List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Worldwide Human Relationships Inferred from Genome-Wide Patterns of Variation. Science, 2008, 319, 1100-1104.	12.6	1,774
2	Personal Omics Profiling Reveals Dynamic Molecular and Medical Phenotypes. Cell, 2012, 148, 1293-1307.	28.9	1,134
3	Estimation of individual admixture: Analytical and study design considerations. Genetic Epidemiology, 2005, 28, 289-301.	1.3	571
4	Genetic Structure, Self-Identified Race/Ethnicity, and Confounding in Case-Control Association Studies. American Journal of Human Genetics, 2005, 76, 268-275.	6.2	513
5	Genetics of blood lipids among ~300,000 multi-ethnic participants of the Million Veteran Program. Nature Genetics, 2018, 50, 1514-1523.	21.4	497
6	Exome-wide association study of plasma lipids in >300,000 individuals. Nature Genetics, 2017, 49, 1758-1766.	21.4	470
7	The Polygenic and Monogenic Basis of Blood Traits and Diseases. Cell, 2020, 182, 1214-1231.e11.	28.9	388
8	Trans-ethnic and Ancestry-Specific Blood-Cell Genetics in 746,667 Individuals from 5 Global Populations. Cell, 2020, 182, 1198-1213.e14.	28.9	353
9	Variation and genetic control of protein abundance in humans. Nature, 2013, 499, 79-82.	27.8	343
10	Characterizing Race/Ethnicity and Genetic Ancestry for 100,000 Subjects in the Genetic Epidemiology Research on Adult Health and Aging (GERA) Cohort. Genetics, 2015, 200, 1285-1295.	2.9	273
11	A Quantitative Proteome Map of the Human Body. Cell, 2020, 183, 269-283.e19.	28.9	243
12	Reconstructing Genetic Ancestry Blocks in Admixed Individuals. American Journal of Human Genetics, 2006, 79, 1-12.	6.2	240
13	Modeling 3D Facial Shape from DNA. PLoS Genetics, 2014, 10, e1004224.	3.5	190
14	Genome-wide Association Analysis of Blood-Pressure Traits in African-Ancestry Individuals Reveals Common Associated Genes in African and Non-African Populations. American Journal of Human Genetics, 2013, 93, 545-554.	6.2	189
15	Harmonizing Genetic Ancestry and Self-identified Race/Ethnicity in Genome-wide Association Studies. American Journal of Human Genetics, 2019, 105, 763-772.	6.2	169
16	Recent Genetic Selection in the Ancestral Admixture of Puerto Ricans. American Journal of Human Genetics, 2007, 81, 626-633.	6.2	168
17	Integrative analysis of RNA, translation, and protein levels reveals distinct regulatory variation across humans. Genome Research, 2015, 25, 1610-1621.	5.5	157
18	Characterizing the admixed African ancestry of African Americans. Genome Biology, 2009, 10, R141.	9.6	145

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19	Genome-Wide Association Study Implicates Chromosome 9q21.31 as a Susceptibility Locus for Asthma in Mexican Children. PLoS Genetics, 2009, 5, e1000623.	3.5	139
20	Genetic Architecture of Skin and Eye Color in an African-European Admixed Population. PLoS Genetics, 2013, 9, e1003372.	3.5	137
21	A Large-Scale Multi-ancestry Genome-wide Study Accounting for Smoking Behavior Identifies Multiple Significant Loci for Blood Pressure. American Journal of Human Genetics, 2018, 102, 375-400.	6.2	123
22	Genome-wide Characterization of Shared and Distinct Genetic Components that Influence Blood Lipid Levels in Ethnically Diverse Human Populations. American Journal of Human Genetics, 2013, 92, 904-916.	6.2	113
23	Multi-ancestry genome-wide gene–smoking interaction study of 387,272 individuals identifies new loci associated with serum lipids. Nature Genetics, 2019, 51, 636-648.	21.4	112
24	Ancestral Components of Admixed Genomes in a Mexican Cohort. PLoS Genetics, 2011, 7, e1002410.	3.5	109
