## FÃ;bio C P Navarro

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5096216/publications.pdf

Version: 2024-02-01

30 papers 7,368 citations

331259 21 h-index 433756 31 g-index

34 all docs

34 docs citations

times ranked

34

16021 citing authors

#	Article	IF	Citations
1	GENCODE reference annotation for the human and mouse genomes. Nucleic Acids Research, 2019, 47, D766-D773.	6.5	2,350
2	Transcriptome-wide isoform-level dysregulation in ASD, schizophrenia, and bipolar disorder. Science, 2018, 362, .	6.0	805
3	Multi-platform discovery of haplotype-resolved structural variation in human genomes. Nature Communications, 2019, 10, 1784.	5.8	636
4	GENCODE 2021. Nucleic Acids Research, 2021, 49, D916-D923.	6.5	633
5	Comprehensive functional genomic resource and integrative model for the human brain. Science, 2018, 362, .	6.0	618
6	Integrative functional genomic analysis of human brain development and neuropsychiatric risks. Science, 2018, 362, .	6.0	516
7	The PsychENCODE project. Nature Neuroscience, 2015, 18, 1707-1712.	7.1	371
8	Pan-cancer analysis of whole genomes identifies driver rearrangements promoted by LINE-1 retrotransposition. Nature Genetics, 2020, 52, 306-319.	9.4	275
9	Transcriptome and epigenome landscape of human cortical development modeled in organoids. Science, 2018, 362, .	6.0	220
10	Diverse human extracellular RNAs are widely detected in human plasma. Nature Communications, 2016, 7, 11106.	5.8	170
11	Sixteen diverse laboratory mouse reference genomes define strain-specific haplotypes and novel functional loci. Nature Genetics, 2018, 50, 1574-1583.	9.4	169
12	Repeat associated mechanisms of genome evolution and function revealed by the <i>Mus caroli</i> and <i>Mus pahari</i> genomes. Genome Research, 2018, 28, 448-459.	2.4	99
13	Gene Copy-Number Polymorphism Caused by Retrotransposition in Humans. PLoS Genetics, 2013, 9, e1003242.	1.5	88
14	Neuronal and glial 3D chromatin architecture informs the cellular etiology of brain disorders. Nature Communications, 2021, 12, 3968.	5.8	48
15	A Genome-Wide Landscape of Retrocopies in Primate Genomes. Genome Biology and Evolution, 2015, 7, 2265-2275.	1.1	46
16	FusorSV: an algorithm for optimally combining data from multiple structural variation detection methods. Genome Biology, 2018, 19, 38.	3.8	46
17	Genomics and data science: an application within an umbrella. Genome Biology, 2019, 20, 109.	3.8	46
18	RCPedia: a database of retrocopied genes. Bioinformatics, 2013, 29, 1235-1237.	1.8	32

#	Article	IF	CITATIONS
19	Mutational analysis of genes coding for cell surface proteins in colorectal cancer cell lines reveal novel altered pathways, druggable mutations and mutated epitopes for targeted therapy. Oncotarget, 2014, 5, 9199-9213.	0.8	31
20	High IL-1R8 expression in breast tumors promotes tumor growth and contributes to impaired antitumor immunity. Oncotarget, 2017, 8, 49470-49483.	0.8	24
21	TeXP: Deconvolving the effects of pervasive and autonomous transcription of transposable elements. PLoS Computational Biology, 2019, 15, e1007293.	1.5	24
22	Prediction of Immunotherapy Response in Melanoma through Combined Modeling of Neoantigen Burden and Immune-Related Resistance Mechanisms. Clinical Cancer Research, 2021, 27, 4265-4276.	3.2	23
23	Distinct patterns of somatic alterations in a lymphoblastoid and a tumor genome derived from the same individual. Nucleic Acids Research, 2011, 39, 6056-6068.	6.5	19
24	Origins and characterization of variants shared between databases of somatic and germline human mutations. BMC Bioinformatics, 2020, 21, 227.	1.2	14
25	Retroposed copies of RET gene: a somatically acquired event in medullary thyroid carcinoma. BMC Medical Genomics, 2019, 12, 104.	0.7	10
26	To mock or not: a comprehensive comparison of mock IP and DNA input for ChIP-seq. Nucleic Acids Research, 2021, 49, e17-e17.	6.5	8
27	SPLOOCE. RNA Biology, 2012, 9, 1339-1343.	1.5	7
28	ICRmax: An optimized approach to detect tumor-specific interchromosomal rearrangements for clinical application. Genomics, 2015, 105, 265-272.	1.3	4
29	STK11/LKB1 Loss of Function Is Associated with Global DNA Hypomethylation and <i>S</i> >-Adenosyl-Methionine Depletion in Human Lung Adenocarcinoma. Cancer Research, 2021, 81, 4194-4204.	0.4	4
30	FANCY: fast estimation of privacy risk in functional genomics data. Bioinformatics, 2021, 36, 5145-5150.	1.8	3