Kari E North

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6655692/publications.pdf

Version: 2024-02-01

400 papers

39,256 citations

80 h-index 4015 176 g-index

417 all docs

 $\begin{array}{c} 417 \\ \text{docs citations} \end{array}$

417 times ranked 44272 citing authors

#	Article	IF	CITATIONS
1	Genetic studies of body mass index yield new insights for obesity biology. Nature, 2015, 518, 197-206.	27.8	3,823
2	Association analyses of 249,796 individuals reveal 18 new loci associated with body mass index. Nature Genetics, 2010, 42, 937-948.	21.4	2,634
3	Defining the role of common variation in the genomic and biological architecture of adult human height. Nature Genetics, 2014, 46, 1173-1186.	21.4	1,818
4	Hundreds of variants clustered in genomic loci and biological pathways affect human height. Nature, 2010, 467, 832-838.	27.8	1,789
5	New genetic loci link adipose and insulin biology to body fat distribution. Nature, 2015, 518, 187-196.	27.8	1,328
6	Sequencing of 53,831 diverse genomes from the NHLBI TOPMed Program. Nature, 2021, 590, 290-299.	27.8	1,069
7	Loss-of-Function Mutations in <i>APOC3,</i> Triglycerides, and Coronary Disease. New England Journal of Medicine, 2014, 371, 22-31.	27.0	936
8	Meta-analysis identifies 13 new loci associated with waist-hip ratio and reveals sexual dimorphism in the genetic basis of fat distribution. Nature Genetics, 2010, 42, 949-960.	21.4	836
9	Large-scale association analyses identify new loci influencing glycemic traits and provide insight into the underlying biological pathways. Nature Genetics, 2012, 44, 991-1005.	21.4	746
10	Genetic analyses of diverse populations improves discovery for complex traits. Nature, 2019, 570, 514-518.	27.8	679
11	Genome-wide meta-analysis identifies 11 new loci for anthropometric traits and provides insights into genetic architecture. Nature Genetics, 2013, 45, 501-512.	21.4	578
12	Mendelian randomization of blood lipids for coronary heart disease. European Heart Journal, 2015, 36, 539-550.	2.2	567
13	Meta-analyses of genome-wide association studies identify multiple loci associated with pulmonary function. Nature Genetics, 2010, 42, 45-52.	21.4	549
14	Rare and low-frequency coding variants alter human adult height. Nature, 2017, 542, 186-190.	27.8	544
15	FTO genotype is associated with phenotypic variability of body mass index. Nature, 2012, 490, 267-272.	27.8	383
16	Inherited causes of clonal haematopoiesis in 97,691 whole genomes. Nature, 2020, 586, 763-768.	27.8	376
17	Sex-stratified Genome-wide Association Studies Including 270,000 Individuals Show Sexual Dimorphism in Genetic Loci for Anthropometric Traits. PLoS Genetics, 2013, 9, e1003500.	3.5	371
18	Genome-wide association and large-scale follow up identifies 16 new loci influencing lung function. Nature Genetics, 2011, 43, 1082-1090.	21.4	367

#	Article	IF	CITATIONS
19	The power of genetic diversity in genome-wide association studies of lipids. Nature, 2021, 600, 675-679.	27.8	353
20	Association of Adolescent Obesity With Risk of Severe Obesity in Adulthood. JAMA - Journal of the American Medical Association, 2010, 304, 2042-7.	7.4	342
21	The trans-ancestral genomic architecture of glycemic traits. Nature Genetics, 2021, 53, 840-860.	21.4	341
22	The Influence of Age and Sex on Genetic Associations with Adult Body Size and Shape: A Large-Scale Genome-Wide Interaction Study. PLoS Genetics, 2015, 11, e1005378.	3.5	331
23	Genetic loci associated with chronic obstructive pulmonary disease overlap with loci for lung function and pulmonary fibrosis. Nature Genetics, 2017, 49, 426-432.	21.4	306
24	Genetic polymorphisms and the progression of liver fibrosis: A critical appraisal. Hepatology, 2003, 37, 493-503.	7.3	298
25	Epigenome-wide association study (EWAS) of BMI, BMI change and waist circumference in African American adults identifies multiple replicated loci. Human Molecular Genetics, 2015, 24, 4464-4479.	2.9	289
26	Protein-altering variants associated with body mass index implicate pathways that control energy intake and expenditure in obesity. Nature Genetics, 2018, 50, 26-41.	21.4	286
27	Genetic Diversity and Association Studies in US Hispanic/Latino Populations: Applications in the Hispanic Community Health Study/Study of Latinos. American Journal of Human Genetics, 2016, 98, 165-184.	6.2	266
28	Fifteen new risk loci for coronary artery disease highlight arterial-wall-specific mechanisms. Nature Genetics, 2017, 49, 1113-1119.	21.4	260
29	Multi-ancestry genetic study of type 2 diabetes highlights the power of diverse populations for discovery and translation. Nature Genetics, 2022, 54, 560-572.	21.4	250
30	Association of Body Mass Index with DNA Methylation and Gene Expression in Blood Cells and Relations to Cardiometabolic Disease: A Mendelian Randomization Approach. PLoS Medicine, 2017, 14, e1002215.	8.4	246
31	New loci for body fat percentage reveal link between adiposity and cardiometabolic disease risk. Nature Communications, 2016, 7, 10495.	12.8	245
32	Genetic evidence of assortative mating in humans. Nature Human Behaviour, 2017, 1, .	12.0	242
33	Generalization and Dilution of Association Results from European GWAS in Populations of Non-European Ancestry: The PAGE Study. PLoS Biology, 2013, 11, e1001661.	5.6	235
34	Genome-wide meta-analysis identifies six novel loci associated with habitual coffee consumption. Molecular Psychiatry, 2015, 20, 647-656.	7.9	235
35	A meta-analysis identifies new loci associated with body mass index in individuals of African ancestry. Nature Genetics, 2013, 45, 690-696.	21.4	232
36	NRXN3 Is a Novel Locus for Waist Circumference: A Genome-Wide Association Study from the CHARGE Consortium. PLoS Genetics, 2009, 5, e1000539.	3.5	230

