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	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
2	Multiple reversible encephalitic attacks: a rare manifestation of neuronal intranuclear inclusion disease. BMC Neurology, 2020, 20, 125.	0.8	16
3	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
4	MRI Lesion Load of Cerebral Small Vessel Disease and Cognitive Impairment in Patients With CADASIL. Frontiers in Neurology, 2018, 9, 862.	1.1	13
5	Heterogeneity of White Matter Hyperintensities in Cognitively Impaired Patients With Cerebral Small Vessel Disease. Frontiers in Immunology, 2021, 12, 803504.	2.2	13
6	Novel Alanyl-tRNA Synthetase 2 Pathogenic Variants in Leukodystrophies. Frontiers in Neurology, 2019, 10, 1321.	1.1	10
7	Sphincter electromyography in diabetes mellitus and multiple system atrophy. Neurourology and Urodynamics, 2015, 34, 669-674.	0.8	8
8	Risk factors of pure leukoaraiosis and the association with preclinical carotid atherosclerosis. Atherosclerosis, 2018, 275, 328-332.	0.4	8
9	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
10	Needle electromyography of the frontalis muscle in patients with amyotrophic lateral sclerosis. Muscle and Nerve, 2016, 54, 1093-1096.	1.0	6
11	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
12	Anti-ganglioside Antibodies in Guillain-Barre Syndrome: A Novel Immunoblotting-Panel Assay. Frontiers in Neurology, 2021, 12, 760889.	1.1	6
13	Clinical Features and Genetic Spectrum of Patients With Clinically Suspected Hereditary Progressive Spastic Paraplegia. Frontiers in Neurology, 2022, 13, 872927.	1.1	6
14	SAMHD1Gene Mutations Are Associated with Cerebral Large-Artery Atherosclerosis. BioMed Research International, 2015, 2015, 1-8.	0.9	5
15	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
16	Dawson's Fingers in Cerebral Small Vessel Disease. Frontiers in Neurology, 2020, 11, 669.	1.1	4
17	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
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	CADASIL with Large Intracranial Arterial Atherosclerotic Stenosis. Radiology, 2019, 292, 538-538.	3.6	3
20	Intracranial Large Artery Abnormalities and Association With Cerebral Small Vessel Disease in CADASIL. Frontiers in Neurology, 2020, 11, 726.	1.1	3
21	Two Novel Myelin Protein Zero Mutations in a Group of Chinese Patients. Frontiers in Neurology, 2021, 12, 734515.	1.1	3
22	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
23	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
24	Age-related characteristics and normative values of F waves in healthy infants. Clinical Neurophysiology, 2020, 131, 1068-1074.	0.7	1
25	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
26	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
27	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
28	Migratory enhancement in brain MRI of rosette-forming glioneuronal tumour over 11Âyears. Acta Neurologica Belgica, 2022, , 1.	0.5	0