Ji-Feng Guo

List of Publications by Year
in descending order

151 papers	3,678 citations	136950 32 h-index	189892 50 g-index
166 all docs	166 docs citations	166 times ranked	4637 citing authors

#	Article	IF	Citations
1	Performance Comparison of Computational Methods for the Prediction of the Function and Pathogenicity of Non-Coding Variants. Genomics, Proteomics and Bioinformatics, 2023, 21, 649-661.	6.9	7
2	Evaluating the role of ARSA in Chinese patients with Parkinson's disease. Neurobiology of Aging, 2022, 109, 269-272.	3.1	5
3	Low-frequency and rare coding variants of NUS1 contribute to susceptibility and phenotype of Parkinson's disease. Neurobiology of Aging, 2022, 110, 106-112.	3.1	2
4	The Phenotypes and Mechanisms of NOTCH2NLC-Related GGC Repeat Expansion Disorders: a Comprehensive Review. Molecular Neurobiology, 2022, 59, 523-534.	4.0	27
5	PINK1 kinase dysfunction triggers neurodegeneration in the primate brain without impacting mitochondrial homeostasis. Protein and Cell, 2022, 13, 26-46.	11.0	32
6	Retinal Microvascular Density Was Associated With the Clinical Progression of Parkinsonâ€™s Disease. Frontiers in Aging Neuroscience, 2022, 14, 818597.	3.4	10
7	Evaluation of common and rare variants of Alzheimer's disease-causal genes in Parkinson's disease. Parkinsonism and Related Disorders, 2022, 97, 8-14.	2.2	4
8	Association Study of TAF1 Variants in Parkinsonâ€™s Disease. Frontiers in Neuroscience, 2022, 16, 846095.	2.8	2
9	Genetic landscape of human mitochondrial genome using whole-genome sequencing. Human Molecular Genetics, 2022, 31, 1747-1761.	2.9	4
10	PSEN1 G417S mutation in a Chinese pedigree causing early-onset parkinsonism with cognitive impairment. Neurobiology of Aging, 2022, 115, 70-76.	3.1	0
11	The association between LIN28A gene rare variants and Parkinsonâ€™s disease in Chinese population. Gene, 2022, 829, 146515.	2.2	2
12	Excimer laser atherectomy combined with drug-coated balloon angioplasty for the treatment of femoropopliteal arteriosclerosis obliterans. Annals of the Royal College of Surgeons of England, 2022, 104, 667-672.	0.6	1
13	Genetic Analysis of Patients With Early-Onset Parkinsonâ€™s Disease in Eastern China. Frontiers in Aging Neuroscience, 2022, 14, .	3.4	6
14	Assessment of <scp>GGC</scp> Repeat Expansion in <scp>GIPC1</scp> in Patients with Parkinson's Disease. Movement Disorders, 2022, 37, 1557-1559.	3.9	8
15	Profiling the Genome-Wide Landscape of Short Tandem Repeats by Long-Read Sequencing. Frontiers in Genetics, 2022, 13, .	2.3	4
16	The Chinese Parkinson's Disease Registry (<scp>CPDR</scp>): Study Design and Baseline Patient Characteristics. Movement Disorders, 2022, 37, 1335-1345.	3.9	13
17	The macular inner plexiform layer thickness as an early diagnostic indicator for Parkinsonâ€™s disease. Npj Parkinson's Disease, 2022, 8, .	5.3	5
18	Genetic Analysis of HSP40/DNAJ Family Genes in Parkinsonâ€™s Disease: a Large Case-Control Study. Molecular Neurobiology, 2022, 59, 5443-5451.	4.0	2

#	Article	IF	Citations
19	Deficiency in endocannabinoid synthase DAGLB contributes to early onset Parkinsonism and murine nigral dopaminergic neuron dysfunction. Nature Communications, 2022, 13, .	12.8	22
20	Biallelic loss-of-function variants in NEMF cause central nervous system impairment and axonal polyneuropathy. Human Genetics, 2021, 140, 579-592.	3.8	14
