Ching-Ching Ng

List of Publications by Year
in descending order

12	101	1478505 6	1372567 10
papers	citations	h-index	g-index

12	12	12	214
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Potential role of regulatory DNA variants in modifying the risk of severe cutaneous reactions induced by aromatic antiâ€seizure medications. Epilepsia, 2022, 63, 936-949.	5.1	5
2	Gene expression profiling and in vitro functional studies reveal RAD54L as a potential therapeutic target in multiple myeloma. Genes and Genomics, 2022, 44, 957-966.	1.4	2
3	A variant e13a3 BCR-ABL1 fusion transcript in refractory adult B-cell acute lymphoblastic leukemia achieving complete remission with CAR-Tcell therapy. Cancer Genetics, 2021, 250-251, 20-24.	0.4	1
4	Whole exome sequencing identifies a novel SCN1A mutation in genetic (idiopathic) generalized epilepsy and juvenile myoclonic epilepsy subtypes. Neurological Sciences, 2020, 41, 591-598.	1.9	6
5	Genetic polymorphisms of ATG16L1 and IRGM genes in Malaysian patients with Crohn's disease. Journal of Digestive Diseases, 2020, 21, 29-37.	1.5	10
6	Pathogenic Variants in CEP85L Cause Sporadic and Familial Posterior Predominant Lissencephaly. Neuron, 2020, 106, 237-245.e8.	8.1	21
7	Association of common genetic variants with vitamin D status in Malaysian children with epilepsy. Seizure: the Journal of the British Epilepsy Association, 2020, 79, 103-111.	2.0	2
8	Determinants of low bone mineral density in children with epilepsy. European Journal of Paediatric Neurology, 2018, 22, 155-163.	1.6	16
9	Molecular Genetic Characterization of Patients With Focal Epilepsy Using a Customized Targeted Resequencing Gene Panel. Frontiers in Neurology, 2018, 9, 515.	2.4	16
10	Ethnic variation of genetic (idiopathic) generalized epilepsy in Malaysia. Seizure: the Journal of the British Epilepsy Association, 2017, 45, 24-27.	2.0	2
11	Association ofNOD1,CXCL16,STAT6andTLR4gene polymorphisms with Malaysian patients with Crohnâ€™s disease. PeerJ, 2016, 4, e1843.	2.0	14
12	R54C Mutation of NOTCH3 Gene in the First Rungus Family with CADASIL. PLoS ONE, 2015, 10, e0135470.	2.5	6