

Zhili Zheng

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

7,026
citations

471509

17
h-index

610901

24
g-index

36
all docs

36
docs citations

36
times ranked

12617
citing authors

#	ARTICLE	IF	CITATIONS
1	Gene discovery and polygenic prediction from a genome-wide association study of educational attainment in 1.1 million individuals. <i>Nature Genetics</i> , 2018, 50, 1112-1121.	21.4	1,835
2	Meta-analysis of genome-wide association studies for height and body mass index in ~ 4.7 million individuals of European ancestry. <i>Human Molecular Genetics</i> , 2018, 27, 3641-3649.	2.9	1,541
3	Causal associations between risk factors and common diseases inferred from GWAS summary data. <i>Nature Communications</i> , 2018, 9, 224.	12.8	629
4	Genome-wide association analyses identify 143 risk variants and putative regulatory mechanisms for type 2 diabetes. <i>Nature Communications</i> , 2018, 9, 2941.	12.8	570
5	Identifying gene targets for brain-related traits using transcriptomic and methylomic data from blood. <i>Nature Communications</i> , 2018, 9, 2282.	12.8	294
6	A resource-efficient tool for mixed model association analysis of large-scale data. <i>Nature Genetics</i> , 2019, 51, 1749-1755.	21.4	294
7	Improved polygenic prediction by Bayesian multiple regression on summary statistics. <i>Nature Communications</i> , 2019, 10, 5086.	12.8	291
8	Integrative analysis of omics summary data reveals putative mechanisms underlying complex traits. <i>Nature Communications</i> , 2018, 9, 918.	12.8	250
9	Genetic signatures of high-altitude adaptation in Tibetans. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, 4189-4194.	7.1	181
10	A generalized linear mixed model association tool for biobank-scale data. <i>Nature Genetics</i> , 2021, 53, 1616-1621.	21.4	168
11	Assessing the contribution of rare variants to complex trait heritability from whole-genome sequence data. <i>Nature Genetics</i> , 2022, 54, 263-273.	21.4	156
12	Genome-wide association study of medication-use and associated disease in the UK Biobank. <i>Nature Communications</i> , 2019, 10, 1891.	12.8	140
13	A Comparison of Ten Polygenic Score Methods for Psychiatric Disorders Applied Across Multiple Cohorts. <i>Biological Psychiatry</i> , 2021, 90, 611-620.	1.3	103
14	Quantifying the mapping precision of genome-wide association studies using whole-genome sequencing data. <i>Genome Biology</i> , 2017, 18, 86.	8.8	84
15	Global genetic differentiation of complex traits shaped by natural selection in humans. <i>Nature Communications</i> , 2018, 9, 1865.	12.8	70
16	Association Between Population Density and Genetic Risk for Schizophrenia. <i>JAMA Psychiatry</i> , 2018, 75, 901.	11.0	67
17	Widespread signatures of natural selection across human complex traits and functional genomic categories. <i>Nature Communications</i> , 2021, 12, 1164.	12.8	50
18	Improved analyses of GWAS summary statistics by reducing data heterogeneity and errors. <i>Nature Communications</i> , 2021, 12, 7117.	12.8	31

#	ARTICLE	IF	CITATIONS
19	Phenotypic covariance across the entire spectrum of relatedness for 86 billion pairs of individuals. <i>Nature Communications</i> , 2021, 12, 1050.	12.8	19
20	Comparison of three monocular methods for measuring accommodative stimulusâ€™response curves. <i>Australasian journal of optometry</i> , The, 2017, 100, 155-161.	1.3	14
21	Tumor Mutational Burden Is Polygenic and Genetically Associated with Complex Traits and Diseases. <i>Cancer Research</i> , 2021, 81, 1230-1239.	0.9	14
22	The effects of spatial frequency on the accommodation responses of myopes and emmetropes under various detection demands. <i>Vision Research</i> , 2015, 115, 1-7.	1.4	8
23	Promoter-anchored chromatin interactions predicted from genetic analysis of epigenomic data. <i>Nature Communications</i> , 2020, 11, 2061.	12.8	8
24	The Effects of Spatial Frequency on the Accommodative Responses of Myopic and Emmetropic Chinese Children. <i>Translational Vision Science and Technology</i> , 2019, 8, 65.	2.2	6
25	F87COMMON GENETIC VARIATION EXPLAINS A HIGH PROPORTION OF THE ELEVATED RISK OF PSYCHIATRIC DISORDERS IN CHILDREN OF YOUNGER MOTHERS. <i>European Neuropsychopharmacology</i> , 2019, 29, S1156-S1157.	0.7	0