Kasper D Hansen

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/12003270/publications.pdf

Version: 2024-02-01

331670 642732 9,266 23 21 23 h-index citations g-index papers 33 33 33 21018 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Human methylome variation across Infinium 450K data on the Gene Expression Omnibus. NAR Genomics and Bioinformatics, 2021, 3, Iqab025.	3.2	12
2	recount3: summaries and queries for large-scale RNA-seq expression and splicing. Genome Biology, 2021, 22, 323.	8.8	103
3	Neuronal brain-region-specific DNA methylation and chromatin accessibility are associated with neuropsychiatric trait heritability. Nature Neuroscience, 2019, 22, 307-316.	14.8	120
4	Preprocessing, normalization and integration of the Illumina HumanMethylationEPIC array with minfi. Bioinformatics, 2017, 33, 558-560.	4.1	583
5	Reproducible RNA-seq analysis using recount2. Nature Biotechnology, 2017, 35, 319-321.	17.5	395
6	A ketogenic diet rescues hippocampal memory defects in a mouse model of Kabuki syndrome. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 125-130.	7.1	102
7	Software for the Integration of Multiomics Experiments in Bioconductor. Cancer Research, 2017, 77, e39-e42.	0.9	80
8	Human splicing diversity and the extent of unannotated splice junctions across human RNA-seq samples on the Sequence Read Archive. Genome Biology, 2016, 17, 266.	8.8	94
9	DNA methylation is stable during replication and cell cycle arrest. Scientific Reports, 2016, 5, 17911.	3.3	44
10	Rail-dbGaP: analyzing dbGaP-protected data in the cloud with Amazon Elastic MapReduce. Bioinformatics, 2016, 32, 2551-2553.	4.1	5
11	Public data and open source tools for multi-assay genomic investigation of disease. Briefings in Bioinformatics, 2016, 17, 603-615.	6.5	46
12	Reconstructing A/B compartments as revealed by Hi-C using long-range correlations in epigenetic data. Genome Biology, 2015, 16, 180.	8.8	232
13	Orchestrating high-throughput genomic analysis with Bioconductor. Nature Methods, 2015, 12, 115-121.	19.0	3,070
14	Coverage recommendations for methylation analysis by whole-genome bisulfite sequencing. Nature Methods, 2015, 12, 230-232.	19.0	248
15	Large-scale hypomethylated blocks associated with Epstein-Barr virus–induced B-cell immortalization. Genome Research, 2014, 24, 177-184.	5.5	130
16	Differential expression analysis of RNA-seq data at single-base resolution. Biostatistics, 2014, 15, 413-426.	1.5	56
17	Removing technical variability in RNA-seq data using conditional quantile normalization. Biostatistics, 2012, 13, 204-216.	1.5	532
18	Social environment is associated with gene regulatory variation in the rhesus macaque immune system. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 6490-6495.	7.1	257

#	Article	IF	CITATIONS
19	BSmooth: from whole genome bisulfite sequencing reads to differentially methylated regions. Genome Biology, 2012, 13, R83.	9.6	650
20	Sequencing technology does not eliminate biological variability. Nature Biotechnology, 2011, 29, 572-573.	17.5	193
21	Evaluation of statistical methods for normalization and differential expression in mRNA-Seq experiments. BMC Bioinformatics, 2010, 11, 94.	2.6	1,421
22	Biases in Illumina transcriptome sequencing caused by random hexamer priming. Nucleic Acids Research, 2010, 38, e131-e131.	14.5	573
23	Cloud-scale RNA-sequencing differential expression analysis with Myrna. Genome Biology, 2010, 11, R83.	9.6	268