

Kin Fai Au

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6808370/publications.pdf>

Version: 2024-02-01

28
papers

4,996
citations

430442

18
h-index

500791

28
g-index

31
all docs

31
docs citations

31
times ranked

7120
citing authors

#	ARTICLE	IF	CITATIONS
1	PacBio Sequencing and Its Applications. <i>Genomics, Proteomics and Bioinformatics</i> , 2015, 13, 278-289.	3.0	1,669
2	Nanopore sequencing technology, bioinformatics and applications. <i>Nature Biotechnology</i> , 2021, 39, 1348-1365.	9.4	521
3	Comprehensive comparison of Pacific Biosciences and Oxford Nanopore Technologies and their applications to transcriptome analysis. <i>F1000Research</i> , 2017, 6, 100.	0.8	366
4	Full-length transcriptome sequences and splice variants obtained by a combination of sequencing platforms applied to different root tissues of <i>Salvia miltiorrhiza</i> and tanshinone biosynthesis. <i>Plant Journal</i> , 2015, 82, 951-961.	2.8	337
5	Characterization of the human ESC transcriptome by hybrid sequencing. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, E4821-30.	3.3	316
6	Detection of splice junctions from paired-end RNA-seq data by SpliceMap. <i>Nucleic Acids Research</i> , 2010, 38, 4570-4578.	6.5	300
7	Improving PacBio Long Read Accuracy by Short Read Alignment. <i>PLoS ONE</i> , 2012, 7, e46679.	1.1	289
8	Gaining comprehensive biological insight into the transcriptome by performing a broad-spectrum RNA-seq analysis. <i>Nature Communications</i> , 2017, 8, 59.	5.8	225
9	Comprehensive comparison of Pacific Biosciences and Oxford Nanopore Technologies and their applications to transcriptome analysis. <i>F1000Research</i> , 2017, 6, 100.	0.8	203
10	The primate-specific noncoding RNA HPAT5 regulates pluripotency during human preimplantation development and nuclear reprogramming. <i>Nature Genetics</i> , 2016, 48, 44-52.	9.4	153
11	Characterization of fusion genes and the significantly expressed fusion isoforms in breast cancer by hybrid sequencing. <i>Nucleic Acids Research</i> , 2015, 43, e116-e116.	6.5	104
12	A comparative evaluation of hybrid error correction methods for error-prone long reads. <i>Genome Biology</i> , 2019, 20, 26.	3.8	86
13	E-C coupling structural protein junctophilin-2 encodes a stress-adaptive transcription regulator. <i>Science</i> , 2018, 362, .	6.0	78
14	Single-molecule long-read sequencing reveals the chromatin basis of gene expression. <i>Genome Research</i> , 2019, 29, 1329-1342.	2.4	46
15	Single-molecule long-read sequencing reveals a conserved intact long RNA profile in sperm. <i>Nature Communications</i> , 2021, 12, 1361.	5.8	43
16	IDP-ASE: haplotyping and quantifying allele-specific expression at the gene and gene isoform level by hybrid sequencing. <i>Nucleic Acids Research</i> , 2017, 45, e32-e32.	6.5	42
17	IDP-denovo: <i>de novo</i> transcriptome assembly and isoform annotation by hybrid sequencing. <i>Bioinformatics</i> , 2018, 34, 2168-2176.	1.8	41
18	Real-time mapping of nanopore raw signals. <i>Bioinformatics</i> , 2021, 37, i477-i483.	1.8	41

#	ARTICLE	IF	CITATIONS
19	Single cell expression analysis of primate-specific retroviruses-derived HPAT lincRNAs in viable human blastocysts identifies embryonic cells co-expressing genetic markers of multiple lineages. <i>Heliyon</i> , 2018, 4, e00667.	1.4	23
20	Hybrid Sequencing of Full-Length cDNA Transcripts of Stems and Leaves in <i>Dendrobium officinale</i> . <i>Genes</i> , 2017, 8, 257.	1.0	20
21	iASPP mediates p53 selectivity through a modular mechanism fine-tuning DNA recognition. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 17470-17479.	3.3	20
22	The transcriptome of human pluripotent stem cells. <i>Current Opinion in Genetics and Development</i> , 2014, 28, 71-77.	1.5	14
23	Performance difference of graph-based and alignment-based hybrid error correction methods for error-prone long reads. <i>Genome Biology</i> , 2020, 21, 14.	3.8	8
24	The blooming of long-read sequencing reforms biomedical research. <i>Genome Biology</i> , 2022, 23, 21.	3.8	8
25	Revealing tumor heterogeneity of breast cancer by utilizing the linkage between somatic and germline mutations. <i>Briefings in Bioinformatics</i> , 2019, 20, 2306-2315.	3.2	4
26	A Statistical Method for Observing Personal Diploid Methylomes and Transcriptomes with Single-Molecule Real-Time Sequencing. <i>Genes</i> , 2018, 9, 460.	1.0	2
27	A network-based computational framework to predict and differentiate functions for gene isoforms using exon-level expression data. <i>Methods</i> , 2021, 189, 54-64.	1.9	2
28	Accurate Mapping of RNA-Seq Data. <i>Methods in Molecular Biology</i> , 2015, 1269, 147-161.	0.4	2