#	Article	IF	Citations
37	Population genetic differentiation of height and body mass index across Europe. Nature Genetics, 2015, 47, 1357-1362.	21.4	227
38	A Bivariate Genome-Wide Approach to Metabolic Syndrome. Diabetes, 2011, 60, 1329-1339.	0.6	226
39	Exploring the genetic basis of chronic periodontitis: a genome-wide association study. Human Molecular Genetics, 2013, 22, 2312-2324.	2.9	210
40	Genome-wide meta-analysis of observational studies shows common genetic variants associated with macronutrient intake. American Journal of Clinical Nutrition, 2013, 97, 1395-1402.	4.7	210
41	Use of >100,000 NHLBI Trans-Omics for Precision Medicine (TOPMed) Consortium whole genome sequences improves imputation quality and detection of rare variant associations in admixed African and Hispanic/Latino populations. PLoS Genetics, 2019, 15, e1008500.	3.5	203
42	Causal Effects of Body Mass Index on Cardiometabolic Traits and Events: A Mendelian Randomization Analysis. American Journal of Human Genetics, 2014, 94, 198-208.	6.2	199
43	Whole-Exome Sequencing Identifies Rare and Low-Frequency Coding Variants Associated with LDL Cholesterol. American Journal of Human Genetics, 2014, 94, 233-245.	6.2	193
44	Meta-analysis of genome-wide association studies from the CHARGE consortium identifies common variants associated with carotid intima media thickness and plaque. Nature Genetics, 2011, 43, 940-947.	21.4	191
45	Genetic and Environmental Contributions to Cardiovascular Disease Risk in American Indians: The Strong Heart Family Study. American Journal of Epidemiology, 2003, 157, 303-314.	3.4	186
46	Genome-wide analysis of dental caries and periodontitis combining clinical and self-reported data. Nature Communications, 2019, 10, 2773.	12.8	183
47	Genome-wide association study identifies multiple risk loci for chronic lymphocytic leukemia. Nature Genetics, 2013, 45, 868-876.	21.4	179
48	Directional dominance on stature and cognition inÂdiverse human populations. Nature, 2015, 523, 459-462.	27.8	173
49	Genetic variation in soluble epoxide hydrolase (EPHX2) and risk of coronary heart disease: The Atherosclerosis Risk in Communities (ARIC) study. Human Molecular Genetics, 2006, 15, 1640-1649.	2.9	171
50	Genome-wide meta-analysis of 241,258 adults accounting for smoking behaviour identifies novel loci for obesity traits. Nature Communications, 2017, 8, 14977.	12.8	169
51	Genome-Wide Association Studies Identify <i>CHRNA5/3</i> and <i>HTR4</i> in the Development of Airflow Obstruction. American Journal of Respiratory and Critical Care Medicine, 2012, 186, 622-632.	5.6	164
52	The Next PAGE in Understanding Complex Traits: Design for the Analysis of Population Architecture Using Genetics and Epidemiology (PAGE) Study. American Journal of Epidemiology, 2011, 174, 849-859.	3.4	161
53	Genome-wide physical activity interactions in adiposity ― A meta-analysis of 200,452 adults. PLoS Genetics, 2017, 13, e1006528.	3.5	158
54	Assessing the contribution of rare variants to complex trait heritability from whole-genome sequence data. Nature Genetics, 2022, 54, 263-273.	21.4	156

#	Article	IF	CITATIONS
55	Genome-wide association study identifies multiple susceptibility loci for diffuse large B cell lymphoma. Nature Genetics, 2014, 46, 1233-1238.	21.4	147
56	Genetic Determinants of Lipid Traits in Diverse Populations from the Population Architecture using Genomics and Epidemiology (PAGE) Study. PLoS Genetics, 2011, 7, e1002138.	3.5	146
57	FTO genetic variants, dietary intake and body mass index: insights from 177 330 individuals. Human Molecular Genetics, 2014, 23, 6961-6972.	2.9	143
58	Metabolic Syndrome and the Development of CKD in American Indians: The Strong Heart Study. American Journal of Kidney Diseases, 2008, 51, 21-28.	1.9	140
59	Genome-wide association analysis identifies six new loci associated with forced vital capacity. Nature Genetics, 2014, 46, 669-677.	21.4	131
60	Genome-Wide Joint Meta-Analysis of SNP and SNP-by-Smoking Interaction Identifies Novel Loci for Pulmonary Function. PLoS Genetics, 2012, 8, e1003098.	3.5	130
61	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
62	Meta-analysis of Dense Genecentric Association Studies Reveals Common and Uncommon Variants Associated with Height. American Journal of Human Genetics, 2011, 88, 6-18.	6.2	122
63	Genome-wide meta-analysis associates HLA-DQA1/DRB1 and LPA and lifestyle factors with human longevity. Nature Communications, 2017, 8, 910.	12.8	118
64	Genome-wide analysis of BMI in adolescents and young adults reveals additional insight into the effects of genetic loci over the life course. Human Molecular Genetics, 2013, 22, 3597-3607.	2.9	116
65	Trans-Ethnic Fine-Mapping of Lipid Loci Identifies Population-Specific Signals and Allelic Heterogeneity That Increases the Trait Variance Explained. PLoS Genetics, 2013, 9, e1003379.	3.5	112
66	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
67	Genome-Wide Association of Body Fat Distribution in African Ancestry Populations Suggests New Loci. PLoS Genetics, 2013, 9, e1003681.	3.5	109
68	Genome-wide association study of biologically informed periodontal complex traits offers novel insights into the genetic basis of periodontal disease. Human Molecular Genetics, 2016, 25, 2113-2129.	2.9	108
69	Heritability of Carotid Artery Structure and Function. Arteriosclerosis, Thrombosis, and Vascular Biology, 2002, 22, 1698-1703.	2.4	107
70	Pleiotropic genes for metabolic syndrome and inflammation. Molecular Genetics and Metabolism, 2014, 112, 317-338.	1.1	107
71	Associations of Mitochondrial and Nuclear Mitochondrial Variants and Genes with Seven Metabolic Traits. American Journal of Human Genetics, 2019, 104, 112-138.	6.2	106
72	Prospective Analysis of Mannose-Binding Lectin Genotypes and Coronary Artery Disease in American Indians. Circulation, 2004, 109, 471-475.	1.6	103

#	Article	IF	Citations
73	Association Between Chromosome 9p21 Variants and the Ankle-Brachial Index Identified by a Meta-Analysis of 21 Genome-Wide Association Studies. Circulation: Cardiovascular Genetics, 2012, 5, 100-112.	5.1	98
74	Discovery and fine-mapping of adiposity loci using high density imputation of genome-wide association studies in individuals of African ancestry: African Ancestry Anthropometry Genetics Consortium. PLoS Genetics, 2017, 13, e1006719.	3.5	98
75	Genome-wide Association Study Identifies Five Susceptibility Loci for Follicular Lymphoma outside the HLA Region. American Journal of Human Genetics, 2014, 95, 462-471.	6.2	96
76	Genetic loci associated with heart rate variability and their effects on cardiac disease risk. Nature Communications, 2017, 8, 15805.	12.8	95
77	Genome-wide association study of lung function decline in adults with and without asthma. Journal of Allergy and Clinical Immunology, 2012, 129, 1218-1228.	2.9	94
78	Meta-analysis of genome-wide association studies discovers multiple loci for chronic lymphocytic leukemia. Nature Communications, 2016, 7, 10933.	12.8	94
79	Gut microbiome composition in the Hispanic Community Health Study/Study of Latinos is shaped by geographic relocation, environmental factors, and obesity. Genome Biology, 2019, 20, 219.	8.8	94
80	Identification, Replication, and Fine-Mapping of Loci Associated with Adult Height in Individuals of African Ancestry. PLoS Genetics, 2011, 7, e1002298.	3.5	93
81	A Phenomics-Based Strategy Identifies Loci on APOC1, BRAP, and PLCG1 Associated with Metabolic Syndrome Phenotype Domains. PLoS Genetics, 2011, 7, e1002322.	3.5	92
82	Linkage Analysis of a Composite Factor for the Multiple Metabolic Syndrome. Diabetes, 2003, 52, 2840-2847.	0.6	89
83	Protein-coding variants implicate novel genes related to lipid homeostasis contributing to body-fat distribution. Nature Genetics, 2019, 51, 452-469.	21.4	89
84	Multiethnic meta-analysis identifies ancestry-specific and cross-ancestry loci for pulmonary function. Nature Communications, 2018, 9, 2976.	12.8	85
85	Multiancestry Genome-Wide Association Study of Lipid Levels Incorporating Gene-Alcohol Interactions. American Journal of Epidemiology, 2019, 188, 1033-1054.	3.4	85
86	Gene $\tilde{A}-$ dietary pattern interactions in obesity: analysis of up to 68 317 adults of European ancestry. Human Molecular Genetics, 2015, 24, 4728-4738.	2.9	84
87	Associations of autozygosity with a broad range of human phenotypes. Nature Communications, 2019, 10, 4957.	12.8	84
88	NPHS2 gene, nephrotic syndrome and focal segmental glomerulosclerosis: A HuGE review. Genetics in Medicine, 2006, 8, 63-75.	2.4	83
89	Meta-analysis of up to 622,409 individuals identifies 40 novel smoking behaviour associated genetic loci. Molecular Psychiatry, 2020, 25, 2392-2409.	7.9	83
90	Race- and Sex-Specific Associations of Obesity Measures With Ischemic Stroke Incidence in the Atherosclerosis Risk in Communities (ARIC) Study. Stroke, 2010, 41, 417-425.	2.0	82

