

## List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Mutational landscape and significance across 12 major cancer types. Nature, 2013, 502, 333-339.	27.8	3,695
2	An integrated map of structural variation in 2,504 human genomes. Nature, 2015, 526, 75-81.	27.8	1,994
3	Pindel: a pattern growth approach to detect break points of large deletions and medium sized insertions from paired-end short reads. Bioinformatics, 2009, 25, 2865-2871.	4.1	1,811
4	A comprehensive catalogue of somatic mutations from a human cancer genome. Nature, 2010, 463, 191-196.	27.8	1,519
5	A Systematic Survey of Loss-of-Function Variants in Human Protein-Coding Genes. Science, 2012, 335, 823-828.	12.6	1,095
6	Mapping copy number variation by population-scale genome sequencing. Nature, 2011, 470, 59-65.	27.8	991
7	Multi-platform discovery of haplotype-resolved structural variation in human genomes. Nature Communications, 2019, 10, 1784.	12.8	636
8	MSIsensor: microsatellite instability detection using paired tumor-normal sequence data. Bioinformatics, 2014, 30, 1015-1016.	4.1	599
9	Genome-wide patterns and properties of de novo mutations in humans. Nature Genetics, 2015, 47, 822-826.	21.4	384
10	Haplotype-resolved diverse human genomes and integrated analysis of structural variation. Science, 2021, 372, .	12.6	358
11	The Genome of the Netherlands: design, and project goals. European Journal of Human Genetics, 2014, 22, 221-227.	2.8	246
12	Patterns and functional implications of rare germline variants across 12 cancer types. Nature Communications, 2015, 6, 10086.	12.8	243
13	The opium poppy genome and morphinan production. Science, 2018, 362, 343-347.	12.6	225
14	Characteristics of de novo structural changes in the human genome. Genome Research, 2015, 25, 792-801.	5.5	115
15	A high-quality human reference panel reveals the complexity and distribution of genomic structural variants. Nature Communications, 2016, 7, 12989.	12.8	99
16	Systematic discovery of complex insertions and deletions in human cancers. Nature Medicine, 2016, 22, 97-104.	30.7	93
17	High-quality Arabidopsis thaliana Genome Assembly with Nanopore and HiFi Long Reads. Genomics, Proteomics and Bioinformatics, 2022, 20, 4-13.	6.9	80
18	Application of Metagenomic Next-Generation Sequencing in the Diagnosis of Pulmonary Infectious Pathogens From Bronchoalveolar Lavage Samples. Frontiers in Cellular and Infection Microbiology, 2021, 11, 541092.	3.9	72

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19	Divergent viral presentation among human tumors and adjacent normal tissues. Scientific Reports, 2016, 6, 28294.	3.3	60
20	One reference genome is not enough. Genome Biology, 2019, 20, 104.	8.8	58
21	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.	6.1	58
22	MSIsensor-pro: Fast, Accurate, and Matched-normal-sample-free Detection of Microsatellite Instability. Genomics, Proteomics and Bioinformatics, 2020, 18, 65-71.	6.9	53
23	Three chromosome-scale Papaver genomes reveal punctuated patchwork evolution of the morphinan and noscapine biosynthesis pathway. Nature Communications, 2021, 12, 6030.	12.8	51
24	A gain of function mutation in <i>TNFRSF11B</i> encoding osteoprotegerin causes osteoarthritis with chondrocalcinosis. Annals of the Rheumatic Diseases, 2015, 74, 1756-1762.	0.9	44
25	Aging as Accelerated Accumulation of Somatic Variants: Whole-Genome Sequencing of Centenarian and Middle-Aged Monozygotic Twin Pairs. Twin Research and Human Genetics, 2013, 16, 1026-1032.	0.6	40
26	A novel CD4+ CTL subtype characterized by chemotaxis and inflammation is involved in the pathogenesis of Graves' orbitopathy. Cellular and Molecular Immunology, 2021, 18, 735-745.	10.5	37
27	A two-entropies analysis to identify functional positions in the transmembrane region of class A G protein-coupled receptors. Proteins: Structure, Function and Bioinformatics, 2006, 63, 1018-1030.	2.6	35
28	Tracing evolutionary pressure. Bioinformatics, 2008, 24, 908-915.	4.1	35
29	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.	9.1	32
30	Multifactorial Deep Learning Reveals Pan-Cancer Genomic Tumor Clusters with Distinct Immunogenomic Landscape and Response to Immunotherapy. Clinical Cancer Research, 2020, 26, 2908-2920.	7.0	30
31	DrugEx v2: de novo design of drug molecules by Pareto-based multi-objective reinforcement learning in polypharmacology. Journal of Cheminformatics, 2021, 13, 85.	6.1	30
32	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.	2.6	29
33	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.	1.8	28
34	Cold atmospheric plasma as a potential tool for multiple myeloma treatment. Oncotarget, 2018, 9, 18002-18017.	1.8	28
35	An efficient, versatile and scalable pattern growth approach to mine frequent patterns in unaligned protein sequences. Bioinformatics, 2007, 23, 687-693.	4.1	23
36	Split-Read Indel and Structural Variant Calling Using PINDEL. Methods in Molecular Biology, 2018, 1833, 95-105.	0.9	20

