

Vikas Bansal

List of Publications by Year in descending order

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Version: 2024-02-01

19
papers

1,337
citations

623574

14
h-index

839398

18
g-index

19
all docs

19
docs citations

19
times ranked

2771
citing authors

#	ARTICLE	IF	CITATIONS
1	Robust and accurate estimation of paralog-specific copy number for duplicated genes using whole-genome sequencing. <i>Nature Communications</i> , 2022, 13, .	5.8	4
2	VarCover. <i>Journal of Molecular Diagnostics</i> , 2020, 22, 123-131.	1.2	2
3	Sensitive alignment using paralogous sequence variants improves long-read mapping and variant calling in segmental duplications. <i>Nucleic Acids Research</i> , 2020, 48, e114-e114.	6.5	12
4	A diploid assembly-based benchmark for variants in the major histocompatibility complex. <i>Nature Communications</i> , 2020, 11, 4794.	5.8	56
5	Perfectionism as a mediator of psychological distress: Implications for addressing underlying vulnerabilities to the mental health of medical students. <i>Medical Teacher</i> , 2020, 42, 1301-1307.	1.0	21
6	Ultralow-input single-tube linked-read library method enables short-read second-generation sequencing systems to routinely generate highly accurate and economical long-range sequencing information. <i>Genome Research</i> , 2020, 30, 898-909.	2.4	68
7	Sequencing Technologies and Analyses: Where Have We Been and Where Are We Going?. <i>IScience</i> , 2019, 18, 37-41.	1.9	31
8	Longshot enables accurate variant calling in diploid genomes from single-molecule long read sequencing. <i>Nature Communications</i> , 2019, 10, 4660.	5.8	156
9	Integrating read-based and population-based phasing for dense and accurate haplotyping of individual genomes. <i>Bioinformatics</i> , 2019, 35, i242-i248.	1.8	23
10	An accurate algorithm for the detection of DNA fragments from dilution pool sequencing experiments. <i>Bioinformatics</i> , 2018, 34, 155-162.	1.8	0
11	Targeted genotyping of variable number tandem repeats with adVNTR. <i>Genome Research</i> , 2018, 28, 1709-1719.	2.4	59
12	Identification of a missense variant in the WFS1 gene that causes a mild form of Wolfram syndrome and is associated with risk for type 2 diabetes in Ashkenazi Jewish individuals. <i>Diabetologia</i> , 2018, 61, 2180-2188.	2.9	38
13	Zika virus infection reprograms global transcription of host cells to allow sustained infection. <i>Emerging Microbes and Infections</i> , 2017, 6, 1-10.	3.0	58
14	A computational method for estimating the PCR duplication rate in DNA and RNA-seq experiments. <i>BMC Bioinformatics</i> , 2017, 18, 43.	1.2	23
15	HapCUT2: robust and accurate haplotype assembly for diverse sequencing technologies. <i>Genome Research</i> , 2017, 27, 801-812.	2.4	285
16	Spectrum of mutations in monogenic diabetes genes identified from high-throughput DNA sequencing of 6888 individuals. <i>BMC Medicine</i> , 2017, 15, 213.	2.3	75
17	InPhaDel: integrative shotgun and proximity-ligation sequencing to phase deletions with single nucleotide polymorphisms. <i>Nucleic Acids Research</i> , 2016, 44, e111-e111.	6.5	1
18	Dynamics of the human and viral m6A RNA methylomes during HIV-1 infection of T cells. <i>Nature Microbiology</i> , 2016, 1, 16011.	5.9	373

#	ARTICLE	IF	CITATIONS
19	Fast individual ancestry inference from DNA sequence data leveraging allele frequencies for multiple populations. BMC Bioinformatics, 2015, 16, 4.	1.2	52