

Erik P Garrison

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

22,488
citations

279798
23
h-index

552781
26
g-index

36
all docs

36
docs citations

36
times ranked

40233
citing authors

#	ARTICLE	IF	CITATIONS
1	The complete sequence of a human genome. <i>Science</i> , 2022, 376, 44-53.	12.6	1,222
2	The Human Pangenome Project: a global resource to map genomic diversity. <i>Nature</i> , 2022, 604, 437-446.	27.8	192
3	ODGI: understanding pangenome graphs. <i>Bioinformatics</i> , 2022, 38, 3319-3326.	4.1	44
4	Efficient dynamic variation graphs. <i>Bioinformatics</i> , 2021, 36, 5139-5144.	4.1	18
5	Towards complete and error-free genome assemblies of all vertebrate species. <i>Nature</i> , 2021, 592, 737-746.	27.8	1,139
6	GRAFIMO: Variant and haplotype aware motif scanning on pangenome graphs. <i>PLoS Computational Biology</i> , 2021, 17, e1009444.	3.2	5
7	Pangenomics enables genotyping of known structural variants in 5202 diverse genomes. <i>Science</i> , 2021, 374, abg8871.	12.6	132
8	Haplotype-aware graph indexes. <i>Bioinformatics</i> , 2020, 36, 400-407.	4.1	59
9	Removing reference bias and improving indel calling in ancient DNA data analysis by mapping to a sequence variation graph. <i>Genome Biology</i> , 2020, 21, 250.	8.8	44
10	Nanopore sequencing and the Shasta toolkit enable efficient de novo assembly of eleven human genomes. <i>Nature Biotechnology</i> , 2020, 38, 1044-1053.	17.5	344
11	Pangenome Graphs. <i>Annual Review of Genomics and Human Genetics</i> , 2020, 21, 139-162.	6.2	148
12	Genotyping structural variants in pangenome graphs using the vg toolkit. <i>Genome Biology</i> , 2020, 21, 35.	8.8	150
13	Genomic diversity and novel genome-wide association with fruit morphology in <i>Capsicum</i> , from 746k polymorphic sites. <i>Scientific Reports</i> , 2019, 9, 10067.	3.3	53
14	Superbubbles, Ultrabubbles, and Cacti. <i>Journal of Computational Biology</i> , 2018, 25, 649-663.	1.6	46
15	Variation graph toolkit improves read mapping by representing genetic variation in the reference. <i>Nature Biotechnology</i> , 2018, 36, 875-879.	17.5	435
16	Genome graphs and the evolution of genome inference. <i>Genome Research</i> , 2017, 27, 665-676.	5.5	264
17	A graph extension of the positional Burrows-Wheeler transform and its applications. <i>Algorithms for Molecular Biology</i> , 2017, 12, 18.	1.2	33
18	Punctuated bursts in human male demography inferred from 1,244 worldwide Y-chromosome sequences. <i>Nature Genetics</i> , 2016, 48, 593-599.	21.4	273

#	ARTICLE	IF	CITATIONS
19	The distribution and mutagenesis of short coding INDELs from 1,128 whole exomes. BMC Genomics, 2015, 16, 143.	2.8	9
20	A global reference for human genetic variation. Nature, 2015, 526, 68-74.	27.8	13,998
21	An integrated map of structural variation in 2,504 human genomes. Nature, 2015, 526, 75-81.	27.8	1,994
22	SpeedSeq: ultra-fast personal genome analysis and interpretation. Nature Methods, 2015, 12, 966-968.	19.0	515
23	MOSAİK: A Hash-Based Algorithm for Accurate Next-Generation Sequencing Short-Read Mapping. PLoS ONE, 2014, 9, e90581.	2.5	249
24	Integrative Annotation of Variants from 1092 Humans: Application to Cancer Genomics. Science, 2013, 342, 1235587.	12.6	341
25	SSW Library: An SIMD Smith-Waterman C/C++ Library for Use in Genomic Applications. PLoS ONE, 2013, 8, e82138.	2.5	175
26	A Comprehensive Map of Mobile Element Insertion Polymorphisms in Humans. PLoS Genetics, 2011, 7, e1002236.	3.5	278