S Cenk Sahinalp

List of Publications by Year in descending order

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46918 18075 18,972 134 47 120 citations h-index g-index papers 157 157 157 34939 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	CYP2C8, CYP2C9, and CYP2C19 Characterization Using Next-Generation Sequencing and Haplotype Analysis. Journal of Molecular Diagnostics, 2022, 24, 337-350.	1.2	23
2	Analytical Validation of a Computational Method for Pharmacogenetic Genotyping from Clinical Whole Exome Sequencing. Journal of Molecular Diagnostics, 2022, 24, 576-585.	1.2	7
3	Characterizing genetic intra-tumor heterogeneity across 2,658 human cancer genomes. Cell, 2021, 184, 2239-2254.e39.	13.5	260
4	Privacy-preserving genotype imputation in a trusted execution environment. Cell Systems, 2021, 12, 983-993.e7.	2.9	13
5	Identification of conserved evolutionary trajectories in tumors. Bioinformatics, 2020, 36, i427-i435.	1.8	9
6	PhISCS-BnB: a fast branch and bound algorithm for the perfect tumor phylogeny reconstruction problem. Bioinformatics, 2020, 36, i169-i176.	1.8	19
7	Tumor Phylogeny Topology Inference via Deep Learning. IScience, 2020, 23, 101655.	1.9	11
8	Sketching algorithms for genomic data analysis and querying in a secure enclave. Nature Methods, 2020, 17, 295-301.	9.0	35
9	Graph Traversal Edit Distance and Extensions. Journal of Computational Biology, 2020, 27, 317-329.	0.8	4
10	The evolutionary history of 2,658 cancers. Nature, 2020, 578, 122-128.	13.7	690
10	The evolutionary history of 2,658 cancers. Nature, 2020, 578, 122-128. Clonal Evolution and Heterogeneity of Osimertinib Acquired Resistance Mechanisms in EGFR Mutant Lung Cancer. Cell Reports Medicine, 2020, 1, 100007.	13.7 3.3	690 78
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11	Clonal Evolution and Heterogeneity of Osimertinib Acquired Resistance Mechanisms in EGFR Mutant Lung Cancer. Cell Reports Medicine, 2020, 1, 100007. lordFAST: sensitive and Fast Alignment Search Tool for LOng noisy Read sequencing Data.	3.3	78
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11 12 13 14	Clonal Evolution and Heterogeneity of Osimertinib Acquired Resistance Mechanisms in EGFR Mutant Lung Cancer. Cell Reports Medicine, 2020, 1, 100007. lordFAST: sensitive and Fast Alignment Search Tool for LOng noisy Read sequencing Data. Bioinformatics, 2019, 35, 20-27. PhISCS: a combinatorial approach for subperfect tumor phylogeny reconstruction via integrative use of single-cell and bulk sequencing data. Genome Research, 2019, 29, 1860-1877. A multi-labeled tree dissimilarity measure for comparing "clonal trees―of tumor progression. Algorithms for Molecular Biology, 2019, 14, 17. Integrative inference of subclonal tumour evolution from single-cell and bulk sequencing data. Nature Communications, 2019, 10, 2750.	3.3 1.8 2.4 0.3	78 23 73 25