#	Article	IF	Citations
91	Longitudinal impact of physical activity on lipid profiles in middle-aged adults: the Atherosclerosis Risk in Communities Study. Journal of Lipid Research, 2009, 50, 1685-1691.	4.2	80
92	Fine Mapping and Identification of BMI Loci in African Americans. American Journal of Human Genetics, 2013, 93, 661-671.	6.2	77
93	A trans-ancestral meta-analysis of genome-wide association studies reveals loci associated with childhood obesity. Human Molecular Genetics, 2019, 28, 3327-3338.	2.9	76
94	Genome-wide association analysis implicates dysregulation of immunity genes in chronic lymphocytic leukaemia. Nature Communications, 2017, 8, 14175.	12.8	75
95	Meta-Analysis Investigating Associations Between Healthy Diet and Fasting Glucose and Insulin Levels and Modification by Loci Associated With Glucose Homeostasis in Data From 15 Cohorts. American Journal of Epidemiology, 2013, 177, 103-115.	3.4	74
96	Association of Functional Polymorphism rs2231142 (Q141K) in the ABCG2 Gene With Serum Uric Acid and Gout in 4 US Populations. American Journal of Epidemiology, 2013, 177, 923-932.	3.4	74
97	A principal component meta-analysis on multiple anthropometric traits identifies novel loci for body shape. Nature Communications, 2016, 7, 13357.	12.8	74
98	Strategies for Enriching Variant Coverage in Candidate Disease Loci on a Multiethnic Genotyping Array. PLoS ONE, 2016, 11, e0167758.	2.5	72
99	Replication of loci influencing ages at menarche and menopause in Hispanic women: the Women's Health Initiative SHARe Study. Human Molecular Genetics, 2012, 21, 1419-1432.	2.9	71
100	Evaluation of the Metabochip Genotyping Array in African Americans and Implications for Fine Mapping of GWAS-Identified Loci: The PAGE Study. PLoS ONE, 2012, 7, e35651.	2.5	71
101	Genome-wide Association Study of Platelet Count Identifies Ancestry-Specific Loci in Hispanic/Latino Americans. American Journal of Human Genetics, 2016, 98, 229-242.	6.2	71
102	Genetic Association Analysis under Complex Survey Sampling: The Hispanic Community Health Study/Study of Latinos. American Journal of Human Genetics, 2014, 95, 675-688.	6.2	70
103	Discovery and Fine Mapping of Serum Protein Loci through Transethnic Meta-analysis. American Journal of Human Genetics, 2012, 91, 744-753.	6.2	69
104	Consumption of meat is associated with higher fasting glucose and insulin concentrations regardless of glucose and insulin genetic risk scores: a meta-analysis of 50,345 Caucasians. American Journal of Clinical Nutrition, 2015, 102, 1266-1278.	4.7	69
105	Exome Chip Meta-analysis Fine Maps Causal Variants and Elucidates the Genetic Architecture of Rare Coding Variants in Smoking and AlcoholÂUse. Biological Psychiatry, 2019, 85, 946-955.	1.3	69
106	Association of Genetic Variants and Incident Coronary Heart Disease in Multiethnic Cohorts. Circulation: Cardiovascular Genetics, 2011, 4, 661-672.	5.1	68
107	Genetic risk factors for BMI and obesity in an ethnically diverse population: Results from the population architecture using genomics and epidemiology (PAGE) study. Obesity, 2013, 21, 835-846.	3.0	68
108	Chocolate consumption is inversely associated with prevalent coronary heart disease: The National Heart, Lung, and Blood Institute Family Heart Study. Clinical Nutrition, 2011, 30, 182-187.	5.0	67

#	Article	IF	CITATIONS
109	Molecular mechanisms underlying variations in lung function: a systems genetics analysis. Lancet Respiratory Medicine, the, 2015, 3, 782-795.	10.7	66
110	Genetic identification of a common collagen disease in Puerto Ricans via identity-by-descent mapping in a health system. ELife, 2017 , 6 , $.$	6.0	65
111	Serum metabolites reflecting gut microbiome alpha diversity predict type 2 diabetes. Gut Microbes, 2020, 11, 1632-1642.	9.8	65
112	A Systematic Mapping Approach of 16q12.2/FTO and BMI in More Than 20,000 African Americans Narrows in on the Underlying Functional Variation: Results from the Population Architecture using Genomics and Epidemiology (PAGE) Study. PLoS Genetics, 2013, 9, e1003171.	3.5	63
113	GWAS for Interleukin- $1\hat{l}^2$ levels in gingival crevicular fluid identifies IL37 variants in periodontal inflammation. Nature Communications, 2018, 9, 3686.	12.8	63
114	DNA Repair Polymorphisms <i>XRCC1 </i> and <i>MGMT</i> and Risk of Adult Gliomas. Neuroepidemiology, 2007, 29, 55-58.	2.3	62
115	Multi-ancestry sleep-by-SNP interaction analysis in 126,926 individuals reveals lipid loci stratified by sleep duration. Nature Communications, 2019, 10, 5121.	12.8	62
116	GWAS for male-pattern baldness identifies 71 susceptibility loci explaining 38% of the risk. Nature Communications, 2017, 8, 1584.	12.8	61
117	Meta-analysis of Gene-Level Associations for Rare Variants Based on Single-Variant Statistics. American Journal of Human Genetics, 2013, 93, 236-248.	6.2	60
118	Association of the FTO Obesity Risk Variant rs8050136 With Percentage of Energy Intake From Fat in Multiple Racial/Ethnic Populations. American Journal of Epidemiology, 2013, 178, 780-790.	3.4	59
119	Angiotensin II type 1 receptor polymorphisms and susceptibility to hypertension: A HuGE review. Genetics in Medicine, 2008, 10, 560-574.	2.4	58
120	A genome-wide association study of marginal zone lymphoma shows association to the HLA region. Nature Communications, 2015, 6, 5751.	12.8	58
121	Methylome-wide association study provides evidence of particulate matter air pollution-associated DNA methylation. Environment International, 2019, 132, 104723.	10.0	58
122	The association of diabetes with breast cancer incidence and mortality in the Long Island Breast Cancer Study Project. Cancer Causes and Control, 2012, 23, 1193-1203.	1.8	57
123	Genotype Imputation of <scp>M</scp> etabochip <scp>SNPs</scp> Using a Studyâ€5pecific Reference Panel of â^1/44,000 Haplotypes in <scp>A</scp> frican <scp>A</scp> mericans From the Women's Health Initiative. Genetic Epidemiology, 2012, 36, 107-117.	1.3	57
124	Gene-based meta-analysis of genome-wide association studies implicates new loci involved in obesity. Human Molecular Genetics, 2015, 24, 6849-6860.	2.9	55
125	Trans-ethnic Meta-analysis and Functional Annotation Illuminates theÂGenetic Architecture of Fasting Glucose and Insulin. American Journal of Human Genetics, 2016, 99, 56-75.	6.2	55
126	GSTM1 and GSTT1 Polymorphisms, Cigarette Smoking, and Risk of Colon Cancer: A Population-based Case-control Study in North Carolina (United States). Cancer Causes and Control, 2006, 17, 385-394.	1.8	54

