Ping Li

List of Publications by Year
in descending order

18	199	1478505 6	1125743 13
papers	citations	h-index	g-index

19	19	19	298
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Syndecan-4 assists Mycobacterium tuberculosis entry into lung epithelial cells by regulating the Cdc42, N-WASP, and Arp2/3 signaling pathways. Microbes and Infection, 2022, , 104931.	1.9	2
2	Functional Characterization of FH Mutation c.557G>A Underlies Uterine Leiomyomas. International Journal of Molecular Sciences, 2022, 23, 1452.	4.1	2
3	Identification and Functional Evaluation of a Novel TBX4 Mutation Underlies Small Patella Syndrome. International Journal of Molecular Sciences, 2022, 23, 2075.	4.1	2
4	The negative charge of the 343 site is essential for maintaining physiological functions of CXCR4. BMC Molecular and Cell Biology, 2021, 22, 8.	2.0	2
5	WDR45 Mutation Impairs the Autophagic Degradation of Transferrin Receptor and Promotes Ferroptosis. Frontiers in Molecular Biosciences, 2021, 8, 645831.	3.5	28
6	Identification of Two Novel Compound Heterozygous EIF2AK3 Mutations Underlying Wolcottâ€“Rallison Syndrome in a Chinese Family. Frontiers in Pediatrics, 2021, 9, 679646.	1.9	3
7	A novel pathogenic splice site variation in STK11 gene results in Peutzâ€“Jeghers syndrome. Molecular Genetics & Genomic Medicine, 2021, 9, e1729.	1.2	5
8	RNAseq and quantitative proteomic analysis of Dictyostelium knock-out cells lacking the core autophagy proteins ATG9 and/or ATG16. BMC Genomics, 2021, 22, 444.	2.8	7
9	Two Novel Hydroxymethylbilane Synthase Splicing Mutations Predispose to Acute Intermittent Porphyria. International Journal of Molecular Sciences, 2021, 22, 11008.	4.1	5
10	Identification and spatiotemporal expression of gpr161 genes in zebrafish. Gene, 2020, 730, 144303.	2.2	1
11	Identification of a novel compound heterozygous IDUA mutation underlies Mucopolysaccharidoses type I in a Chinese pedigree. Molecular Genetics & Genomic Medicine, 2020, 8, e1058.	1.2	2
12	A novel splicing pathogenic variant in COL1A1 causing osteogenesis imperfecta (OI) type I in a Chinese family. Molecular Genetics & Genomic Medicine, 2020, 8, e1366.	1.2	4
13	Functional evaluation of a novel GLA causative mutation in Fabry disease. Molecular Genetics & Genomic Medicine, 2019, 7, e864.	1.2	5
14	Functional evidence for a de novo mutation in WDR45 leading to BPAN in a Chinese girl. Molecular Genetics & Genomic Medicine, 2019, 7, e858.	1.2	9
15	Bacteria Exploit Autophagy For Their Own Benefit . Infection and Drug Resistance, 2019, Volume 12, 3205-3215.	2.7	10
16	A Novel Î±-Galactosidase A Splicing Mutation Predisposes to Fabry Disease. Frontiers in Genetics, 2019, 10, 60.	2.3	15
17	The Role of ATG16 in Autophagy and The Ubiquitin Proteasome System. Cells, 2019, 8, 2.	4.1	48
18	Inner nuclear envelope protein SUN1 plays a prominent role in mammalian mRNA export. Nucleic Acids Research, 2015, 43, gkv1058.	14.5	48