

Kai Ye

List of Publications by Year in descending order

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Version: 2024-02-01

58
papers

15,170
citations

212478

28
h-index

169272

56
g-index

72
all docs

72
docs citations

72
times ranked

33452
citing authors

#	ARTICLE	IF	CITATIONS
1	Homotopic Convex Transformation: A New Landscape Smoothing Method for the Traveling Salesman Problem. <i>IEEE Transactions on Cybernetics</i> , 2022, 52, 495-507.	6.2	7
2	Mako: A Graph-based Pattern Growth Approach to Detect Complex Structural Variants. <i>Genomics, Proteomics and Bioinformatics</i> , 2022, 20, 205-218.	3.0	6
3	High-quality <i>Arabidopsis thaliana</i> Genome Assembly with Nanopore and HiFi Long Reads. <i>Genomics, Proteomics and Bioinformatics</i> , 2022, 20, 4-13.	3.0	80
4	A global survey of the transcriptome of the opium poppy (<i>Papaver somniferum</i>) based on single-molecule long-read isoform sequencing. <i>Plant Journal</i> , 2022, 110, 607-620.	2.8	5
5	JAX-CNV: A Whole-genome Sequencing-based Algorithm for Copy Number Detection at Clinical Grade Level. <i>Genomics, Proteomics and Bioinformatics</i> , 2022, 20, 1197-1206.	3.0	3
6	Integrating bulk and single-cell RNA sequencing reveals cellular heterogeneity and immune infiltration in hepatocellular carcinoma. <i>Molecular Oncology</i> , 2022, 16, 2195-2213.	2.1	16
7	IAGS: Inferring Ancestor Genome Structure under a Wide Range of Evolutionary Scenarios. <i>Molecular Biology and Evolution</i> , 2022, 39, .	3.5	5
8	Cellular heterogeneity and transcriptomic profiles during intrahepatic cholangiocarcinoma initiation and progression. <i>Hepatology</i> , 2022, 76, 1302-1317.	3.6	13
9	Haplotype-resolved Chinese male genome assembly based on high-fidelity sequencing. <i>Fundamental Research</i> , 2022, 2, 946-953.	1.6	11
10	A novel CD4+ CTL subtype characterized by chemotaxis and inflammation is involved in the pathogenesis of Graves' orbitopathy. <i>Cellular and Molecular Immunology</i> , 2021, 18, 735-745.	4.8	37
11	Application of Metagenomic Next-Generation Sequencing in the Diagnosis of Pulmonary Infectious Pathogens From Bronchoalveolar Lavage Samples. <i>Frontiers in Cellular and Infection Microbiology</i> , 2021, 11, 541092.	1.8	72
12	Haplotype-resolved diverse human genomes and integrated analysis of structural variation. <i>Science</i> , 2021, 372, .	6.0	358
13	Three chromosome-scale <i>Papaver</i> genomes reveal punctuated patchwork evolution of the morphinan and noscapine biosynthesis pathway. <i>Nature Communications</i> , 2021, 12, 6030.	5.8	51
14	DrugEx v2: de novo design of drug molecules by Pareto-based multi-objective reinforcement learning in polypharmacology. <i>Journal of Cheminformatics</i> , 2021, 13, 85.	2.8	30
15	Cerebrospinal Fluid from Healthy Pregnant Women Does Not Harbor a Detectable Microbial Community. <i>Microbiology Spectrum</i> , 2021, 9, e0076921.	1.2	5
16	Multifactorial Deep Learning Reveals Pan-Cancer Genomic Tumor Clusters with Distinct Immunogenomic Landscape and Response to Immunotherapy. <i>Clinical Cancer Research</i> , 2020, 26, 2908-2920.	3.2	30
17	Chromosome-Scale Genome Assembly of <i>Talaromyces rugulosus</i> W13939, a Mycoparasitic Fungus and Promising Biocontrol Agent. <i>Molecular Plant-Microbe Interactions</i> , 2020, 33, 1446-1450.	1.4	4
18	Predicting the early risk of ophthalmopathy in Graves' disease patients using TCR repertoire. <i>Clinical and Translational Medicine</i> , 2020, 10, e218.	1.7	2

