Andrea Ganna

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6294168/publications.pdf

Version: 2024-02-01

64 papers

13,342 citations

35 h-index 65 g-index

85 all docs 85 docs citations

85 times ranked

32197 citing authors

#	Article	IF	CITATIONS
1	Effect of General Adiposity and Central Body Fat Distribution on the Circulating Metabolome: A Multicohort Nontargeted Metabolomics Observational and Mendelian Randomization Study. Diabetes, 2022, 71, 329-339.	0.3	14
2	Fast estimation of genetic correlation for biobank-scale data. American Journal of Human Genetics, 2022, 109, 24-32.	2.6	11
3	Mendelian randomization of genetically independent aging phenotypes identifies LPA and VCAM1 as biological targets for human aging. Nature Aging, 2022, 2, 19-30.	5.3	17
4	Multi-ancestry fine mapping implicates OAS1 splicing in risk of severe COVID-19. Nature Genetics, 2022, 54, 125-127.	9.4	75
5	Rare coding variants in ten genes confer substantial risk for schizophrenia. Nature, 2022, 604, 509-516.	13.7	326
6	Genome-wide risk prediction of common diseases across ancestries in one million people. Cell Genomics, 2022, 2, 100118.	3.0	34
7	The human genetic epidemiology of COVID-19. Nature Reviews Genetics, 2022, 23, 533-546.	7.7	64
8	Novel genes and sex differences in COVID-19 severity. Human Molecular Genetics, 2022, 31, 3789-3806.	1.4	38
9	The 22q11.2 region regulates presynaptic gene-products linked to schizophrenia. Nature Communications, 2022, 13, .	5.8	22
10	Detailed stratified GWAS analysis for severe COVID-19 in four European populations. Human Molecular Genetics, 2022, 31, 3945-3966.	1.4	46
11	Sex-dimorphic genetic effects and novel loci for fasting glucose and insulin variability. Nature Communications, 2021, 12, 24.	5.8	87
12	Response to Comment on "Large-scale GWAS reveals insights into the genetic architecture of same-sex sexual behavior― Science, 2021, 371, .	6.0	5
13	Genetic analyses identify widespread sex-differential participation bias. Nature Genetics, 2021, 53, 663-671.	9.4	124
14	COVID-19 tissue atlases reveal SARS-CoV-2 pathology and cellular targets. Nature, 2021, 595, 107-113.	13.7	537
15	Hematopoietic mosaic chromosomal alterations increase the risk for diverse types of infection. Nature Medicine, 2021, 27, 1012-1024.	15.2	109
16	Mapping the human genetic architecture of COVID-19. Nature, 2021, 600, 472-477.	13.7	640
17	Clinical Conditions and Their Impact on Utility of Genetic Scores for Prediction of Acute Coronary Syndrome. Circulation Genomic and Precision Medicine, 2021, 14, e003283.	1.6	4
18	Using symptom-based case predictions to identify host genetic factors that contribute to COVID-19 susceptibility. PLoS ONE, 2021, 16, e0255402.	1.1	6

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19	Addendum: The mutational constraint spectrum quantified from variation in 141,456 humans. Nature, 2021, 597, E3-E4.	13.7	45
20	Age-dependent impact of the major common genetic risk factor for COVID-19 on severity and mortality. Journal of Clinical Investigation, 2021, 131, .	3.9	72
21	Polygenic risk score for general cognitive function is associated with measures of cognition in South Asians from the LASI-DAD Study Alzheimer's and Dementia, 2021, 17 Suppl 3, e053977.	0.4	О
22	Common and rare variants in topologically associated domains for cognitive function in South Asians from the LASI-DAD Study Alzheimer's and Dementia, 2021, 17 Suppl 3, e054029.	0.4	0
23	A Multi-Cohort Metabolomics Analysis Discloses Sphingomyelin (32:1) Levels to be Inversely Related to Incident Ischemic Stroke. Journal of Stroke and Cerebrovascular Diseases, 2020, 29, 104476.	0.7	14
24	Non-targeted urine metabolomics and associations with prevalent and incident type 2 diabetes. Scientific Reports, 2020, 10, 16474.	1.6	11
25	Development and validation of risk prediction models for multiple cardiovascular diseases and Type 2 diabetes. PLoS ONE, 2020, 15, e0235758.	1.1	13
26	Association Between Episodic Memory and Genetic Risk Factors for Alzheimer's Disease in South Asians from the Longitudinal Aging Study in India–Diagnostic Assessment of Dementia (LASIâ€DAD). Journal of the American Geriatrics Society, 2020, 68, S45-S53.	1.3	10
27	Common and rare variants in Alzheimer's disease genes are associated with episodic memory in South Asians from the LASIâ€DAD study. Alzheimer's and Dementia, 2020, 16, e045189.	0.4	0
28	Proteomic profiles before and during weight loss: Results from randomized trial of dietary intervention. Scientific Reports, 2020, 10, 7913.	1.6	22
29	The mutational constraint spectrum quantified from variation in 141,456 humans. Nature, 2020, 581, 434-443.	13.7	6,140
30	Discriminating bipolar depression from major depressive disorder with polygenic risk scores. Psychological Medicine, 2020, 51, 1-8.	2.7	12
31	The plasma protein profile and cardiovascular risk differ between intima-media thickness of the common carotid artery and the bulb: A meta-analysis and a longitudinal evaluation. Atherosclerosis, 2020, 295, 25-30.	0.4	18
32	Large-scale GWAS reveals insights into the genetic architecture of same-sex sexual behavior. Science, 2019, 365, .	6.0	245
33	The metabolites urobilin and sphingomyelin (30:1) are associated with incident heart failure in the general population. ESC Heart Failure, 2019, 6, 764-773.	1.4	23
34	Polygenic adaptation on height is overestimated due to uncorrected stratification in genome-wide association studies. ELife, 2019, 8, .	2.8	276
35	Genome studies must account for history—Response. Science, 2019, 366, 1461-1462.	6.0	4
36	Identification of metabolic profiles associated with human exposure to perfluoroalkyl substances. Journal of Exposure Science and Environmental Epidemiology, 2019, 29, 196-205.	1.8	55