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37	Integrating bulk and singleâ€cell RNA sequencing reveals cellular heterogeneity and immune infiltration in hepatocellular carcinoma. Molecular Oncology, 2022, 16, 2195-2213.	4.6	16
38	Precision Medicine: What Challenges Are We Facing?. Genomics, Proteomics and Bioinformatics, 2016, 14, 253-261.	6.9	15
39	Cellular heterogeneity and transcriptomic profiles during intrahepatic cholangiocarcinoma initiation and progression. Hepatology, 2022, 76, 1302-1317.	7.3	13
40	An improved burden-test pipeline for identifying associations from rare germline and somatic variants. BMC Genomics, 2017, 18, 753.	2.8	11
41	Haplotype-resolved Chinese male genome assembly based on high-fidelity sequencing. Fundamental Research, 2022, 2, 946-953.	3.3	11
42	MEpurity: estimating tumor purity using DNA methylation data. Bioinformatics, 2019, 35, 5298-5300.	4.1	8
43	Dynamic network inference and association computation discover gene modules regulating virulence, mycotoxin and sexual reproduction in Fusarium graminearum. BMC Genomics, 2020, 21, 179.	2.8	8
44	Homotopic Convex Transformation: A New Landscape Smoothing Method for the Traveling Salesman Problem. IEEE Transactions on Cybernetics, 2022, 52, 495-507.	9.5	7
45	PVTree: A Sequential Pattern Mining Method for Alignment Independent Phylogeny Reconstruction. Genes, 2019, 10, 73.	2.4	6
46	PRESM: personalized reference editor for somatic mutation discovery in cancer genomics. Bioinformatics, 2019, 35, 1445-1452.	4.1	6
47	Mako: A Graph-based Pattern Growth Approach to Detect Complex Structural Variants. Genomics, Proteomics and Bioinformatics, 2022, 20, 205-218.	6.9	6
48	A global survey of the transcriptome of the opium poppy ( <i>Papaver somniferum</i> ) based on singleâ€molecule longâ€read isoform sequencing. Plant Journal, 2022, 110, 607-620.	5.7	5
49	IAGS: Inferring Ancestor Genome Structure under a Wide Range of Evolutionary Scenarios. Molecular Biology and Evolution, 2022, 39, .	8.9	5
50	Cerebrospinal Fluid from Healthy Pregnant Women Does Not Harbor a Detectable Microbial Community. Microbiology Spectrum, 2021, 9, e0076921.	3.0	5
51	Chromosome-Scale Genome Assembly of <i>Talaromyces rugulosus</i> W13939, a Mycoparasitic Fungus and Promising Biocontrol Agent. Molecular Plant-Microbe Interactions, 2020, 33, 1446-1450.	2.6	4
52	Transportation, germs, culture: a dynamic graph model of COVIDâ€19 outbreak. Quantitative Biology, 2020, 8, 238-244.	0.5	4
53	Mining Unique-m Substrings from Genomes. Journal of Proteomics and Bioinformatics, 2010, 03, 099-103.	0.4	4

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55	JAX-CNV: A Whole-genome Sequencing-based Algorithm for Copy Number Detection at Clinical Grade Level. Genomics, Proteomics and Bioinformatics, 2022, 20, 1197-1206.	6.9	3
56	Predicting the early risk of ophthalmopathy in Graves' disease patients using TCR repertoire. Clinical and Translational Medicine, 2020, 10, e218.	4.0	2
57	Mapping Genome Variants Sheds Light on Genetic and Phenotypic Differentiation in Chinese. Genomics, Proteomics and Bioinformatics, 2019, 17, 226-228.	6.9	1
58	gCAnno: a graph-based single cell type annotation method. BMC Genomics, 2020, 21, 823.	2.8	0