Zhan Qi

List of Publications by Year
in descending order

		1162367	1125271
17	198	8	13
papers	citations	h-index	g-index

17	17	17	356
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Toll-like receptor 1(TLR1) Gene SNP rs5743618 is associated with increased risk for tuberculosis in Han Chinese children. Tuberculosis, 2015, 95, 197-203.	0.8	33
2	Genetic Analysis and Follow-Up of 25 Neonatal Diabetes Mellitus Patients in China. Journal of Diabetes Research, 2016, 2016, 1-9.	1.0	28
3	Whole-exome sequencing identified compound heterozygous variants in MMKS in a Chinese pedigree with Bardet-Biedl syndrome. Science China Life Sciences, 2017, 60, 739-745.	2.3	21
4	Newborn screening with targeted sequencing: a multicenter investigation and a pilot clinical study in China. Journal of Genetics and Genomics, 2022, 49, 13-19.	1.7	19
5	Identification of AAAS gene mutation in Allgrove syndrome: A report of three cases. Experimental and Therapeutic Medicine, 2015, 10, 1277-1282.	0.8	16
6	rs2243268 and rs2243274 of Interleukin-4 (IL-4) gene are associated with reduced risk for extrapulmonary and severe tuberculosis in Chinese Han children. Infection, Genetics and Evolution, 2014, 23, 121-128.	1.0	14
7	Instability of BLOCâ€2 and BLOCâ€3 in Chinese patients with Hermanskyâ€Pudlak syndrome. Pigment Cell and Melanoma Research, 2019, 32, 373-380.	1.5	12
8	Genetic variants and mutational spectrum of Chinese Hermanskyâ€“Pudlak syndrome patients. Pigment Cell and Melanoma Research, 2021, 34, 111-121.	1.5	9
9	Clinical feature and waveform in infantile nystagmus syndrome in children with FRMD7 gene mutations. Science China Life Sciences, 2017, 60, 707-713.	2.3	8
10	Parallel Tests of Whole Exome Sequencing and Copy Number Variant Sequencing Increase the Diagnosis Yields of Rare Pediatric Disorders. Frontiers in Genetics, 2020, 11, 473.	1.1	8
11	The first Hermanskyâ€“Pudlak syndrome type 9 patient with two novel variants in Chinese population. Journal of Dermatology, 2021, 48, 676-680.	0.6	7
12	Clinical Application of Whole Exome Sequencing for Monogenic Disorders in PICU of China. Frontiers in Genetics, 2021, 12, 677699.	1.1	6
13	Spectrum Analysis of Albinism Genes in a Large Cohort of Chinese Index Patients. Journal of Investigative Dermatology, 2022, 142, 1752-1755.e3.	0.3	6
14	Neonates are armed with deviated immune cell proportion and cytokine reduction but higher T cell proliferation potentiality. Acta Biochimica Et Biophysica Sinica, 2018, 50, 934-937.	0.9	5
15	Application of multiplex ligation-dependent probe amplification in the genetic testing of oculocutaneous albinism. Chinese Medical Journal, 2019, 132, 2011-2012.	0.9	3
16	Exome sequencing as the firstâ€tier test for pediatric respiratory diseases: A singleâ€center study. Human Mutation, 2021, 42, 891-900.	1.1	2
17	Simpsonâ€“Golabiâ€“Behmel syndrome with 46,XY disorders of sex development: A case report. American Journal of Medical Genetics, Part A, 2019, 179, 285-289.	0.7	1