

Daniel Cameron

List of Publications by Year in descending order

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20
papers

1,071
citations

758635

12
h-index

752256

20
g-index

29
all docs

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docs citations

29
times ranked

2358
citing authors

#	ARTICLE	IF	CITATIONS
1	StructuralVariantAnnotation: a R/Bioconductor foundation for a caller-agnostic structural variant software ecosystem. <i>Bioinformatics</i> , 2022, 38, 2046-2048.	1.8	4
2	Unscrambling cancer genomes via integrated analysis of structural variation and copy number. <i>Cell Genomics</i> , 2022, 2, 100112.	3.0	34
3	A multi-platform reference for somatic structural variation detection. <i>Cell Genomics</i> , 2022, 2, 100139.	3.0	10
4	An international virtual hackathon to build tools for the analysis of structural variants within species ranging from coronaviruses to vertebrates. <i>F1000Research</i> , 2021, 10, 246.	0.8	3
5	Evolution of late-stage metastatic melanoma is dominated by aneuploidy and whole genome doubling. <i>Nature Communications</i> , 2021, 12, 1434.	5.8	46
6	VIRUSBreakend: Viral Integration Recognition Using Single Breakends. <i>Bioinformatics</i> , 2021, 37, 3115-3119.	1.8	15
7	GRIDSS2: comprehensive characterisation of somatic structural variation using single breakend variants and structural variant phasing. <i>Genome Biology</i> , 2021, 22, 202.	3.8	73
8	An international virtual hackathon to build tools for the analysis of structural variants within species ranging from coronaviruses to vertebrates. <i>F1000Research</i> , 2021, 10, 246.	0.8	2
9	Relationship between CD4 T cell turnover, cellular differentiation and HIV persistence during ART. <i>PLoS Pathogens</i> , 2021, 17, e1009214.	2.1	25
10	GA4GH: International policies and standards for data sharing across genomic research and healthcare. <i>Cell Genomics</i> , 2021, 1, 100029.	3.0	94
11	The GA4GH Variation Representation Specification: A computational framework for variation representation and federated identification. <i>Cell Genomics</i> , 2021, 1, 100027.	3.0	18
12	Comprehensive evaluation and characterisation of short read general-purpose structural variant calling software. <i>Nature Communications</i> , 2019, 10, 3240.	5.8	184
13	Barcoding reveals complex clonal behavior in patient-derived xenografts of metastatic triple negative breast cancer. <i>Nature Communications</i> , 2019, 10, 766.	5.8	99
14	Optical mapping reveals a higher level of genomic architecture of chained fusions in cancer. <i>Genome Research</i> , 2018, 28, 726-738.	2.4	41
15	TERT structural rearrangements in metastatic pheochromocytomas. <i>Endocrine-Related Cancer</i> , 2018, 25, 1-9.	1.6	45
16	Detection of clinically relevant early genomic lesions in B cell malignancies from circulating tumour DNA using a single hybridisation-based next generation sequencing assay. <i>British Journal of Haematology</i> , 2018, 183, 146-149.	1.2	8
17	GRIDSS: sensitive and specific genomic rearrangement detection using positional de Bruijn graph assembly. <i>Genome Research</i> , 2017, 27, 2050-2060.	2.4	255
18	Digital PCR of Genomic Rearrangements for Monitoring Circulating Tumour DNA. <i>Advances in Experimental Medicine and Biology</i> , 2016, 924, 139-146.	0.8	6

#	ARTICLE	IF	CITATIONS
19	Bioinformatics Analysis of Sequence Data. , 2016, , 317-333.		0
20	Knowledge-Based Autonomous Dynamic Colour Calibration. Lecture Notes in Computer Science, 2004, , 226-237.	1.0	24