21	No relationship between SRY variants and risk of Parkinson's disease in Chinese population. Neurobiology of Aging, 2021, 100, 119.e3-119.e6.	3.1	5
22	Rare variant analysis of essential tremorâ€associated genes in earlyâ€onset Parkinsonâ€™s disease. Annals of Clinical and Translational Neurology, 2021, 8, 119-125.	3.7	4
23	Assessment of the association between NUS1 variants and essential tremor. Neuroscience Letters, 2021, 740, 135441.	2.1	3
24	Reply: Assessing the NOTCH2NLC GGC expansion in essential tremor patients from eastern China. Brain, 2021, 144, e2-e2.	7.6	0
25	Association of rare heterozygous PLA2G6 variants with the risk of Parkinson's disease. Neurobiology of Aging, 2021, 101, 297.e5-297.e8.	3.1	6
26	GPCards: An integrated database of genotypeâ€“phenotype correlations in human genetic diseases. Computational and Structural Biotechnology Journal, 2021, 19, 1603-1611.	4.1	5
27	Preliminary Study of hsa-mir-626 Change in the Cerebrospinal Fluid in Parkinson's Disease. Neurology India, 2021, 69, 115.	0.4	9
28	ATP10B variants in Parkinsonâ€™s disease: a large cohort study in Chinese mainland population. Acta Neuropathologica, 2021, 141, 805-806.	7.7	8
29	UQCRC1 variants in Parkinsonâ€™s disease: a large cohort study in Chinese mainland population. Brain, 2021, 144, e54-e54.	7.6	5
30	Contribution of coding/non-coding variants in NUS1 to late-onset sporadic Parkinson's disease. Parkinsonism and Related Disorders, 2021, 84, 29-34.	2.2	2
31	Characterizing the Expression Patterns of Parkinsonâ€™s Disease Associated Genes. Frontiers in Neuroscience, 2021, 15, 629156.	2.8	6
32	Gene4PD: A Comprehensive Genetic Database of Parkinsonâ€™s Disease. Frontiers in Neuroscience, 2021, 15, 679568.	2.8	16
33	Gene4MND: An Integrative Genetic Database and Analytic Platform for Motor Neuron Disease. Frontiers in Molecular Neuroscience, 2021, 14, 644202.	2.9	1
34	Evaluation of Peripheral Immune Activation in Amyotrophic Lateral Sclerosis. Frontiers in Neurology, 2021, 12, 628710.	2.4	3
35	Genotype and phenotype distribution of 435 patients with Charcotâ€“Marieâ€“Tooth disease from central south China. European Journal of Neurology, 2021, 28, 3774-3783.	3.3	19
36	PSAP variants in Parkinsonâ€™s disease: a large cohort study in Chinese mainland population. Brain, 2021, 144, e25-e25.	7.6	7

#	Article	IF	Citations
37	The Association Between Lysosomal Storage Disorder Genes and Parkinsonâ€™s Disease: A Large Cohort Study in Chinese Mainland Population. Frontiers in Aging Neuroscience, 2021, 13, 749109.	3.4	11
38	Characteristics of Autonomic Dysfunction in Parkinsonâ€™s Disease: A Large Chinese Multicenter Cohort Study. Frontiers in Aging Neuroscience, 2021, 13, 761044.	3.4	15
39	One PMP22/MPZ and Three MFN2/GDAP1 Concomitant Variants Occurred in a Cohort of 189 Chinese Charcot-Marie-Tooth Families. Frontiers in Neurology, 2021, 12, 736704.	2.4	1
40	Constructing Prediction Models for Freezing of Gait by Nomogram and Machine Learning: A Longitudinal Study. Frontiers in Neurology, 2021, 12, 684044.	2.4	6
41	A presenilin-1 mutation causes Alzheimer disease without affecting Notch signaling. Molecular Psychiatry, 2020, 25, 603-613.	7.9	37
42	Expansion of GGC repeat in the human-specific NOTCH2NLC gene is associated with essential tremor. Brain, 2020, 143, 222-233.	7.6	139
43	Ethnicity-specific and overlapping alterations of brain hydroxymethylome in Alzheimerâ€™s disease. Human Molecular Genetics, 2020, 29, 149-158.	2.9	11
