

# Davis James McCarthy

## List of Publications by Year in descending order

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Version: 2024-02-01

36  
papers

47,084  
citations

270111

25  
h-index

371746

37  
g-index

53  
all docs

53  
docs citations

53  
times ranked

89021  
citing authors

#	ARTICLE	IF	CITATIONS
1	<code>edgeR</code> : a Bioconductor package for differential expression analysis of digital gene expression data. <i>Bioinformatics</i> , 2010, 26, 139-140.	1.8	32,955
2	Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. <i>Nucleic Acids Research</i> , 2012, 40, 4288-4297.	6.5	4,474
3	Scater: pre-processing, quality control, normalization and visualization of single-cell RNA-seq data in R. <i>Bioinformatics</i> , 2017, 33, 1179-1186.	1.8	1,283
4	Count-based differential expression analysis of RNA sequencing data using R and Bioconductor. <i>Nature Protocols</i> , 2013, 8, 1765-1786.	5.5	1,124
5	The genetic architecture of type 2 diabetes. <i>Nature</i> , 2016, 536, 41-47.	13.7	952
6	A step-by-step workflow for low-level analysis of single-cell RNA-seq data with Bioconductor. <i>F1000Research</i> , 2016, 5, 2122.	0.8	822
7	Eleven grand challenges in single-cell data science. <i>Genome Biology</i> , 2020, 21, 31.	3.8	742
8	A step-by-step workflow for low-level analysis of single-cell RNA-seq data. <i>F1000Research</i> , 2016, 5, 2122.	0.8	654
9	Classification of low quality cells from single-cell RNA-seq data. <i>Genome Biology</i> , 2016, 17, 29.	3.8	572
10	Testing significance relative to a fold-change threshold is a TREAT. <i>Bioinformatics</i> , 2009, 25, 765-771.	1.8	570
11	Common genetic variation drives molecular heterogeneity in human iPSCs. <i>Nature</i> , 2017, 546, 370-375.	13.7	491
12	Benchmarking single-cell RNA-sequencing protocols for cell atlas projects. <i>Nature Biotechnology</i> , 2020, 38, 747-755.	9.4	313
13	Factors influencing success of clinical genome sequencing across a broad spectrum of disorders. <i>Nature Genetics</i> , 2015, 47, 717-726.	9.4	310
14	Detecting Differential Expression in RNA-sequence Data Using Quasi-likelihood with Shrunken Dispersion Estimates. <i>Statistical Applications in Genetics and Molecular Biology</i> , 2012, 11, .	0.2	282
15	Single-cell RNA-sequencing of differentiating iPSC cells reveals dynamic genetic effects on gene expression. <i>Nature Communications</i> , 2020, 11, 810.	5.8	235
16	Tutorial: guidelines for the computational analysis of single-cell RNA sequencing data. <i>Nature Protocols</i> , 2021, 16, 1-9.	5.5	169
17	Choice of transcripts and software has a large effect on variant annotation. <i>Genome Medicine</i> , 2014, 6, 26.	3.6	158
18	Vireo: Bayesian demultiplexing of pooled single-cell RNA-seq data without genotype reference. <i>Genome Biology</i> , 2019, 20, 273.	3.8	152

#	ARTICLE	IF	CITATIONS
19	f-sLVM: scalable and versatile factor analysis for single-cell RNA-seq. <i>Genome Biology</i> , 2017, 18, 212.	3.8	119
20	NOX1 loss-of-function genetic variants in patients with inflammatory bowel disease. <i>Mucosal Immunology</i> , 2018, 11, 562-574.	2.7	71
21	Properties of structural variants and short tandem repeats associated with gene expression and complex traits. <i>Nature Communications</i> , 2020, 11, 2927.	5.8	67
22	Visualization of Biomedical Data. <i>Annual Review of Biomedical Data Science</i> , 2018, 1, 275-304.	2.8	63
23	Combined single-cell profiling of expression and DNA methylation reveals splicing regulation and heterogeneity. <i>Genome Biology</i> , 2019, 20, 30.	3.8	61
24	Cardelino: computational integration of somatic clonal substructure and single-cell transcriptomes. <i>Nature Methods</i> , 2020, 17, 414-421.	9.0	48
25	Optimizing expression quantitative trait locus mapping workflows for single-cell studies. <i>Genome Biology</i> , 2021, 22, 188.	3.8	36
26	Erythrocytosis associated with a novel missense mutation in the BPGM gene. <i>Haematologica</i> , 2014, 99, e201-e204.	1.7	35
27	Method to Synchronize Cell Cycle of Human Pluripotent Stem Cells without Affecting Their Fundamental Characteristics. <i>Stem Cell Reports</i> , 2019, 12, 165-179.	2.3	35
28	Sequence data and association statistics from 12,940 type 2 diabetes cases and controls. <i>Scientific Data</i> , 2017, 4, 170179.	2.4	31
29	MOZ and BMI1 play opposing roles during <i>Hox</i> gene activation in ES cells and in body segment identity specification in vivo. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 5437-5442.	3.3	28
30	Aliskiren increases bradykinin and tissue kallikrein mRNA levels in the heart. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2011, 38, 623-631.	0.9	23
31	Discovery and quality analysis of a comprehensive set of structural variants and short tandem repeats. <i>Nature Communications</i> , 2020, 11, 2928.	5.8	22
32	Key signaling networks are dysregulated in patients with the adipose tissue disorder, lipedema. <i>International Journal of Obesity</i> , 2022, 46, 502-514.	1.6	15
33	Personalized genome structure via single gamete sequencing. <i>Genome Biology</i> , 2021, 22, 112.	3.8	10
34	Case Report: Hypoglycemia Due to a Novel Activating Glucokinase Variant in an Adult – a Molecular Approach. <i>Frontiers in Endocrinology</i> , 2022, 13, 842937.	1.5	8
35	Differential Expression for RNA Sequencing (RNA-Seq) Data: Mapping, Summarization, Statistical Analysis, and Experimental Design. , 2012, , 169-190.		4
36	splatPop: simulating population scale single-cell RNA sequencing data. <i>Genome Biology</i> , 2021, 22, 341.	3.8	4