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19	BAP1 haploinsufficiency predicts a distinct immunogenic class of malignant peritoneal mesothelioma. Genome Medicine, 2019, 11, 8.	3.6	88
20	Structural variation and fusion detection using targeted sequencing data from circulating cell free DNA. Nucleic Acids Research, 2019, 47, e38-e38.	6.5	17
21	SubGraph2Vec: Highly-Vectorized Tree-like Subgraph Counting. , 2019, , .		3
22	Genomic Data Compression. , 2019, , 783-783.		1
23	Protecting Genomic Data Privacy with Probabilistic Modeling. Pacific Symposium on Biocomputing Pacific Symposium on Biocomputing, 2019, 24, 403-414.	0.7	1
24	GTED: Graph Traversal Edit Distance. Lecture Notes in Computer Science, 2018, , 37-53.	1.0	1
25	MechRNA: prediction of lncRNA mechanisms from RNA–RNA and RNA–protein interactions. Bioinformatics, 2018, 34, 3101-3110.	1.8	48
26	Optimal compressed representation of high throughput sequence data via light assembly. Nature Communications, 2018, 9, 566.	5.8	14
27	Computational identification of micro-structural variations and their proteogenomic consequences in cancer. Bioinformatics, 2018, 34, 1672-1681.	1.8	8
28	Allelic decomposition and exact genotyping of highly polymorphic and structurally variant genes. Nature Communications, 2018, 9, 828.	5.8	67
29	Ultra High-Dimensional Nonlinear Feature Selection for Big Biological Data. IEEE Transactions on Knowledge and Data Engineering, 2018, 30, 1352-1365.	4.0	48
30	Stromal Gene Expression is Predictive for Metastatic Primary Prostate Cancer. European Urology, 2018, 73, 524-532.	0.9	60
31	Analysis of the androgen receptor–regulated IncRNA landscape identifies a role for ARLNC1 in prostate cancer progression. Nature Genetics, 2018, 50, 814-824.	9.4	196
32	The long noncoding RNA landscape of neuroendocrine prostate cancer and its clinical implications. GigaScience, 2018, 7, .	3.3	54
33	Preface: Selected Articles from RECOMB 2017. Journal of Computational Biology, 2018, 25, 623-623.	0.8	0
34	Protecting Genomic Data Privacy with Probabilistic Modeling. , 2018, , .		5
35	SRRM4 Drives Neuroendocrine Transdifferentiation of Prostate Adenocarcinoma Under Androgen Receptor Pathway Inhibition. European Urology, 2017, 71, 68-78.	0.9	136
36	Genomic hallmarks of localized, non-indolent prostate cancer. Nature, 2017, 541, 359-364.	13.7	462

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37	PRINCESS: Privacy-protecting Rare disease International Network Collaboration via Encryption through Software guard extensionS. Bioinformatics, 2017, 33, 871-878.	1.8	7 5
38	Clonality Inference from Single Tumor Samples Using Low-Coverage Sequence Data. Journal of Computational Biology, 2017, 24, 515-523.	0.8	20
39	Mutational Analysis of Gene Fusions Predicts Novel MHC Class I–Restricted T-Cell Epitopes and Immune Signatures in a Subset of Prostate Cancer. Clinical Cancer Research, 2017, 23, 7596-7607.	3.2	14
40	HIT'nDRIVE: patient-specific multidriver gene prioritization for precision oncology. Genome Research, 2017, 27, 1573-1588.	2.4	78
41	SiNVICT: ultra-sensitive detection of single nucleotide variants and indels in circulating tumour DNA. Bioinformatics, 2017, 33, 26-34.	1.8	48
42	PRESAGE: PRivacy-preserving gEnetic testing via SoftwAre Guard Extension. BMC Medical Genomics, 2017, 10, 48.	0.7	32
43	ReMixT: clone-specific genomic structure estimation in cancer. Genome Biology, 2017, 18, 140.	3.8	29
44	Abstract 378: The Cancer Genome Collaboratory. Cancer Research, 2017, 77, 378-378.	0.4	1
45	Edit Distance Under Block Operations. , 2016, , 611-614.		0
46	Enabling Privacy-Preserving GWASs in Heterogeneous Human Populations. Cell Systems, 2016, 3, 54-61.	2.9	62
47	CoLoRMap: Correcting Long Reads by Mapping short reads. Bioinformatics, 2016, 32, i545-i551.	1.8	49
48	Comparison of high-throughput sequencing data compression tools. Nature Methods, 2016, 13, 1005-1008.	9.0	91
49	Spatial genomic heterogeneity within localized, multifocal prostate cancer. Nature Genetics, 2015, 47, 736-745.	9.4	395
50	Cypiripi: exact genotyping of <i>CYP2D6</i> using high-throughput sequencing data. Bioinformatics, 2015, 31, i27-i34.	1.8	37
51	Robustness of Massively Parallel Sequencing Platforms. PLoS ONE, 2015, 10, e0138259.	1.1	3
52	mrsFAST-Ultra: a compact, SNP-aware mapper for high performance sequencing applications. Nucleic Acids Research, 2014, 42, W494-W500.	6.5	54
53	ORMAN: Optimal resolution of ambiguous RNA-Seq multimappings in the presence of novel isoforms. Bioinformatics, 2014, 30, 644-651.	1.8	17
54	Whole genome sequencing of Turkish genomes reveals functional private alleles and impact of genetic interactions with Europe, Asia and Africa. BMC Genomics, 2014, 15, 963.	1.2	46

