

Quan Long

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/8997906/quan-long-publications-by-year.pdf>

Version: 2024-04-29

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

27
papers

5,499
citations

11
h-index

29
g-index

29
ext. papers

7,045
ext. citations

9.3
avg. IF

4.65
L-index

#	Paper	IF	Citations
27	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
26	Nucleocytoplasmic transport of the RNA-binding protein CELF2 regulates neural stem cell fates. <i>Cell Reports</i> , 2021 , 35, 109226	10.6	6
25	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
24	WgLink: reconstructing whole-genome viral haplotypes using $\ell_0+\ell_1$ -regularization. <i>Bioinformatics</i> , 2021 ,	7.2	1
23	Power analysis of transcriptome-wide association study: Implications for practical protocol choice. <i>PLoS Genetics</i> , 2021 , 17, e1009405	6	5
22	Genetic variations of DNA bindings of FOXA1 and co-factors in breast cancer susceptibility. <i>Nature Communications</i> , 2021 , 12, 5318	17.4	1
21	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
20	Evaluation of A Phylogenetic Pipeline to Examine Transmission Networks in A Canadian HIV Cohort. <i>Microorganisms</i> , 2020 , 8,	4.9	5
19	An Individualized Approach for Somatic Variant Discovery. <i>Methods in Molecular Biology</i> , 2020 , 2120, 11-36	1.4	
18	OCMA: Fast, Memory-Efficient Factorization of Prohibitively Large Relationship Matrices. <i>G3: Genes, Genomes, Genetics</i> , 2019 , 9, 13-19	3.2	1
17	PRESM: personalized reference editor for somatic mutation discovery in cancer genomics. <i>Bioinformatics</i> , 2019 , 35, 1445-1452	7.2	3
16	SimPEL: Simulation-based power estimation for sequencing studies of low-prevalence conditions. <i>Genetic Epidemiology</i> , 2018 , 42, 480-487	2.6	
15	Computational Haplotype Inference from Pooled Samples. <i>Methods in Molecular Biology</i> , 2017 , 1551, 309-319	1.4	3
14	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
13	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
12	Human genomics. The Genotype-Tissue Expression (GTEx) pilot analysis: multitissue gene regulation in humans. <i>Science</i> , 2015 , 348, 648-60	33.3	3242
11	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

10	An extended Tajima's D neutrality test incorporating SNP calling and imputation uncertainties. <i>Statistics and Its Interface</i> , 2015 , 8, 447-456	0.4	2
9	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
8	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
7	Massive genomic variation and strong selection in <i>Arabidopsis thaliana</i> lines from Sweden. <i>Nature Genetics</i> , 2013 , 45, 884-890	36.3	264
6	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
5	PoolHap: inferring haplotype frequencies from pooled samples by next generation sequencing. <i>PLoS ONE</i> , 2011 , 6, e15292	3.7	28
4	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
3	HI: haplotype improver using paired-end short reads. <i>Bioinformatics</i> , 2009 , 25, 2436-7	7.2	12
2	Detecting disease-associated genotype patterns. <i>BMC Bioinformatics</i> , 2009 , 10 Suppl 1, S75	3.6	19
1	Reconstruction of microbial haplotypes by integration of statistical and physical linkage in scaffolding		1