

Wan-Ping Lee

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4935642/publications.pdf>

Version: 2024-02-01

26
papers

30,279
citations

759233

12
h-index

642732

23
g-index

32
all docs

32
docs citations

32
times ranked

54067
citing authors

#	ARTICLE	IF	CITATIONS
1	A global reference for human genetic variation. <i>Nature</i> , 2015, 526, 68-74.	27.8	13,998
2	A map of human genome variation from population-scale sequencing. <i>Nature</i> , 2010, 467, 1061-1073.	27.8	7,209
3	An integrated map of genetic variation from 1,092 human genomes. <i>Nature</i> , 2012, 491, 56-65.	27.8	7,199
4	Multi-platform discovery of haplotype-resolved structural variation in human genomes. <i>Nature Communications</i> , 2019, 10, 1784.	12.8	636
5	A Comprehensive Map of Mobile Element Insertion Polymorphisms in Humans. <i>PLoS Genetics</i> , 2011, 7, e1002236.	3.5	278
6	MOSAİK: A Hash-Based Algorithm for Accurate Next-Generation Sequencing Short-Read Mapping. <i>PLoS ONE</i> , 2014, 9, e90581.	2.5	249
7	SSW Library: An SIMD Smith-Waterman C/C++ Library for Use in Genomic Applications. <i>PLoS ONE</i> , 2013, 8, e82138.	2.5	175
8	Fast and accurate genomic analyses using genome graphs. <i>Nature Genetics</i> , 2019, 51, 354-362.	21.4	167
9	Expectations and blind spots for structural variation detection from long-read assemblies and short-read genome sequencing technologies. <i>American Journal of Human Genetics</i> , 2021, 108, 919-928.	6.2	72
10	One reference genome is not enough. <i>Genome Biology</i> , 2019, 20, 104.	8.8	58
11	Tangram: a comprehensive toolbox for mobile element insertion detection. <i>BMC Genomics</i> , 2014, 15, 795.	2.8	54
12	FusorSV: an algorithm for optimally combining data from multiple structural variation detection methods. <i>Genome Biology</i> , 2018, 19, 38.	8.8	46
13	Polygenic Risk Scores in Alzheimer's Disease Genetics: Methodology, Applications, Inclusion, and Diversity. <i>Journal of Alzheimer's Disease</i> , 2022, 89, 1-12.	2.6	17
14	Voltage-Island Partitioning and Floorplanning Under Timing Constraints. <i>IEEE Transactions on Computer-Aided Design of Integrated Circuits and Systems</i> , 2009, 28, 690-702.	2.7	13
15	Comprehensive Analysis of Alternative Splicing in Gastric Cancer Identifies Epithelial-Mesenchymal Transition Subtypes Associated with Survival. <i>Cancer Research</i> , 2022, 82, 543-555.	0.9	12
16	Voltage Island Aware Floorplanning for Power and Timing Optimization. <i>IEEE/ACM International Conference on Computer-Aided Design, Digest of Technical Papers</i> , 2006, , .	0.0	10
17	Post-floorplanning power/ground ring synthesis for multiple-supply-voltage designs. , 2009, , .		9
18	Toolbox for Mobile-Element Insertion Detection on Cancer Genomes. <i>Cancer Informatics</i> , 2014, 13s4, CIN.S13979.	1.9	4

#	ARTICLE	IF	CITATIONS
19	Toolbox for Mobile-Element Insertion Detection on Cancer Genomes. <i>Cancer Informatics</i> , 2015, 14s1, CIN.S24657.	1.9	4
20	Copy Number Variation Identification on 3,800 Alzheimer's Disease Whole Genome Sequencing Data from the Alzheimer's Disease Sequencing Project. <i>Frontiers in Genetics</i> , 2021, 12, 752390.	2.3	4
21	JAX-CNV: A Whole-genome Sequencing-based Algorithm for Copy Number Detection at Clinical Grade Level. <i>Genomics, Proteomics and Bioinformatics</i> , 2022, 20, 1197-1206.	6.9	3
22	An ILP algorithm for post-floorplanning voltage-island generation considering power-network planning. <i>IEEE/ACM International Conference on Computer-Aided Design, Digest of Technical Papers</i> , 2007, , .	0.0	2
23	SEAGLE: A Scalable Exact Algorithm for Large-Scale Set-Based Gene-Environment Interaction Tests in Biobank Data. <i>Frontiers in Genetics</i> , 2021, 12, 710055.	2.3	2
24	Sensitivity-based multiple-Vt cell swapping for leakage power reduction. , 2008, , .		1
25	NIA genetics of Alzheimer's disease data storage site (NIAGADS): 2021 update.. <i>Alzheimer's and Dementia</i> , 2021, 17 Suppl 3, e052258.	0.8	0
26	Copy number variation (CNV) identification and association study on 3,928 Alzheimer's disease whole genome sequencing data from the Alzheimer's Disease Sequencing Project (ADSP).. <i>Alzheimer's and Dementia</i> , 2021, 17 Suppl 3, e052721.	0.8	0