Zaiqiang Zhang

List of Publications by Year
in descending order

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28	172	1477746 6	1372195 10
papers	citations	h-index	g-index

30	30	30	280
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A Novel Mutation in COL4A1 Gene in a Chinese Family with Pontine Autosomal Dominant Microangiopathy and Leukoencephalopathy. Translational Stroke Research, 2022, 13, 238-244.	2.3	8
2	Clinical and pathological study of SORD-related distal motor neuropathy caused by novel compound heterozygous mutations in a Chinese patient. Clinical Neurology and Neurosurgery, 2022, 213, 107118.	0.6	1
3	Migratory enhancement in brain MRI of rosette-forming glioneuronal tumour over 11Âyears. Acta Neurologica Belgica, 2022, , 1.	0.5	0
4	Genotypic and phenotypic characteristics of juvenile/adult onset vanishing white matter: a series of 14 Chinese patients. Neurological Sciences, 2022, 43, 4961-4977.	0.9	1
5	Clinical Features and Genetic Spectrum of Patients With Clinically Suspected Hereditary Progressive Spastic Paraplegia. Frontiers in Neurology, 2022, 13, 872927.	1.1	6
6	Genotypic and Phenotypic Characteristics of Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy from China. European Neurology, 2021, 84, 237-245.	0.6	2
7	A patient with SCA17 featuring 41 CAG repeats presents with spastic paraplegia and involuntary movement. Parkinsonism and Related Disorders, 2021, 89, 87-89.	1.1	0
8	Anti-ganglioside Antibodies in Guillain-Barre Syndrome: A Novel Immunoblotting-Panel Assay. Frontiers in Neurology, 2021, 12, 760889.	1.1	6
9	Two Novel Myelin Protein Zero Mutations in a Group of Chinese Patients. Frontiers in Neurology, 2021, 12, 734515.	1.1	3
10	Heterogeneity of White Matter Hyperintensities in Cognitively Impaired Patients With Cerebral Small Vessel Disease. Frontiers in Immunology, 2021, 12, 803504.	2.2	13
11	Dawson's Fingers in Cerebral Small Vessel Disease. Frontiers in Neurology, 2020, 11, 669.	1.1	4
12	Intracranial Large Artery Abnormalities and Association With Cerebral Small Vessel Disease in CADASIL. Frontiers in Neurology, 2020, 11, 726.	1.1	3
13	Age-related characteristics and normative values of F waves in healthy infants. Clinical Neurophysiology, 2020, 131, 1068-1074.	0.7	1
14	Multiple reversible encephalitic attacks: a rare manifestation of neuronal intranuclear inclusion disease. BMC Neurology, 2020, 20, 125.	0.8	16
15	Three novel mutations in a group of Chinese patients with X-linked Charcot-Marie-Tooth disease. Clinical Neurology and Neurosurgery, 2019, 184, 105430.	0.6	2
16	CADASIL with Large Intracranial Arterial Atherosclerotic Stenosis. Radiology, 2019, 292, 538-538.	3.6	3
17	Novel Alanyl-tRNA Synthetase 2 Pathogenic Variants in Leukodystrophies. Frontiers in Neurology, 2019, 10, 1321.	1.1	10
18	A novel homozygous NDRG1 mutation in a Chinese patient with Charcot-Marie-Tooth disease 4D. Journal of Clinical Neuroscience, 2018, 53, 231-234.	0.8	3

#	Article	IF	Citations
19	Clinical, electrophysiological, genetic, and imaging features of six Chinese Han patients with hereditary neuropathy with liability to pressure palsies (HNPP). Journal of Clinical Neuroscience, 2018, 48, 133-137.	0.8	6
20	MRI Lesion Load of Cerebral Small Vessel Disease and Cognitive Impairment in Patients With CADASIL. Frontiers in Neurology, 2018, 9, 862.	1.1	13
21	A Novel Mutation of GARS in a Chinese Family With Distal Hereditary Motor Neuropathy Type V. Frontiers in Neurology, 2018, 9, 571.	1.1	5
22	Risk factors of pure leukoaraiosis and the association with preclinical carotid atherosclerosis. Atherosclerosis, 2018, 275, 328-332.	0.4	8
23	A Novel KRIT1/CCM1 Gene Insertion Mutation Associated with Cerebral Cavernous Malformations in a Chinese Family. Journal of Molecular Neuroscience, 2017, 61, 221-226.	1.1	13
24	Needle electromyography of the frontalis muscle in patients with amyotrophic lateral sclerosis. Muscle and Nerve, 2016, 54, 1093-1096.	1.0	6
25	Toll-like receptor 2 and -4 are involved in the pathogenesis of the Guillain-BarrÃ© syndrome. Molecular Medicine Reports, 2015, 12, 3207-3213.	1.1	20
26	Sphincter electromyography in diabetes mellitus and multiple system atrophy. Neurourology and Urodynamics, 2015, 34, 669-674.	0.8	8
27	SAMHD1Gene Mutations Are Associated with Cerebral Large-Artery Atherosclerosis. BioMed Research International, 2015, 2015, 1-8.	0.9	5
28	A novel compound heterozygous mutation in a Chinese boy with L-2-hydroxyglutaric aciduria: a case study. BMC Neurology, 2015, 15, 117.	0.8	3