Yan-ming Xu

List of Publications by Year
in descending order

76	943	623574	580701
papers	citations	14 h-index	25 g-index

79	79	79	1308
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Development and validation of a nomogram for freezing of gait in patients with Parkinson's Disease. Acta Neurologica Scandinavica, 2022, 145, 658-668.	1.0	2
2	Exploring the dual character of metformin in Alzheimer's disease. Neuropharmacology, 2022, 207, 108966.	2.0	10
3	Fatigue in Chinese Patients With Amyotrophic Lateral Sclerosis: Associated Factors and Impact on Quality of Life. Frontiers in Neurology, 2022, 13, 806577.	1.1	3
4	Oculopharyngodistal myopathy with CGG repeat expansions in GIPC1: the first report from southwestern China. Neurological Sciences, 2022, 43, 3989-3993.	0.9	1
5	Rare missense variants in the PPP2R5D gene associated with Parkinsonâ€™s disease in the Han Chinese population. Neuroscience Letters, 2022, 776, 136564.	1.0	2
6	Prevalence of restless legs syndrome in people with diabetes mellitus: A pooling analysis of observational studies. EClinicalMedicine, 2022, 46, 101357.	3.2	7
7	Transcranial direct current stimulation for migraine: a systematic review and metaâ€analysis of randomized controlled trials. CNS Neuroscience and Therapeutics, 2022, 28, 992-998.	1.9	8
8	An Exponential Curve Relationship Between Serum Urate and Migraine: A Cross-Section Study From NHANES. Frontiers in Neurology, 2022, 13, 871783.	1.1	7
9	Effect of shift work on fatigue, reaction time and accuracy of nurses in the Department of Neurology: A crossâ€sectional observational study. Journal of Nursing Management, 2022, 30, 2074-2083.	1.4	3
10	Inhibition of miR-421 Preserves Mitochondrial Function and Protects against Parkinsonâ€™s Disease Pathogenesis via Pink1/Parkin-Dependent Mitophagy. Disease Markers, 2022, 2022, 1-13.	0.6	7
11	Association Between Metformin and Alzheimerâ€™s Disease: A Systematic Review and Meta-Analysis of Clinical Observational Studies. Journal of Alzheimer's Disease, 2022, 88, 1311-1323.	1.2	9
12	Clinical value of C-reactive protein/albumin ratio in Guillain-BarrÃ© syndrome. Neurological Sciences, 2021, 42, 3275-3283.	0.9	8
13	Non-motor symptoms are associated with REM sleep behavior disorder in Parkinsonâ€™s disease: a systematic review and meta-analysis. Neurological Sciences, 2021, 42, 47-60.	0.9	18
14	LRP10 Mutations May Correlate with Sporadic Parkinsonâ€™s Disease in China. Molecular Neurobiology, 2021, 58, 1212-1216.	1.9	7
15	Acute cognitive disorder as the initial manifestation of nitrous oxide abusing: a case report. Neurological Sciences, 2021, 42, 755-756.	0.9	8
16	Association of Single Nucleotide Polymorphism at rs2275294 in the ZNF512B Gene with Prognosis in Amyotrophic Lateral Sclerosis. NeuroMolecular Medicine, 2021, 23, 242-246.	1.8	7
17	A deep learning algorithm for automatic detection and classification of acute intracranial hemorrhages in head CT scans. NeuroImage: Clinical, 2021, 32, 102785.	1.4	62
18	Genetic Polymorphisms of Delta-Like 1 Homolog Influence the Susceptibility to Antituberculosis Drug-Induced Hepatotoxicity. DNA and Cell Biology, 2021, 40, 231-238.	0.9	0

