## Yun Li

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1635882/publications.pdf

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37	2,900	14	36
papers	citations	h-index	g-index
55	55	55	7475
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Generalized multiâ€SNP mediation intersection–union test. Biometrics, 2022, 78, 364-375.	1.4	7
2	Single-cell dual-omics reveals the transcriptomic and epigenomic diversity of cardiac non-myocytes. Cardiovascular Research, 2022, 118, 1548-1563.	3.8	31
3	eSCAN: scan regulatory regions for aggregate association testing using whole-genome sequencing data. Briefings in Bioinformatics, 2022, 23, .	6.5	5
4	Do adverse childhood experiences and genetic obesity risk interact in relation to body mass index in young adulthood? Findings from the National Longitudinal Study of Adolescent to Adult Health. Pediatric Obesity, 2022, 17, e12885.	2.8	4
5	THUNDER: A reference-free deconvolution method to infer cell type proportions from bulk Hi-C data. PLoS Genetics, 2022, 18, e1010102.	3.5	9
6	A systematic evaluation of Hi-C data enhancement methods for enhancing PLAC-seq and HiChIP data. Briefings in Bioinformatics, 2022, 23, .	6.5	3
7	SnapHiC2: A computationally efficient loop caller for single cell Hi-C data. Computational and Structural Biotechnology Journal, 2022, 20, 2778-2783.	4.1	7
8	CUE: CpG impUtation ensemble for DNA methylation levels across the human methylation450 (HM450) and EPIC (HM850) BeadChip platforms. Epigenetics, 2021, 16, 851-861.	2.7	1
9	SMNN: batch effect correction for single-cell RNA-seq data via supervised mutual nearest neighbor detection. Briefings in Bioinformatics, 2021, 22, .	6.5	17
10	TWOâ€SIGMA: A novel twoâ€component single cell modelâ€based association method for singleâ€cell RNAâ€seq data. Genetic Epidemiology, 2021, 45, 142-153.	1.3	11
11	Maternal gut microbiota reflecting poor diet quality is associated with spontaneous preterm birth in a prospective cohort study. American Journal of Clinical Nutrition, 2021, 113, 602-611.	4.7	19
12	MOSTWAS: Multi-Omic Strategies for Transcriptome-Wide Association Studies. PLoS Genetics, 2021, 17, e1009398.	3.5	46
13	MRLocus: Identifying causal genes mediating a trait through Bayesian estimation of allelic heterogeneity. PLoS Genetics, 2021, 17, e1009455.	3.5	24
14	Genetic correlations between COVID-19 and a variety of traits and diseases. Innovation(China), 2021, 2, 100112.	9.1	7
15	ExpressHeart: Web Portal to Visualize Transcriptome Profiles of Non-Cardiomyocyte Cells. International Journal of Molecular Sciences, 2021, 22, 8943.	4.1	3
16	HPRep: Quantifying Reproducibility in HiChIP and PLAC-Seq Datasets. Current Issues in Molecular Biology, 2021, 43, 1156-1170.	2.4	4
17	Soluble Urokinase Plasminogen Activator Receptor: Genetic Variation and Cardiovascular Disease Risk in Black Adults. Circulation Genomic and Precision Medicine, 2021, 14, CIRCGEN121003421.	3.6	7
18	Age-related DNA hydroxymethylation is enriched for gene expression and immune system processes in human peripheral blood. Epigenetics, 2020, 15, 294-306.	2.7	8

#	Article	IF	CITATIONS
19	SAME-clustering: Single-cell Aggregated Clustering via Mixture Model Ensemble. Nucleic Acids Research, 2020, 48, 86-95.	14.5	55
20	Genome-Wide Association of Kidney Traits in Hispanics/Latinos Using Dense Imputed Whole-Genome Sequencing Data. Circulation Genomic and Precision Medicine, 2020, 13, e002891.	3.6	6
21	Bacterial colonization reprograms the neonatal gut metabolome. Nature Microbiology, 2020, 5, 838-847.	13.3	70
22	MAPS: Model-based analysis of long-range chromatin interactions from PLAC-seq and HiChIP experiments. PLoS Computational Biology, 2019, 15, e1006982.	3.2	94
23	Coexpression network analysis identifies transcriptional modules associated with genomic alterations in neuroblastoma. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2018, 1864, 2341-2348.	3.8	35
24	FXR-Dependent Modulation of the Human Small Intestinal Microbiome by the Bile Acid Derivative Obeticholic Acid. Gastroenterology, 2018, 155, 1741-1752.e5.	1.3	82
25	299 - FXR-Dependent Modification of the Human Small Intestinal Microbiome. Gastroenterology, 2018, 154, S-1084.	1.3	0
26	Weighted Gene Co-Expression Network Analysis Reveals Dysregulation of Mitochondrial Oxidative Phosphorylation in Eating Disorders. Genes, 2018, 9, 325.	2.4	14
27	A Multiancestral Genome-Wide Exome Array Study of Alzheimer Disease, Frontotemporal Dementia, and Progressive Supranuclear Palsy. JAMA Neurology, 2015, 72, 414.	9.0	37
28	An Epigenetic Signature in Peripheral Blood Associated with the Haplotype on 17q21.31, a Risk Factor for Neurodegenerative Tauopathy. PLoS Genetics, 2014, 10, e1004211.	3.5	65
29	Dynamic modular architecture of protein-protein interaction networks beyond the dichotomy of $\hat{a} \in \hat{a}$ and $\hat{a} \in \hat{a}$ hubs. Scientific Reports, 2013, 3, 1691.	3.3	71
30	A cross-species analysis method to analyze animal models' similarity to human's disease state. BMC Systems Biology, 2012, 6, S18.	3.0	7
31	Rare-Variant Association Testing for Sequencing Data with the Sequence Kernel Association Test. American Journal of Human Genetics, 2011, 89, 82-93.	6.2	2,060
32	EcoBrowser: a web-based tool for visualizing transcriptome data of Escherichia coli. BMC Research Notes, 2011, 4, 405.	1.4	3
33	ASSOCIATION OF FEATURE GENE EXPRESSION WITH STRUCTURAL FINGERPRINTS OF CHEMICAL COMPOUNDS. Journal of Bioinformatics and Computational Biology, 2011, 09, 503-519.	0.8	4
34	GEOGLE: context mining tool for the correlation between gene expression and the phenotypic distinction. BMC Bioinformatics, 2009, 10, 264.	2.6	4
35	MPSQ: a web tool for protein-state searching. Bioinformatics, 2008, 24, 2412-2413.	4.1	4
36	Gene expression module-based chemical function similarity search. Nucleic Acids Research, 2008, 36, e137-e137.	14.5	23

# ARTICLE IF CITATIONS

37 Tree of Life Based on Genome Context Networks. PLoS ONE, 2008, 3, e3357. 2.5 26