Ryan D Hernandez

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/119406/publications.pdf

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51 papers 9,729 citations

34 h-index 53 g-index

68 all docs 68
docs citations

68 times ranked 15541 citing authors

#	Article	IF	CITATIONS
1	Inferring the Joint Demographic History of Multiple Populations from Multidimensional SNP Frequency Data. PLoS Genetics, 2009, 5, e1000695.	3.5	1,522
2	Sequencing of 53,831 diverse genomes from the NHLBI TOPMed Program. Nature, 2021, 590, 290-299.	27.8	1,069
3	Genome-Wide Survey of SNP Variation Uncovers the Genetic Structure of Cattle Breeds. Science, 2009, 324, 528-532.	12.6	746
4	Assessing the Evolutionary Impact of Amino Acid Mutations in the Human Genome. PLoS Genetics, 2008, 4, e1000083.	3.5	586
5	selscan: An Efficient Multithreaded Program to Perform EHH-Based Scans for Positive Selection. Molecular Biology and Evolution, 2014, 31, 2824-2827.	8.9	555
6	Classic Selective Sweeps Were Rare in Recent Human Evolution. Science, 2011, 331, 920-924.	12.6	432
7	Genetic Ancestry and Natural Selection Drive Population Differences in Immune Responses to Pathogens. Cell, 2016, 167, 657-669.e21.	28.9	419
8	Genome-Wide Patterns of Nucleotide Polymorphism in Domesticated Rice. PLoS Genetics, 2007, 3, e163.	3.5	406
9	Inherited causes of clonal haematopoiesis in 97,691 whole genomes. Nature, 2020, 586, 763-768.	27.8	376
10	Proportionally more deleterious genetic variation in European than in African populations. Nature, 2008, 451, 994-997.	27.8	365
11	Simultaneous inference of selection and population growth from patterns of variation in the human genome. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 7882-7887.	7.1	310
12	Global environmental drivers of influenza. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 13081-13086.	7.1	239
13	A flexible forward simulator for populations subject to selection and demography. Bioinformatics, 2008, 24, 2786-2787.	4.1	197
14	Context Dependence, Ancestral Misidentification, and Spurious Signatures of Natural Selection. Molecular Biology and Evolution, 2007, 24, 1792-1800.	8.9	162
15	Demographic Histories and Patterns of Linkage Disequilibrium in Chinese and Indian Rhesus Macaques. Science, 2007, 316, 240-243.	12.6	161
16	Assessing the contribution of rare variants to complex trait heritability from whole-genome sequence data. Nature Genetics, 2022, 54, 263-273.	21.4	156
17	Dynamic incorporation of multiple in silico functional annotations empowers rare variant association analysis of large whole-genome sequencing studies at scale. Nature Genetics, 2020, 52, 969-983.	21.4	146
18	A continuum of admixture in the Western Hemisphere revealed by the African Diaspora genome. Nature Communications, 2016, 7, 12522.	12.8	136

