

# Alexis Battle

## List of Publications by Year in descending order

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Version: 2024-02-01

51  
papers

20,686  
citations

117625

34  
h-index

175258

52  
g-index

71  
all docs

71  
docs citations

71  
times ranked

38931  
citing authors

#	ARTICLE	IF	CITATIONS
1	Redefining tissue specificity of genetic regulation of gene expression in the presence of allelic heterogeneity. <i>American Journal of Human Genetics</i> , 2022, 109, 223-239.	6.2	26
2	Single-cell sequencing reveals lineage-specific dynamic genetic regulation of gene expression during human cardiomyocyte differentiation. <i>PLoS Genetics</i> , 2022, 18, e1009666.	3.5	28
3	Human embryoid bodies as a novel system for genomic studies of functionally diverse cell types. <i>ELife</i> , 2022, 11, .	6.0	7
4	Where Are the Disease-Associated eQTLs?. <i>Trends in Genetics</i> , 2021, 37, 109-124.	6.7	163
5	Transcriptional profile of platelets and iPSC-derived megakaryocytes from whole-genome and RNA sequencing. <i>Blood</i> , 2021, 137, 959-968.	1.4	21
6	Linear and Nonlinear Mendelian Randomization Analyses of the Association Between Diastolic Blood Pressure and Cardiovascular Events. <i>Circulation</i> , 2021, 143, 895-906.	1.6	73
7	InÂvivo CD8+ TÂcell CRISPR screening reveals control by Fli1 in infection and cancer. <i>Cell</i> , 2021, 184, 1262-1280.e22.	28.9	107
8	Population-scale tissue transcriptomics maps long non-coding RNAs to complex disease. <i>Cell</i> , 2021, 184, 2633-2648.e19.	28.9	94
9	Coexpression network architecture reveals the brain-wide and multiregional basis of disease susceptibility. <i>Nature Neuroscience</i> , 2021, 24, 1313-1323.	14.8	44
10	Large-scale cis- and trans-eQTL analyses identify thousands of genetic loci and polygenic scores that regulate blood gene expression. <i>Nature Genetics</i> , 2021, 53, 1300-1310.	21.4	590
11	DNA methylation signatures reveal that distinct combinations of transcription factors specify human immune cell epigenetic identity. <i>Immunity</i> , 2021, 54, 2465-2480.e5.	14.3	31
12	GBAT: a gene-based association test for robust detection of trans-gene regulation. <i>Genome Biology</i> , 2020, 21, 211.	8.8	12
13	Transcriptomic signatures across human tissues identify functional rare genetic variation. <i>Science</i> , 2020, 369, .	12.6	89
14	The impact of sex on gene expression across human tissues. <i>Science</i> , 2020, 369, .	12.6	329
15	A vast resource of allelic expression data spanning human tissues. <i>Genome Biology</i> , 2020, 21, 234.	8.8	68
16	sn-spMF: matrix factorization informs tissue-specific genetic regulation of gene expression. <i>Genome Biology</i> , 2020, 21, 235.	8.8	18
17	Genome-wide association and multi-omic analyses reveal ACTN2 as a gene linked to heart failure. <i>Nature Communications</i> , 2020, 11, 1122.	12.8	57
18	ACE inhibition and cardiometabolic risk factors, lung<i>ACE2</i>and<i>TMPRSS2</i>gene expression, and plasma ACE2 levels: a Mendelian randomization study. <i>Royal Society Open Science</i> , 2020, 7, 200958.	2.4	12

