

Daniel R Zerbino

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/167508/daniel-r-zerbino-publications-by-year.pdf>

Version: 2024-04-10

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

53 papers	37,217 citations	33 h-index	57 g-index
57 ext. papers	48,548 ext. citations	17.4 avg, IF	8.11 L-index

#	Paper	IF	Citations
53	Ensembl 2022. <i>Nucleic Acids Research</i> , 2021 ,	20.1	72
52	The gene regulation knowledge commons: the action area of GREEKC. <i>Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms</i> , 2021 , 1865, 194768	6	1
51	Recommendations for the FAIRification of genomic track metadata. <i>F1000Research</i> , 2021 , 10,	3.6	1
50	Population-scale tissue transcriptomics maps long non-coding RNAs to complex disease. <i>Cell</i> , 2021 , 184, 2633-2648.e19	56.2	20
49	The FAANG Data Portal: Global, Open-Access, "FAIR", and Richly Validated Genotype to Phenotype Data for High-Quality Functional Annotation of Animal Genomes. <i>Frontiers in Genetics</i> , 2021 , 12, 639238	4.5	0
48	Ensembl 2021. <i>Nucleic Acids Research</i> , 2021 , 49, D884-D891	20.1	324
47	GENCODE 2021. <i>Nucleic Acids Research</i> , 2021 , 49, D916-D923	20.1	82
46	Transcription and DNA Methylation Patterns of Blood-Derived CD8 T Cells Are Associated With Age and Inflammatory Bowel Disease But Do Not Predict Prognosis. <i>Gastroenterology</i> , 2021 , 160, 232-244.e7	13.3	17
45	The Ensembl COVID-19 resource: ongoing integration of public SARS-CoV-2 data. <i>Nucleic Acids Research</i> , 2021 ,	20.1	3
44	A compendium of uniformly processed human gene expression and splicing quantitative trait loci. <i>Nature Genetics</i> , 2021 , 53, 1290-1299	36.3	28
43	Sequence Ontology terminology for gene regulation. <i>Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms</i> , 2021 , 1864, 194745	6	3
42	Progress, Challenges, and Surprises in Annotating the Human Genome. <i>Annual Review of Genomics and Human Genetics</i> , 2020 , 21, 55-79	9.7	6
41	The open targets post-GWAS analysis pipeline. <i>Bioinformatics</i> , 2020 , 36, 2936-2937	7.2	13
40	Ensembl 2020. <i>Nucleic Acids Research</i> , 2020 , 48, D682-D688	20.1	645
39	Perspectives on ENCODE. <i>Nature</i> , 2020 , 583, 693-698	50.4	61
38	Sequence tube maps: making graph genomes intuitive to commuters. <i>Bioinformatics</i> , 2019 , 35, 5318-5320	9.2	16
37	GENCODE reference annotation for the human and mouse genomes. <i>Nucleic Acids Research</i> , 2019 , 47, D766-D773	20.1	1140