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19	Lymphocyte-based ratios for predicting respiratory failure in Guillain-BarrÃ© syndrome. Journal of Neuroimmunology, 2021, 353, 577504.	1.1	12
20	Cerebrospinal Fluid Biomarkers in Multiple System Atrophy Relative to Parkinsonâ€™s Disease: A Meta-Analysis. Behavioural Neurology, 2021, 2021, 1-9.	1.1	4
21	Fatigue prevalence and associated factors in patients with multiple system atrophy. Acta Neurologica Scandinavica, 2021, 144, 553-558.	1.0	4
22	The Evaluation of Pain with Nociceptive and Neuropathic Characteristics from Three Different Perspectives in Amyotrophic Lateral Sclerosis Patients: A Case Controlled Observational Study in Southwestern China. Neural Plasticity, 2021, 2021, 1-6.	1.0	3
23	Blockade of macrophage-associated programmed death 1 inhibits the pyroptosis signalling pathway in sepsis. Inflammation Research, 2021, 70, 993-1004.	1.6	10
24	Retinal Flow Density Changes in Early-stage Parkinsonâ€™s Disease Investigated by Swept-Source Optical Coherence Tomography Angiography. Current Eye Research, 2021, 46, 1886-1891.	0.7	17
25	Novel deletion in the ACTA1 gene associated with milder phenotype of nemaline myopathy in Chinese patient: a case report. Neurological Sciences, 2021, 42, 5401-5405.	0.9	0
26	Excessive Daytime Sleepiness Is Associated With Non-motor Symptoms of Multiple System Atrophy: A Cross-Sectional Study in China. Frontiers in Neurology, 2021, 12, 798771.	1.1	4
27	Rare, low-frequency and common coding variants of ARHGEF28 gene and their association with sporadic amyotrophic lateral sclerosis. Neurobiology of Aging, 2020, 87, 138.e1-138.e6.	1.5	6
28	Lack of association between appendectomy and Parkinsonâ€™s disease: a systematic review and meta-analysis. Aging Clinical and Experimental Research, 2020, 32, 2201-2209.	1.4	13
29	Association between REM sleep behavior disorder and impulsiveâ€“compulsive behaviors in Parkinsonâ€™s disease: a systematic review and meta-analysis of observational studies. Journal of Neurology, 2020, 267, 331-340.	1.8	14
30	Clinical features of multiple system atrophy with or without rapid eye movement behavior disorder: a cross-sectional study in southwest China. Clinical Autonomic Research, 2020, 30, 239-245.	1.4	3
31	Clinical characteristics of patients with essential tremor or essential tremor plus. Acta Neurologica Scandinavica, 2020, 141, 335-341.	1.0	22
32	Nonâ€motor symptoms are associated with midline tremor in essential tremor. Acta Neurologica Scandinavica, 2020, 142, 501-510.	1.0	6
33	A nomogram to predict mechanical ventilation in Guillainâ€BarrÃ© syndrome patients. Acta Neurologica Scandinavica, 2020, 142, 466-474.	1.0	15
34	Acute Flaccid Paralysis as the Initial Manifestation of Japanese Encephalitis: a Case Report. Japanese Journal of Infectious Diseases, 2020, 73, 381-382.	0.5	8
35	The Gene Polymorphism of VMAT2 Is Associated with Risk of Schizophrenia in Male Han Chinese. Psychiatry Investigation, 2020, 17, 1073-1078.	0.7	4
36	Is obstructive sleep apnea related to postoperative delirium, postoperative, coma or both?. Journal of Thoracic Disease, 2019, 11, S1320-S1321.	0.6	0

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37	Association between suicide and multiple sclerosis: An updated meta-analysis. Multiple Sclerosis and Related Disorders, 2019, 34, 83-90.	0.9	20
38	Alpha-synuclein gene polymorphism affects risk of dementia in Han Chinese with Parkinson's disease. Neuroscience Letters, 2019, 706, 146-150.	1.0	6
39	Frequency and factors related to drooling in Chinese patients with multiple system atrophy: a cross-sectional study. Clinical Autonomic Research, 2019, 29, 595-601.	1.4	6
40	Systematic review of the prognostic role of body mass index in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 356-367.	1.1	14
41	Prevalence and Risk Factors of Depression and Anxiety in Essential Tremor Patients: A Cross-Sectional Study in Southwest China. Frontiers in Neurology, 2019, 10, 1194.	1.1	21
42	Leukoencephalopathy in mitochondrial neurogastrointestinal encephalomyopathy-like syndrome with polymerase-gamma mutations. Annals of Indian Academy of Neurology, 2019, 22, 325.	0.2	3
43	Prevalence of restless legs syndrome in Parkinson's disease: a systematic review and meta-analysis of observational studies. Sleep Medicine, 2018, 43, 40-46.	0.8	48
44	Recurrent stroke-like episodes of Wilson disease with a novel Val176fs mutation. Neurological Sciences, 2018, 39, 973-974.	0.9	2
45	Meta-analysis of the association between ZNF512B polymorphism rs2275294 and risk of amyotrophic lateral sclerosis. Neurological Sciences, 2018, 39, 1261-1266.	0.9	7
46	Absence of association of the Ala58Val (rs17571) CTSD gene variant with Parkinsonâ€™s disease or amyotrophic lateral sclerosis in a Han Chinese population. Neuroscience Letters, 2018, 662, 181-184.	1.0	2
47	Lack of association between the P413L variant of chromogranin B and ALS risk or age at onset: a meta-analysis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2018, 19, 80-86.	1.1	0
48	Systematic review and meta-analysis of observational studies to understand the prevalence of restless legs syndrome in multiple sclerosis: an update. Sleep Medicine, 2018, 50, 97-104.	0.8	21
49	Prevalence of pre-diagnostic symptoms did not differ between LRRK2-related, GBA-related and idiopathic patients with Parkinson's disease. Parkinsonism and Related Disorders, 2018, 57, 72-76.	1.1	6
50	Rare Co-occurrence of Ocular Myasthenia Gravis and Thyroid-Associated Orbitopathy (Ophthalmopathy) in an Individual With Hypothyroidism. Frontiers in Endocrinology, 2018, 9, 801.	1.5	5
51	HLA-DRA/HLA-DRB5 polymorphism affects risk of sporadic ALS and survival in a southwest Chinese cohort. Journal of the Neurological Sciences, 2017, 373, 124-128.	0.3	18
52	Association between gene polymorphism and depression in Parkinson's disease: A case-control study. Journal of the Neurological Sciences, 2017, 375, 231-234.	0.3	18
53	Onset of bladder and motor symptoms in multiple system atrophy: differences according to phenotype. Clinical Autonomic Research, 2017, 27, 103-106.	1.4	6
54	Association between a heme oxygenase-2 genetic variant and risk of Parkinsonâ€™s disease in Han Chinese. Neuroscience Letters, 2017, 642, 119-122.	1.0	5

