Guohua Zhao

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7465187/publications.pdf

Version: 2024-02-01

24	270	933447 10	996975 15
papers	citations	h-index	g-index

26	26	26	374
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Risk factors of impulsive-compulsive behaviors in PD patients: a meta-analysis. Journal of Neurology, 2022, 269, 1298-1315.	3.6	16
2	Artificial Intelligent Olfactory System for the Diagnosis of Parkinsonâ€™s Disease. ACS Omega, 2022, 7, 4001-4010.	3.5	11
3	Novel SPG11 Mutation in Hereditary Spastic Paraplegia with Thin Corpus Callosum. Neurology India, 2022, 70, 424.	0.4	0
4	Neuroimaging evidence of glymphatic system dysfunction in possible REM sleep behavior disorder and Parkinsonâ€™s disease. Npj Parkinson's Disease, 2022, 8, 54.	5.3	42
5	Assessing the NOTCH2NLC GGC expansion in essential tremor patients from eastern China. Brain, 2021, 144, e1-e1.	7.6	15
6	Recommendations for the diagnosis and treatment of paroxysmal kinesigenic dyskinesia: an expert consensus in China. Translational Neurodegeneration, 2021, 10, 7.	8.0	19
7	NOTCH2NLC-related repeat expansion disorders: an expanding group of neurodegenerative disorders. Neurological Sciences, 2021, 42, 4055-4062.	1.9	21
8	Molecular Features of Parkinson's Disease in Patientâ€Derived Midbrain Dopaminergic Neurons. Movement Disorders, 2021, , .	3.9	4
9	Hesperetin inhibits KSHV reactivation and is reversed by HIF1Î± overexpression. Journal of General Virology, 2021, 102, .	2.9	4
10	Validation of the Chinese Version of the Questionnaire for Impulsive-Compulsive Disorders in Parkinson's Disease. Frontiers in Neurology, 2021, 12, 731552.	2.4	3
11	TGM6 L517W is not a pathogenic variant for spinocerebellar ataxia type 35. Neurology: Genetics, 2020, 6, e424.	1.9	2
12	Novel mutation of the PRRT2 gene in two cases of paroxysmal kinesigenic dyskinesia: Two case reports. Biomedical Reports, 2020, 12, 309-312.	2.0	0
13	ATL3 gene mutation in a Chinese family with hereditary sensory neuropathy type 1F. Journal of the Peripheral Nervous System, 2019, 24, 150-155.	3.1	12
14	Neurologists' attitudes toward driving among persons with epilepsy in China: A pilot electronic survey. Epilepsy and Behavior, 2019, 94, 47-51.	1.7	4
15	Novel mutations in the SPAST gene cause hereditary spastic paraplegia. Parkinsonism and Related Disorders, 2019, 69, 125-133.	2.2	10
16	A novel mutation of LRSAM1 in a Chinese family with Charcotâ€Marieâ€Tooth disease. Journal of the Peripheral Nervous System, 2018, 23, 55-59.	3.1	8
17	Feature-based information filtering in visual working memory is impaired in Parkinson's disease. Neuropsychologia, 2018, 111, 317-323.	1.6	9
18	PRRT2 mutations in a cohort of Chinese families with paroxysmal kinesigenic dyskinesia and genotypeâ€“phenotype correlation reanalysis in literatures. International Journal of Neuroscience, 2018, 128, 751-760.	1.6	13

#	Article	IF	Citations
19	Novel KRIT1/CCM1 and MGC4607/CCM2 Gene Variants in Chinese Families With Cerebral Cavernous Malformations. Frontiers in Neurology, 2018, 9, 1128.	2.4	5
20	Identification of a novel SPG4 tandem base substitution in a Chinese hereditary spastic paraplegia family. Neurological Sciences, 2017, 38, 903-905.	1.9	2
21	Exon 8â€“17 deletions of SPAST in a Chinese family with hereditary spastic paraplegia: A case report and literature review. Journal of the Neurological Sciences, 2015, 357, 282-284.	0.6	5
22	Genetic analysis of SPG4 and SPG3A genes in a cohort of Chinese patients with hereditary spastic paraplegia. Journal of the Neurological Sciences, 2014, 347, 368-371.	0.6	16
23	Clinical Features and Cu/Zn Superoxide Dismutase Gene Mutations in Two Mainland Chinese Families With Amyotrophic Lateral Sclerosis. International Journal of Neuroscience, 2011, 121, 191-195.	1.6	2
24	Clinical Manifestations and Management of Acute Thallium Poisoning. European Neurology, 2008, 60, 292-297.	1.4	45