Yongzhuang Liu

List of Publications by Year in descending order

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840776 940533 20 482 11 16 citations h-index g-index papers 22 22 22 967 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Long-read-based human genomic structural variation detection with cuteSV. Genome Biology, 2020, 21, 189.	8.8	164
2	A Bayesian framework for <i>de novo</i> mutation calling in parents-offspring trios. Bioinformatics, 2015, 31, 1375-1381.	4.1	87
3	A gradient-boosting approach for filtering <i>de novo</i> mutations in parent–offspring trios. Bioinformatics, 2014, 30, 1830-1836.	4.1	33
4	A deep learning approach for filtering structural variants in short read sequencing data. Briefings in Bioinformatics, 2021 , 22 , .	6.5	32
5	A pipeline for RNA-seq based eQTL analysis with automated quality control procedures. BMC Bioinformatics, 2021, 22, 403.	2.6	27
6	InteGO2: a web tool for measuring and visualizing gene semantic similarities using Gene Ontology. BMC Genomics, 2016, 17, 530.	2.8	26
7	Disease Module Identification Based on Representation Learning of Complex Networks Integrated From GWAS, eQTL Summaries, and Human Interactome. Frontiers in Bioengineering and Biotechnology, 2020, 8, 418.	4.1	22
8	abPOA: an SIMD-based C library for fast partial order alignment using adaptive band. Bioinformatics, 2021, 37, 2209-2211.	4.1	20
9	Joint detection of copy number variations in parent-offspring trios. Bioinformatics, 2016, 32, 1130-1137.	4.1	18
10	eQTLMAPT: Fast and Accurate eQTL Mediation Analysis With Efficient Permutation Testing Approaches. Frontiers in Genetics, 2019, 10, 1309.	2.3	17
11	Sexually Dimorphic Gene Expression Associated with Growth and Reproduction of Tongue Sole (Cynoglossus semilaevis) Revealed by Brain Transcriptome Analysis. International Journal of Molecular Sciences, 2016, 17, 1402.	4.1	15
12	Enhancing discoveries of molecular QTL studies with small sample size using summary statistic imputation. Briefings in Bioinformatics, 2022, 23, .	6.5	15
13	MGMIN: A Normalization Method for Correcting Probe Design Bias in Illumina Infinium HumanMethylation450 BeadChips. Frontiers in Genetics, 2020, 11, 538492.	2.3	2
14	Fast and Accurate Classification of Meta-Genomics Long Reads With deSAMBA. Frontiers in Cell and Developmental Biology, 2021, 9, 643645.	3.7	1
15	PocaCNV: A Tool to Detect Copy Number Variants from Population-Scale Genome Sequencing Data. , 2021, , .		1
16	Joint detection of germline and somatic copy number events in matched tumor–normal sample pairs. Bioinformatics, 2019, 35, 4955-4961.	4.1	0
17	DNMFilter_Indel: Filtering de novo Indels in Parent-Offspring Trios. , 2019, , .		0
18	An integrated approach for copy number variation discovery in parent–offspring trios. Briefings in Bioinformatics, 2021, 22, .	6.5	0

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#	#	Article	IF	CITATIONS
1	19	Filtering de novo indels in parent-offspring trios. BMC Bioinformatics, 2020, 21, 547.	2.6	0
2	20	Ontology-based annotation and retrieval for large-scale VCF data., 2021,,.		0