

Jakob Skou Pedersen

List of Publications by Year in descending order

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Version: 2024-02-01

66
papers

21,834
citations

66250

44
h-index

124990

64
g-index

81
all docs

81
docs citations

81
times ranked

37974
citing authors

#	ARTICLE	IF	CITATIONS
1	Error Characterization and Statistical Modeling Improves Circulating Tumor DNA Detection by Droplet Digital PCR. <i>Clinical Chemistry</i> , 2022, 68, 657-667.	1.5	9
2	The transcriptional landscape and biomarker potential of circular RNAs in prostate cancer. <i>Genome Medicine</i> , 2022, 14, 8.	3.6	19
3	miRNA activity inferred from single cell mRNA expression. <i>Scientific Reports</i> , 2021, 11, 9170.	1.6	16
4	The landscape and driver potential of site-specific hotspots across cancer genomes. <i>Npj Genomic Medicine</i> , 2021, 6, 33.	1.7	8
5	Transcriptome-wide profiles of circular RNA and RNA-binding protein interactions reveal effects on circular RNA biogenesis and cancer pathway expression. <i>Genome Medicine</i> , 2020, 12, 112.	3.6	106
6	Molecular correlates of cisplatin-based chemotherapy response in muscle invasive bladder cancer by integrated multi-omics analysis. <i>Nature Communications</i> , 2020, 11, 4858.	5.8	124
7	Retrospective evaluation of whole exome and genome mutation calls in 746 cancer samples. <i>Nature Communications</i> , 2020, 11, 4748.	5.8	27
8	Epigenetic and transcriptomic consequences of excess X chromosome material in 47,XXX females. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2020, 184, 279-293.	0.7	21
9	Passenger Mutations in More Than 2,500 Cancer Genomes: Overall Molecular Functional Impact and Consequences. <i>Cell</i> , 2020, 180, 915-927.e16.	13.5	98
10	Pathway and network analysis of more than 2500 whole cancer genomes. <i>Nature Communications</i> , 2020, 11, 729.	5.8	73
11	Analyses of non-coding somatic drivers in 2,658 cancer whole genomes. <i>Nature</i> , 2020, 578, 102-111.	13.7	424
12	Cancer LncRNA Census reveals evidence for deep functional conservation of long noncoding RNAs in tumorigenesis. <i>Communications Biology</i> , 2020, 3, 56.	2.0	140
13	ncdDetect2: improved models of the site-specific mutation rate in cancer and driver detection with robust significance evaluation. <i>Bioinformatics</i> , 2019, 35, 189-199.	1.8	6
14	Deficiency of nucleotide excision repair is associated with mutational signature observed in cancer. <i>Genome Research</i> , 2019, 29, 1067-1077.	2.4	66
15	EBADIMEX: an empirical Bayes approach to detect joint differential expression and methylation and to classify samples. <i>Statistical Applications in Genetics and Molecular Biology</i> , 2019, 18, .	0.2	0
16	Pan-cancer screen for mutations in non-coding elements with conservation and cancer specificity reveals correlations with expression and survival. <i>Npj Genomic Medicine</i> , 2018, 3, 1.	1.7	79
17	Optimized targeted sequencing of cell-free plasma DNA from bladder cancer patients. <i>Scientific Reports</i> , 2018, 8, 1917.	1.6	50
18	A site specific model and analysis of the neutral somatic mutation rate in whole-genome cancer data. <i>BMC Bioinformatics</i> , 2018, 19, 147.	1.2	11

