Lin Han

List of Publications by Year
in descending order

10	58	1936888 4	1719596 7
papers	citations	h-index	g-index

11	11	11	75
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Identification of Two Novel Mutations in COG5 Causing Congenital Disorder of Glycosylation. Frontiers in Genetics, 2020, 11, 168.	1.1	11
2	Frequency detection of BRAF V600E mutation in a cohort of pediatric langerhans cell histiocytosis patients by next-generation sequencing. Orphanet Journal of Rare Diseases, 2021, 16, 272.	1.2	11
3	A somatic mutation in PIK3CD unravels a novel candidate gene for lymphatic malformation. Orphanet Journal of Rare Diseases, 2021, 16, 208.	1.2	8
4	Autosomal dominant hereditary spastic paraplegia caused by mutation of UBAP1. Neurogenetics, 2020, 21, 169-177.	0.7	7
5	Novel Compound Heterozygous Variants of ETHE1 Causing Ethylmalonic Encephalopathy in a Chinese Patient: A Case Report. Frontiers in Genetics, 2020, 11, 341.	1.1	6
6	Case Report: Identification of a Novel Homozygous Mutation in GPD1 Gene of a Chinese Child With Transient Infantile Hypertriglyceridemia. Frontiers in Genetics, 2021, 12, 726116.	1.1	4
7	A Novel CCM2 Missense Variant Caused Cerebral Cavernous Malformations in a Chinese Family. Frontiers in Neuroscience, 2020, 14, 604350.	1.4	4
8	Screening of the TMEM151A Gene in Patients With Paroxysmal Kinesigenic Dyskinesia and Other Movement Disorders. Frontiers in Neurology, 0, 13, .	1.1	3
9	Case Report: Novel Compound-Heterozygous Variants of SKIV2L Gene that Cause Trichohepatoenteric Syndrome 2. Frontiers in Genetics, 2021, 12, 756451.	1.1	2
10	A novel SCN9A gene variant identified in a Chinese girl with paroxysmal extreme pain disorder (PEPD): a rare case report. BMC Medical Genomics, 2022, 15, .	0.7	2