Lifang Dai

List of Publications by Year
in descending order

8	138	1683354 5	1473754 9
papers	citations	h-index	g-index

10	10	10	365
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical, biochemical and genetic spectrum of 70 patients with ACAD9 deficiency: is riboflavin supplementation effective?. Orphanet Journal of Rare Diseases, 2018, 13, 120.	1.2	61
2	Altered inhibitory synapses in de novo GABRA5 and GABRA1 mutations associated with early onset epileptic encephalopathies. Brain, 2019, 142, 1938-1954.	3.7	32
3	An inherited KMT2B duplication variant in a Chinese family with dystonia and/or development delay. Parkinsonism and Related Disorders, 2019, 63, 227-228.	1.1	15
4	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.	0.6	10
5	Case Report: Autoimmune Encephalitis Associated With Anti-glutamic Acid Decarboxylase Antibodies: A Pediatric Case Series. Frontiers in Neurology, 2021, 12, 641024.	1.1	9
6	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.	0.7	5
7	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.	0.9	4
8	Clinical Attributes and Electroencephalogram Analysis of Patients With Varying Alpersâ€™ Syndrome Genotypes. Frontiers in Pharmacology, 2021, 12, 669516.	1.6	1