Yi Zhao

List of Publications by Year
in descending order

31 papers	488 citations	687363 13 h-index	713466 21 g-index
31 all docs	31 docs citations	31 times ranked	1022 citing authors

#	Article	IF	Citations
1	Longitudinal Study of SNCA Rep1 Polymorphism on Executive Function inÂEarly Parkinsonâ€™s Disease. Journal of Parkinson's Disease, 2022, , 1-6.	2.8	0
2	Delayed Diagnosis, Difficult Decisions: Novel Gene Deletion Causing X-Linked Hypophosphatemia in a Middle-Aged Man with Achondroplastic Features and Tertiary Hyperparathyroidism. Case Reports in Endocrinology, 2021, 2021, 1-7.	0.4	1
3	Epidermolytic ichthyosis in a child and systematized epidermolytic nevi in the mosaic parent associated with a KRT1 variant. European Journal of Medical Genetics, 2021, 64, 104324.	1.3	3
4	Phenotypic bases of <scp>NOTCH2NLC GGC</scp> expansion positive neuronal intranuclear inclusion disease in a Southeast Asian cohort. Clinical Genetics, 2020, 98, 274-281.	2.0	25
5	SNCA Rep1 microsatellite length influences non-motor symptoms in early Parkinson’s disease. Aging, 2020, 12, 20880-20887.	3.1	2
6	The Therapeutic Implications of Tea Polyphenols Against Dopamine (DA) Neuron Degeneration in Parkinsonâ€™s Disease (PD). Cells, 2019, 8, 911.	4.1	69
7	LRRK2 N551K and R1398H variants are protective in Malays and Chinese in Malaysia: A caseâ€“control association study for Parkinson's disease. Molecular Genetics & Genomic Medicine, 2019, 7, e604.	1.2	11
8	SNCA Rep1 promoter variability influences cognition in Parkinson's disease. Movement Disorders, 2019, 34, 1232-1236.	3.9	13
9	PD-linked CHCHD2 mutations impair CHCHD10 and MICOS complex leading to mitochondria dysfunction. Human Molecular Genetics, 2019, 28, 1100-1116.	2.9	48
10	Traditional clinical criteria outperform high-sensitivity C-reactive protein for the screening of hepatic nuclear factor 1 alpha maturity-onset diabetes of the young among young Asians with diabetes. Therapeutic Advances in Endocrinology and Metabolism, 2018, 9, 271-282.	3.2	6
11	Reprogramming of a human induced pluripotent stem cell (iPSC) line from a Parkinson's disease patient with a R1628P variant in the LRRK2 gene. Stem Cell Research, 2017, 18, 45-47.	0.7	7
12	Development of a human induced pluripotent stem cell (iPSC) line from a Parkinson's disease patient carrying the N551K variant in LRRK2 gene. Stem Cell Research, 2017, 18, 51-53.	0.7	10
13	Derivation of human induced pluripotent stem cell (iPSC) line with LRRK2 gene R1398H variant in Parkinson's disease. Stem Cell Research, 2017, 18, 48-50.	0.7	6
14	Generation of a human induced pluripotent stem cell (iPSC) line carrying the Parkinsonâ€™s disease linked LRRK2 variant S1647T. Stem Cell Research, 2017, 18, 54-56.	0.7	10
15	Varied pathological and therapeutic response effects associated withCHCHD2mutant and risk variants. Human Mutation, 2017, 38, 978-987.	2.5	21
16	GWAS-linked PPARGC1A variant in Asian patients with essential tremor. Brain, 2017, 140, e24-e24.	7.6	18
17	Screening for TMEM230 mutations in young-onset Parkinson's disease. Neurobiology of Aging, 2017, 58, 239.e9-239.e10.	3.1	8
18	Real world experience with pegylated interferon and ribavirin in hepatitis C genotype 1 population with favourableIL28Bpolymorphism. Gastroenterology Report, 2016, 5, gow033.	1.3	5

#	Article	IF	Citations
19	PARK16 is associated with PD in the Malaysian population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2016, 171, 839-847.	1.7	11
20	Methylation of serum gene is an independent prognostic marker in colorectal cancer. American Journal of Cancer Research, 2016, 6, 2098-2108.	1.4	23
21	Lrrk2 R1628P variant is a risk factor for essential tremor. Scientific Reports, 2015, 5, 9029.	3.3	18
22	Nonsynonymous variants in MC1R are rare in Chinese Parkinson disease cases. Annals of Neurology, 2015, 78, 152-153.	5.3	9
23	Whole Exome Sequencing Identifies a Novel and a Recurrent Mutation inBBS2Gene in a Family with Bardet-Biedl Syndrome. BioMed Research International, 2015, 2015, 1-5.	1.9	14
24	Association Analysis of COQ2 Variant in Dementia and Essential Tremor. Parkinson's Disease, 2015, 2015, 1-4.	1.1	1
25	LRRK2 G2385R and R1628P Mutations Are Associated with an Increased Risk of Parkinsonâ€™s Disease in the Malaysian Population. BioMed Research International, 2014, 2014, 1-4.	1.9	26
26	In vivo evidence of pathogenicity of VPS35 mutations in the Drosophila. Molecular Brain, 2014, 7, 73.	2.6	35
27	Analysis of EIF4G1 in Parkinson's disease among Asians. Neurobiology of Aging, 2013, 34, 1311.e5-1311.e6.	3.1	11
28	LRRK2 variant associated with Alzheimer's disease. Neurobiology of Aging, 2011, 32, 1990-1993.	3.1	26
29	IgA nephropathy: effects of clinical indices, ACEI/ATRA therapy and ACE gene polymorphism on disease progression. Nephrology, 2002, 7, S166.	1.6	2
30	Topoisomerase-I- and Alu-mediated genomic deletions of the APC gene in familial adenomatous polyposis. Human Genetics, 2001, 108, 436-442.	3.8	25
31	Two Chinese Families with Pendredâ€™s Syndromeâ€”Radiological Imaging of the Ear and Molecular Analysis of the Pendrin Gene. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 3907-3911.	3.6	24