44	Study on the Clinical Features of Parkinson's Disease With Probable Rapid Eye Movement Sleep Behavior Disorder. Frontiers in Neurology, 2020, 11, 979.	2.4	12
45	Olfactory Dysfunction and Its Relationship With Clinical Features of Parkinson's Disease. Frontiers in Neurology, 2020, 11, 526615.	2.4	4
46	Different iron deposition patterns in akinetic/rigid-dominant and tremor-dominant Parkinson's disease. Clinical Neurology and Neurosurgery, 2020, 198, 106181.	1.4	12
47	GCH1 variants contribute to the risk and earlier age-at-onset of Parkinsonâ€™s disease: a two-cohort case-control study. Translational Neurodegeneration, 2020, 9, 31.	8.0	30
48	Genetic and Clinical Features in 24 Chinese Distal Hereditary Motor Neuropathy Families. Frontiers in Neurology, 2020, 11, 603003.	2.4	8
49	Olfactory Dysfunction Predicts Disease Progression in Parkinsonâ€™s Disease: A Longitudinal Study. Frontiers in Neuroscience, 2020, 14, 569777.	2.8	25
50	Reply: Assessing the NOTCH2NLC GGC expansion in European patients with essential tremor. Brain, 2020, 143, e90-e90.	7.6	0
51	Association Between REM Sleep Behavior Disorder and Cognitive Dysfunctions in Parkinson's Disease: A Systematic Review and Meta-Analysis of Observational Studies. Frontiers in Neurology, 2020, 11, 577874.	2.4	10
52	Generation of an induced pluripotent stem cell line (GIBHi004-A) from a Parkinsonâ€™s disease patient with mutant DJ-1/PARK7 (p.L10P). Stem Cell Research, 2020, 46, 101845.	0.7	3
53	<scp>PINK</scp> 1 phosphorylates Drp1 ^{S616} to regulate mitophagyâ€independent mitochondrial dynamics. EMBO Reports, 2020, 21, e48686.	4.5	112
54	Research advances on neurite outgrowth inhibitor B receptor. Journal of Cellular and Molecular Medicine, 2020, 24, 7697-7705.	3.6	13

#	Article	IF	Citations
55	Lee Silverman Voice Treatment for dysarthria in patients with Parkinsonâ€™s disease: a systematic review and metaâ€analysis. European Journal of Neurology, 2020, 27, 1957-1970.	3.3	30
56	The Discriminative Power of Different Olfactory Domains in Parkinson's Disease. Frontiers in Neurology, 2020, 11, 420.	2.4	8
57	The role of genetics in Parkinsonâ€™s disease: a large cohort study in Chinese mainland population. Brain, 2020, 143, 2220-2234.	7.6	97
58	Genetic analysis of N6-methyladenosine modification genes in Parkinson's disease. Neurobiology of Aging, 2020, 93, 143.e9-143.e13.	3.1	35
59	Microglial autophagy defect causes parkinson disease-like symptoms by accelerating inflammasome activation in mice. Autophagy, 2020, 16, 2193-2205.	9.1	134
60	Identification of Alzheimerâ€™s diseaseâ€“associated rare coding variants in the ECE2 gene. JCI Insight, 2020, 5, .	5.0	19
61	Factors Associated With Dyskinesia in Parkinson's Disease in Mainland China. Frontiers in Neurology, 2019, 10, 477.	2.4	15
62	Expansion of Human-Specific GGC Repeat in Neuronal Intranuclear Inclusion Disease-Related Disorders. American Journal of Human Genetics, 2019, 105, 166-176.	6.2	212
63	Association of HIF1A and Parkinsonâ€™s disease in a Han Chinese population demonstrated by molecular inversion probe analysis. Neurological Sciences, 2019, 40, 1927-1931.	1.9	15
64	Clinical Features and Correlates of Excessive Daytime Sleepiness in Parkinson's Disease. Frontiers in Neurology, 2019, 10, 121.	2.4	38