36	Ensembl 2019. <i>Nucleic Acids Research</i> , 2019 , 47, D745-D751	20.1	554
35	Ensembl 2018. <i>Nucleic Acids Research</i> , 2018 , 46, D754-D761	20.1	1822
34	Ensembl 2017. <i>Nucleic Acids Research</i> , 2017 , 45, D635-D642	20.1	404
33	Ensembl core software resources: storage and programmatic access for DNA sequence and genome annotation. <i>Database: the Journal of Biological Databases and Curation</i> , 2017 , 2017,	5	35
32	CHiCAGO: robust detection of DNA looping interactions in Capture Hi-C data. <i>Genome Biology</i> , 2016 , 17, 127	18.3	220
31	Lineage-Specific Genome Architecture Links Enhancers and Non-coding Disease Variants to Target Gene Promoters. <i>Cell</i> , 2016 , 167, 1369-1384.e19	56.2	556
30	The International Human Epigenome Consortium: A Blueprint for Scientific Collaboration and Discovery. <i>Cell</i> , 2016 , 167, 1145-1149	56.2	232
29	Representing and decomposing genomic structural variants as balanced integer flows on sequence graphs. <i>BMC Bioinformatics</i> , 2016 , 17, 400	3.6	6
28	Ensembl 2016. <i>Nucleic Acids Research</i> , 2016 , 44, D710-6	20.1	1094
27	Ensembl regulation resources. <i>Database: the Journal of Biological Databases and Curation</i> , 2016 , 2016,	5	33
26	Punctuated bursts in human male demography inferred from 1,244 worldwide Y-chromosome sequences. <i>Nature Genetics</i> , 2016 , 48, 593-9	36.3	204
25	The ensembl regulatory build. <i>Genome Biology</i> , 2015 , 16, 56	18.3	255
24	A global reference for human genetic variation. <i>Nature</i> , 2015 , 526, 68-74	50.4	8599
23	Ensembl 2015. <i>Nucleic Acids Research</i> , 2015 , 43, D662-9	20.1	1013
22	Building a pan-genome reference for a population. <i>Journal of Computational Biology</i> , 2015 , 22, 387-401	1.7	41
21	A unifying model of genome evolution under parsimony. <i>BMC Bioinformatics</i> , 2014 , 15, 206	3.6	8
20	Ensembl 2014. <i>Nucleic Acids Research</i> , 2014 , 42, D749-55	20.1	1087
19	WiggleTools: parallel processing of large collections of genome-wide datasets for visualization and statistical analysis. <i>Bioinformatics</i> , 2014 , 30, 1008-9	7.2	44

18	Building a Pangenome Reference for a Population. <i>Lecture Notes in Computer Science</i> , 2014 , 207-221	0.9	4
17	Assessment of transcript reconstruction methods for RNA-seq. <i>Nature Methods</i> , 2013 , 10, 1177-84	21.6	477
16	The NGS WikiBook: a dynamic collaborative online training effort with long-term sustainability. <i>Briefings in Bioinformatics</i> , 2013 , 14, 548-55	13.4	7
15	HAL: a hierarchical format for storing and analyzing multiple genome alignments. <i>Bioinformatics</i> , 2013 , 29, 1341-2	7.2	78
14	Oases: robust de novo RNA-seq assembly across the dynamic range of expression levels. <i>Bioinformatics</i> , 2012 , 28, 1086-92	7.2	1129
13	Comprehensive molecular characterization of human colon and rectal cancer. <i>Nature</i> , 2012 , 487, 330-7	50.4	5640
12	Comprehensive genomic characterization of squamous cell lung cancers. <i>Nature</i> , 2012 , 489, 519-25	50.4	2820
11	Assemblathon 1: a competitive assessment of de novo short read assembly methods. <i>Genome Research</i> , 2011 , 21, 2224-41	9.7	364
10	Cactus: Algorithms for genome multiple sequence alignment. <i>Genome Research</i> , 2011 , 21, 1512-28	9.7	170
9	Improvements to services at the European Nucleotide Archive. <i>Nucleic Acids Research</i> , 2010 , 38, D39-45	20.1	47
8	A new strategy for genome assembly using short sequence reads and reduced representation libraries. <i>Genome Research</i> , 2010 , 20, 249-56	9.7	24
7	Using the Velvet de novo assembler for short-read sequencing technologies. <i>Current Protocols in Bioinformatics</i> , 2010 , Chapter 11, Unit 11.5	24.2	358
6	Pebble and rock band: heuristic resolution of repeats and scaffolding in the velvet short-read de novo assembler. <i>PLoS ONE</i> , 2009 , 4, e8407	3.7	151
5	Velvet: algorithms for de novo short read assembly using de Bruijn graphs. <i>Genome Research</i> , 2008 , 18, 821-9	9.7	7156
4	An efficient conformational sampling method for homology modeling. <i>Proteins: Structure, Function and Bioinformatics</i> , 2008 , 71, 175-88	4.2	20
3	An analysis of core deformations in protein superfamilies. <i>Biophysical Journal</i> , 2005 , 88, 1291-9	2.9	91
2	Core deformations in protein families: a physical perspective. <i>Biophysical Chemistry</i> , 2005 , 115, 125-8	3.5	9
1	eQTL Catalogue: a compendium of uniformly processed human gene expression and splicing QTLs		33