25	Genome-wide Association and Population Genetic Analysis of C-Reactive Protein in African American and Hispanic American Women. American Journal of Human Genetics, 2012, 91, 502-512.	6.2	107
26	Genome-Wide Association Studies of Quantitatively Measured Skin, Hair, and Eye Pigmentation in Four European Populations. PLoS ONE, 2012, 7, e48294.	2.5	94
27	Novel genetic associations for blood pressure identified via gene-alcohol interaction in up to 570K individuals across multiple ancestries. PLoS ONE, 2018, 13, e0198166.	2.5	94
28	Single-trait and multi-trait genome-wide association analyses identify novel loci for blood pressure in African-ancestry populations. PLoS Genetics, 2017, 13, e1006728.	3.5	88
29	Racial admixture and its impact on BMI and blood pressure in African and Mexican Americans. Human Genetics, 2006, 119, 624-633.	3.8	81
30	Frequentist Estimation of Coalescence Times From Nucleotide Sequence Data Using a Tree-Based Partition. Genetics, 2002, 161, 447-459.	2.9	76
31	Leveraging Multi-ethnic Evidence for Risk Assessment of Quantitative Traits in Minority Populations. American Journal of Human Genetics, 2017, 101, 218-226.	6.2	75
32	Leveraging population admixture to characterize the heritability of complex traits. Nature Genetics, 2014, 46, 1356-1362.	21.4	69
33	Multi-ancestry study of blood lipid levels identifies four loci interacting with physical activity. Nature Communications, 2019, 10, 376.	12.8	64
34	Meta-analysis of lipid-traits in Hispanics identifies novel loci, population-specific effects and tissue-specific enrichment of eQTLs. Scientific Reports, 2016, 6, 19429.	3.3	63
35	Trans-ethnic meta-analysis of white blood cell phenotypes. Human Molecular Genetics, 2014, 23, 6944-6960.	2.9	60
36	Genome-wide Trans-ethnic Meta-analysis Identifies Seven Genetic Loci Influencing Erythrocyte Traits and a Role for RBPMS in Erythropoiesis. American Journal of Human Genetics, 2017, 100, 51-63.	6.2	45

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37	Joint testing of genotype and ancestry association in admixed families. Genetic Epidemiology, 2010, 34, 783-791.	1.3	43
38	Regulation of gene expression and RNA editing in Drosophila adapting to divergent microclimates. Nature Communications, 2017, 8, 1570.	12.8	43
39	Locating Regions of Differential Variability in DNA and Protein Sequences. Genetics, 1999, 153, 485-495.	2.9	43
40	Dissecting Complex Diseases in Complex Populations: Asthma in Latino Americans. Proceedings of the American Thoracic Society, 2007, 4, 226-233.	3.5	41
41	A multi-ancestry genome-wide study incorporating gene–smoking interactions identifies multiple new loci for pulse pressure and mean arterial pressure. Human Molecular Genetics, 2019, 28, 2615-2633.	2.9	31
42	Inference on the Genetic Basis of Eye and Skin Color in an Admixed Population via Bayesian Linear Mixed Models. Genetics, 2017, 206, 1113-1126.	2.9	30
43	Whole-genome sequencing association analysis of quantitative red blood cell phenotypes: The NHLBI TOPMed program. American Journal of Human Genetics, 2021, 108, 874-893.	6.2	28
44	Genome-wide association and admixture analysis of glaucoma in the Women's Health Initiative. Human Molecular Genetics, 2014, 23, 6634-6643.	2.9	22
45	Leveraging Multi-ethnic Evidence for Mapping Complex Traits in Minority Populations: An Empirical Bayes Approach. American Journal of Human Genetics, 2015, 96, 740-752.	6.2	22
46	Rare transmission of commensal and pathogenic bacteria in the gut microbiome of hospitalized adults. Nature Communications, 2022, 13, 586.	12.8	21