65	Genetic analysis of DNA methylation and hydroxymethylation genes in Parkinson's disease. Neurobiology of Aging, 2019, 84, 242.e13-242.e16.	3.1	10
66	Impaired iPLA2Î² activity affects iron uptake and storage without iron accumulation: An in vitro study excluding decreased iPLA2Î² activity as the cause of iron deposition in PLAN. Brain Research, 2019, 1712, 25-33.	2.2	6
67	The Effects of SNCA rs894278 on Resting-State Brain Activity in Parkinsonâ€™s Disease. Frontiers in Neuroscience, 2019, 13, 47.	2.8	12
68	Systematically analyzing rare variants of autosomal-dominant genes for sporadic Parkinson's disease in a Chinese cohort. Neurobiology of Aging, 2019, 76, 215.e1-215.e7.	3.1	17
69	MicroRNA-26a/Death-Associated Protein KinaseÂ1 Signaling Induces Synucleinopathy andÂDopaminergic Neuron Degeneration in Parkinsonâ€™s Disease. Biological Psychiatry, 2019, 85, 769-781.	1.3	92
70	Identification of CHCHD10 variants in Chinese patients with Parkinson's disease. Parkinsonism and Related Disorders, 2018, 47, 96-97.	2.2	3
71	Familial paroxysmal kinesigenic dyskinesia is associated with mutations in the KCNA1 gene. Human Molecular Genetics, 2018, 27, 625-637.	2.9	43
72	Alzheimerâ€™s disease susceptibility genes modify the risk of Parkinson disease and Parkinsonâ€™s disease-associated cognitive impairment. Neuroscience Letters, 2018, 677, 55-59.	2.1	6

#	Article	IF	Citations
73	Hsp70 participates in PINK1-mediated mitophagy by regulating the stability of PINK1. Neuroscience Letters, 2018, 662, 264-270.	2.1	24
74	A Comprehensive Analysis of the Association Between SNCA Polymorphisms and the Risk of Parkinson's Disease. Frontiers in Molecular Neuroscience, 2018, 11, 391.	2.9	31
75	Genetic Impact on Clinical Features in Parkinsonâ€™s Disease: A Study on SNCA-rs11931074. Parkinson's Disease, 2018, 2018, 1-4.	1.1	2
76	PLA2G6-Associated Neurodegeneration (PLAN): Review of Clinical Phenotypes and Genotypes. Frontiers in Neurology, 2018, 9, 1100.	2.4	75
77	Clinical Heterogeneity Among LRRK2 Variants in Parkinson's Disease: A Meta-Analysis. Frontiers in Aging Neuroscience, 2018, 10, 283.	3.4	40
78	A Meta-Analysis of GBA-Related Clinical Symptoms in Parkinsonâ€™s Disease. Parkinson's Disease, 2018, 2018, 1-7.	1.1	29
79	Recent Advances in Biomarkers for Parkinsonâ€™s Disease. Frontiers in Aging Neuroscience, 2018, 10, 305.	3.4	120
80	Coding mutations inNUS1contribute to Parkinsonâ€™s disease. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 11567-11572.	7.1	78
81	Identifying mild-moderate Parkinsonâ€™s disease using whole-brain functional connectivity. Clinical Neurophysiology, 2018, 129, 2507-2516.	1.5	14
82	Integrated Genetic Analysis of Racial Differences of Common GBA Variants in Parkinson's Disease: A Meta-Analysis. Frontiers in Molecular Neuroscience, 2018, 11, 43.	2.9	71
83	Piperine ameliorates SCA17 neuropathology by reducing ER stress. Molecular Neurodegeneration, 2018, 13, 4.	10.8	29
84	Clinical characteristics of PD patients with LRRK2 G2385R and R1628P variants. Neuroscience Letters, 2018, 685, 185-189.	2.1	8
85	SNCA REP1 and Parkinsonâ€™s disease. Neuroscience Letters, 2018, 682, 79-84.	2.1	5
86	Influence of cognitive function on cerebrovascular disease among the elderly. Acta Neurologica Scandinavica, 2017, 135, 308-315.	2.1	1
87	Novel variants in PAX6 gene caused congenital aniridia in two Chinese families. Eye, 2017, 31, 956-961.	2.1	14