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55	DeeZ: reference-based compression by local assembly. Nature Methods, 2014, 11, 1082-1084.	9.0	45
56	HIT'nDRIVE: Multi-driver Gene Prioritization Based on Hitting Time. Lecture Notes in Computer Science, 2014, , 293-306.	1.0	35
57	Barnacle: detecting and characterizing tandem duplications and fusions in transcriptome assemblies. BMC Genomics, 2013, 14, 550.	1.2	12
58	Abstract B129: Clinical implications of inter- and intra- prostatic heterogeneity , 2013, , .		0
59	nFuse: Discovery of complex genomic rearrangements in cancer using high-throughput sequencing. Genome Research, 2012, 22, 2250-2261.	2.4	67
60	SCALCE: boosting sequence compression algorithms using locally consistent encoding. Bioinformatics, 2012, 28, 3051-3057.	1.8	129
61	Mirroring co-evolving trees in the light of their topologies. Bioinformatics, 2012, 28, 1202-1208.	1.8	4
62	Dissect: detection and characterization of novel structural alterations in transcribed sequences. Bioinformatics, 2012, 28, i179-i187.	1.8	13
63	Polyâ€gene fusion transcripts and chromothripsis in prostate cancer. Genes Chromosomes and Cancer, 2012, 51, 1144-1153.	1.5	46
64	Integrated genome and transcriptome sequencing identifies a novel form of hybrid and aggressive prostate cancer. Journal of Pathology, 2012, 227, 53-61.	2.1	63
65	From sequence to molecular pathology, and a mechanism driving the neuroendocrine phenotype in prostate cancer. Journal of Pathology, 2012, 227, 286-297.	2.1	161
66	CLIIQ: Accurate Comparative Detection and Quantification of Expressed Isoforms in a Population. Lecture Notes in Computer Science, 2012, , 178-189.	1.0	32
67	Proteome Network Emulating Models. , 2012, , 69-95.		0
68	Mapping the Protein Interaction Network in Methicillin-Resistant <i>Staphylococcus aureus</i> Journal of Proteome Research, 2011, 10, 1139-1150.	1.8	55
69	Demographic history and rare allele sharing among human populations. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 11983-11988.	3.3	589
70	Comrad: detection of expressed rearrangements by integrated analysis of RNA-Seq and low coverage genome sequence data. Bioinformatics, 2011, 27, 1481-1488.	1.8	39
71	<i>Alu</i> repeat discovery and characterization within human genomes. Genome Research, 2011, 21, 840-849.	2.4	94
72	Optimally discriminative subnetwork markers predict response to chemotherapy. Bioinformatics, 2011, 27, i205-i213.	1.8	81