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37	Exploring Comorbidity Within Mental Disorders Among a Danish National Population. JAMA Psychiatry, 2019, 76, 259.	6.0	374
38	Functional architecture of low-frequency variants highlights strength of negative selection across coding and non-coding annotations. Nature Genetics, 2018, 50, 1600-1607.	9.4	132
39	Genetic predictors of testosterone and their associations with cardiovascular disease and risk factors: A Mendelian randomization investigation. International Journal of Cardiology, 2018, 267, 171-176.	0.8	49
40	Quantifying the Impact of Rare and Ultra-rare Coding Variation across the Phenotypic Spectrum. American Journal of Human Genetics, 2018, 102, 1204-1211.	2.6	102
41	Deep coverage whole genome sequences and plasma lipoprotein(a) in individuals of European and African ancestries. Nature Communications, 2018, 9, 2606.	5.8	79
42	Glucose challenge metabolomics implicates medium-chain acylcarnitines in insulin resistance. Scientific Reports, 2018, 8, 8691.	1.6	47
43	Insights into the genetic epidemiology of Crohn's and rare diseases in the Ashkenazi Jewish population. PLoS Genetics, 2018, 14, e1007329.	1.5	66
44	Loss of Cardioprotective Effects at the <i>ADAMTS7</i> Locus as a Result of Gene-Smoking Interactions. Circulation, 2017, 135, 2336-2353.	1.6	51
45	The impact of rare variation on gene expression across tissues. Nature, 2017, 550, 239-243.	13.7	229
46	Heterogeneous contribution of microdeletions in the development of common generalised and focal epilepsies. Journal of Medical Genetics, 2017, 54, 598-606.	1.5	22
47	Nonparametric estimation of the rediscovery rate. Statistics in Medicine, 2016, 35, 3203-3212.	0.8	2
48	Effects of cigarette smoking on cardiovascular-related protein profiles in two community-based cohort studies. Atherosclerosis, 2016, 254, 52-58.	0.4	18
49	Ultra-rare disruptive and damaging mutations influence educational attainment in the general population. Nature Neuroscience, 2016, 19, 1563-1565.	7.1	90
50	Prediction impact curve is a new measure integrating intervention effects in the evaluation of risk models. Journal of Clinical Epidemiology, 2016, 69, 89-95.	2.4	8
51	The metabolic fingerprint of p,p′-DDE and HCB exposure in humans. Environment International, 2016, 88, 60-66.	4.8	61
52	Large-scale non-targeted metabolomic profiling in three human population-based studies. Metabolomics, 2016, 12, 1.	1.4	29
53	Effect of Insulin Resistance on Monounsaturated Fatty Acid Levels: A Multi-cohort Non-targeted Metabolomics and Mendelian Randomization Study. PLoS Genetics, 2016, 12, e1006379.	1.5	20
54	5 year mortality predictors in $498\hat{a}\in 103$ UK Biobank participants: a prospective population-based study. Lancet, The, 2015, 386, 533-540.	6.3	319

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55	Gene × dietary pattern interactions in obesity: analysis of up to 68 317 adults of European ancestry. Human Molecular Genetics, 2015, 24, 4728-4738.	1.4	84
56	Gene-based meta-analysis of genome-wide association studies implicates new loci involved in obesity. Human Molecular Genetics, 2015, 24, 6849-6860.	1.4	55
57	Genome-wide meta-analysis identifies six novel loci associated with habitual coffee consumption. Molecular Psychiatry, 2015, 20, 647-656.	4.1	235
58	Large-scale Metabolomic Profiling Identifies Novel Biomarkers for Incident Coronary Heart Disease. PLoS Genetics, 2014, 10, e1004801.	1.5	225
59	Parent-of-origin-specific allelic associations among 106 genomic loci for age at menarche. Nature, 2014, 514, 92-97.	13.7	548
60	Genetic determinants of mortality. Can findings from genome-wide association studies explain variation in human mortality? Human Genetics, 2013, 132, 553-561.	1.8	29
61	Sex-stratified Genome-wide Association Studies Including 270,000 Individuals Show Sexual Dimorphism in Genetic Loci for Anthropometric Traits. PLoS Genetics, 2013, 9, e1003500.	1.5	371
62	Multilocus Genetic Risk Scores for Coronary Heart Disease Prediction. Arteriosclerosis, Thrombosis, and Vascular Biology, 2013, 33, 2267-2272.	1.1	138
63	Utilizing Twins as Controls for Non-Twin Case-Materials in Genome Wide Association Studies. PLoS ONE, 2013, 8, e83101.	1.1	6
64	Risk Prediction Measures for Case-Cohort and Nested Case-Control Designs: An Application to Cardiovascular Disease. American Journal of Epidemiology, 2012, 175, 715-724.	1.6	75