Xiaoqing Zhang

List of Publications by Year
in descending order

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19 papers	243 citations	1307594 7 h-index	996975 15 g-index
21 all docs	21 docs citations	21 times ranked	587 citing authors

#	Article	IF	Citations
1	Clinical features, laboratory findings and persistence of virus in 10 children with coronavirus disease 2019 (COVID-19). Biomedical Journal, 2021, 44, 94-100.	3.1	8
2	Identification of the genetic basis of sporadic polydactyly in China by targeted sequencing. Computational and Structural Biotechnology Journal, 2021, 19, 3482-3490.	4.1	0
3	FGD5-AS1 Is a Hub lncRNA ceRNA in Hearts With Tetralogy of Fallot Which Regulates Congenital Heart Disease Genes Transcriptionally and Epigenetically. Frontiers in Cell and Developmental Biology, 2021, 9, 630634.	3.7	13
4	Long Noncoding RNA lnc-TSSK2-8 Activates Canonical Wnt/Î²-Catenin Signaling Through Small Heat Shock Proteins HSPA6 and CRYAB. Frontiers in Cell and Developmental Biology, 2021, 9, 660576.	3.7	7
5	SORBS2 is a genetic factor contributing to cardiac malformation of 4q deletion syndrome patients. ELife, 2021, 10, .	6.0	9
6	Generation of a homozygous CRISPR/Cas9-mediated knockout human iPSC line for PTCH1 gene. Stem Cell Research, 2021, 56, 102517.	0.7	0
7	Copy number variation analysis in Chinese children with complete atrioventricular canal and single ventricle. BMC Medical Genomics, 2021, 14, 243.	1.5	0
8	A Novel Nonsense GLI3 Variant Is Associated With Polydactyly and Syndactyly in a Family by Blocking the Sonic Hedgehog Signaling Pathway. Frontiers in Genetics, 2020, 11, 542004.	2.3	5
9	Rare Copy Number Variations Might Not be Involved in the Molecular Pathogenesis of PAâ€“IVS in an Unselected Chinese Cohort. Pediatric Cardiology, 2019, 40, 762-767.	1.3	3
10	Intracellular osteopontin negatively regulates toll-like receptor 4-mediated inflammatory response via regulating GSK3Î² and 4EBP1 phosphorylation. Cytokine, 2018, 108, 89-95.	3.2	13
11	A novel dNTP-limited PCR and HRM assay to detect Williams-Beuren syndrome. Clinica Chimica Acta, 2018, 481, 171-176.	1.1	1
12	A loss-of-function mutation p.T52S in RIPPLY3 is a potential predisposing genetic risk factor for Chinese Han conotruncal heart defect patients without the 22q11.2 deletion/duplication. Journal of Translational Medicine, 2018, 16, 260.	4.4	3
13	Accurate diagnosis of spinal muscular atrophy and 22q11.2 deletion syndrome using limited deoxynucleotide triphosphates and high-resolution melting. BMC Genomics, 2018, 19, 485.	2.8	7
14	A Novel Multiplex HRM Assay to Detect Clopidogrel Resistance. Scientific Reports, 2017, 7, 16021.	3.3	6
15	A Novel Missense Mutation of GATA4 in a Chinese Family with Congenital Heart Disease. PLoS ONE, 2016, 11, e0158904.	2.5	19
16	Three novel missense mutations in the filamin B gene are associated with isolated congenital talipes equinovarus. Human Genetics, 2016, 135, 1181-1189.	3.8	22
17	A novel missense mutation of TNNI2 in a Chinese family cause distal arthrogryposis type 1. American Journal of Medical Genetics, Part A, 2016, 170, 135-141.	1.2	8
18	A modified multiplex ligation-dependent probe amplification method for the detection of 22q11.2 copy number variations in patients with congenital heart disease. BMC Genomics, 2015, 16, 364.	2.8	44

#	ARTICLE	IF	CITATIONS
19	Chromosome microarray testing for patients with congenital heart defects reveals novel disease causing loci and high diagnostic yield. BMC Genomics, 2014, 15, 1127.	2.8	74