

Lifang Dai

List of Publications by Year in Descending Order

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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.

8

papers

81

citations

4

h-index

9

g-index

10

ext. papers

109

ext. citations

4.2

avg, IF

2.08

L-index

#	Paper	IF	Citations
8	Clinical Attributes and Electroencephalogram Analysis of Patients With Varying Alpers Syndrome Genotypes. <i>Frontiers in Pharmacology</i> , 2021 , 12, 669516	5.6	0
7	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
6	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
5	Altered inhibitory synapses in de novo GABRA5 and GABRA1 mutations associated with early onset epileptic encephalopathies. <i>Brain</i> , 2019 , 142, 1938-1954	11.2	16
4	A novel DDC gene deletion mutation in two Chinese mainland siblings with aromatic l-amino acid decarboxylase deficiency. <i>Brain and Development</i> , 2019 , 41, 205-209	2.2	6
3	Two Chinese siblings with two novel KCTD7 mutations have dystonia or seizures and epileptic discharge on electroencephalograms. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019 , 70, 27-29	3.2	1
2	An inherited KMT2B duplication variant in a Chinese family with dystonia and/or development delay. <i>Parkinsonism and Related Disorders</i> , 2019 , 63, 227-228	3.6	14
1	Clinical, biochemical and genetic spectrum of 70 patients with ACAD9 deficiency: is riboflavin supplementation effective?. <i>Orphanet Journal of Rare Diseases</i> , 2018 , 13, 120	4.2	41