Chen-Shan Chin

List of Publications by Year in descending order

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Version: 2024-02-01

43 papers

17,688 citations

33 h-index 243625 44 g-index

67 all docs

67 docs citations

67 times ranked

23257 citing authors

#	Article	IF	CITATIONS
1	Curated variation benchmarks for challenging medically relevant autosomal genes. Nature Biotechnology, 2022, 40, 672-680.	17.5	90
2	Ten simple rules for large-scale data processing. PLoS Computational Biology, 2022, 18, e1009757.	3.2	1
3	A complete reference genome improves analysis of human genetic variation. Science, 2022, 376, eabl3533.	12.6	144
4	The complete sequence of a human genome. Science, 2022, 376, 44-53.	12.6	1,222
5	Benchmarking challenging small variants with linked and long reads. Cell Genomics, 2022, 2, 100128.	6.5	77
6	Chromosome-scale, haplotype-resolved assembly of human genomes. Nature Biotechnology, 2021, 39, 309-312.	17.5	109
7	AnÂinternationalÂvirtualÂhackathon toÂbuildÂtools for theÂanalysis ofÂstructuralÂvariants withinÂspeciesÂranging fromÂcoronaviruses toÂvertebrates. F1000Research, 2021, 10, 246.	1.6	3
8	A draft sequence reference of the Psilocybe cubensis genome. F1000Research, 2021, 10, 281.	1.6	1
9	A draft reference assembly of the Psilocybe cubensis genome. F1000Research, 2021, 10, 281.	1.6	1
10	AnÂinternationalÂvirtualÂhackathon toÂbuildÂtools for theÂanalysis ofÂstructuralÂvariants withinÂspeciesÂranging fromÂcoronaviruses toÂvertebrates. F1000Research, 2021, 10, 246.	1.6	2
11	Amplification-free long-read sequencing reveals unforeseen CRISPR-Cas9 off-target activity. Genome Biology, 2020, 21, 290.	8.8	35
12	A diploid assembly-based benchmark for variants in the major histocompatibility complex. Nature Communications, 2020, 11, 4794.	12.8	56
13	Effect of sequence depth and length in long-read assembly of the maize inbred NC358. Nature Communications, 2020, 11, 2288.	12.8	39
14	Accurate circular consensus long-read sequencing improves variant detection and assembly of a human genome. Nature Biotechnology, 2019, 37, 1155-1162.	17.5	1,010
15	Multi-platform discovery of haplotype-resolved structural variation in human genomes. Nature Communications, 2019, 10, 1784.	12.8	636
16	Complex rearrangements and oncogene amplifications revealed by long-read DNA and RNA sequencing of a breast cancer cell line. Genome Research, 2018, 28, 1126-1135.	5.5	142
17	Comprehensive analysis of single molecule sequencing-derived complete genome and whole transcriptome of Hyposidra talaca nuclear polyhedrosis virus. Scientific Reports, 2018, 8, 8924.	3.3	8
18	Evaluation of GRCh38 and de novo haploid genome assemblies demonstrates the enduring quality of the reference assembly. Genome Research, 2017, 27, 849-864.	5.5	728

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19	Heterogeneous resistance to quizartinib in acute myeloid leukemia revealed by single-cell analysis. Blood, 2017, 130, 48-58.	1.4	143
20	Improved maize reference genome with single-molecule technologies. Nature, 2017, 546, 524-527.	27.8	1,113
21	De novo PacBio long-read and phased avian genome assemblies correct and add to reference genes generated with intermediate and short reads. GigaScience, 2017, 6, 1-16.	6.4	165
22	Scaffolding of long read assemblies using long range contact information. BMC Genomics, 2017, 18, 527.	2.8	194
23	Chromosomal-Level Assembly of the Asian Seabass Genome Using Long Sequence Reads and Multi-layered Scaffolding. PLoS Genetics, 2016, 12, e1005954.	3.5	105
24	Long-read sequence assembly of the gorilla genome. Science, 2016, 352, aae0344.	12.6	368
25	Phased diploid genome assembly with single-molecule real-time sequencing. Nature Methods, 2016, 13, 1050-1054.	19.0	1,658
26	HLA Typing for the Next Generation. PLoS ONE, 2015, 10, e0127153.	2.5	125
27	Assembling large genomes with single-molecule sequencing and locality-sensitive hashing. Nature Biotechnology, 2015, 33, 623-630.	17.5	877
28	Assembly and diploid architecture of an individual human genome via single-molecule technologies. Nature Methods, 2015, 12, 780-786.	19.0	465
29	Extending reference assembly models. Genome Biology, 2015, 16, 13.	8.8	139
30	Long-read, whole-genome shotgun sequence data for five model organisms. Scientific Data, 2014, 1, 140045.	5.3	138
31	Nonhybrid, finished microbial genome assemblies from long-read SMRT sequencing data. Nature Methods, 2013, 10, 563-569.	19.0	4,029
32	A hybrid approach for the automated finishing of bacterial genomes. Nature Biotechnology, 2012, 30, 701-707.	17.5	178
33	Validation of ITD mutations in FLT3 as a therapeutic target in human acute myeloid leukaemia. Nature, 2012, 485, 260-263.	27.8	641
34	Origins of the <i>E. coli</i> Strain Causing an Outbreak of Hemolyticâ€"Uremic Syndrome in Germany. New England Journal of Medicine, 2011, 365, 709-717.	27.0	778
35	The Origin of the Haitian Cholera Outbreak Strain. New England Journal of Medicine, 2011, 364, 33-42.	27.0	676
36	A flexible and efficient template format for circular consensus sequencing and SNP detection. Nucleic Acids Research, 2010, 38, e159-e159.	14.5	377

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37	Dynamics and Design Principles of a Basic Regulatory Architecture Controlling Metabolic Pathways. PLoS Biology, 2008, 6, e146.	5.6	43
38	Global identification of noncoding RNAs in Saccharomyces cerevisiae by modulating an essential RNA processing pathway. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 4192-4197.	7.1	69
39	Genome-wide regulatory complexity in yeast promoters: Separation of functionally conserved and neutral sequence. Genome Research, 2005, 15, 205-213.	5. 5	43
40	Comparing genomic expression patterns across species identifies shared transcriptional profile in aging. Nature Genetics, 2004, 36, 197-204.	21.4	434
41	Passive random walkers and riverlike networks on growing surfaces. Physical Review E, 2002, 66, 021104.	2.1	22
42	Reconstructed rough growing interfaces: Ridge-line trapping of domain walls. Physical Review E, 2001, 64, 031606.	2.1	1
43	Stationary-state skewness in two-dimensional Kardar-Parisi-Zhang type growth. Physical Review E, 1999, 59, 2633-2641.	2.1	40