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73	deFuse: An Algorithm for Gene Fusion Discovery in Tumor RNA-Seq Data. PLoS Computational Biology, 2011, 7, e1001138.	1.5	477
74	Simultaneous structural variation discovery among multiple paired-end sequenced genomes. Genome Research, 2011, 21, 2203-2212.	2.4	60
75	Sensitive and fast mapping of di-base encoded reads. Bioinformatics, 2011, 27, 1915-1921.	1.8	16
76	Simultaneous Structural Variation Discovery in Multiple Paired-End Sequenced Genomes. Lecture Notes in Computer Science, 2011, , 104-105.	1.0	4
77	Applicability Domains for Classification Problems: Benchmarking of Distance to Models for Ames Mutagenicity Set. Journal of Chemical Information and Modeling, 2010, 50, 2094-2111.	2.5	202
78	Fast prediction of RNA-RNA interaction. Algorithms for Molecular Biology, 2010, 5, 5.	0.3	44
79	A map of human genome variation from population-scale sequencing. Nature, 2010, 467, 1061-1073.	13.7	7,209
80	mrsFAST: a cache-oblivious algorithm for short-read mapping. Nature Methods, 2010, 7, 576-577.	9.0	248
81	Detection and characterization of novel sequence insertions using paired-end next-generation sequencing. Bioinformatics, 2010, 26, 1277-1283.	1.8	98
82	Sparsification of RNA structure prediction including pseudoknots. Algorithms for Molecular Biology, 2010, 5, 39.	0.3	17
83	Towards Improved Assessment of Functional Similarity in Large-Scale Screens: A Study on Indel Length. Journal of Computational Biology, 2010, 17, 1-20.	0.8	15
84	Protein-Protein Interaction Network Evaluation for Identifying Potential Drug Targets. Journal of Computational Biology, 2010, 17, 669-684.	0.8	30
85	Next-generation VariationHunter: combinatorial algorithms for transposon insertion discovery. Bioinformatics, 2010, 26, i350-i357.	1.8	190
86	Periodicity testing with sublinear samples and space. ACM Transactions on Algorithms, 2010, 6, 1-14.	0.9	3
87	PSORTb 3.0: improved protein subcellular localization prediction with refined localization subcategories and predictive capabilities for all prokaryotes. Bioinformatics, 2010, 26, 1608-1615.	1.8	2,044
88	Time and Space Efficient RNA-RNA Interaction Prediction via Sparse Folding. Lecture Notes in Computer Science, 2010, , 473-490.	1.0	18
89	Sparsification of RNA Structure Prediction Including Pseudoknots. Lecture Notes in Computer Science, 2010, , 40-51.	1.0	3
90	Pair HMM Based Gap Statistics for Re-evaluation of Indels in Alignments with Affine Gap Penalties. Lecture Notes in Computer Science, 2010, , 350-361.	1.0	0

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91	A partition function algorithm for interacting nucleic acid strands. Bioinformatics, 2009, 25, i365-i373.	1.8	85
92	Personalized copy number and segmental duplication maps using next-generation sequencing. Nature Genetics, 2009, 41, 1061-1067.	9.4	656
93	Combinatorial algorithms for structural variation detection in high-throughput sequenced genomes. Genome Research, 2009, 19, 1270-1278.	2.4	266
94	The Effect of Insertions and Deletions on Wirings in Protein-Protein Interaction Networks: A Large-Scale Study. Journal of Computational Biology, 2009, 16, 159-167.	0.8	25
95	Fast Prediction of RNA-RNA Interaction. Lecture Notes in Computer Science, 2009, , 261-272.	1.0	6
96	biRNA: Fast RNA-RNA Binding Sites Prediction. Lecture Notes in Computer Science, 2009, , 25-36.	1.0	26
97	smyRNA: A Novel Ab Initio ncRNA Gene Finder. PLoS ONE, 2009, 4, e5433.	1.1	13
98	Combinatorial Algorithms for Structural Variation Detection in High Throughput Sequenced Genomes. Lecture Notes in Computer Science, 2009, , 218-219.	1.0	3
99	Quantifying Systemic Evolutionary Changes by Color Coding Confidence-Scored PPI Networks. Lecture Notes in Computer Science, 2009, , 37-48.	1.0	4
100	COMPUTATIONAL STUDIES OF NON-CODING RNAS – Session Introduction. , 2009, , 54-56.		0
101	PERSONAL GENOMICS – Session Introduction. , 2009, , 302-304.		0
102	Biomolecular network motif counting and discovery by color coding. Bioinformatics, 2008, 24, i241-i249.	1.8	155
103	Comparative analysis of the small RNA transcriptomes of Pinus contorta and Oryza sativa. Genome Research, 2008, 18, 571-584.	2.4	305
104	Conifers have a unique small RNA silencing signature. Rna, 2008, 14, 1508-1515.	1.6	108
105	Optimal pooling for genome re-sequencing with ultra-high-throughput short-read technologies. Bioinformatics, 2008, 24, i32-i40.	1.8	16
106	Edit Distance Under Block Operations. , 2008, , 265-267.		2
107	The Relation between Indel Length and Functional Divergence: A Formal Study. Lecture Notes in Computer Science, 2008, , 330-341.	1.0	1
108	Not All Scale-Free Networks Are Born Equal: The Role of the Seed Graph in PPI Network Evolution. PLoS Computational Biology, 2007, 3, e118.	1,5	77