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19	gCAnno: a graph-based single cell type annotation method. BMC Genomics, 2020, 21, 823.	1.2	0
20	A Chromosome-Scale Genome Assembly for the <i>Fusarium oxysporum</i> Strain Fo5176 To Establish a Model <i>Arabidopsis</i> -Fungal Pathosystem. G3: Genes, Genomes, Genetics, 2020, 10, 3549-3555.	0.8	28
21	Transportation, germs, culture: a dynamic graph model of COVID-19 outbreak. Quantitative Biology, 2020, 8, 238-244.	0.3	4
22	MSIsensor-pro: Fast, Accurate, and Matched-normal-sample-free Detection of Microsatellite Instability. Genomics, Proteomics and Bioinformatics, 2020, 18, 65-71.	3.0	53
23	Chromosome-Scale Genome Assembly of <i>Fusarium oxysporum</i> Strain Fo47, a Fungal Endophyte and Biocontrol Agent. Molecular Plant-Microbe Interactions, 2020, 33, 1108-1111.	1.4	29
24	Dynamic network inference and association computation discover gene modules regulating virulence, mycotoxin and sexual reproduction in <i>Fusarium graminearum</i> . BMC Genomics, 2020, 21, 179.	1.2	8
25	What Is Required to Prevent a Second Major Outbreak of SARS-CoV-2 upon Lifting Quarantine in Wuhan City, China. Innovation(China), 2020, 1, 100006.	5.2	32
26	From Innovations to Prospects. , 2020, , .		3
27	MEpurity: estimating tumor purity using DNA methylation data. Bioinformatics, 2019, 35, 5298-5300.	1.8	8
28	Mapping Genome Variants Sheds Light on Genetic and Phenotypic Differentiation in Chinese. Genomics, Proteomics and Bioinformatics, 2019, 17, 226-228.	3.0	1
29	One reference genome is not enough. Genome Biology, 2019, 20, 104.	3.8	58
30	An exploration strategy improves the diversity of de novo ligands using deep reinforcement learning: a case for the adenosine A2A receptor. Journal of Cheminformatics, 2019, 11, 35.	2.8	58
31	Multi-platform discovery of haplotype-resolved structural variation in human genomes. Nature Communications, 2019, 10, 1784.	5.8	636
32	PVTree: A Sequential Pattern Mining Method for Alignment Independent Phylogeny Reconstruction. Genes, 2019, 10, 73.	1.0	6
33	PRESM: personalized reference editor for somatic mutation discovery in cancer genomics. Bioinformatics, 2019, 35, 1445-1452.	1.8	6
34	The opium poppy genome and morphinan production. Science, 2018, 362, 343-347.	6.0	225
35	Split-Read Indel and Structural Variant Calling Using PINDEL. Methods in Molecular Biology, 2018, 1833, 95-105.	0.4	20
36	Cold atmospheric plasma as a potential tool for multiple myeloma treatment. Oncotarget, 2018, 9, 18002-18017.	0.8	28

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37	An improved burden-test pipeline for identifying associations from rare germline and somatic variants. <i>BMC Genomics</i> , 2017, 18, 753.	1.2	11
38	Divergent viral presentation among human tumors and adjacent normal tissues. <i>Scientific Reports</i> , 2016, 6, 28294.	1.6	60
39	Precision Medicine: What Challenges Are We Facing?. <i>Genomics, Proteomics and Bioinformatics</i> , 2016, 14, 253-261.	3.0	15
40	A high-quality human reference panel reveals the complexity and distribution of genomic structural variants. <i>Nature Communications</i> , 2016, 7, 12989.	5.8	99
41	Systematic discovery of complex insertions and deletions in human cancers. <i>Nature Medicine</i> , 2016, 22, 97-104.	15.2	93
42	Patterns and functional implications of rare germline variants across 12 cancer types. <i>Nature Communications</i> , 2015, 6, 10086.	5.8	243
43	Characteristics of de novo structural changes in the human genome. <i>Genome Research</i> , 2015, 25, 792-801.	2.4	115
44	Genome-wide patterns and properties of de novo mutations in humans. <i>Nature Genetics</i> , 2015, 47, 822-826.	9.4	384
45	An integrated map of structural variation in 2,504 human genomes. <i>Nature</i> , 2015, 526, 75-81.	13.7	1,994
46	A gain of function mutation in <i>TNFRSF11B</i> encoding osteoprotegerin causes osteoarthritis with chondrocalcinosis. <i>Annals of the Rheumatic Diseases</i> , 2015, 74, 1756-1762.	0.5	44
47	The Genome of the Netherlands: design, and project goals. <i>European Journal of Human Genetics</i> , 2014, 22, 221-227.	1.4	246
48	MSIsensor: microsatellite instability detection using paired tumor-normal sequence data. <i>Bioinformatics</i> , 2014, 30, 1015-1016.	1.8	599
49	Mutational landscape and significance across 12 major cancer types. <i>Nature</i> , 2013, 502, 333-339.	13.7	3,695
50	Aging as Accelerated Accumulation of Somatic Variants: Whole-Genome Sequencing of Centenarian and Middle-Aged Monozygotic Twin Pairs. <i>Twin Research and Human Genetics</i> , 2013, 16, 1026-1032.	0.3	40
51	A Systematic Survey of Loss-of-Function Variants in Human Protein-Coding Genes. <i>Science</i> , 2012, 335, 823-828.	6.0	1,095
52	Mapping copy number variation by population-scale genome sequencing. <i>Nature</i> , 2011, 470, 59-65.	13.7	991
53	A comprehensive catalogue of somatic mutations from a human cancer genome. <i>Nature</i> , 2010, 463, 191-196.	13.7	1,519
54	Mining Unique-m Substrings from Genomes. <i>Journal of Proteomics and Bioinformatics</i> , 2010, 03, 099-103.	0.4	4

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55	Pindel: a pattern growth approach to detect break points of large deletions and medium sized insertions from paired-end short reads. <i>Bioinformatics</i> , 2009, 25, 2865-2871.	1.8	1,811
56	Tracing evolutionary pressure. <i>Bioinformatics</i> , 2008, 24, 908-915.	1.8	35
57	An efficient, versatile and scalable pattern growth approach to mine frequent patterns in unaligned protein sequences. <i>Bioinformatics</i> , 2007, 23, 687-693.	1.8	23
58	A two-entropies analysis to identify functional positions in the transmembrane region of class A G protein-coupled receptors. <i>Proteins: Structure, Function and Bioinformatics</i> , 2006, 63, 1018-1030.	1.5	35