#	Article	IF	Citations
127	Association of rs780094 in GCKR with Metabolic Traits and Incident Diabetes and Cardiovascular Disease: The ARIC Study. PLoS ONE, 2010, 5, e11690.	2.5	54
128	The Genetics of Obesity and the Metabolic Syndrome. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2010, 10, 86-108.	1.2	54
129	Using genetics to test the causal relationship of total adiposity and periodontitis: Mendelian randomization analyses in the Gene-Lifestyle Interactions and Dental Endpoints (GLIDE) Consortium. International Journal of Epidemiology, 2015, 44, 638-650.	1.9	54
130	Multiethnic Exome-Wide Association Study of Subclinical Atherosclerosis. Circulation: Cardiovascular Genetics, 2016, 9, 511-520.	5.1	54
131	Profiles of internalizing and externalizing symptoms associated with bullying victimization. Journal of Adolescence, 2018, 65, 101-110.	2.4	54
132	PAH–DNA Adducts, Cigarette Smoking, <i>GST</i> Polymorphisms, and Breast Cancer Risk. Environmental Health Perspectives, 2009, 117, 552-558.	6.0	53
133	Genome-wide association study of red blood cell traits in Hispanics/Latinos: The Hispanic Community Health Study/Study of Latinos. PLoS Genetics, 2017, 13, e1006760.	3.5	53
134	Genome-wide association study of age at menarche in African-American women. Human Molecular Genetics, 2013, 22, 3329-3346.	2.9	52
135	Large-Scale Genome-Wide Association Studies and Meta-Analyses of Longitudinal Change in Adult Lung Function. PLoS ONE, 2014, 9, e100776.	2.5	52
136	Genetically predicted longer telomere length is associated with increased risk of B-cell lymphoma subtypes. Human Molecular Genetics, 2016, 25, 1663-1676.	2.9	52
137	Association of Adiposity Genetic Variants With Menarche Timing in 92,105 Women of European Descent. American Journal of Epidemiology, 2013, 178, 451-460.	3.4	51
138	The ERLIN1-CHUK-CWF19L1 gene cluster influences liver fat deposition and hepatic inflammation in the NHLBI Family Heart Study. Atherosclerosis, 2013, 228, 175-180.	0.8	50
139	Melanocortin 4 Receptor Pathway Dysfunction in Obesity: Patient Stratification Aimed at MC4R Agonist Treatment. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 2601-2612.	3.6	50
140	Consistent Directions of Effect for Established Type 2 Diabetes Risk Variants Across Populations. Diabetes, 2012, 61, 1642-1647.	0.6	49
141	Use of Nonsteroidal Antiinflammatory Drugs and Risk of Colon Cancer in a Population-based, Case-Control Study of African Americans and Whites. American Journal of Epidemiology, 2005, 162, 548-558.	3.4	48
142	The association of genetic variants of type 2 diabetes with kidney function. Kidney International, 2012, 82, 220-225.	5.2	48
143	Novel Loci Associated With PR Interval in a Genome-Wide Association Study of 10 African American Cohorts. Circulation: Cardiovascular Genetics, 2012, 5, 639-646.	5.1	48
144	A Genome-wide Association Study Discovers 46 Loci of the Human Metabolome in the Hispanic Community Health Study/Study of Latinos. American Journal of Human Genetics, 2020, 107, 849-863.	6.2	48

#	Article	IF	CITATIONS
145	Higher Magnesium Intake Is Associated with Lower Fasting Glucose and Insulin, with No Evidence of Interaction with Select Genetic Loci, in a Meta-Analysis of 15 CHARGE Consortium Studies. Journal of Nutrition, 2013, 143, 345-353.	2.9	47
146	Associations of Obesity Measures with Subtypes of Ischemic Stroke in the ARIC Study. Journal of Epidemiology, 2010, 20, 347-354.	2.4	46
147	Association of dietary folate and vitamin B-12 intake with genome-wide DNA methylation in blood: a large-scale epigenome-wide association analysis in 5841 individuals. American Journal of Clinical Nutrition, 2019, 110, 437-450.	4.7	46
148	Serum Adiponectin in Relation to Body Mass Index and Other Correlates in Black and White Women. Annals of Epidemiology, 2011, 21, 86-94.	1.9	45
149	Genome-Wide Detection of Allele Specific Copy Number Variation Associated with Insulin Resistance in African Americans from the HyperGEN Study. PLoS ONE, 2011, 6, e24052.	2.5	45
150	A whole genome association study of mother-to-child transmission of HIV in Malawi. Genome Medicine, 2010, 2, 17.	8.2	44
151	Mapping adipose and muscle tissue expression quantitative trait loci in African Americans to identify genes for type 2 diabetes and obesity. Human Genetics, 2016, 135, 869-880.	3.8	44
152	Rare Exome Sequence Variants in <i>CLCN6</i> Reduce Blood Pressure Levels and Hypertension Risk. Circulation: Cardiovascular Genetics, 2016, 9, 64-70.	5.1	44
153	Genome-wide meta-analysis of macronutrient intake of 91,114 European ancestry participants from the cohorts for heart and aging research in genomic epidemiology consortium. Molecular Psychiatry, 2019, 24, 1920-1932.	7.9	44
154	TCF7L2 Variants Associate with CKD Progression and Renal Function in Population-Based Cohorts. Journal of the American Society of Nephrology: JASN, 2008, 19, 1989-1999.	6.1	43
155	Analysis of Metabolic Syndrome Components in >15 000 African Americans Identifies Pleiotropic Variants. Circulation: Cardiovascular Genetics, 2014, 7, 505-513.	5.1	43
156	COX-2 Polymorphism, Use of Nonsteroidal Anti-Inflammatory Drugs, and Risk of Colon Cancer in African Americans (United States). Cancer Causes and Control, 2006, 17, 257-266.	1.8	41
157	Gene-environment interactions and obesity traits among postmenopausal African-American and Hispanic women in the Women's Health Initiative SHARe Study. Human Genetics, 2013, 132, 323-336.	3.8	41
158	Multi-ethnic fine-mapping of 14 central adiposity loci. Human Molecular Genetics, 2014, 23, 4738-4744.	2.9	41
159	A powerful statistical framework for generalization testing in GWAS, with application to the HCHS/SOL. Genetic Epidemiology, 2017, 41, 251-258.	1.3	41
160	Association of Birth Weight With Type 2 Diabetes and Glycemic Traits. JAMA Network Open, 2019, 2, e1910915.	5.9	41
161	Chocolate consumption is inversely associated with calcified atherosclerotic plaque in the coronary arteries: The NHLBI Family Heart Studyâ [†] t. Clinical Nutrition, 2011, 30, 38-43.	5.0	39
162	Association of dietary omega-3 fatty acids with prevalence of metabolic syndrome: The National Heart, Lung, and Blood Institute Family Heart Study. Clinical Nutrition, 2013, 32, 966-969.	5.0	39

