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	The Therapeutic Implications of Tea Polyphenols Against Dopamine (DA) Neuron Degeneration in Parkinsonâ€™s Disease (PD). Cells, 2019, 8, 911.	4.1	69
2	PD-linked CHCHD2 mutations impair CHCHD10 and MICOS complex leading to mitochondria dysfunction. Human Molecular Genetics, 2019, 28, 1100-1116.	2.9	48
3	In vivo evidence of pathogenicity of VPS35 mutations in the Drosophila. Molecular Brain, 2014, 7, 73.	2.6	35
4	LRRK2 variant associated with Alzheimer's disease. Neurobiology of Aging, 2011, 32, 1990-1993.	3.1	26
5	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
6	Topoisomerase-I- and Alu-mediated genomic deletions of the APC gene in familial adenomatous polyposis. Human Genetics, 2001, 108, 436-442.	3.8	25
7	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
8	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
9	Methylation of serum gene is an independent prognostic marker in colorectal cancer. American Journal of Cancer Research, 2016, 6, 2098-2108.	1.4	23
10	Varied pathological and therapeutic response effects associated withCHCHD2mutant and risk variants. Human Mutation, 2017, 38, 978-987.	2.5	21
11	Lrrk2 R1628P variant is a risk factor for essential tremor. Scientific Reports, 2015, 5, 9029.	3.3	18
12	GWAS-linked PPARGC1A variant in Asian patients with essential tremor. Brain, 2017, 140, e24-e24.	7.6	18
13	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
14	SNCA Rep1 promoter variability influences cognition in Parkinson's disease. Movement Disorders, 2019, 34, 1232-1236.	3.9	13
15	Analysis of EIF4G1 in Parkinson's disease among Asians. Neurobiology of Aging, 2013, 34, 1311.e5-1311.e6.	3.1	11
16	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
17	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
18	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

#	Article	IF	Citations
19	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
20	Nonsynonymous variants in MC1R are rare in Chinese Parkinson disease cases. Annals of Neurology, 2015, 78, 152-153.	5.3	9
21	Screening for TMEM230 mutations in young-onset Parkinson's disease. Neurobiology of Aging, 2017, 58, 239.e9-239.e10.	3.1	8
22	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
23	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
24	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
25	Real world experience with pegylated interferon and ribavirin in hepatitis C genotype 1 population with favourableIL28Bpolymorphism. Gastroenterology Report, 2016, 5, gow033.	1.3	5
26	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
27	IgA nephropathy: effects of clinical indices, ACEI/ATRA therapy and ACE gene polymorphism on disease progression. Nephrology, 2002, 7, S166.	1.6	2
28	SNCA Rep1 microsatellite length influences non-motor symptoms in early Parkinson’s disease. Aging, 2020, 12, 20880-20887.	3.1	2
29	Association Analysis of COQ2 Variant in Dementia and Essential Tremor. Parkinson's Disease, 2015, 2015, 1-4.	1.1	1
30	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
31	Longitudinal Study of SNCA Rep1 Polymorphism on Executive Function inÂEarly Parkinsonâ€™s Disease. Journal of Parkinson's Disease, 2022, , 1-6.	2.8	0