Pin Fee Chong

List of Publications by Year
in descending order

15 papers	392 citations	1163117 8 h-index	1199594 12 g-index
15 all docs	15 docs citations	15 times ranked	636 citing authors

#	Article	IF	Citations
1	Acute flaccid myelitis: cause, diagnosis, and management. Lancet, The, 2021, 397, 334-346.	13.7	88
2	Disseminated cortical and subcortical lesions in neonatal enterovirus 71 encephalitis. Journal of NeuroVirology, 2020, 26, 790-792.	2.1	3
3	Acute Flaccid Myelitis With Neuroradiological Finding of Brachial Plexus Swelling. Pediatric Neurology, 2020, 109, 85-88.	2.1	5
4	Letter to the Editor. Journal of Paediatrics and Child Health, 2020, 56, 348-349.	0.8	0
5	Long surviving classical Menkes disease treated with weekly intravenous copper therapy. Journal of Trace Elements in Medicine and Biology, 2019, 54, 172-174.	3.0	8
6	Serial MRI findings of acute flaccid myelitis during an outbreak of enterovirus D68 infection in Japan. Brain and Development, 2019, 41, 443-451.	1.1	31
7	Cytotoxic lesion of the corpus callosum exclusively at the genu in a case of callosal hypogenesis. Journal of Neuroradiology, 2019, 46, 222-223.	1.1	1
8	Mulberries in the urine: a tellâ€tale sign of Fabry disease. Journal of Inherited Metabolic Disease, 2018, 41, 745-746.	3.6	4
9	Clinical Features of Acute Flaccid Myelitis Temporally Associated With an Enterovirus D68 Outbreak: Results of a Nationwide Survey of Acute Flaccid Paralysis in Japan, Augustâ€"December 2015. Clinical Infectious Diseases, 2018, 66, 653-664.	5.8	110
10	Deletions of SCN2A and SCN3A genes in a patient with West syndrome and autistic spectrum disorder. Seizure: the Journal of the British Epilepsy Association, 2018, 60, 91-93.	2.0	10
11	Ineffective quinidine therapy in early onset epileptic encephalopathy with <scp>KCNT</scp>1 mutation. Annals of Neurology, 2016, 79, 502-503.	5.3	68
12	Challenges in detecting genomic copy number aberrations using next-generation sequencing data and the eXome Hidden Markov Model: a clinical exome-first diagnostic approach. Human Genome Variation, 2016, 3, 16025.	0.7	38
13	Water Immersion-Induced Skin Wrinkling Test in Complex Regional Pain Syndrome. Pediatric Neurology, 2015, 52, 649-650.	2.1	0
14	Early onset of moyamoya syndrome in a Down syndrome patient with the genetic variant RNF213 p.R4810K. Brain and Development, 2015, 37, 822-824.	1.1	13
15	A case of pontine tegmental cap dysplasia with comorbidity of oculoauriculovertebral spectrum. Brain and Development, 2015, 37, 171-174.	1.1	13