#	Article	IF	CITATIONS
163	Impact of Ancestry and Common Genetic Variants on QT Interval in African Americans. Circulation: Cardiovascular Genetics, 2012, 5, 647-655.	5.1	38
164	Genome-wide association of white blood cell counts in Hispanic/Latino Americans: the Hispanic Community Health Study/Study of Latinos. Human Molecular Genetics, 2017, 26, 1193-1204.	2.9	38
165	Fine-mapping, novel loci identification, and SNP association transferability in a genome-wide association study of QRS duration in African Americans. Human Molecular Genetics, 2016, 25, 4350-4368.	2.9	37
166	Functional COMT Val158Met Polymorphism, Risk of Acute Coronary Events and Serum Homocysteine: The Kuopio Ischaemic Heart Disease Risk Factor Study. PLoS ONE, 2007, 2, e181.	2.5	36
167	<i>IGF-I</i> and <i>IGFBP-3</i> Polymorphisms in Relation to Circulating Levels among African American and Caucasian Women. Cancer Epidemiology Biomarkers and Prevention, 2009, 18, 954-966.	2.5	36
168	Imputation of coding variants in African Americans: better performance using data from the exome sequencing project. Bioinformatics, 2013, 29, 2744-2749.	4.1	36
169	Mendelian randomization supports bidirectional causality between telomere length and clonal hematopoiesis of indeterminate potential. Science Advances, 2022, 8, eabl6579.	10.3	36
170	Genetic Determination of Acute Phase Reactant Levels: The Strong Heart Study. Human Heredity, 2004, 58, 112-116.	0.8	35
171	Pleiotropic Associations of Risk Variants Identified for Other Cancers With Lung Cancer Risk: The PAGE and TRICL Consortia. Journal of the National Cancer Institute, 2014, 106, dju061.	6.3	35
172	Genome scan of glomerular filtration rate and albuminuria: the HyperGEN study. Nephrology Dialysis Transplantation, 2007, 22, 763-771.	0.7	34
173	Lack of associations of ten candidate coronary heart disease risk genetic variants and subclinical atherosclerosis in four U.S. populations: The Population Architecture using Genomics and Epidemiology (PAGE) study. Atherosclerosis, 2013, 228, 390-399.	0.8	33
174	An epigenome-wide study of obesity in African American youth and young adults: novel findings, replication in neutrophils, and relationship with gene expression. Clinical Epigenetics, 2018, 10, 3.	4.1	33
175	The Future of Genomic Studies Must Be Globally Representative: Perspectives from PAGE. Annual Review of Genomics and Human Genetics, 2019, 20, 181-200.	6.2	33
176	Distinct Loci in the <i>CHRNA5</i> / <i>CHRNA3</i> / <i>CHRNB4</i> Gene Cluster Are Associated With Onset of Regular Smoking. Genetic Epidemiology, 2013, 37, 846-859.	1.3	32
177	Association of Cardiometabolic Genes with Arsenic Metabolism Biomarkers in American Indian Communities: The Strong Heart Family Study (SHFS). Environmental Health Perspectives, 2017, 125, 15-22.	6.0	32
178	Sugar-sweetened beverage intake associations with fasting glucose and insulin concentrations are not modified by selected genetic variants in a ChREBP-FGF21 pathway: a meta-analysis. Diabetologia, 2018, 61, 317-330.	6.3	32
179	Heritability and Preliminary Genome-Wide Linkage Analysis of Arsenic Metabolites in Urine. Environmental Health Perspectives, 2013, 121, 345-351.	6.0	31
180	Multiancestral Analysis of Inflammation-Related Genetic Variants and C-Reactive Protein in the Population Architecture Using Genomics and Epidemiology Study. Circulation: Cardiovascular Genetics, 2014, 7, 178-188.	5.1	31

#	Article	IF	Citations
181	Trans-ethnic fine-mapping of genetic loci for body mass index in the diverse ancestral populations of the Population Architecture using Genomics and Epidemiology (PAGE) Study reveals evidence for multiple signals at established loci. Human Genetics, 2017, 136, 771-800.	3.8	31
182	Metabolomics Identifies Novel Blood Biomarkers of Pulmonary Function and COPD in the General Population. Metabolites, 2019, 9, 61.	2.9	30
183	Pleiotropic QTL on chromosome 19q13 for triglycerides and adiposity: The HERITAGE family study. Atherosclerosis, 2006, 185, 426-432.	0.8	29
184	Admixture mapping of quantitative trait loci for blood lipids in African-Americans. Human Molecular Genetics, 2009, 18, 2091-2098.	2.9	29
185	The Influence of Obesity-Related Single Nucleotide Polymorphisms on BMI Across the Life Course: The PAGE Study. Diabetes, 2013, 62, 1763-1767.	0.6	29
186	Fine-mapping of lipid regions in global populations discovers ethnic-specific signals and refines previously identified lipid loci. Human Molecular Genetics, 2016, 25, 5500-5512.	2.9	29
187	Genetics of Chronic Kidney Disease Stages Across Ancestries: The PAGE Study. Frontiers in Genetics, 2019, 10, 494.	2.3	29
188	A System for Phenotype Harmonization in the National Heart, Lung, and Blood Institute Trans-Omics for Precision Medicine (TOPMed) Program. American Journal of Epidemiology, 2021, 190, 1977-1992.	3.4	29
189	Genetic determinants of telomere length from 109,122 ancestrally diverse whole-genome sequences in TOPMed. Cell Genomics, 2022, 2, 100084.	6.5	29
190	Quantitative effects of common genetic variations in the $3\hat{a}\in^2$ UTR of the human LDL-receptor gene and their associations with plasma lipid levels in the Atherosclerosis Risk in Communities study. Human Genetics, 2007, 121, 421-431.	3.8	28
191	Transcription Factor 7-Like 2 (TCF7L2) Polymorphism and Context-Specific Risk of Type 2 Diabetes in African American and Caucasian Adults: The Atherosclerosis Risk in Communities Study. Diabetes, 2009, 58, 285-289.	0.6	28
192	Gene-centric meta-analyses for central adiposity traits in up to 57 412 individuals of European descent confirm known loci and reveal several novel associations. Human Molecular Genetics, 2014, 23, 2498-2510.	2.9	28
193	Integrative pathway genomics of lung function and airflow obstruction. Human Molecular Genetics, 2015, 24, 6836-6848.	2.9	28
194	Objectively Measured Physical Activity, Sedentary Behavior, and Genetic Predisposition to Obesity in U.S. Hispanics/Latinos: Results From the Hispanic Community Health Study/Study of Latinos (HCHS/SOL). Diabetes, 2017, 66, 3001-3012.	0.6	28
195	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
196	Genetic Influences on Aortic Root Size in American Indians. Arteriosclerosis, Thrombosis, and Vascular Biology, 2002, 22, 1008-1011.	2.4	27
197	Genetics of Obesity in Diverse Populations. Current Diabetes Reports, 2018, 18, 145.	4.2	27
198	Analyses of biomarker traits in diverse UK biobank participants identify associations missed by European-centric analysis strategies. Journal of Human Genetics, 2022, 67, 87-93.	2.3	27

