Serghei Mangul

List of Publications by Year in descending order

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361413 345221 1,785 41 20 36 citations h-index g-index papers 63 63 63 4133 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Correction for Rando et al., "Pathogenesis, Symptomatology, and Transmission of SARS-CoV-2 through Analysis of Viral Genomics and Structure― MSystems, 2022, , e0144721.	3.8	2
2	Unlocking capacities of genomics for the COVID-19 response and future pandemics. Nature Methods, 2022, 19, 374-380.	19.0	35
3	Virtual meetings promise to eliminate geographical and administrative barriers and increase accessibility, diversity and inclusivity. Nature Biotechnology, 2022, 40, 133-137.	17.5	30
4	A comprehensive benchmarking of WGS-based deletion structural variant callers. Briefings in Bioinformatics, 2022, 23, .	6.5	9
5	The Gene Expression Deconvolution Interactive Tool (GEDIT): accurate cell type quantification from gene expression data. GigaScience, 2021, 10, .	6.4	33
6	Improving the completeness of public metadata accompanying omics studies. Genome Biology, 2021, 22, 106.	8.8	22
7	Detection of viral gene expression in riskâ€stratified biopsies reveals no active HPV in cutaneous squamous cell carcinoma. Experimental Dermatology, 2021, 30, 1711-1716.	2.9	4
8	Diversity in immunogenomics: the value and the challenge. Nature Methods, 2021, 18, 588-591.	19.0	40
9	Accurate assembly of minority viral haplotypes from next-generation sequencing through efficient noise reduction. Nucleic Acids Research, 2021, 49, e102-e102.	14.5	36
10	Rare loss-of-function variants in type I IFN immunity genes are not associated with severe COVID-19. Journal of Clinical Investigation, 2021, 131, .	8.2	56
11	Technology dictates algorithms: recent developments in read alignment. Genome Biology, 2021, 22, 249.	8.8	51
12	Systematic evaluation of transcriptomics-based deconvolution methods and references using thousands of clinical samples. Briefings in Bioinformatics, 2021, 22, .	6.5	10
13	Pathogenesis, Symptomatology, and Transmission of SARS-CoV-2 through Analysis of Viral Genomics and Structure. MSystems, 2021, 6, e0009521.	3.8	26
14	Ancestral diversity is limited in published T cell receptor sequencing studies. Immunity, 2021, 54, 2177-2179.	14.3	3
15	Integrating big data computational skills in education to facilitate reproducibility and transparency in pharmaceutical sciences. JACCP Journal of the American College of Clinical Pharmacy, 2021, 4, 1263-1266.	1.0	O
16	PUMAA: A Platform for Accessible Microbiome Analysis in the Undergraduate Classroom. Frontiers in Microbiology, 2020, 11, 584699.	3.5	10
17	Cell type–specific genetic regulation of gene expression across human tissues. Science, 2020, 369, .	12.6	210
18	Metalign: efficient alignment-based metagenomic profiling via containment min hash. Genome Biology, 2020, 21, 242.	8.8	29

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19	Recommendations to enhance rigor and reproducibility in biomedical research. GigaScience, 2020, 9, .	6.4	83
20	Intratumoral CD4+ T Cells Mediate Anti-tumor Cytotoxicity in Human Bladder Cancer. Cell, 2020, 181, 1612-1625.e13.	28.9	436
21	Benchmarking of computational error-correction methods for next-generation sequencing data. Genome Biology, 2020, 21, 71.	8.8	26
22	Profiling immunoglobulin repertoires across multiple human tissues using RNA sequencing. Nature Communications, 2020, 11, 3126.	12.8	44
23	Telescope: an interactive tool for managing large-scale analysis from mobile devices. GigaScience, 2020, 9, .	6.4	0
24	Challenges and recommendations to improve the installability and archival stability of omics computational tools. PLoS Biology, 2019, 17, e3000333.	5.6	54
25	Transcriptional profiling of single fiber cells in a transgenic paradigm of an inherited childhood cataract reveals absence of molecular heterogeneity. Journal of Biological Chemistry, 2019, 294, 13530-13544.	3.4	5
26	Interpreting and integrating big data in the life sciences. Emerging Topics in Life Sciences, 2019, 3, 335-341.	2.6	2
27	MiCoP: microbial community profiling method for detecting viral and fungal organisms in metagenomic samples. BMC Genomics, 2019, 20, 423.	2.8	22
28	Improving the usability and archival stability of bioinformatics software. Genome Biology, 2019, 20, 47.	8.8	62
29	How bioinformatics and open data can boost basic science in countries and universities with limited resources. Nature Biotechnology, 2019, 37, 324-326.	17.5	25
30	Systematic benchmarking of omics computational tools. Nature Communications, 2019, 10, 1393.	12.8	111
31	Involving undergraduates in genomics research to narrow the education–research gap. Nature Biotechnology, 2018, 36, 369-371.	17.5	2
32	Spatial Analysis of Single Fiber Cells of the Developing Ocular Lens Reveals Regulated Heterogeneity of Gene Expression. IScience, 2018, 10, 66-79.	4.1	15
33	A Comprehensive Map of Genetic Variation in the World's Largest Ethnic Group—Han Chinese. Molecular Biology and Evolution, 2018, 35, 2736-2750.	8.9	86
34	ROP: dumpster diving in RNA-sequencing to find the source of 1 trillion reads across diverse adult human tissues. Genome Biology, 2018, $19, 36$.	8.8	42
35	Long Single-Molecule Reads Can Resolve the Complexity of the Influenza Virus Composed of Rare, Closely Related Mutant Variants. Journal of Computational Biology, 2017, 24, 558-570.	1.6	14
36	Addressing the Digital Divide in Contemporary Biology: Lessons from Teaching UNIX. Trends in Biotechnology, 2017, 35, 901-903.	9.3	22

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#	Article	IF	CITATIONS
37	Reference-free comparison of microbial communities via de Bruijn graphs. , 2016, , .		3
38	Discovering Single Nucleotide Polymorphisms Regulating Human Gene Expression Using Allele Specific Expression from RNA-seq Data. Genetics, 2016, 204, 1057-1064.	2.9	17
39	Accurate viral population assembly from ultra-deep sequencing data. Bioinformatics, 2014, 30, i329-i337.	4.1	48
40	Transcriptome assembly and quantification from Ion Torrent RNA-Seq data. , 2013, , .		0
41	Monte-Carlo Regression algorithm for isoform frequency estimation from RNA-Seq data. , 2013, , .		1