Quan Long

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

5,499 27 11 29 g-index h-index citations papers 4.65 29 7,045 9.3 avg, IF L-index ext. papers ext. citations

#	Paper	IF	Citations
27	Human genomics. The Genotype-Tissue Expression (GTEx) pilot analysis: multitissue gene regulation in humans. <i>Science</i> , 2015 , 348, 648-60	33.3	3242
26	Pindel: a pattern growth approach to detect break points of large deletions and medium sized insertions from paired-end short reads. <i>Bioinformatics</i> , 2009 , 25, 2865-71	7.2	1423
25	A mixed-model approach for genome-wide association studies of correlated traits in structured populations. <i>Nature Genetics</i> , 2012 , 44, 1066-71	36.3	273
24	Massive genomic variation and strong selection in Arabidopsis thaliana lines from Sweden. <i>Nature Genetics</i> , 2013 , 45, 884-890	36.3	264
23	Synchronized age-related gene expression changes across multiple tissues in human and the link to complex diseases. <i>Scientific Reports</i> , 2015 , 5, 15145	4.9	128
22	PoolHap: inferring haplotype frequencies from pooled samples by next generation sequencing. <i>PLoS ONE</i> , 2011 , 6, e15292	3.7	28
21	Patterns of Polymorphism at the Self-Incompatibility Locus in 1,083 Arabidopsis thaliana Genomes. <i>Molecular Biology and Evolution</i> , 2017 , 34, 1878-1889	8.3	25
20	Inter-tissue coexpression network analysis reveals DPP4 as an important gene in heart to blood communication. <i>Genome Medicine</i> , 2016 , 8, 15	14.4	20
19	Detecting disease-associated genotype patterns. <i>BMC Bioinformatics</i> , 2009 , 10 Suppl 1, S75	3.6	19
18	AprioriGWAS, a new pattern mining strategy for detecting genetic variants associated with disease through interaction effects. <i>PLoS Computational Biology</i> , 2014 , 10, e1003627	5	18
17	HI: haplotype improver using paired-end short reads. <i>Bioinformatics</i> , 2009 , 25, 2436-7	7.2	12
16	JAWAMix5: an out-of-core HDF5-based java implementation of whole-genome association studies using mixed models. <i>Bioinformatics</i> , 2013 , 29, 1220-2	7.2	8
15	Nucleocytoplasmic transport of the RNA-binding protein CELF2 regulates neural stem cell fates. <i>Cell Reports</i> , 2021 , 35, 109226	10.6	6
14	Evaluation of A Phylogenetic Pipeline to Examine Transmission Networks in A Canadian HIV Cohort. <i>Microorganisms</i> , 2020 , 8,	4.9	5
13	kTWAS: integrating kernel machine with transcriptome-wide association studies improves statistical power and reveals novel genes. <i>Briefings in Bioinformatics</i> , 2021 , 22,	13.4	5
12	Power analysis of transcriptome-wide association study: Implications for practical protocol choice. <i>PLoS Genetics</i> , 2021 , 17, e1009405	6	5
11	Reconstruction of Microbial Haplotypes by Integration of Statistical and Physical Linkage in Scaffolding. <i>Molecular Biology and Evolution</i> , 2021 , 38, 2660-2672	8.3	4

LIST OF PUBLICATIONS

10	Computational Haplotype Inference from Pooled Samples. <i>Methods in Molecular Biology</i> , 2017 , 1551, 309-319	1.4	3
9	PRESM: personalized reference editor for somatic mutation discovery in cancer genomics. <i>Bioinformatics</i> , 2019 , 35, 1445-1452	7.2	3
8	An extended Tajima & D neutrality test incorporating SNP calling and imputation uncertainties. Statistics and Its Interface, 2015, 8, 447-456	0.4	2
7	OCMA: Fast, Memory-Efficient Factorization of Prohibitively Large Relationship Matrices. <i>G3: Genes, Genomes, Genetics</i> , 2019 , 9, 13-19	3.2	1
6	Reconstruction of microbial haplotypes by integration of statistical and physical linkage in scaffolding		1
5	WgLink: reconstructing whole-genome viral haplotypes using ${ m L}0+{ m L}1$ -regularization. <i>Bioinformatics</i> , 2021 ,	7.2	1
4	Genetic variations of DNA bindings of FOXA1 and co-factors in breast cancer susceptibility. <i>Nature Communications</i> , 2021 , 12, 5318	17.4	1
3	Reconstructing SARS-CoV-2 infection dynamics through the phylogenetic inference of unsampled sources of infection <i>PLoS ONE</i> , 2021 , 16, e0261422	3.7	1
2	An Individualized Approach for Somatic Variant Discovery. <i>Methods in Molecular Biology</i> , 2020 , 2120, 11-36	1.4	
1	SimPEL: Simulation-based power estimation for sequencing studies of low-prevalence conditions. <i>Genetic Epidemiology</i> , 2018 , 42, 480-487	2.6	