

Zhang Weihua

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

17
papers

64
citations

4
h-index

7
g-index

18
ext. papers

118
ext. citations

4.2
avg, IF

1.91
L-index

#	Paper	IF	Citations
17	Long-term efficacy of mycophenolate mofetil in myelin oligodendrocyte glycoprotein antibody-associated disorders: A prospective study. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2020 , 7,	9.1	28
16	Neurochondrin Antibody Serum Positivity in Three Cases of Autoimmune Cerebellar Ataxia. <i>Cerebellum</i> , 2019 , 18, 1137-1142	4.3	11
15	A systematic review and meta-analysis on the efficacy and safety of traditional Chinese patent medicine Jinqi Jiangtang Tablet in the treatment of type 2 diabetes. <i>Complementary Therapies in Medicine</i> , 2019 , 47, 102021	3.5	4
14	Study of B Cell Repertoire in Patients With Anti-N-Methyl-D-Aspartate Receptor Encephalitis. <i>Frontiers in Immunology</i> , 2020 , 11, 1539	8.4	4
13	Compound Heterozygous Gene Mutations of a Large Deletion and a Missense Variant in a Chinese Patient With Severe Congenital Myasthenic Syndrome With Episodic Apnea. <i>Frontiers in Pharmacology</i> , 2019 , 10, 259	5.6	3
12	Effect of a modified Banxia Xiexin decoction plus chemotherapy on stage III colon cancer. <i>Journal of Traditional Chinese Medicine</i> , 2019 , 39, 251-257	1.1	3
11	Leigh syndrome: a study of 209 patients at the Beijing Children's Hospital.. <i>Annals of Neurology</i> , 2022 ,	9.4	2
10	Whole exome sequencing identifies a novel homozygous MECP2 mutation in a Chinese patient with childhood-onset dystonia and basal ganglia abnormalities, without optic atrophy. <i>Mitochondrion</i> , 2021 , 57, 222-229	4.9	2
9	Case Report: Autoimmune Encephalitis Associated With Anti-glutamic Acid Decarboxylase Antibodies: A Pediatric Case Series. <i>Frontiers in Neurology</i> , 2021 , 12, 641024	4.1	2
8	Clinical Features and Outcomes of Anti-N-Methyl-d-Aspartate Receptor Encephalitis in Infants and Toddlers. <i>Pediatric Neurology</i> , 2021 , 119, 27-33	2.9	2
7	Expanding the mutational spectrum of Rahman syndrome: A rare disorder with severe intellectual disability and particular facial features in two Chinese patients.. <i>Molecular Genetics & Genomic Medicine</i> , 2022 , e1825	2.3	1
6	Pediatric anti-N-methyl-D-aspartate receptor encephalitis with MOG-Ab co-existence: Relapse propensity and treatability.. <i>Multiple Sclerosis and Related Disorders</i> , 2021 , 58, 103447	4	0
5	Phenotypes and genotypes of mitochondrial diseases with mtDNA variations in Chinese children: a multi-center study. <i>Mitochondrion</i> , 2021 , 62, 139-139	4.9	0
4	Immunotherapies for Anti-N-M-methyl-D-aspartate Receptor Encephalitis: Multicenter Retrospective Pediatric Cohort Study in China. <i>Frontiers in Pediatrics</i> , 2021 , 9, 691599	3.4	0
3	Age-dependent characteristics and prognostic factors of pediatric anti-N-methyl-d-aspartate receptor encephalitis in a Chinese single-center study. <i>European Journal of Paediatric Neurology</i> , 2021 , 34, 67-73	3.8	0
2	Chinese patients with p.Arg756 mutations of : Clinical manifestations, treatment, and follow-up.. <i>Pediatric Investigation</i> , 2022 , 6, 5-10	1.3	0
1	A 5-year-old child presenting with tumor-like primary angiitis of the central nervous system. <i>Pediatric Investigation</i> ,	1.3	0