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19	Regmex: a statistical tool for exploring motifs in ranked sequence lists from genomics experiments. <i>Algorithms for Molecular Biology</i> , 2018, 13, 17.	0.3	5
20	DNA hypermethylation and differential gene expression associated with Klinefelter syndrome. <i>Scientific Reports</i> , 2018, 8, 13740.	1.6	75
21	APD-Containing Cyclolipodepsipeptides Target Mitochondrial Function in Hypoxic Cancer Cells. <i>Cell Chemical Biology</i> , 2018, 25, 1337-1349.e12.	2.5	27
22	epiG: statistical inference and profiling of DNA methylation from whole-genome bisulfite sequencing data. <i>Genome Biology</i> , 2017, 18, 38.	3.8	6
23	Circular RNA expression is abundant and correlated to aggressiveness in early-stage bladder cancer. <i>Npj Genomic Medicine</i> , 2017, 2, 36.	1.7	105
24	Significance evaluation in factor graphs. <i>BMC Bioinformatics</i> , 2017, 18, 199.	1.2	2
25	RHCG and TCAF1 promoter hypermethylation predicts biochemical recurrence in prostate cancer patients treated by radical prostatectomy. <i>Oncotarget</i> , 2017, 8, 5774-5788.	0.8	22
26	Non-coding cancer driver candidates identified with a sample- and position-specific model of the somatic mutation rate. <i>ELife</i> , 2017, 6, .	2.8	36
27	SNHG16 is regulated by the Wnt pathway in colorectal cancer and affects genes involved in lipid metabolism. <i>Molecular Oncology</i> , 2016, 10, 1266-1282.	2.1	151
28	SNHG5 promotes colorectal cancer cell survival by counteracting STAU1-mediated mRNA destabilization. <i>Nature Communications</i> , 2016, 7, 13875.	5.8	170
29	ProbFold: a probabilistic method for integration of probing data in RNA secondary structure prediction. <i>Bioinformatics</i> , 2016, 32, 2626-2635.	1.8	9
30	Paired Exome Analysis Reveals Clonal Evolution and Potential Therapeutic Targets in Urothelial Carcinoma. <i>Cancer Research</i> , 2016, 76, 5894-5906.	0.4	87
31	Fast, Accurate and Automatic Ancient Nucleosome and Methylation Maps with epiPALEOMIX. <i>Molecular Biology and Evolution</i> , 2016, 33, 3284-3298.	3.5	53
32	miR-625-3p regulates oxaliplatin resistance by targeting MAP2K6-p38 signalling in human colorectal adenocarcinoma cells. <i>Nature Communications</i> , 2016, 7, 12436.	5.8	82
33	Widespread DNA hypomethylation and differential gene expression in Turner syndrome. <i>Scientific Reports</i> , 2016, 6, 34220.	1.6	106
34	Comprehensive Transcriptional Analysis of Early-Stage Urothelial Carcinoma. <i>Cancer Cell</i> , 2016, 30, 27-42.	7.7	486
35	Analysis of circulating tumour DNA to monitor disease burden following colorectal cancer surgery. <i>Cut</i> , 2016, 65, 625-634.	6.1	381
36	PINCAGE: probabilistic integration of cancer genomics data for perturbed gene identification and sample classification. <i>Bioinformatics</i> , 2016, 32, 1353-1365.	1.8	12