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55	No association of PARK10 polymorphism with Parkinson's disease in Han Chinese population. Parkinsonism and Related Disorders, 2017, 42, 105-106.	1.1	2
56	Sequencing TMEM230 in Chinese patients with sporadic or familial Parkinson's disease. Movement Disorders, 2017, 32, 800-802.	2.2	12
57	Polymorphism in MIR4697 but not VPS13C, GCH1, or SIPA1L2 is associated with risk of Parkinsonâ€™s disease in a Han Chinese population. Neuroscience Letters, 2017, 650, 8-11.	1.0	10
58	Sequence TMEM230 gene in patients with multiple system atrophy in a southwest Chinese population: A pilot study. Journal of the Neurological Sciences, 2017, 375, 264-265.	0.3	8
59	Sequence CLCN1 and SCN4A in patients with Nondystrophic myotonias in Chinese populations: Genetic and pedigree analysis of 10 families and review of the literature. Channels, 2017, 11, 55-65.	1.5	13
60	Levofloxacin-induced transient musculospiral paralysis. American Journal of Emergency Medicine, 2017, 35, 375.e1-375.e2.	0.7	5
61	Festination Correlates with SNCA Polymorphism in Chinese Patients with Parkinsonâ€™s Disease. Parkinson's Disease, 2017, 2017, 1-4.	0.6	8
62	Association of the COQ2 V393A variant with risk of multiple system atrophy in East Asians: a caseâ€“control study and meta-analysis of the literature. Neurological Sciences, 2016, 37, 423-430.	0.9	44
63	SNP rs1805874 of the Calbindin1 Gene Is Associated with Parkinson's Disease in Han Chinese. Genetic Testing and Molecular Biomarkers, 2016, 20, 753-757.	0.3	4
64	Mutational analysis of CHCHD2 in Chinese patients with multiple system atrophy and amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 2016, 368, 389-391.	0.3	8
65	Letter by Yang et al Regarding Article, â€œUse of Oral Anticoagulants for Stroke Prevention in Patients With Atrial Fibrillation Who Have a History of Intracranial Hemorrhageâ€. Circulation, 2016, 134, e228-9.	1.6	0
66	Identification of TMEM230 mutations in familial Parkinson's disease. Nature Genetics, 2016, 48, 733-739.	9.4	146
67	Mutational scanning of the CHCHD2 gene in Han Chinese patients with Parkinsonâ€™s disease and meta-analysis of the literature. Parkinsonism and Related Disorders, 2016, 29, 42-46.	1.1	17
68	Association of the functional SNP rs2275294 in ZNF512B with risk of amyotrophic lateral sclerosis and Parkinson's disease in Han Chinese. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2016, 17, 142-147.	1.1	12
69	Association of Histamine N-Methyltransferase Thr105Ile Polymorphism with Parkinsonâ€™s Disease and Schizophrenia in Han Chinese: A Case-Control Study. PLoS ONE, 2015, 10, e0119692.	1.1	19
70	Association of the COQ2 V393A Variant with Parkinson's Disease: A Case-Control Study and Meta-Analysis. PLoS ONE, 2015, 10, e0130970.	1.1	8
71	Polymorphism in theVesicular Monoamine Transporter 2Gene Decreases the Risk of Parkinsonâ€™s Disease in Han Chinese Men. Parkinson's Disease, 2015, 2015, 1-8.	0.6	9
72	Lack of association between theATP13A2A746T variant and Parkinson's disease susceptibility in Han Chinese: a meta-analysis. International Journal of Neuroscience, 2015, 126, 1-7.	0.8	2

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73	Linkage analysis and whole-exome sequencing exclude extra mutations responsible for the parkinsonian phenotype of spinocerebellar ataxia-2. Neurobiology of Aging, 2015, 36, 545.e1-545.e7.	1.5	14
74	Mutations in theATP13A2Gene and Parkinsonism: A Preliminary Review. BioMed Research International, 2014, 2014, 1-9.	0.9	48
75	SNP rs7684318 of the Î±-synuclein gene is associated with Parkinson's disease in the Han Chinese population. Brain Research, 2010, 1346, 262-265.	1.1	30
76	Pain-Related Factors and Their Impact on Quality of Life in Chinese Patients With Amyotrophic Lateral Sclerosis. Frontiers in Neuroscience, 0, 16, .	1.4	2