David A Knowles

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5940664/publications.pdf

Version: 2024-02-01

27 papers

4,862 citations

20 h-index 27 g-index

42 all docs 42 docs citations

times ranked

42

12091 citing authors

#	Article	IF	CITATIONS
1	An integrated approach to identify environmental modulators of genetic risk factors for complex traits. American Journal of Human Genetics, 2021, 108, 1866-1879.	6.2	9
2	Welch-weighted Egger regression reduces false positives due to correlated pleiotropy in Mendelian randomization. American Journal of Human Genetics, 2021, 108, 2319-2335.	6.2	8
3	Distinct Classes of Complex Structural Variation Uncovered across Thousands of Cancer Genome Graphs. Cell, 2020, 183, 197-210.e32.	28.9	141
4	A human lung tumor microenvironment interactome identifies clinically relevant cell-type cross-talk. Genome Biology, 2020, 21, 107.	8.8	33
5	Molecular Choreography of Acute Exercise. Cell, 2020, 181, 1112-1130.e16.	28.9	261
6	Genetic regulation of gene expression and splicing during a 10-year period of human aging. Genome Biology, 2019, 20, 230.	8.8	57
7	Sparse discriminative latent characteristics for predicting cancer drug sensitivity from genomic features. PLoS Computational Biology, 2019, 15, e1006743.	3.2	4
8	Opportunities and challenges for transcriptome-wide association studies. Nature Genetics, 2019, 51, 592-599.	21.4	592
9	Landscape of stimulation-responsive chromatin across diverse human immune cells. Nature Genetics, 2019, 51, 1494-1505.	21.4	196
10	Annotation-free quantification of RNA splicing using LeafCutter. Nature Genetics, 2018, 50, 151-158.	21.4	520
11	Interactions between genetic variation and cellular environment in skeletal muscle gene expression. PLoS ONE, 2018, 13, e0195788.	2.5	18
12	Determining the genetic basis of anthracycline-cardiotoxicity by molecular response QTL mapping in induced cardiomyocytes. ELife, 2018, 7, .	6.0	94
13	Therapeutic reduction of ataxin-2 extends lifespan and reduces pathology in TDP-43 mice. Nature, 2017, 544, 367-371.	27.8	422
14	Allele-specific expression reveals interactions between genetic variation and environment. Nature Methods, 2017, 14, 699-702.	19.0	135
15	Batch effects and the effective design of single-cell gene expression studies. Scientific Reports, 2017, 7, 39921.	3.3	275
16	Inferring Relevant Cell Types for Complex Traits by Using Single-Cell Gene Expression. American Journal of Human Genetics, 2017, 101, 686-699.	6.2	102
17	Small RNA Sequencing in Cells and Exosomes Identifies eQTLs and 14q32 as a Region of Active Export. G3: Genes, Genomes, Genetics, 2017, 7, 31-39.	1.8	16
18	Impact of the X Chromosome and sex on regulatory variation. Genome Research, 2016, 26, 768-777.	5 . 5	88

#	Article	IF	CITATIONS
19	RNA splicing is a primary link between genetic variation and disease. Science, 2016, 352, 600-604.	12.6	574
20	An Efficient Multiple-Testing Adjustment for eQTL Studies that Accounts for Linkage Disequilibrium between Variants. American Journal of Human Genetics, 2016, 98, 216-224.	6.2	91
21	Relational Learning and Network Modelling Using Infinite Latent Attribute Models. IEEE Transactions on Pattern Analysis and Machine Intelligence, 2015, 37, 462-474.	13.9	8
22	Pitman Yor Diffusion Trees for Bayesian Hierarchical Clustering. IEEE Transactions on Pattern Analysis and Machine Intelligence, 2015, 37, 271-289.	13.9	11
23	Allelic Expression of Deleterious Protein-Coding Variants across Human Tissues. PLoS Genetics, 2014, 10, e1004304.	3.5	60
24	Transcriptome Sequencing of a Large Human Family Identifies the Impact of Rare Noncoding Variants. American Journal of Human Genetics, 2014, 95, 245-256.	6.2	63
25	Fixed-Form Variational Posterior Approximation through Stochastic Linear Regression. Bayesian Analysis, 2013, 8, .	3.0	65
26	Mapping cis- and trans-regulatory effects across multiple tissues in twins. Nature Genetics, 2012, 44, 1084-1089.	21.4	701
27	Distinct Epigenomic Features in End-Stage Failing Human Hearts. Circulation, 2011, 124, 2411-2422.	1.6	245