Gaik-Siew Ch'ng

List of Publications by Year
in descending order

9 papers	237 citations	1684188 5 h-index	1720034 7 g-index
10 all docs	10 docs citations	10 times ranked	572 citing authors

#	Article	IF	Citations
1	Mutations in KEOPS-complex genes cause nephrotic syndrome with primary microcephaly. Nature Genetics, 2017, 49, 1529-1538.	21.4	164
2	De Novo Truncating Variants in the Last Exon of SEMA6B Cause Progressive Myoclonic Epilepsy. American Journal of Human Genetics, 2020, 106, 549-558.	6.2	32
3	Efficient detection of copyâ€number variations using exome data: Batchâ€and sexâ€based analyses. Human Mutation, 2021, 42, 50-65.	2.5	18
4	Identification of two novel mutations, PSEN1 E280K and PRNP G127S, in a Malaysian family. Neuropsychiatric Disease and Treatment, 2015, 11, 2315.	2.2	9
5	Hyperexcretion of homocitrulline in a Malaysian patient with lysinuric protein intolerance. European Journal of Pediatrics, 2013, 172, 1277-1281.	2.7	6
6	Cost-Effectiveness of Colorectal Cancer Genetic Testing. International Journal of Environmental Research and Public Health, 2021, 18, 8330.	2.6	6
7	A Pathogenic Presenilin-1 Val96Phe Mutation from a Malaysian Family. Brain Sciences, 2021, 11, 1328.	2.3	2
8	P3-015: Novel PSEN1 mutation (E280K) in a malaysian family. , 2015, 11, P625-P625.		0
9	P4â€039: APP, PSEN1, AND PSEN2 MUTATIONS IN EARLYâ€ONSET ALZHEIMER'S DISEASE DISCOVERED IN ASIAN COUNTRIES: A GENETIC SCREENING STUDY OF FAMILIAL AND SPORADIC CASES. Alzheimer's and Dementia, 2018, 14, P1447.	0.8	0