

Jin Yu

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3827639/publications.pdf>

Version: 2024-02-01

26
papers

31,371
citations

430442

18
h-index

610482

24
g-index

31
all docs

31
docs citations

31
times ranked

54923
citing authors

#	ARTICLE	IF	CITATIONS
1	A global reference for human genetic variation. <i>Nature</i> , 2015, 526, 68-74.	13.7	13,998
2	A map of human genome variation from population-scale sequencing. <i>Nature</i> , 2010, 467, 1061-1073.	13.7	7,209
3	An integrated map of genetic variation from 1,092 human genomes. <i>Nature</i> , 2012, 491, 56-65.	13.7	7,199
4	Genome-wide association meta-analysis in 269,867 individuals identifies new genetic and functional links to intelligence. <i>Nature Genetics</i> , 2018, 50, 912-919.	9.4	893
5	Study of 300,486 individuals identifies 148 independent genetic loci influencing general cognitive function. <i>Nature Communications</i> , 2018, 9, 2098.	5.8	484
6	The 1000 Genomes Project: data management and community access. <i>Nature Methods</i> , 2012, 9, 459-462.	9.0	308
7	An integrative variant analysis suite for whole exome next-generation sequencing data. <i>BMC Bioinformatics</i> , 2012, 13, 8.	1.2	252
8	The functional spectrum of low-frequency coding variation. <i>Genome Biology</i> , 2011, 12, R84.	13.9	173
9	Whole-genome sequence-based analysis of high-density lipoprotein cholesterol. <i>Nature Genetics</i> , 2013, 45, 899-901.	9.4	132
10	Schizophrenia Polygenic Risk Score as a Predictor of Antipsychotic Efficacy in First-Episode Psychosis. <i>American Journal of Psychiatry</i> , 2019, 176, 21-28.	4.0	127
11	Large-Scale Cognitive GWAS Meta-Analysis Reveals Tissue-Specific Neural Expression and Potential Nootropic Drug Targets. <i>Cell Reports</i> , 2017, 21, 2597-2613.	2.9	103
12	An integrative variant analysis pipeline for accurate genotype/haplotype inference in population NGS data. <i>Genome Research</i> , 2013, 23, 833-842.	2.4	93
13	Pleiotropic Meta-Analysis of Cognition, Education, and Schizophrenia Differentiates Roles of Early Neurodevelopmental and Adult Synaptic Pathways. <i>American Journal of Human Genetics</i> , 2019, 105, 334-350.	2.6	86
14	Association of a Schizophrenia Risk Variant at the <i>DRD2</i> Locus With Antipsychotic Treatment Response in First-Episode Psychosis. <i>Schizophrenia Bulletin</i> , 2015, 41, 1248-1255.	2.3	64
15	Next-generation sequencing study finds an excess of rare, coding single-nucleotide variants of <i>ADAMTS13</i> in patients with deep vein thrombosis. <i>Journal of Thrombosis and Haemostasis</i> , 2013, 11, 1228-1239.	1.9	52
16	Atlas2 Cloud: a framework for personal genome analysis in the cloud. <i>BMC Genomics</i> , 2012, 13, S19.	1.2	42
17	Expanded genetic screening panel for the Ashkenazi Jewish population. <i>Genetics in Medicine</i> , 2016, 18, 522-528.	1.1	33
18	Identification of genetic risk variants for deep vein thrombosis by multiplexed next-generation sequencing of 186 hemostatic/pro-inflammatory genes. <i>BMC Medical Genomics</i> , 2012, 5, 7.	0.7	32

#	ARTICLE	IF	CITATIONS
19	High-depth whole genome sequencing of an Ashkenazi Jewish reference panel: enhancing sensitivity, accuracy, and imputation. <i>Human Genetics</i> , 2018, 137, 343-355.	1.8	24
20	Novel ultra-rare exonic variants identified in a founder population implicate cadherins in schizophrenia. <i>Neuron</i> , 2021, 109, 1465-1478.e4.	3.8	21
21	Population Genomic Analysis of 962 Whole Genome Sequences of Humans Reveals Natural Selection in Non-Coding Regions. <i>PLoS ONE</i> , 2015, 10, e0121644.	1.1	13
22	Identifying nootropic drug targets via large-scale cognitive GWAS and transcriptomics. <i>Neuropsychopharmacology</i> , 2021, 46, 1788-1801.	2.8	12
23	Multi-Trait Analysis of GWAS and Biological Insights Into Cognition: A Response to Hill (2018). <i>Twin Research and Human Genetics</i> , 2018, 21, 394-397.	0.3	3
24	Cloud processing of 1000 genomes sequencing data using Amazon Web Service. , 2013, , .		2
25	Identification of DEEP Vein Thrombosis GENETIC RISK Variants by NEXT GENERATION Sequencing of Hemostatic Genes. <i>Blood</i> , 2011, 118, 710-710.	0.6	0
26	Rare Coding Single Nucleotide Variants of ADAMTS13 Are Associated with Deep Vein Thrombosis in a Next-Generation Sequencing Association Study. <i>Blood</i> , 2012, 120, 107-107.	0.6	0