

Lin S Chen

List of Publications by Year in descending order

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Version: 2024-02-01

30
papers

2,108
citations

471509

17
h-index

434195

31
g-index

33
all docs

33
docs citations

33
times ranked

4423
citing authors

#	ARTICLE	IF	CITATIONS
1	Integrated Proteogenomic Characterization of Clear Cell Renal Cell Carcinoma. <i>Cell</i> , 2019, 179, 964-983.e31.	28.9	430
2	Proteogenomic and metabolomic characterization of human glioblastoma. <i>Cancer Cell</i> , 2021, 39, 509-528.e20.	16.8	327
3	Determinants of telomere length across human tissues. <i>Science</i> , 2020, 369, .	12.6	257
4	Proteogenomic insights into the biology and treatment of HPV-negative head and neck squamous cell carcinoma. <i>Cancer Cell</i> , 2021, 39, 361-379.e16.	16.8	189
5	Insights into Colon Cancer Etiology via a Regularized Approach to Gene Set Analysis of GWAS Data. <i>American Journal of Human Genetics</i> , 2010, 86, 860-871.	6.2	130
6	Harnessing naturally randomized transcription to infer regulatory relationships among genes. <i>Genome Biology</i> , 2007, 8, R219.	9.6	109
7	Mediation Analysis Demonstrates That Trans-eQTLs Are Often Explained by Cis-Mediation: A Genome-Wide Analysis among 1,800 South Asians. <i>PLoS Genetics</i> , 2014, 10, e1004818.	3.5	88
8	Characterization of Nigerian breast cancer reveals prevalent homologous recombination deficiency and aggressive molecular features. <i>Nature Communications</i> , 2018, 9, 4181.	12.8	77
9	Identifying cis-mediators for trans-eQTLs across many human tissues using genomic mediation analysis. <i>Genome Research</i> , 2017, 27, 1859-1871.	5.5	72
10	Co-occurring expression and methylation QTLs allow detection of common causal variants and shared biological mechanisms. <i>Nature Communications</i> , 2018, 9, 804.	12.8	66
11	Imputing Gene Expression in Uncollected Tissues Within and Beyond GTEx. <i>American Journal of Human Genetics</i> , 2016, 98, 697-708.	6.2	51
12	Genome-wide association studies in women of African ancestry identified 3q26.21 as a novel susceptibility locus for oestrogen receptor negative breast cancer. <i>Human Molecular Genetics</i> , 2016, 25, ddw305.	2.9	50
13	An Exponential Combination Procedure for Set-Based Association Tests in Sequencing Studies. <i>American Journal of Human Genetics</i> , 2012, 91, 977-986.	6.2	34
14	Genome-wide association study of telomere length among South Asians identifies a second RTEL1 association signal. <i>Journal of Medical Genetics</i> , 2018, 55, 64-71.	3.2	33
15	Primo: integration of multiple GWAS and omics QTL summary statistics for elucidation of molecular mechanisms of trait-associated SNPs and detection of pleiotropy in complex traits. <i>Genome Biology</i> , 2020, 21, 236.	8.8	26
16	The contribution of parent-to-offspring transmission of telomeres to the heritability of telomere length in humans. <i>Human Genetics</i> , 2019, 138, 49-60.	3.8	24
17	Insights into Impact of DNA Copy Number Alteration and Methylation on the Proteogenomic Landscape of Human Ovarian Cancer via a Multi-omics Integrative Analysis. <i>Molecular and Cellular Proteomics</i> , 2019, 18, S52-S65.	3.8	23
18	IGREX for quantifying the impact of genetically regulated expression on phenotypes. <i>NAR Genomics and Bioinformatics</i> , 2020, 2, lqaa010.	3.2	15

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19	A Unified Set-Based Test with Adaptive Filtering for Gene-Environment Interaction Analyses. <i>Biometrics</i> , 2016, 72, 629-638.	1.4	14
20	The association between telomere length and mortality in Bangladesh. <i>Aging</i> , 2017, 9, 1537-1551.	3.1	12
21	Breast cancer risk after full-term pregnancies among African women from Nigeria, Cameroon, and Uganda. <i>Cancer</i> , 2015, 121, 2237-2243.	4.1	11
22	A robust two-sample transcriptome-wide Mendelian randomization method integrating GWAS with multi-tissue eQTL summary statistics. <i>Genetic Epidemiology</i> , 2021, 45, 353-371.	1.3	11
23	A mixed-effects model for incomplete data from labeling-based quantitative proteomics experiments. <i>Annals of Applied Statistics</i> , 2017, 11, 114-138.	1.1	9
24	Germline Variation and Breast Cancer Incidence: A Gene-Based Association Study and Whole-Genome Prediction of Early-Onset Breast Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2018, 27, 1057-1064.	2.5	9
25	Rare, Protein-Altering Variants in <i>AS3MT</i> and Arsenic Metabolism Efficiency: A Multi-Population Association Study. <i>Environmental Health Perspectives</i> , 2021, 129, 47007.	6.0	9
26	Integrative Proteo-genomic Analysis to Construct CNA-protein Regulatory Map in Breast and Ovarian Tumors. <i>Molecular and Cellular Proteomics</i> , 2019, 18, S66-S81.	3.8	6
27	A meta-analysis approach with filtering for identifying gene-level gene-environment interactions. <i>Genetic Epidemiology</i> , 2018, 42, 434-446.	1.3	5
28	Using eQTLs to Reconstruct Gene Regulatory Networks. <i>Methods in Molecular Biology</i> , 2012, 871, 175-189.	0.9	4
29	CCmed: cross-condition mediation analysis for identifying replicable trans-associations mediated by cis-gene expression. <i>Bioinformatics</i> , 2021, 37, 2513-2520.	4.1	4
30	Using multivariate mixed-effects selection models for analyzing batch-processed proteomics data with non-ignorable missingness. <i>Biostatistics</i> , 2019, 20, 648-665.	1.5	3