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109	Organization and Evolution of Primate Centromeric DNA from Whole-Genome Shotgun Sequence Data. PLoS Computational Biology, 2007, 3, e181.	1.5	80
110	taveRNA: a web suite for RNA algorithms and applications. Nucleic Acids Research, 2007, 35, W325-W329.	6.5	14
111	Optimal spaced seeds for faster approximate string matching. Journal of Computer and System Sciences, 2007, 73, 1035-1044.	0.9	17
112	Novel approaches for small biomolecule classification and structural similarity search. SIGKDD Explorations: Newsletter of the Special Interest Group (SIG) on Knowledge Discovery & Data Mining, 2007, 9, 14-21.	3.2	7
113	Comparative QSAR- and Fragments Distribution Analysis of Drugs, Druglikes, Metabolic Substances, and Antimicrobial Compounds. Journal of Chemical Information and Modeling, 2006, 46, 2167-2182.	2.5	50
114	Distance based algorithms for small biomolecule classification and structural similarity search. Bioinformatics, 2006, 22, e243-e251.	1.8	41
115	RNA–RNA Interaction Prediction and Antisense RNA Target Search. Journal of Computational Biology, 2006, 13, 267-282.	0.8	106
116	RNA Secondary Structure Prediction Via Energy Density Minimization. Lecture Notes in Computer Science, 2006, , 130-142.	1.0	9
117	Oblivious string embeddings and edit distance approximations. , 2006, , .		46
118	COMPARATIVE QSAR ANALYSIS OF BACTERIAL, FUNGAL PLANT AND HUMAN METABOLITES. , 2006, , .		0
119	Not All Scale Free Networks Are Born Equal: The Role of the Seed Graph in PPI Network Emulation. Lecture Notes in Computer Science, 2006, , $1-13$.	1.0	0
120	Locally Consistent Parsing and Applications to Approximate String Comparisons. Lecture Notes in Computer Science, 2005, , 22-35.	1.0	2
121	Manipulating multiple sequence alignments via MaM and WebMaM. Nucleic Acids Research, 2005, 33, W295-W298.	6. 5	5
122	Improved Duplication Models for Proteome Network Evolution., 2005,, 119-137.		7
123	IDENTIFYING UNIFORMLY MUTATED SEGMENTS WITHIN REPEATS. Journal of Bioinformatics and Computational Biology, 2004, 02, 657-668.	0.3	0
124	The Role of Unequal Crossover in Alpha-Satellite DNA Evolution: A Computational Analysis. Journal of Computational Biology, 2004, $11,933-944$.	0.8	20
125	The structure and evolution of centromeric transition regions within the human genome. Nature, 2004, 430, 857-864.	13.7	179
126	Sublinear Methods for Detecting Periodic Trends in Data Streams. Lecture Notes in Computer Science, 2004, , 16-28.	1.0	11

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127	Hardness of String Similarity Search and Other Indexing Problems. Lecture Notes in Computer Science, 2004, , 1080-1098.	1.0	4
128	Analysis of Primate Genomic Variation Reveals a Repeat-Driven Expansion of the Human Genome. Genome Research, 2003, 13, 358-368.	2.4	127
129	Simple and Practical Sequence Nearest Neighbors with Block Operations. Lecture Notes in Computer Science, 2002, , 262-278.	1.0	9
130	The Complexity of Gene Placement. Journal of Algorithms, 2001, 41, 225-243.	0.9	3
131	Divergent Origins and Concerted Expansion of Two Segmental Duplications on Chromosome 16. Journal of Heredity, 2001, 92, 462-468.	1.0	25
132	Permutation Editing and Matching via Embeddings. Lecture Notes in Computer Science, 2001, , 481-492.	1.0	14
133	Approximate nearest neighbors and sequence comparison with block operations. , 2000, , .		58
134	On a parallel-algorithms method for string matching problems (overview). Lecture Notes in Computer Science, 1994, , 22-32.	1.0	6