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19	Dynamic genetic regulation of gene expression during cellular differentiation. <i>Science</i> , 2019, 364, 1287-1290.	12.6	142
20	Identification of rare-disease genes using blood transcriptome sequencing and large control cohorts. <i>Nature Medicine</i> , 2019, 25, 911-919.	30.7	221
21	Addressing confounding artifacts in reconstruction of gene co-expression networks. <i>Genome Biology</i> , 2019, 20, 94.	8.8	68
22	Imputed gene associations identify replicable <i>trans</i> -acting genes enriched in transcription pathways and complex traits. <i>Genetic Epidemiology</i> , 2019, 43, 596-608.	1.3	19
23	Retinal transcriptome and eQTL analyses identify genes associated with age-related macular degeneration. <i>Nature Genetics</i> , 2019, 51, 606-610.	21.4	201
24	False positives in trans-eQTL and co-expression analyses arising from RNA-sequencing alignment errors. <i>F1000Research</i> , 2018, 7, 1860.	1.6	51
25	False positives in trans-eQTL and co-expression analyses arising from RNA-sequencing alignment errors. <i>F1000Research</i> , 2018, 7, 1860.	1.6	45
26	Population- and individual-specific regulatory variation in Sardinia. <i>Nature Genetics</i> , 2017, 49, 700-707.	21.4	38
27	Allele-specific expression reveals interactions between genetic variation and environment. <i>Nature Methods</i> , 2017, 14, 699-702.	19.0	135
28	The impact of structural variation on human gene expression. <i>Nature Genetics</i> , 2017, 49, 692-699.	21.4	334
29	The impact of rare variation on gene expression across tissues. <i>Nature</i> , 2017, 550, 239-243.	27.8	229
30	Genetic effects on gene expression across human tissues. <i>Nature</i> , 2017, 550, 204-213.	27.8	3,500
31	Co-expression networks reveal the tissue-specific regulation of transcription and splicing. <i>Genome Research</i> , 2017, 27, 1843-1858.	5.5	139
32	FIRE: functional inference of genetic variants that regulate gene expression. <i>Bioinformatics</i> , 2017, 33, 3895-3901.	4.1	30
33	Incorporation of Biological Knowledge Into the Study of Gene-Environment Interactions. <i>American Journal of Epidemiology</i> , 2017, 186, 771-777.	3.4	23
34	Identifying global expression patterns and key regulators in epithelial to mesenchymal transition through multi-study integration. <i>BMC Cancer</i> , 2017, 17, 447.	2.6	26
35	Impact of the X Chromosome and sex on regulatory variation. <i>Genome Research</i> , 2016, 26, 768-777.	5.5	88
36	Genetic variation in MHC proteins is associated with T cell receptor expression biases. <i>Nature Genetics</i> , 2016, 48, 995-1002.	21.4	151

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37	An Efficient Multiple-Testing Adjustment for eQTL Studies that Accounts for Linkage Disequilibrium between Variants. <i>American Journal of Human Genetics</i> , 2016, 98, 216-224.	6.2	91
38	EIF3G is associated with narcolepsy across ethnicities. <i>European Journal of Human Genetics</i> , 2015, 23, 1573-1580.	2.8	21
39	Sharing and Specificity of Co-expression Networks across 35 Human Tissues. <i>PLoS Computational Biology</i> , 2015, 11, e1004220.	3.2	158
40	The Genotype-Tissue Expression (GTEx) pilot analysis: Multitissue gene regulation in humans. <i>Science</i> , 2015, 348, 648-660.	12.6	4,659
41	Impact of regulatory variation from RNA to protein. <i>Science</i> , 2015, 347, 664-667.	12.6	399
42	High-Resolution Transcriptome Analysis with Long-Read RNA Sequencing. <i>PLoS ONE</i> , 2014, 9, e108095.	2.5	47
43	Type I interferon signaling genes in recurrent major depression: increased expression detected by whole-blood RNA sequencing. <i>Molecular Psychiatry</i> , 2014, 19, 1267-1274.	7.9	151
44	Transcriptome Sequencing of a Large Human Family Identifies the Impact of Rare Noncoding Variants. <i>American Journal of Human Genetics</i> , 2014, 95, 245-256.	6.2	63
45	Determining causality and consequence of expression quantitative trait loci. <i>Human Genetics</i> , 2014, 133, 727-735.	3.8	58
46	Characterizing the genetic basis of transcriptome diversity through RNA-sequencing of 922 individuals. <i>Genome Research</i> , 2014, 24, 14-24.	5.5	547
47	Transcriptome Analysis Reveals Differential Splicing Events in IPF Lung Tissue. <i>PLoS ONE</i> , 2014, 9, e92111.	2.5	73
48	The Genotype-Tissue Expression (GTEx) project. <i>Nature Genetics</i> , 2013, 45, 580-585.	21.4	6,815
49	Normalizing RNA-Sequencing Data by Modeling Hidden Covariates with Prior Knowledge. <i>PLoS ONE</i> , 2013, 8, e68141.	2.5	68
50	Automated identification of pathways from quantitative genetic interaction data. <i>Molecular Systems Biology</i> , 2010, 6, 379.	7.2	70
51	Probabilistic Discovery of Overlapping Cellular Processes and Their Regulation. <i>Journal of Computational Biology</i> , 2005, 12, 909-927.	1.6	24