Lifang Dai

List of Publications by Citations

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, biochemical and genetic spectrum of 70 patients with ACAD9 deficiency: is riboflavin supplementation effective?. *Orphanet Journal of Rare Diseases*, **2018**, 13, 120	4.2	41
7	Altered inhibitory synapses in de novo GABRA5 and GABRA1 mutations associated with early onset epileptic encephalopathies. *Brain*, **2019**, 142, 1938-1954	11.2	16
6	An inherited KMT2B duplication variant in a Chinese family with dystonia and/or development delay. *Parkinsonism and Related Disorders*, **2019**, 63, 227-228	3.6	14
5	A novel DDC gene deletion mutation in two Chinese mainland siblings with aromatic l-amino acid decarboxylase deficiency. *Brain and Development*, **2019**, 41, 205-209	2.2	6
4	Case Report: Autoimmune Encephalitis Associated With Anti-glutamic Acid Decarboxylase Antibodies: A Pediatric Case Series. *Frontiers in Neurology*, **2021**, 12, 641024	4.1	2
3	Two Chinese siblings with two novel KCTD7 mutations have dystonia or seizures and epileptic discharge on electroencephalograms. *Seizure: the Journal of the British Epilepsy Association*, **2019**, 70, 27-29	3.2	1
2	Clinical Attributes and Electroencephalogram Analysis of Patients With Varying AlpersfSyndrome Genotypes. *Frontiers in Pharmacology*, **2021**, 12, 669516	5.6	0
1	Age-dependent characteristics and prognostic factors of pediatric anti-N-methyl-d-aspartate receptor encephalitis in a Chinese single-center study. *European Journal of Paediatric Neurology*, **2021**, 34, 67-73	3.8	0