

Ching-Ching Ng

List of Publications by Year in descending order

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Version: 2024-02-01

12
papers

101
citations

1478505

6
h-index

1372567

10
g-index

12
all docs

12
docs citations

12
times ranked

214
citing authors

#	ARTICLE	IF	CITATIONS
1	Pathogenic Variants in CEP85L Cause Sporadic and Familial Posterior Predominant Lissencephaly. <i>Neuron</i> , 2020, 106, 237-245.e8.	8.1	21
2	Determinants of low bone mineral density in children with epilepsy. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 155-163.	1.6	16
3	Molecular Genetic Characterization of Patients With Focal Epilepsy Using a Customized Targeted Resequencing Gene Panel. <i>Frontiers in Neurology</i> , 2018, 9, 515.	2.4	16
4	Association of <i>NOD1</i> , <i>CXCL16</i> , <i>STAT6</i> and <i>TLR4</i> gene polymorphisms with Malaysian patients with Crohn's disease. <i>PeerJ</i> , 2016, 4, e1843.	2.0	14
5	Genetic polymorphisms of <i>ATG16L1</i> and <i>IRGM</i> genes in Malaysian patients with Crohn's disease. <i>Journal of Digestive Diseases</i> , 2020, 21, 29-37.	1.5	10
6	R54C Mutation of NOTCH3 Gene in the First Rungus Family with CADASIL. <i>PLoS ONE</i> , 2015, 10, e0135470.	2.5	6
7	Whole exome sequencing identifies a novel SCN1A mutation in genetic (idiopathic) generalized epilepsy and juvenile myoclonic epilepsy subtypes. <i>Neurological Sciences</i> , 2020, 41, 591-598.	1.9	6
8	Potential role of regulatory DNA variants in modifying the risk of severe cutaneous reactions induced by aromatic anti-epilepsy medications. <i>Epilepsia</i> , 2022, 63, 936-949.	5.1	5
9	Ethnic variation of genetic (idiopathic) generalized epilepsy in Malaysia. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2017, 45, 24-27.	2.0	2
10	Association of common genetic variants with vitamin D status in Malaysian children with epilepsy. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2020, 79, 103-111.	2.0	2
11	Gene expression profiling and in vitro functional studies reveal RAD54L as a potential therapeutic target in multiple myeloma. <i>Genes and Genomics</i> , 2022, 44, 957-966.	1.4	2
12	A variant e13a3 BCR-ABL1 fusion transcript in refractory adult B-cell acute lymphoblastic leukemia achieving complete remission with CAR-Tcell therapy. <i>Cancer Genetics</i> , 2021, 250-251, 20-24.	0.4	1