#	Article	IF	CITATIONS
199	Prospective Associations of Coronary Heart Disease Loci in African Americans Using the MetaboChip: The PAGE Study. PLoS ONE, 2014, 9, e113203.	2.5	27
200	Evidence for a Gene on Chromosome 13 Influencing Postural Systolic Blood Pressure Change and Body Mass Index. Hypertension, 2004, 43, 780-784.	2.7	26
201	Joint Effects of Alcohol Consumption and Polymorphisms in Alcohol and Oxidative Stress Metabolism Genes on Risk of Head and Neck Cancer. Cancer Epidemiology Biomarkers and Prevention, 2011, 20, 2438-2449.	2.5	26
202	Genome-wide association study of 1,5-anhydroglucitol identifies novel genetic loci linked to glucose metabolism. Scientific Reports, 2017, 7, 2812.	3.3	26
203	Genetic Studies of Leptin Concentrations Implicate Leptin in the Regulation of Early Adiposity. Diabetes, 2020, 69, 2806-2818.	0.6	26
204	Linkage analysis of LDL cholesterol in American Indian populations: the Strong Heart Family Study. Journal of Lipid Research, 2006, 47, 59-66.	4.2	25
205	Peroxisome proliferator-activated receptor-alpha (PPARA) genetic polymorphisms and breast cancer risk: a Long Island ancillary study. Carcinogenesis, 2008, 29, 1944-1949.	2.8	25
206	Hip Circumference and Incident Metabolic Risk Factors in Chinese Men and Women: The People's Republic of China Study. Metabolic Syndrome and Related Disorders, 2011, 9, 55-62.	1.3	25
207	Genetic determinants of the ankle-brachial index: A meta-analysis of a cardiovascular candidate gene 50K SNP panel in the candidate gene association resource (CARe) consortium. Atherosclerosis, 2012, 222, 138-147.	0.8	25
208	Sex-influenced association of nonalcoholic fatty liver disease with coronary heart disease. Atherosclerosis, 2013, 227, 420-424.	0.8	25
209	The genetic underpinnings of variation in ages at menarche and natural menopause among women from the multi-ethnic Population Architecture using Genomics and Epidemiology (PAGE) Study: A trans-ethnic meta-analysis. PLoS ONE, 2018, 13, e0200486.	2.5	25
210	DRD4 gene variant associated with body mass: The National Longitudinal Study of Adolescent Health. Human Mutation, 2006, 27, 236-241.	2.5	24
211	Linkage Analysis of Albuminuria. Journal of the American Society of Nephrology: JASN, 2009, 20, 1597-1606.	6.1	24
212	The association of the MYH9 gene and kidney outcomes in American Indians: the Strong Heart Family Study. Human Genetics, 2010, 127, 295-301.	3.8	24
213	Genetic variants associated with fasting glucose and insulin concentrations in an ethnically diverse population: results from the Population Architecture using Genomics and Epidemiology (PAGE) study. BMC Medical Genetics, 2013, 14, 98.	2.1	24
214	Polymorphisms in DNA repair genes, recreational physical activity and breast cancer risk. International Journal of Cancer, 2014, 134, 654-663.	5.1	24
215	Both Rare and Common Variants in PCSK9 Influence Plasma Low-Density Lipoprotein Cholesterol Level in American Indians. Journal of Clinical Endocrinology and Metabolism, 2015, 100, E345-E349.	3.6	24
216	Variant Discovery and Fine Mapping of Genetic Loci Associated with Blood Pressure Traits in Hispanics and African Americans. PLoS ONE, 2016, 11, e0164132.	2.5	24

#	Article	IF	CITATIONS
217	Exploring the genomic basis of early childhood caries: a pilot study. International Journal of Paediatric Dentistry, 2018, 28, 217-225.	1.8	24
218	No Interactions Between Previously Associated 2-Hour Glucose Gene Variants and Physical Activity or BMI on 2-Hour Glucose Levels. Diabetes, 2012, 61, 1291-1296.	0.6	23
219	Replication of the effect of SLC2A9 genetic variation on serum uric acid levels in American Indians. European Journal of Human Genetics, 2014, 22, 938-943.	2.8	23
220	Association of Cancer Susceptibility Variants with Risk of Multiple Primary Cancers: The Population Architecture using Genomics and Epidemiology Study. Cancer Epidemiology Biomarkers and Prevention, 2014, 23, 2568-2578.	2.5	23
221	Genetic discovery and risk characterization in type 2 diabetes across diverse populations. Human Genetics and Genomics Advances, 2021, 2, 100029.	1.7	23
222	Linkage analysis of glomerular filtration rate in American Indians. Kidney International, 2008, 74, 1185-1191.	5.2	22
223	Genetic Epidemiology of BMI and Body Mass Change From Adolescence to Young Adulthood. Obesity, 2010, 18, 1474-1476.	3.0	22
224	Evidence of Heterogeneity by Race/Ethnicity in Genetic Determinants of QT Interval. Epidemiology, 2014, 25, 790-798.	2.7	22
225	Targeting epigenetic mechanisms in periodontal diseases. Periodontology 2000, 2018, 78, 174-184.	13.4	22
226	A high throughput, functional screen of human Body Mass Index GWAS loci using tissue-specific RNAi Drosophila melanogaster crosses. PLoS Genetics, 2018, 14, e1007222.	3.5	22
227	CCR5 Haplotypes and Mother-to-Child HIV Transmission in Malawi. PLoS ONE, 2007, 2, e838.	2.5	22
228	Evidence for Genetic Factors Underlying the Insulin Resistance Syndrome in American Indians. Obesity, 2003, 11, 1444-1448.	4.0	21
229	Evidence for a gene influencing fasting LDL cholesterol and triglyceride levels on chromosome 21q. Atherosclerosis, 2005, 179, 119-125.	0.8	21
230	Genotype-by-sex interaction in the aetiology of type 2 diabetes mellitus: support for sex-specific quantitative trait loci in Hypertension Genetic Epidemiology Network participants. Diabetologia, 2006, 49, 2329-2336.	6.3	21
231	Sex and race differences in the prevalence of fatty liver disease as measured by computed tomography liver attenuation in European American and African American participants of the NHLBI family heart study. European Journal of Gastroenterology and Hepatology, 2012, 24, 9-16.	1.6	21
232	Exome-Derived Adiponectin-Associated Variants Implicate Obesity and Lipid Biology. American Journal of Human Genetics, 2019, 105, 15-28.	6.2	21
233	A Quantitative Trait Loci-Specific Gene-by-Sex Interaction on Systolic Blood Pressure Among American Indians. Hypertension, 2006, 48, 266-270.	2.7	20
234	Interaction of Folate Intake and the Paraoxonase Q192R Polymorphism with Risk of Incident Coronary Heart Disease and Ischemic Stroke: TheÂAtherosclerosis Risk in Communities Study. Annals of Epidemiology, 2011, 21, 815-823.	1.9	20

