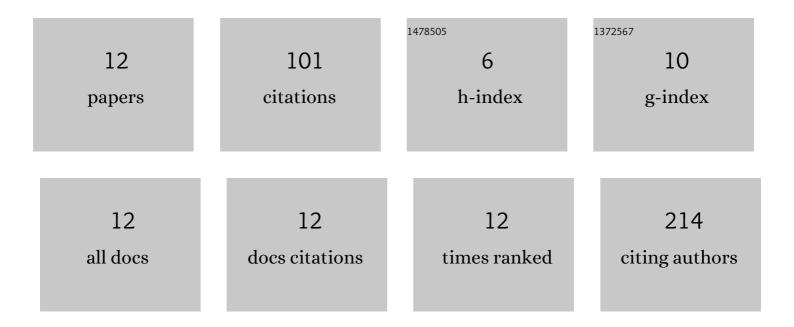
## Ching-Ching Ng

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4864334/publications.pdf Version: 2024-02-01



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#	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 <i>ATG16L1</i> and <i>IRGM</i> 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 of <i>NOD1</i> , <i>CXCL16</i> , <i>STAT6</i> and <i>TLR4</i> gene polymorphisms with Malaysian patients with Crohn'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