#	Article	IF	Citations
19	Evolutionary Processes Acting on Candidate cis-Regulatory Regions in Humans Inferred from Patterns of Polymorphism and Divergence. PLoS Genetics, 2009, 5, e1000592.	3.5	123
20	A genome-wide association study of bronchodilator response in Latinos implicates rare variants. Journal of Allergy and Clinical Immunology, 2014, 133, 370-378.e15.	2.9	105
21	Ultrarare variants drive substantial cis heritability of human gene expression. Nature Genetics, 2019, 51, 1349-1355.	21.4	98
22	A <i>Cutibacterium acnes </i> antibiotic modulates human skin microbiota composition in hair follicles. Science Translational Medicine, 2020, 12, .	12.4	83
23	Human demographic history has amplified the effects of background selection across the genome. PLoS Genetics, 2018, 14, e1007387.	3.5	71
24	Association study in African-admixed populations across the Americas recapitulates asthma risk loci in non-African populations. Nature Communications, 2019, 10, 880.	12.8	71
25	De novo mutations across 1,465 diverse genomes reveal mutational insights and reductions in the Amish founder population. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 2560-2569.	7.1	71
26	A high-resolution HLA reference panel capturing global population diversity enables multi-ancestry fine-mapping in HIV host response. Nature Genetics, 2021, 53, 1504-1516.	21.4	69
27	Selection and explosive growth alter genetic architecture and hamper the detection of causal rare variants. Genome Research, 2016, 26, 863-873.	5.5	63
28	Functional Segregation of Overlapping Genes in HIV. Cell, 2016, 167, 1762-1773.e12.	28.9	58
29	Cancer-associated arginine-to-histidine mutations confer a gain in pH sensing to mutant proteins. Science Signaling, 2017, 10, .	3.6	54
30	A genome-wide association and admixture mapping study of bronchodilator drug response in African Americans with asthma. Pharmacogenomics Journal, 2019, 19, 249-259.	2.0	54
31	Population Genetics of Rare Variants and Complex Diseases. Human Heredity, 2012, 74, 118-128.	0.8	53
32	Genome-wide association study and admixture mapping reveal new loci associated with total IgE levels in Latinos. Journal of Allergy and Clinical Immunology, 2015, 135, 1502-1510.	2.9	52
33	Context-Dependent Mutation Rates May Cause Spurious Signatures of a Fixation Bias Favoring Higher GC-Content in Humans. Molecular Biology and Evolution, 2007, 24, 2196-2202.	8.9	50
34	Population sequencing data reveal a compendium of mutational processes in the human germ line. Science, 2021, 373, 1030-1035.	12.6	43
35	ROP: dumpster diving in RNA-sequencing to find the source of 1 trillion reads across diverse adult human tissues. Genome Biology, 2018, 19, 36.	8.8	42
36	Statistical inference of a convergent antibody repertoire response to influenza vaccine. Genome Medicine, 2016, 8, 60.	8.2	41

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37	Ancestry-Dependent Enrichment of Deleterious Homozygotes in Runs of Homozygosity. American Journal of Human Genetics, 2019, 105, 747-762.	6.2	36
38	An admixture mapping meta-analysis implicates genetic variation at $18q21$ with asthma susceptibility in Latinos. Journal of Allergy and Clinical Immunology, 2019, 143, 957-969.	2.9	33
39	Genome sequencing unveils a regulatory landscape of platelet reactivity. Nature Communications, 2021, 12, 3626.	12.8	29
40	Prominent features of the amino acid mutation landscape in cancer. PLoS ONE, 2017, 12, e0183273.	2.5	26
41	Robust Forward Simulations of Recurrent Hitchhiking. Genetics, 2014, 197, 221-236.	2.9	24
42	The Temporal Dynamics of Background Selection in Nonequilibrium Populations. Genetics, 2020, 214, 1019-1030.	2.9	23
43	Genetic Simulation Tools for Postâ€Genome Wide Association Studies of Complex Diseases. Genetic Epidemiology, 2015, 39, 11-19.	1.3	22
44	Population Genetic Simulations of Complex Phenotypes with Implications for Rare Variant Association Tests. Genetic Epidemiology, 2015, 39, 35-44.	1.3	16
45	Whole-genome association analyses of sleep-disordered breathing phenotypes in the NHLBI TOPMed program. Genome Medicine, 2021, 13, 136.	8.2	16
46	Recent shifts in the genomic ancestry of Mexican Americans may alter the genetic architecture of biomedical traits. ELife, 2020, 9, .	6.0	15
47	CauseMap: fast inference of causality from complex time series. PeerJ, 2015, 3, e824.	2.0	12
48	Rock, Paper, Scissors: Harnessing Complementarity in Ortholog Detection Methods Improves Comparative Genomic Inference. G3: Genes, Genomes, Genetics, 2015, 5, 629-638.	1.8	10
49	Pooled Sequencing of Candidate Genes Implicates Rare Variants in the Development of Asthma Following Severe RSV Bronchiolitis in Infancy. PLoS ONE, 2015, 10, e0142649.	2.5	10
50	Population genetic simulation study of power in association testing across genetic architectures and study designs. Genetic Epidemiology, 2020, 44, 90-103.	1.3	7
51	Singleton Variants Dominate the Genetic Architecture of Human Gene Expression. SSRN Electronic Journal, 2018, , .	0.4	4