

Jin Yu

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

27
papers

22,677
citations

18
h-index

31
g-index

31
ext. papers

28,128
ext. citations

15.3
avg, IF

5.9
L-index

#	Paper	IF	Citations
27	Novel ultra-rare exonic variants identified in a founder population implicate cadherins in schizophrenia. <i>Neuron</i> , 2021 , 109, 1465-1478.e4	13.9	8
26	Identifying nootropic drug targets via large-scale cognitive GWAS and transcriptomics. <i>Neuropsychopharmacology</i> , 2021 , 46, 1788-1801	8.7	1
25	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	11.1	37
24	Schizophrenia Polygenic Risk Score as a Predictor of Antipsychotic Efficacy in First-Episode Psychosis. <i>American Journal of Psychiatry</i> , 2019 , 176, 21-28	11.9	68
23	High-depth whole genome sequencing of an Ashkenazi Jewish reference panel: enhancing sensitivity, accuracy, and imputation. <i>Human Genetics</i> , 2018 , 137, 343-355	6.3	16
22	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	2.2	2
21	Study of 300,486 individuals identifies 148 independent genetic loci influencing general cognitive function. <i>Nature Communications</i> , 2018 , 9, 2098	17.4	254
20	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	36.3	475
19	Large-Scale Cognitive GWAS Meta-Analysis Reveals Tissue-Specific Neural Expression and Potential Nootropic Drug Targets. <i>Cell Reports</i> , 2017 , 21, 2597-2613	10.6	71
18	Expanded genetic screening panel for the Ashkenazi Jewish population. <i>Genetics in Medicine</i> , 2016 , 18, 522-8	8.1	24
17	A global reference for human genetic variation. <i>Nature</i> , 2015 , 526, 68-74	50.4	8599
16	Association of a Schizophrenia Risk Variant at the DRD2 Locus With Antipsychotic Treatment Response in First-Episode Psychosis. <i>Schizophrenia Bulletin</i> , 2015 , 41, 1248-55	1.3	52
15	Population genomic analysis of 962 whole genome sequences of humans reveals natural selection in non-coding regions. <i>PLoS ONE</i> , 2015 , 10, e0121644	3.7	12
14	An integrative variant analysis pipeline for accurate genotype/haplotype inference in population NGS data. <i>Genome Research</i> , 2013 , 23, 833-42	9.7	69
13	Whole-genome sequence-based analysis of high-density lipoprotein cholesterol. <i>Nature Genetics</i> , 2013 , 45, 899-901	36.3	117
12	Next-generation sequencing study finds an excess of rare, coding single-nucleotide variants of ADAMTS13 in patients with deep vein thrombosis. <i>Journal of Thrombosis and Haemostasis</i> , 2013 , 11, 1228-39	15.4	41
11	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	3.7	26

10	The 1000 Genomes Project: data management and community access. <i>Nature Methods</i> , 2012 , 9, 459-62	21.6	202
9	An integrative variant analysis suite for whole exome next-generation sequencing data. <i>BMC Bioinformatics</i> , 2012 , 13, 8	3.6	211
8	An integrated map of genetic variation from 1,092 human genomes. <i>Nature</i> , 2012 , 491, 56-65	50.4	6049
7	Atlas2 Cloud: a framework for personal genome analysis in the cloud. <i>BMC Genomics</i> , 2012 , 13 Suppl 6, S19	4.5	34
6	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	2.2	
5	The functional spectrum of low-frequency coding variation. <i>Genome Biology</i> , 2011 , 12, R84	18.3	161
4	Identification of DEEP Vein Thrombosis GENETIC RISK Variants by NEXT GENERATION Sequencing of Hemostatic Genes. <i>Blood</i> , 2011 , 118, 710-710	2.2	
3	A map of human genome variation from population-scale sequencing. <i>Nature</i> , 2010 , 467, 1061-73	50.4	6142
2	Novel Ultra-Rare Exonic Variants Identified in a Founder Population Implicate Cadherins in Schizophrenia		1
1	Pleiotropic Meta-Analysis of Cognition, Education, and Schizophrenia Differentiates Roles of Early Neurodevelopmental and Adult Synaptic Pathways		2