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	Inner nuclear envelope protein SUN1 plays a prominent role in mammalian mRNA export. Nucleic Acids Research, 2015, 43, gkv1058.	14.5	48
2	The Role of ATG16 in Autophagy and The Ubiquitin Proteasome System. Cells, 2019, 8, 2.	4.1	48
3	WDR45 Mutation Impairs the Autophagic Degradation of Transferrin Receptor and Promotes Ferroptosis. Frontiers in Molecular Biosciences, 2021, 8, 645831.	3.5	28
4	A Novel Î±-Galactosidase A Splicing Mutation Predisposes to Fabry Disease. Frontiers in Genetics, 2019, 10, 60.	2.3	15
5	Bacteria Exploit Autophagy For Their Own Benefit . Infection and Drug Resistance, 2019, Volume 12, 3205-3215.	2.7	10
6	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
7	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
8	Functional evaluation of a novel GLA causative mutation in Fabry disease. Molecular Genetics & Genomic Medicine, 2019, 7, e864.	1.2	5
9	A novel pathogenic splice site variation in STK11 gene results in Peutzâ€“Jeghers syndrome. Molecular Genetics & Genomic Medicine, 2021, 9, e1729.	1.2	5
10	Two Novel Hydroxymethylbilane Synthase Splicing Mutations Predispose to Acute Intermittent Porphyria. International Journal of Molecular Sciences, 2021, 22, 11008.	4.1	5
11	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
12	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
13	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
14	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
15	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
16	Functional Characterization of FH Mutation c.557G>A Underlies Uterine Leiomyomas. International Journal of Molecular Sciences, 2022, 23, 1452.	4.1	2
17	Identification and Functional Evaluation of a Novel TBX4 Mutation Underlies Small Patella Syndrome. International Journal of Molecular Sciences, 2022, 23, 2075.	4.1	2
18	Identification and spatiotemporal expression of gpr161 genes in zebrafish. Gene, 2020, 730, 144303.	2.2	1