88	Identifying the presence of Parkinsonâ€™s disease using low-frequency fluctuations in BOLD signals. Neuroscience Letters, 2017, 645, 1-6.	2.1	59
89	Novel mutations in ADSL for Adenylosuccinate Lyase Deficiency identified by the combination of Trio-WES and constantly updated guidelines. Scientific Reports, 2017, 7, 1625.	3.3	12
90	Rare GCH1 heterozygous variants contributing to Parkinsonâ€™s disease. Brain, 2017, 140, e41-e41.	7.6	21

#	Article	IF	Citations
91	Synergistic Toxicity of Polyglutamine-Expanded TATA-Binding Protein in Glia and Neuronal Cells: Therapeutic Implications for Spinocerebellar Ataxia 17. Journal of Neuroscience, 2017, 37, 9101-9115.	3.6	18
92	Assessment of Three New Loci from Genome-wide Association Study in Essential Tremor in Chinese population. Scientific Reports, 2017, 7, 7981.	3.3	14
93	Relationship between GWAS-linked three new loci in Essential tremor and risk of Parkinson's disease in Chinese population. Parkinsonism and Related Disorders, 2017, 43, 124-126.	2.2	8
94	TMEM230 mutation analysis in Parkinson's disease in a Chinese population. Neurobiology of Aging, 2017, 49, 219.e1-219.e3.	3.1	34
95	Identification of Ser465 as a novel PINK1 autophosphorylation site. Translational Neurodegeneration, 2017, 6, 34.	8.0	4
96	Altered Functional Brain Connectomes between Sporadic and Familial Parkinson's Patients. Frontiers in Neuroanatomy, 2017, 11, 99.	1.7	7
97	BAG5 Interacts with DJ-1 and Inhibits the Neuroprotective Effects of DJ-1 to Combat Mitochondrial Oxidative Damage. Oxidative Medicine and Cellular Longevity, 2017, 2017, 1-10.	4.0	24
98	Genetic Analysis of LRRK2 R1628P in Parkinsonâ€™s Disease in Asian Populations. Parkinson's Disease, 2017, 2017, 1-6.	1.1	16
99	The Association betweenC9orf72Repeats and Risk of Alzheimerâ€™s Disease and Amyotrophic Lateral Sclerosis: A Meta-Analysis. Parkinson's Disease, 2016, 2016, 1-8.	1.1	13
100	Parkinsonâ€™s Disease and Cognitive Impairment. Parkinson's Disease, 2016, 2016, 1-8.	1.1	50
101	The Progress of Induced Pluripotent Stem Cells as Models of Parkinsonâ€™s Disease. Stem Cells International, 2016, 2016, 1-6.	2.5	19
102	Prediction of orthostatic hypotension in multiple system atrophy and Parkinson disease. Scientific Reports, 2016, 6, 21649.	3.3	13
103	Genome-wide analysis of DNA methylation during antagonism of DMOG to MnCl2-induced cytotoxicity in the mouse substantia nigra. Scientific Reports, 2016, 6, 28933.	3.3	15
104	Lack of association between IL-10 and IL-18 gene promoter polymorphisms and Parkinsonâ€™s disease with cognitive impairment in a Chinese population. Scientific Reports, 2016, 6, 19021.	3.3	35
105	RAB39B gene mutations are not linked to familial Parkinsonâ€™s disease in China. Scientific Reports, 2016, 6, 34502.	3.3	9
106	Mutations analysis of RAB39B gene in Chinese early-onset Parkinson's disease. Parkinsonism and Related Disorders, 2016, 28, 157-158.	2.2	7
107	The GBA, DYRK1A and MS4A6A polymorphisms influence the age at onset of Chinese Parkinson patients. Neuroscience Letters, 2016, 621, 133-136.	2.1	16
108	The rs3756063 polymorphism is associated with SNCA methylation in the Chinese Han population. Journal of the Neurological Sciences, 2016, 367, 11-14.	0.6	24

#	Article	IF	Citations
109	BAG2 structure, function and involvement in disease. Cellular and Molecular Biology Letters, 2016, 21, 18.	7.0	54