47	<i>Trans</i> -ancestry Fine Mapping and Molecular Assays Identify Regulatory Variants at the <i>ANGPTL8</i> HDL-C GWAS Locus. G3: Genes, Genomes, Genetics, 2017, 7, 3217-3227.	1.8	19
48	Variants for HDL-C, LDL-C, and Triglycerides Identified from Admixture Mapping and Fine-Mapping Analysis in African American Families. Circulation: Cardiovascular Genetics, 2015, 8, 106-113.	5.1	18
49	Functional and structural basis of extreme conservation in vertebrate 5′ untranslated regions. Nature Genetics, 2021, 53, 729-741.	21.4	17
50	Identification of putative causal loci in whole-genome sequencing data via knockoff statistics. Nature Communications, 2021, 12, 3152.	12.8	17
51	The Association of the Vanin-1 N131S Variant with Blood Pressure Is Mediated by Endoplasmic Reticulum-Associated Degradation and Loss of Function. PLoS Genetics, 2014, 10, e1004641.	3.5	16
52	Whole-genome sequencing in diverse subjects identifies genetic correlates of leukocyte traits: The NHLBI TOPMed program. American Journal of Human Genetics, 2021, 108, 1836-1851.	6.2	14
53	Gene by Environment Investigation of Incident Lung Cancer Risk in African-Americans. EBioMedicine, 2016, 4, 153-161.	6.1	12
54	Genome-wide analysis of common and rare variants via multiple knockoffs at biobank scale, with an application to Alzheimer disease genetics. American Journal of Human Genetics, 2021, 108, 2336-2353.	6.2	12

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55	Joint genotype―and ancestryâ€based genomeâ€wide association studies in admixed populations. Genetic Epidemiology, 2017, 41, 555-566.	1.3	11
56	PREMIX: PRivacy-preserving EstiMation of Individual admiXture. AMIA Annual Symposium proceedings, 2016, 2016, 1747-1755.	0.2	11
57	PLOS Genetics Data Sharing Policy: In Pursuit of Functional Utility. PLoS Genetics, 2015, 11, e1005716.	3.5	10
58	Robust identification of temporal biomarkers in longitudinal omics studies. Bioinformatics, 2022, 38, 3802-3811.	4.1	10
59	Genomeâ€wide survey in African Americans demonstrates potential epistasis of fitness in the human genome. Genetic Epidemiology, 2017, 41, 122-135.	1.3	9
60	Skin color variation in Africa. Science, 2017, 358, 867-868.	12.6	8
61	Controlling for false positive findings of trans-hubs in expression quantitative trait loci mapping. BMC Proceedings, 2007, 1, S157.	1.6	6
62	A Poisson Log-Normal Model for Constructing Gene Covariation Network Using RNA-seq Data. Journal of Computational Biology, 2017, 24, 721-731.	1.6	4
63	Detecting fitness epistasis in recently admixed populations with genome-wide data. BMC Genomics, 2020, 21, 476.	2.8	4
64	Combining multiple family-based association studies. BMC Proceedings, 2007, 1, S162.	1.6	3
65	Response—How the Gray Wolf Got Its Color. Science, 2009, 325, 34-34.	12.6	3
66	RobNorm: model-based robust normalization method for labeled quantitative mass spectrometry proteomics data. Bioinformatics, 2021, 37, 815-821.	4.1	3
67	Evaluating the strength of genetic results: Risks and responsibilities. PLoS Genetics, 2019, 15, e1008437.	3.5	1
68	GENETIC STRUCTURE OF HUMAN POPULATION. , 2015, , 937-960.		0
69	Doubling down on forensic twin studies. PLoS Genetics, 2018, 14, e1007831.	3.5	0
70	Advances and challenges in quantitative delineation of the genetic architecture of complex traits. Quantitative Biology, 2021, 9, 168-184.	0.5	0
71	ldentification Of The Disease-Causing Mutation In Autosomal Dominant Familial Immune Thrombocytopenia By Genome-Wide Linkage Analysis and Whole Genome Sequencing. Blood, 2013, 122, 565-565.	1.4	0
72	Expanding human variation at PLOS Genetics. PLoS Genetics. 2022, 18. e1010070.	3.5	0