#	Article	IF	Citations
235	A genome-wide association study meta-analysis of clinical fracture in 10,012 African American women. Bone Reports, 2016, 5, 233-242.	0.4	20
236	Transethnic insight into the genetics of glycaemic traits: fine-mapping results from the Population Architecture using Genomics and Epidemiology (PAGE) consortium. Diabetologia, 2017, 60, 2384-2398.	6.3	20
237	Dairy Intake and Body Composition and Cardiometabolic Traits among Adults: Mendelian Randomization Analysis of 182041 Individuals from 18 Studies. Clinical Chemistry, 2019, 65, 751-760.	3.2	20
238	Cohort Profile: ZOE 2.0—A Community-Based Genetic Epidemiologic Study of Early Childhood Oral Health. International Journal of Environmental Research and Public Health, 2020, 17, 8056.	2.6	20
239	HDL cholesterol in females in the Framingham Heart Study is linked to a region of chromosome 2q. BMC Genetics, 2003, 4, S98.	2.7	19
240	Linkage Analysis of Diabetes Status Among Hypertensive Families: The Hypertension Genetic Epidemiology Network Study. Diabetes, 2004, 53, 3307-3312.	0.6	19
241	Genotype-by-Sex Interaction on Fasting Insulin Concentration: The HyperGEN Study. Diabetes, 2007, 56, 137-142.	0.6	19
242	A gene-centric association scan for Coagulation Factor VII levels in European and African Americans: the Candidate Gene Association Resource (CARe) Consortium. Human Molecular Genetics, 2011, 20, 3525-3534.	2.9	19
243	Fine mapping of QT interval regions in global populations refines previously identified QT interval loci and identifies signals unique to African and Hispanic descent populations. Heart Rhythm, 2017, 14, 572-580.	0.7	19
244	Importance of Genetic Studies of Cardiometabolic Disease in Diverse Populations. Circulation Research, 2020, 126, 1816-1840.	4. 5	19
245	Meta-analysis of exome array data identifies six novel genetic loci for lung function. Wellcome Open Research, 2018, 3, 4.	1.8	19
246	Val153Met Polymorphism of Catechol-O-Methyltransferase and Prevalence of Uterine Leiomyomata. Reproductive Sciences, 2007, 14, 117-120.	2.5	18
247	Polymorphisms in oxidative stress genes, physical activity, and breast cancer risk. Cancer Causes and Control, 2012, 23, 1949-1958.	1.8	18
248	Sequencing of 2 Subclinical Atherosclerosis Candidate Regions in 3669 Individuals. Circulation: Cardiovascular Genetics, 2014, 7, 359-364.	5.1	18
249	Genetic architecture of lipid traits in the Hispanic community health study/study of Latinos. Lipids in Health and Disease, 2017, 16, 200.	3.0	18
250	Discovery and fine-mapping of height loci via high-density imputation of GWASs in individuals of African ancestry. American Journal of Human Genetics, 2021, 108, 564-582.	6.2	18
251	Associations of body mass index with incident hypertension in American white, American black and Chinese Asian adults in early and middle adulthood: the Coronary Artery Risk Development in Young Adults (CARDIA) study, the Atherosclerosis Risk in Communities (ARIC) study and the People's Republic of China (PRC) study. Asia Pacific Journal of Clinical Nutrition, 2013, 22, 626-34.	0.4	18
252	Racial Differences in the Association of Coronary Calcified Plaque With Left Ventricular Hypertrophy: The National Heart, Lung, and Blood Institute Family Heart Study and Hypertension Genetic Epidemiology Network. American Journal of Cardiology, 2006, 97, 1441-1448.	1.6	17

#	Article	IF	Citations
253	Fasting Insulin and Obesity-Related Phenotypes Are Linked to Chromosome 2p: The Strong Heart Family Study. Diabetes, 2006, 55, 1874-1878.	0.6	17
254	Relation of Albuminuria to Left Ventricular Mass (from the HyperGEN Study). American Journal of Cardiology, 2008, 101, 212-216.	1.6	17
255	Genome-wide Linkage Analysis of Pulse Pressure in American Indians: The Strong Heart Study. American Journal of Hypertension, 2008, 21, 194-199.	2.0	17
256	Regulation of CCR5 Expression in Human Placenta: Insights from a Study of Mother-to-Child Transmission of HIV in Malawi. PLoS ONE, 2010, 5, e9212.	2.5	17
257	Ultraconserved Elements in the Human Genome: Association and Transmission Analyses of Highly Constrained Single-Nucleotide Polymorphisms. Genetics, 2012, 192, 253-266.	2.9	17
258	Genetic Determinants of Pelvic Organ Prolapse among African American and Hispanic Women in the Women's Health Initiative. PLoS ONE, 2015, 10, e0141647.	2.5	17
259	Gene-educational attainment interactions in a multi-ancestry genome-wide meta-analysis identify novel blood pressure loci. Molecular Psychiatry, 2020, 26, 2111-2125.	7.9	17
260	Minority-centric meta-analyses of blood lipid levels identify novel loci in the Population Architecture using Genomics and Epidemiology (PAGE) study. PLoS Genetics, 2020, 16, e1008684.	3.5	17
261	Host genetic effects in pneumonia. American Journal of Human Genetics, 2021, 108, 194-201.	6.2	17
262	Chromosome Xq23 is associated with lower atherogenic lipid concentrations and favorable cardiometabolic indices. Nature Communications, 2021, 12, 2182.	12.8	17
263	A Population Association Study of Angiotensinogen Polymorphisms and Haplotypes With Left Ventricular Phenotypes. Hypertension, 2005, 46, 1294-1299.	2.7	16
264	Metaâ€Analysis of Genomeâ€Wide Association Studies with Correlated Individuals: Application to the Hispanic Community Health Study/Study of Latinos (HCHS/SOL). Genetic Epidemiology, 2016, 40, 492-501.	1.3	16
265	Discovery, fine-mapping, and conditional analyses of genetic variants associated with C-reactive protein in multiethnic populations using the Metabochip in the Population Architecture using Genomics and Epidemiology (PAGE) study. Human Molecular Genetics, 2018, 27, 2940-2953.	2.9	16
266	Associations of maternal arsenic exposure with adult fasting glucose and insulin resistance in the Strong Heart Study and Strong Heart Family Study. Environment International, 2020, 137, 105531.	10.0	16
267	Diabetes-specific genetic effects on obesity traits in American Indian populations: the Strong Heart Family Study. BMC Medical Genetics, 2008, 9, 90.	2.1	15
268	Lupus-related single nucleotide polymorphisms and risk of diffuse large B-cell lymphoma. Lupus Science and Medicine, 2017, 4, e000187.	2.7	15
269	Two high-risk susceptibility loci at 6p25.3 and 14q32.13 for Waldenström macroglobulinemia. Nature Communications, 2018, 9, 4182.	12.8	15
270	A phenome-wide association study (PheWAS) in the Population Architecture using Genomics and Epidemiology (PAGE) study reveals potential pleiotropy in African Americans. PLoS ONE, 2019, 14, e0226771.	2.5	15