110	Relationship between Alzheimer's disease GWAS-linked top hits and risk of Parkinson's disease with or without cognitive decline: a Chinese population-based study. Neurobiology of Aging, 2016, 39, 217.e9-217.e11.	3.1	12
111	UBA5 Mutations Cause a New Form of Autosomal Recessive Cerebellar Ataxia. PLoS ONE, 2016, 11, e0149039.	2.5	68
112	Association between 1p13.3 genomic markers and coronary artery disease: a meta-analysis involving patients and controls. Genetics and Molecular Research, 2015, 14, 9092-9102.	0.2	4
113	Effect ofGBAMutations on Phenotype of Parkinsonâ€™s Disease: A Study on Chinese Population and aMeta-Analysis. Parkinson's Disease, 2015, 2015, 1-10.	1.1	38
114	Polygenic determinants of Parkinson's disease in a Chinese population. Neurobiology of Aging, 2015, 36, 1765.e1-1765.e6.	3.1	73
115	Analysis of several loci from genome-wide association studies in Parkinsonâ€™s disease in mainland China. Neuroscience Letters, 2015, 587, 68-71.	2.1	15
116	Assessment of RIT2 rs12456492 association with Parkinson's disease in Mainland China. Neurobiology of Aging, 2015, 36, 1600.e9-1600.e11.	3.1	13
117	The single nucleotide polymorphism Rs12817488 is associated with Parkinsonâ€™s disease in the Chinese population. Journal of Clinical Neuroscience, 2015, 22, 1002-1004.	1.5	11
118	A neurophysiological profile in Parkinsonâ€™s disease with mild cognitive impairment and dementia in China. Journal of Clinical Neuroscience, 2015, 22, 981-985.	1.5	23
119	Exon dosage analysis of parkin gene in Chinese sporadic Parkinsonâ€™s disease. Neuroscience Letters, 2015, 604, 47-51.	2.1	17
120	Large Polyglutamine Repeats Cause Muscle Degeneration in SCA17 Mice. Cell Reports, 2015, 13, 196-208.	6.4	39
121	LRRK2 A419V variant is a risk factor for Parkinson's disease in Asian population. Neurobiology of Aging, 2015, 36, 2908.e11-2908.e15.	3.1	35
122	Mutation analysis of CHCHD2 gene in Chinese familial Parkinson's disease. Neurobiology of Aging, 2015, 36, 3117.e7-3117.e8.	3.1	22
123	The contribution of GIGYF2 to Parkinsonâ€™s disease: a meta-analysis. Neurological Sciences, 2015, 36, 2073-2079.	1.9	26
124	The BAG2 and BAG5 proteins inhibit the ubiquitination of pathogenic ataxin3-80Q. International Journal of Neuroscience, 2015, 125, 390-394.	1.6	8
125	Genetic Identification Is Critical for the Diagnosis of Parkinsonism: A Chinese Pedigree with Early Onset of Parkinsonism. PLoS ONE, 2015, 10, e0136245.	2.5	9
126	Association study between SMPD1 p.L302P and sporadic Parkinson's disease in ethnic Chinese population. International Journal of Clinical and Experimental Medicine, 2015, 8, 13869-73.	1.3	1

#	Article	IF	Citations
127	Involvement of Bcl-2-associated athanogene (BAG)-family proteins in the neuroprotection by rasagiline. International Journal of Clinical and Experimental Medicine, 2015, 8, 18158-64.	1.3	6
128	Genetic analysis of P387L mutation in SLC18A2 gene in sporadic Parkinson's disease in Chinese Han population. Journal of Central South University (Medical Sciences), 2015, 40, 825-8.	0.1	0
129	L10P mutation in DJ-1 gene induced oxidative stress and mitochondrial disfunction. Journal of Central South University (Medical Sciences), 2015, 40, 1285-91.	0.1	1
130	BAG5 Protects against Mitochondrial Oxidative Damage through Regulating PINK1 Degradation. PLoS ONE, 2014, 9, e86276.	2.5	56
131	Genetic Diagnosis of Two Dopa-Responsive Dystonia Families by Exome Sequencing. PLoS ONE, 2014, 9, e106388.	2.5	9
