis of gray and white matter volume abnormalities in spinocerebellar ataxia type 2. <i>Brain and Behavior</i> , 2018 , 8, e01099	3.4	6
108	Mutation screening of the TIA1 gene in Chinese patients with amyotrophic lateral sclerosis/frontotemporal dementia. <i>Neurobiology of Aging</i> , 2018 , 68, 161.e1-161.e3	5.6	4

107	Impaired topographic organization in cognitively unimpaired drug-naïve patients with rigidity-dominant Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2018 , 56, 52-57	3.6	17
106	Mutation Screening of the CHCHD10 Gene in Chinese Patients with Amyotrophic Lateral Sclerosis. <i>Molecular Neurobiology</i> , 2017 , 54, 3189-3194	6.2	14
105	Nonmotor symptoms in primary adult-onset cervical dystonia and blepharospasm. <i>Brain and Behavior</i> , 2017 , 7, e00592	3.4	32
104	Serum uric acid levels and freezing of gait in Parkinson's disease. <i>Neurological Sciences</i> , 2017 , 38, 955-960	3.5	10
103	Ubiquitin-related network underlain by (CAG) _n loci modulate age at onset in Machado-Joseph disease. <i>Brain</i> , 2017 , 140, e25	11.2	7
102	Association analysis of polymorphisms in VMAT2 and TMEM106B genes for Parkinson's disease, amyotrophic lateral sclerosis and multiple system atrophy. <i>Journal of the Neurological Sciences</i> , 2017 , 377, 65-71	3.2	5
101	White matter microstructure damage in tremor-dominant Parkinson's disease patients. <i>Neuroradiology</i> , 2017 , 59, 691-698	3.2	18
100	Corrigendum to "Rotigotine transdermal patch in Chinese patients with early Parkinson's disease: A randomized, double-blind, placebo-controlled pivotal study" [Park. Relat. Disord. 28 (2016) 49-55]. <i>Parkinsonism and Related Disorders</i> , 2017 , 38, 108	3.6	
99	Analysis of SOD1 mutations in a Chinese population with amyotrophic lateral sclerosis: a case-control study and literature review. <i>Scientific Reports</i> , 2017 , 7, 44606	4.9	16
98	Non-motor symptoms and the quality of life in multiple system atrophy with different subtypes. <i>Parkinsonism and Related Disorders</i> , 2017 , 35, 63-68	3.6	27
97	Encephalitis with antibodies against the GABA receptor: seizures as the most common presentation at admission. <i>Neurological Research</i> , 2017 , 39, 973-980	2.7	24
96	Blood hemoglobin A1c levels and amyotrophic lateral sclerosis survival. <i>Molecular Neurodegeneration</i> , 2017 , 12, 69	19	20
95	Tacrolimus in the treatment of myasthenia gravis in patients with an inadequate response to glucocorticoid therapy: randomized, double-blind, placebo-controlled study conducted in China. <i>Therapeutic Advances in Neurological Disorders</i> , 2017 , 10, 315-325	6.6	22
94	Apathy in drug-naïve patients with Parkinson's disease. <i>Parkinsonism and Related Disorders</i> , 2017 , 44, 28-32	3.6	9
93	Default-mode network connectivity in cognitively unimpaired drug-naïve patients with rigidity-dominant Parkinson's disease. <i>Journal of Neurology</i> , 2017 , 264, 152-160	5.5	23
92	Patients' self-perceived burden, caregivers' burden and quality of life for amyotrophic lateral sclerosis patients: a cross-sectional study. <i>Journal of Clinical Nursing</i> , 2017 , 26, 3188-3199	3.2	19
91	Teaching Neuro: Reversible brain MRI lesions in adult-onset type II citrullinemia. <i>Neurology</i> , 2017 , 89, e115	6.5	1
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