Changhong Ding

List of Publications by Citations

17	76	5	8
papers	citations	h-index	g-index
20	122	4.4	2.33
ext. papers	ext. citations	avg, IF	L-index

#	Paper	IF	Citations
17	The clinical and genetic characteristics in children with mitochondrial disease in China. *Science China Life Sciences*, **2017**, 60, 746-757	8.5	20
16	Altered inhibitory synapses in de novo GABRA5 and GABRA1 mutations associated with early onset epileptic encephalopathies. *Brain*, **2019**, 142, 1938-1954	11.2	16
15	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
14	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
13	Molecular spectrum of excision repair cross-complementation group 8 gene defects in Chinese patients with Cockayne syndrome type A. *Scientific Reports*, **2017**, 7, 13686	4.9	5
12	Compound Heterozygous Gene Mutations of a Large Deletion and a Missense Variant in a Chinese Patient With Severe Congenital Myasthenic Syndrome With Episodic Apnea. *Frontiers in Pharmacology*, **2019**, 10, 259	5.6	3
11	Leigh syndrome: a study of 209 patients at the Beijing Childrenæs Hospital.. *Annals of Neurology*, **2022**,	9.4	2
10	Clinical Features and Outcomes of Anti-N-Methyl-d-Aspartate Receptor Encephalitis in Infants and Toddlers. *Pediatric Neurology*, **2021**, 119, 27-33	2.9	2
9	Genotype-phenotype correlation of CACNA1A variants in children with epilepsy. *Developmental Medicine and Child Neurology*, **2021**,	3.3	2
8	Novel truncating mutations in ASXL1 identified in two boys with Bohring-Opitz syndrome. *European Journal of Medical Genetics*, **2021**, 64, 104155	2.6	2
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	3.2	1
6	Expanding the mutational spectrum of Rahman syndrome: A rare disorder with severe intellectual disability and particular facial features in two Chinese patients.. *Molecular Genetics & Genomic Medicine*, **2022**, e1825	2.3	1
5	Phenotypes and genotypes of mitochondrial diseases with mtDNA variations in Chinese children: a multi-center study. *Mitochondrion*, **2021**, 62, 139-139	4.9	0
4	Clinical Attributes and Electroencephalogram Analysis of Patients With Varying AlpersaSyndrome Genotypes. *Frontiers in Pharmacology*, **2021**, 12, 669516	5.6	0
3	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
2	Chinese patients with p.Arg756 mutations of : Clinical manifestations, treatment, and follow-up.. *Pediatric Investigation*, **2022**, 6, 5-10	1.3	0
1	Novel Loss-of-Function Variants in CHD2 Cause Childhood-Onset Epileptic Encephalopathy in Chinese Patients. *Genes*, **2022**, 13, 908	4.2	