132	Role of periodically rotated overlapping parallel lines with enhanced reconstruction diffusion-weighted imaging in correcting distortion and evaluating head and neck masses using 3 T MRI. Clinical Radiology, 2014, 69, 403-409.	1.1	22
133	Investigation of TREM2, PLD3, and UNC5C variants in patients with Alzheimer's disease from mainland China. Neurobiology of Aging, 2014, 35, 2422.e9-2422.e11.	3.1	61
134	Association analysis of STK39, MCCC1/LAMP3 and sporadic PD in the Chinese Han population. Neuroscience Letters, 2014, 566, 206-209.	2.1	9
135	Hypomethylation of SNCA in blood of patients with sporadic Parkinson's disease. Journal of the Neurological Sciences, 2014, 337, 123-128.	0.6	90
136	The BAG2 protein stabilises PINK1 by decreasing its ubiquitination. Biochemical and Biophysical Research Communications, 2013, 441, 488-492.	2.1	32
137	Association study between SNP rs150689919 in the DNA demethylation gene, TET1, and Parkinsonâ€™s disease in Chinese Han population. BMC Neurology, 2013, 13, 196.	1.8	7
138	C9orf72 mutation is rare in Alzheimer's disease, Parkinson's disease, and essential tremor in China. Frontiers in Cellular Neuroscience, 2013, 7, 164.	3.7	35
139	VPS35 gene variants are not associated with Parkinson's disease in the mainland Chinese population. Parkinsonism and Related Disorders, 2012, 18, 983-985.	2.2	15
140	Mutation analysis of LRRK2, SCNA, UCHL1, HtrA2 and GIGYF2 genes in Chinese patients with autosomal dorminant Parkinson's disease. Neuroscience Letters, 2012, 516, 207-211.	2.1	14
141	Variant in the 3â€² region of SNCA associated with Parkinsonâ€™s disease and serum Î±-synuclein levels. Journal of Neurology, 2012, 259, 497-504.	3.6	37
142	Clinical features and [11C]-CFT PET analysis of PARK2, PARK6, PARK7-linked autosomal recessive early onset Parkinsonism. Neurological Sciences, 2011, 32, 35-40.	1.9	24
143	Mutation analysis of Parkin, PINK1 and DJ-1 genes in Chinese patients with sporadic early onset parkinsonism. Journal of Neurology, 2010, 257, 1170-1175.	3.6	46
144	R492X mutation in PTEN-induced putative kinase 1 induced cellular mitochondrial dysfunction and oxidative stress. Brain Research, 2010, 1351, 229-237.	2.2	16

#	Article	IF	Citations
145	Glucocerebrosidase Gene L444P mutation is a risk factor for Parkinson's disease in Chinese population. Movement Disorders, 2010, 25, 1005-1011.	3.9	50
146	A novel presenilin 1 mutation (Ser169del) in a Chinese family with early-onset Alzheimer's disease. Neuroscience Letters, 2010, 468, 34-37.	2.1	23
147	A novel LRRK2 mutation in a mainland Chinese patient with familial Parkinson's disease. Neuroscience Letters, 2010, 468, 198-201.	2.1	22
148	Screening for two SNPs of LINGO1 gene in patients with essential tremor or sporadic Parkinson's disease in Chinese population. Neuroscience Letters, 2010, 481, 69-72.	2.1	35
149	Analysis of SCA2 and SCA3/MJD repeats in Parkinson's disease in mainland China: Genetic, clinical, and positron emission tomography findings. Movement Disorders, 2009, 24, 2007-2011.	3.9	43
150	Mutation analysis of Parkin, PINK1, DJâ€1 and ATP13A2 genes in Chinese patients with autosomal recessive earlyâ€onset Parkinsonism. Movement Disorders, 2008, 23, 2074-2079.	3.9	61
151	Genetic Analysis of Six Transmembrane Protein Family Genes in Parkinsonâ€™s Disease in a Large Chinese Cohort. Frontiers in Aging Neuroscience, 0, 14, .	3.4	8