Aaron K Wong

List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/5886205/aaron-k-wong-publications-by-year.pdf

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

38 41 2,335 20 h-index g-index citations papers 4.66 41 19 3,237 L-index ext. citations avg, IF ext. papers

#	Paper	IF	Citations
38	Tissue-specific enhancer functional networks for associating distal regulatory regions to disease. <i>Cell Systems</i> , 2021 , 12, 353-362.e6	10.6	6
37	Modeling transcriptional regulation of model species with deep learning. <i>Genome Research</i> , 2021 , 31, 1097-1105	9.7	2
36	Machine learning methods to model multicellular complexity and tissue specificity. <i>Nature Reviews Materials</i> , 2021 , 6, 717-729	73.3	O
35	Genome-wide landscape of RNA-binding protein target site dysregulation reveals a major impact on psychiatric disorder risk. <i>Nature Genetics</i> , 2021 , 53, 166-173	36.3	10
34	Spatial transcriptional mapping of the human nephrogenic program. <i>Developmental Cell</i> , 2021 , 56, 238 ²	1- <u>23.9</u> 8	.e%
33	Decoding disease: from genomes to networks to phenotypes. <i>Nature Reviews Genetics</i> , 2021 , 22, 774-79	99 0.1	7
32	SARS-CoV-2 receptor networks in diabetic and COVID-19 associated kidney disease 2020 ,		2
31	SARS-CoV-2 receptor networks in diabetic and COVID-19-associated kidney disease. <i>Kidney International</i> , 2020 , 98, 1502-1518	9.9	33
30	Whole-genome deep-learning analysis identifies contribution of noncoding mutations to autism risk. <i>Nature Genetics</i> , 2019 , 51, 973-980	36.3	122
29	Deep learning sequence-based ab initio prediction of variant effects on expression and disease risk. <i>Nature Genetics</i> , 2018 , 50, 1171-1179	36.3	155
28	Enabling Precision Medicine through Integrative Network Models. <i>Journal of Molecular Biology</i> , 2018 , 430, 2913-2923	6.5	11
27	Interpretation of an individual functional genomics experiment guided by massive public data. <i>Nature Methods</i> , 2018 , 15, 1049-1052	21.6	2
26	An integrative tissue-network approach to identify and test human disease genes. <i>Nature Biotechnology</i> , 2018 ,	44.5	31
25	GIANT 2.0: genome-scale integrated analysis of gene networks in tissues. <i>Nucleic Acids Research</i> , 2018 , 46, W65-W70	20.1	33
24	GIANT API: an application programming interface for functional genomics. <i>Nucleic Acids Research</i> , 2016 , 44, W587-92	20.1	2
23	Network-based analysis of genetic variants associated with hippocampal volume in Alzheimers disease: a study of ADNI cohorts. <i>BioData Mining</i> , 2016 , 9, 3	4.3	21
22	Genome-wide prediction and functional characterization of the genetic basis of autism spectrum disorder. <i>Nature Neuroscience</i> , 2016 , 19, 1454-1462	25.5	214

21	Targeted exploration and analysis of large cross-platform human transcriptomic compendia. <i>Nature Methods</i> , 2015 , 12, 211-4, 3 p following 214	21.6	92	
20	Understanding multicellular function and disease with human tissue-specific networks. <i>Nature Genetics</i> , 2015 , 47, 569-76	36.3	473	
19	Tissue-aware data integration approach for the inference of pathway interactions in metazoan organisms. <i>Bioinformatics</i> , 2015 , 31, 1093-101	7.2	63	
18	Low-variance RNAs identify Parkinsons disease molecular signature in blood. <i>Movement Disorders</i> , 2015 , 30, 813-21	7	14	
17	IMP 2.0: a multi-species functional genomics portal for integration, visualization and prediction of protein functions and networks. <i>Nucleic Acids Research</i> , 2015 , 43, W128-33	20.1	44	
16	Interactive Big Data Resource to Elucidate Human Immune Pathways and Diseases. <i>Immunity</i> , 2015 , 43, 605-14	32.3	41	
15	FNTM: a server for predicting functional networks of tissues in mouse. <i>Nucleic Acids Research</i> , 2015 , 43, W182-7	20.1	16	
14	Functional knowledge transfer for high-accuracy prediction of under-studied biological processes. <i>PLoS Computational Biology</i> , 2013 , 9, e1002957	5	44	
13	Marker panels for genealogy-based mapping, breed demographics, and inference-of-ancestry in the dog. <i>Animal Biotechnology</i> , 2012 , 23, 241-52	1.4	1	
12	Tissue-specific functional networks for prioritizing phenotype and disease genes. <i>PLoS Computational Biology</i> , 2012 , 8, e1002694	5	114	
11	IMP: a multi-species functional genomics portal for integration, visualization and prediction of protein functions and networks. <i>Nucleic Acids Research</i> , 2012 , 40, W484-90	20.1	76	
10	An ADAMTSL2 founder mutation causes Musladin-Lueke Syndrome, a heritable disorder of beagle dogs, featuring stiff skin and joint contractures. <i>PLoS ONE</i> , 2010 , 5, e12817	3.7	22	
9	Tracking footprints of artificial selection in the dog genome. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010 , 107, 1160-5	11.5	222	
8	A comprehensive linkage map of the dog genome. <i>Genetics</i> , 2010 , 184, 595-605	4	71	
7	Coat variation in the domestic dog is governed by variants in three genes. <i>Science</i> , 2009 , 326, 150-3	33.3	226	
6	Breed distribution and history of canine mdr1-1Delta, a pharmacogenetic mutation that marks the emergence of breeds from the collie lineage. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004 , 101, 11725-30	11.5	139	
5	Tribe: The collaborative platform for reproducible web-based analysis of gene sets		1	
4	DeepArk: modeling cis-regulatory codes of model species with deep learning		1	

2

Spatial Transcriptional Mapping of the Human Nephrogenic Program

Genome-wide characterization of genetic and functional dysregulation in autism spectrum disorder

2

A sequence-based global map of regulatory activity for deciphering human genetics