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37	The lncRNA MIR31HG regulates p16INK4A expression to modulate senescence. <i>Nature Communications</i> , 2015, 6, 6967.	5.8	161
38	Next-Generation Sequencing of RNA and DNA Isolated from Paired Fresh-Frozen and Formalin-Fixed Paraffin-Embedded Samples of Human Cancer and Normal Tissue. <i>PLoS ONE</i> , 2014, 9, e98187.	1.1	284
39	SNPest: a probabilistic graphical model for estimating genotypes. <i>BMC Research Notes</i> , 2014, 7, 698.	0.6	6
40	Genome-wide nucleosome map and cytosine methylation levels of an ancient human genome. <i>Genome Research</i> , 2014, 24, 454-466.	2.4	161
41	Identification of expressed and conserved human noncoding RNAs. <i>Rna</i> , 2014, 20, 236-251.	1.6	47
42	Cellular Disposal of miR23b by RAB27-Dependent Exosome Release Is Linked to Acquisition of Metastatic Properties. <i>Cancer Research</i> , 2014, 74, 5758-5771.	0.4	237
43	A Dual Program for Translation Regulation in Cellular Proliferation and Differentiation. <i>Cell</i> , 2014, 158, 1281-1292.	13.5	414
44	Mutational Context and Diverse Clonal Development in Early and Late Bladder Cancer. <i>Cell Reports</i> , 2014, 7, 1649-1663.	2.9	128
45	Abstract 3153: Whole genome and transcriptome analysis reveals novel genomic alterations in bladder cancer.. , 2013, , .		0
46	A high-resolution map of human evolutionary constraint using 29 mammals. <i>Nature</i> , 2011, 478, 476-482.	13.7	1,016
47	Tumor-specific usage of alternative transcription start sites in colorectal cancer identified by genome-wide exon array analysis. <i>BMC Genomics</i> , 2011, 12, 505.	1.2	57
48	Locating protein-coding sequences under selection for additional, overlapping functions in 29 mammalian genomes. <i>Genome Research</i> , 2011, 21, 1916-1928.	2.4	83
49	New families of human regulatory RNA structures identified by comparative analysis of vertebrate genomes. <i>Genome Research</i> , 2011, 21, 1929-1943.	2.4	100
50	Ancient human genome sequence of an extinct Palaeo-Eskimo. <i>Nature</i> , 2010, 463, 757-762.	13.7	750
51	Posttranscriptional Crossregulation between Drosha and DGCR8. <i>Cell</i> , 2009, 136, 75-84.	13.5	380
52	The UCSC genome browser database: update 2007. <i>Nucleic Acids Research</i> , 2007, 35, D668-D673.	6.5	260
53	Structured RNAs in the ENCODE selected regions of the human genome. <i>Genome Research</i> , 2007, 17, 852-864.	2.4	150
54	The UCSC Genome Browser Database: 2008 update. <i>Nucleic Acids Research</i> , 2007, 36, D773-D779.	6.5	459

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55	Editing modifies the GABAA receptor subunit $\alpha 3$. <i>Rna</i> , 2007, 13, 698-703.	1.6	184
56	Identification and analysis of functional elements in 1% of the human genome by the ENCODE pilot project. <i>Nature</i> , 2007, 447, 799-816.	13.7	4,709
57	Discovery of functional elements in 12 <i>Drosophila</i> genomes using evolutionary signatures. <i>Nature</i> , 2007, 450, 219-232.	13.7	573
58	Evolution of genes and genomes on the <i>Drosophila</i> phylogeny. <i>Nature</i> , 2007, 450, 203-218.	13.7	1,886
59	An RNA gene expressed during cortical development evolved rapidly in humans. <i>Nature</i> , 2006, 443, 167-172.	13.7	884
60	The UCSC Genome Browser Database: update 2006. <i>Nucleic Acids Research</i> , 2006, 34, D590-D598.	6.5	1,156
61	Identification and Classification of Conserved RNA Secondary Structures in the Human Genome. <i>PLoS Computational Biology</i> , 2006, 2, e33.	1.5	439
62	Forces Shaping the Fastest Evolving Regions in the Human Genome. <i>PLoS Genetics</i> , 2006, 2, e168.	1.5	399
63	Evolutionarily conserved elements in vertebrate, insect, worm, and yeast genomes. <i>Genome Research</i> , 2005, 15, 1034-1050.	2.4	3,517
64	An Evolutionary Model for Protein-Coding Regions with Conserved RNA Structure. <i>Molecular Biology and Evolution</i> , 2004, 21, 1913-1922.	3.5	36
65	A comparative method for finding and folding RNA secondary structures within protein-coding regions. <i>Nucleic Acids Research</i> , 2004, 32, 4925-4936.	6.5	82
66	Gene finding with a hidden Markov model of genome structure and evolution. <i>Bioinformatics</i> , 2003, 19, 219-227.	1.8	96