Jiebiao Wang

List of Publications by Year in descending order

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LIERIAO MANG

#	Article	IF	CITATIONS
1	Large-Scale Exome Sequencing Study Implicates Both Developmental and Functional Changes in the Neurobiology of Autism. Cell, 2020, 180, 568-584.e23.	28.9	1,422
2	Characterization of Nigerian breast cancer reveals prevalent homologous recombination deficiency and aggressive molecular features. Nature Communications, 2018, 9, 4181.	12.8	77
3	Transcriptional Alterations in Dorsolateral Prefrontal Cortex and Nucleus Accumbens Implicate Neuroinflammation and Synaptic Remodeling in Opioid Use Disorder. Biological Psychiatry, 2021, 90, 550-562.	1.3	76
4	Large eQTL meta-analysis reveals differing patterns between cerebral cortical and cerebellar brain regions. Scientific Data, 2020, 7, 340.	5.3	75
5	Identifying <i>cis</i> -mediators for <i>trans</i> -eQTLs across many human tissues using genomic mediation analysis. Genome Research, 2017, 27, 1859-1871.	5.5	72
6	An interactome perturbation framework prioritizes damaging missense mutations for developmental disorders. Nature Genetics, 2018, 50, 1032-1040.	21.4	64
7	Global landscape and genetic regulation of RNA editing in cortical samples from individuals with schizophrenia. Nature Neuroscience, 2019, 22, 1402-1412.	14.8	63
8	Imputing Gene Expression in Uncollected Tissues Within and Beyond GTEx. American Journal of Human Genetics, 2016, 98, 697-708.	6.2	51
9	Bayesian estimation of cell type–specific gene expression with prior derived from single-cell data. Genome Research, 2021, 31, 1807-1818.	5.5	40
10	Prospective Development and Validation of the Computerized Adaptive Screen for Suicidal Youth. JAMA Psychiatry, 2021, 78, 540.	11.0	30
11	Using multiple measurements of tissue to estimate subject- and cell-type-specific gene expression. Bioinformatics, 2020, 36, 782-788.	4.1	28
12	ESCO: single cell expression simulation incorporating gene co-expression. Bioinformatics, 2021, 37, 2374-2381.	4.1	21
13	De novo missense variants disrupting protein–protein interactions affect risk for autism through gene co-expression and protein networks in neuronal cell types. Molecular Autism, 2020, 11, 76.	4.9	19
14	Small nucleolar RNAs in plasma extracellular vesicles and their discriminatory power as diagnostic biomarkers of Alzheimer's disease. Neurobiology of Disease, 2021, 159, 105481.	4.4	17
15	Transcriptome alterations are enriched for synapse-associated genes in the striatum of subjects with obsessive-compulsive disorder. Translational Psychiatry, 2021, 11, 171.	4.8	13
16	Large-Scale Exome Sequencing Study Implicates Both Developmental and Functional Changes in the Neurobiology of Autism. SSRN Electronic Journal, 0, , .	0.4	12
17	Robust and accurate estimation of cellular fraction from tissue omics data via ensemble deconvolution. Bioinformatics, 2022, 38, 3004-3010.	4.1	10
18	A mixed-effects model for incomplete data from labeling-based quantitative proteomics experiments. Annals of Applied Statistics, 2017, 11, 114-138.	1.1	9

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19	Identification of cell-type-specific marker genes from co-expression patterns in tissue samples. Bioinformatics, 2021, 37, 3228-3234.	4.1	9
20	Medications and Suicide: High Dimensional Empirical Bayes Screening (iDEAS). , 0, , .		7
21	A metaâ€analysis approach with filtering for identifying geneâ€level gene–environment interactions. Genetic Epidemiology, 2018, 42, 434-446.	1.3	5
22	CCmed: cross-condition mediation analysis for identifying replicable trans-associations mediated by cis-gene expression. Bioinformatics, 2021, 37, 2513-2520.	4.1	4
23	Using multivariate mixed-effects selection models for analyzing batch-processed proteomics data with non-ignorable missingness. Biostatistics, 2019, 20, 648-665.	1.5	3
24	Gene Expression Deconvolution Implicates Cell-Type-Specific Gene Expression and Co-Expression in Autism. Biological Psychiatry, 2020, 87, S60-S61.	1.3	1
25	MAD1L1 Harbors Schizophrenia-Associated Differential Methylation and Methylation/Transcription Quantitative Trait Loci that Colocalize With Genetic Risk for Schizophrenia. Biological Psychiatry, 2021, 89, S153.	1.3	0
26	Abstract 4494: Whole genome sequencing reveals different patterns of mutational mechanisms in breast tumors between women of African and European descent. , 2016, , .		0