Daniel J Gaffney

List of Publications by Year in descending order

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117571 206029 9,286 45 34 48 citations g-index h-index papers 63 63 63 19215 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	A survey of best practices for RNA-seq data analysis. Genome Biology, 2016, 17, 13.	3.8	1,898
2	DNA methylation patterns associate with genetic and gene expression variation in HapMap cell lines. Genome Biology, 2011, 12, R10.	3.8	754
3	DNase l sensitivity QTLs are a major determinant of human expression variation. Nature, 2012, 482, 390-394.	13.7	608
4	Accurate inference of transcription factor binding from DNA sequence and chromatin accessibility data. Genome Research, 2011, 21, 447-455.	2.4	501
5	Common genetic variation drives molecular heterogeneity in human iPSCs. Nature, 2017, 546, 370-375.	13.7	491
6	DNA Sequence-Dependent Compartmentalization and Silencing of Chromatin at the Nuclear Lamina. Cell, 2012, 149, 1474-1487.	13.5	405
7	Genetic Background Drives Transcriptional Variation in Human Induced Pluripotent Stem Cells. PLoS Genetics, 2014, 10, e1004432.	1.5	260
8	Genome-wide meta-analysis, fine-mapping and integrative prioritization implicate new Alzheimer's disease risk genes. Nature Genetics, 2021, 53, 392-402.	9.4	258
9	Controls of Nucleosome Positioning in the Human Genome. PLoS Genetics, 2012, 8, e1003036.	1.5	255
10	Shared genetic effects on chromatin and gene expression indicate a role for enhancer priming in immune response. Nature Genetics, 2018, 50, 424-431.	9.4	253
11	Dense fine-mapping study identifies new susceptibility loci for primary biliary cirrhosis. Nature Genetics, 2012, 44, 1137-1141.	9.4	251
12	Epithelial IL-22RA1-Mediated Fucosylation Promotes Intestinal Colonization Resistance to an Opportunistic Pathogen. Cell Host and Microbe, 2014, 16, 504-516.	5.1	237
13	Souporcell: robust clustering of single-cell RNA-seq data by genotype without reference genotypes. Nature Methods, 2020, 17, 615-620.	9.0	232
14	Genome-wide association study of primary sclerosing cholangitis identifies new risk loci and quantifies the genetic relationship with inflammatory bowel disease. Nature Genetics, 2017, 49, 269-273.	9.4	230
15	Fine-mapping cellular QTLs with RASQUAL and ATAC-seq. Nature Genetics, 2016, 48, 206-213.	9.4	199
16	Quantifying the Slightly Deleterious Mutation Model of Molecular Evolution. Molecular Biology and Evolution, 2002, 19, 2142-2149.	3.5	191
17	Molecular and functional variation in iPSC-derived sensory neurons. Nature Genetics, 2018, 50, 54-61.	9.4	191
18	Dissecting the regulatory architecture of gene expression QTLs. Genome Biology, 2012, 13, R7.	13.9	188

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19	N6-methyladenosine regulates the stability of RNA:DNA hybrids in human cells. Nature Genetics, 2020, 52, 48-55.	9.4	147
20	Population-scale single-cell RNA-seq profiling across dopaminergic neuron differentiation. Nature Genetics, 2021, 53, 304-312.	9.4	146
21	Functional constraints and frequency of deleterious mutations in noncoding DNA of rodents. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 13402-13406.	3.3	120
22	Early maturation and distinct tau pathology in induced pluripotent stem cell-derived neurons from patients with <i>MAPT </i> mutations. Brain, 2015, 138, 3345-3359.	3.7	116
23	Activin/Nodal signaling and NANOG orchestrate human embryonic stem cell fate decisions by controlling the H3K4me3 chromatin mark. Genes and Development, 2015, 29, 702-717.	2.7	115
24	A map of transcriptional heterogeneity and regulatory variation in human microglia. Nature Genetics, 2021, 53, 861-868.	9.4	115
25	The Contribution of RNA Decay Quantitative Trait Loci to Inter-Individual Variation in Steady-State Gene Expression Levels. PLoS Genetics, 2012, 8, e1003000.	1.5	104
26	Transcriptional profiling of macrophages derived from monocytes and iPS cells identifies a conserved response to LPS and novel alternative transcription. Scientific Reports, 2015, 5, 12524.	1.6	94
27	High-resolution genetic mapping of putative causal interactions between regions of open chromatin. Nature Genetics, 2019, 51, 128-137.	9.4	80
28	The scale of mutational variation in the murid genome. Genome Research, 2005, 15, 1086-1094.	2.4	75
29	False positive peaks in ChIP-seq and other sequencing-based functional assays caused by unannotated high copy number regions. Bioinformatics, 2011, 27, 2144-2146.	1.8	74
30	Genomic Selective Constraints in Murid Noncoding DNA. PLoS Genetics, 2006, 2, e204.	1.5	60
31	Global Properties and Functional Complexity of Human Gene Regulatory Variation. PLoS Genetics, 2013, 9, e1003501.	1.5	55
32	Genetic effects on promoter usage are highly context-specific and contribute to complex traits. ELife, 2019, 8, .	2.8	53
33	Loss of IL-10 signaling in macrophages limits bacterial killing driven by prostaglandin E2. Journal of Experimental Medicine, 2020, 217, .	4.2	51
34	Evolutionary constraints in conserved nongenic sequences of mammals. Genome Research, 2005, 15, 1373-1378.	2.4	50
35	Cardelino: computational integration of somatic clonal substructure and single-cell transcriptomes. Nature Methods, 2020, 17, 414-421.	9.0	48
36	Population-scale proteome variation in human induced pluripotent stem cells. ELife, 2020, 9, .	2.8	40

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37	Locus-specific expression of transposable elements in single cells with CELLO-seq. Nature Biotechnology, 2022, 40, 546-554.	9.4	38
38	DNA Sequence Error Rates in Genbank Records Estimated using the Mouse Genome as a Reference. DNA Sequence, 2004, 15, 362-364.	0.7	37
39	Unexpected conserved non-coding DNA blocks in mammals. Trends in Genetics, 2004, 20, 332-337.	2.9	22
40	Cell reprogramming shapes the mitochondrial DNA landscape. Nature Communications, 2021, 12, 5241.	5.8	21
41	Selective Constraints in Experimentally Defined Primate Regulatory Regions. PLoS Genetics, 2008, 4, e1000157.	1.5	20
42	Exon-Specific QTLs Skew the Inferred Distribution of Expression QTLs Detected Using Gene Expression Array Data. PLoS ONE, 2012, 7, e30629.	1.1	18
43	Effect of the assignment of ancestral CpG state on the estimation of nucleotide substitution rates in mammals. BMC Evolutionary Biology, 2008, 8, 265.	3.2	15
44	Robust temporal map of human in vitro myelopoiesis using single-cell genomics. Nature Communications, 2022, 13, .	5.8	13
45	Mapping and predicting gene–enhancer interactions. Nature Genetics, 2019, 51, 1662-1663.	9.4	11