#	Article	IF	Citations
271	Epigenetic reprogramming in periodontal disease: Dynamic crosstalk with potential impact in oncogenesis. Periodontology 2000, 2020, 82, 157-172.	13.4	15
272	Multi-ethnic GWAS and fine-mapping of glycaemic traits identify novel loci in the PAGE Study. Diabetologia, 2022, 65, 477-489.	6.3	15
273	Pulmonary Function and Blood DNA Methylation: A Multiancestry Epigenome-Wide Association Meta-analysis. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 321-336.	5.6	15
274	Analysis of metabolic syndrome phenotypes in Framingham Heart Study families from Genetic Analysis Workshop 13. Genetic Epidemiology, 2003, 25, S78-S89.	1.3	14
275	The Importance of Geneâ€"Environment Interaction. Sociological Methods and Research, 2008, 37, 164-200.	6.8	14
276	Genome-wide linkage scan for plasma high density lipoprotein cholesterol, apolipoprotein A-1 and triglyceride variation among American Indian populations: the Strong Heart Family Study. Journal of Medical Genetics, 2009, 46, 472-479.	3.2	14
277	Transcription factor 7â€like 2 (<i>TCF7L2</i>) polymorphism and contextâ€specific risk of impaired fasting glucose in African American and Caucasian adults: the atherosclerosis risk in communities (ARIC) study. Diabetes/Metabolism Research and Reviews, 2010, 26, 371-377.	4.0	14
278	The 9p21 genetic variant is additive to carotid intima media thickness and plaque in improving coronary heart disease risk prediction in white participants of the Atherosclerosis Risk in Communities (ARIC) Study. Atherosclerosis, 2012, 222, 135-137.	0.8	14
279	Effects of polymorphisms in alcohol metabolism and oxidative stress genes on survival from head and neck cancer. Cancer Epidemiology, 2013, 37, 479-491.	1.9	14
280	Whole-exome imputation of sequence variants identified two novel alleles associated with adult body height in African Americans. Human Molecular Genetics, 2014, 23, 6607-6615.	2.9	14
281	Linkage Analysis of Urine Arsenic Species Patterns in the Strong Heart Family Study. Toxicological Sciences, 2015, 148, 89-100.	3.1	14
282	The Novel <i>ASIC2</i> Locus Is Associated with Severe Gingival Inflammation. JDR Clinical and Translational Research, 2016, 1, 163-170.	1.9	14
283	Protocols, Methods, and Tools for Genome-Wide Association Studies (GWAS) of Dental Traits. Methods in Molecular Biology, 2019, 1922, 493-509.	0.9	14
284	Omega-3 Fatty Acids and Genome-Wide Interaction Analyses Reveal <i>DPP10–</i> Pulmonary Function Association. American Journal of Respiratory and Critical Care Medicine, 2019, 199, 631-642.	5.6	14
285	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
286	Genome-wide linkage analysis replicates susceptibility locus for fasting plasma triglycerides: NHLBI Family Heart Study. Human Genetics, 2004, 115, 468-474.	3.8	13
287	Linkage Analysis of Factors Underlying Insulin Resistance: Strong Heart Family Study. Obesity, 2005, 13, 1877-1884.	4.0	13
288	Hysterectomy prevalence and cardiovascular disease risk factors in American Indian women. Maturitas, 2005, 52, 328-336.	2.4	13

#	Article	IF	Citations
289	Evaluation of Serum Immunoglobulins among Individuals Living Near Six SuperfundSites. Environmental Health Perspectives, 2006, 114, 1065-1071.	6.0	13
290	QTL-specific genotype-by-smoking interaction and burden of calcified coronary atherosclerosis: The NHLBI Family Heart Study. Atherosclerosis, 2007, 193, 11-19.	0.8	13
291	Comparison of genome-wide variation between Malawians and African ancestry HapMap populations. Journal of Human Genetics, 2010, 55, 366-374.	2.3	13
292	Fine-Mapping and Initial Characterization of QT Interval Loci in African Americans. PLoS Genetics, 2012, 8, e1002870.	3.5	13
293	Genetic polymorphisms of diabetesâ€related genes, their interaction with diabetes status, and breast cancer incidence and mortality: The Long Island Breast Cancer Study Project. Molecular Carcinogenesis, 2019, 58, 436-446.	2.7	13
294	Genetic variants affecting bone mineral density and bone mineral content at multiple skeletal sites in Hispanic children. Bone, 2020, 132, 115175.	2.9	13
295	Association Between Midlife Obesity and Kidney Function Trajectories: The Atherosclerosis Risk in Communities (ARIC) Study. American Journal of Kidney Diseases, 2021, 77, 376-385.	1.9	13
296	Multi-ancestry genome-wide gene–sleep interactions identify novel loci for blood pressure. Molecular Psychiatry, 2021, 26, 6293-6304.	7.9	13
297	Meta-analysis of genome-wide linkage studies for quantitative lipid traits in African Americans. Human Molecular Genetics, 2005, 14, 3955-3962.	2.9	12
298	Linkage study of fibrinogen levels: the Strong Heart Family Study. BMC Medical Genetics, 2008, 9, 77.	2.1	12
299	A multi-population phenome-wide association study of genetically-predicted height in the Million Veteran Program. PLoS Genetics, 2022, 18, e1010193.	3.5	12
300	Insights From a Large-Scale Whole-Genome Sequencing Study of Systolic Blood Pressure, Diastolic Blood Pressure, and Hypertension. Hypertension, 2022, 79, 1656-1667.	2.7	12
301	Genetic meta-analysis of 15,901 African Americans identifies variation in EXOC3L1 is associated with HDL concentration. Journal of Lipid Research, 2015, 56, 1781-1786.	4.2	11
302	Identifying gene–gene interactions that are highly associated with four quantitative lipid traits across multiple cohorts. Human Genetics, 2017, 136, 165-178.	3.8	11
303	Transcriptome-Wide Association Study of Blood Cell Traits in African Ancestry and Hispanic/Latino Populations. Genes, 2021, 12, 1049.	2.4	11
304	Interaction of smoking and obesity susceptibility loci on adolescent BMI: The National Longitudinal Study of Adolescent to Adult Health. BMC Genetics, 2015, 16, 131.	2.7	10
305	Analysis of Sequence Data Under Multivariate Trait-Dependent Sampling. Journal of the American Statistical Association, 2015, 110, 560-572.	3.1	10
306	Testing the role of predicted gene knockouts in human anthropometric trait variation. Human Molecular Genetics, 2016, 25, 2082-2092.	2.9	10

#	Article	IF	CITATIONS
307	Obesity Duration, Severity, and Distribution Trajectories and Cardiovascular Disease Risk in the Atherosclerosis Risk in Communities Study. Journal of the American Heart Association, 2021, 10, e019946.	3.7	10
308	Evidence for distinct genetic effects on obesity and lipid-related CVD risk factors in diabetic compared to nondiabetic American Indians: the Strong Heart Family Study. Diabetes/Metabolism Research and Reviews, 2003, 19, 140-147.	4.0	9
309	DNA methylation of imprinted gene control regions in the regression of lowâ€grade cervical lesions. International Journal of Cancer, 2018, 143, 552-560.	5.1	9
310	Genomeâ€Wide Interactions with Dairy Intake for Body Mass Index in Adults of European Descent. Molecular Nutrition and Food Research, 2018, 62, 1700347.	3.3	9
311	Meta-analysis across Cohorts for Heart and Aging Research in Genomic Epidemiology (CHARGE) consortium provides evidence for an association of serum vitamin D with pulmonary function. British Journal of Nutrition, 2018, 120, 1159-1170.	2.3	9
312	Associations between SLC16A11 variants and diabetes in the Hispanic Community Health Study/Study of Latinos (HCHS/SOL). Scientific Reports, 2019, 9, 843.	3.3	9
313	Mendelian randomization analysis does not support causal associations of birth weight with hypertension risk and blood pressure in adulthood. European Journal of Epidemiology, 2020, 35, 685-697.	5.7	9
314	Social- and Behavioral-Specific Genetic Effects on Blood Pressure Traits. Circulation: Cardiovascular Genetics, 2009, 2, 396-401.	5.1	8
315	<i>ADAM19</i> and <i>HTR4</i> Variants and Pulmonary Function. Circulation: Cardiovascular Genetics, 2014, 7, 350-358.	5.1	8
316	Sequence Variation in $\langle i \rangle$ TMEM18 $\langle i \rangle$ in Association With Body Mass Index. Circulation: Cardiovascular Genetics, 2014, 7, 344-349.	5.1	8
317	Genome-wide association of trajectories of systolic blood pressure change. BMC Proceedings, 2016, 10, 321-327.	1.6	8
318	Validation in a Brazilian population of gene markers of periodontitis previously investigated by GWAS and bioinformatic studies. Journal of Periodontology, 2021, 92, 689-703.	3.4	8
319	Genome-wide association study of body fat distribution traits in Hispanics/Latinos from the HCHS/SOL. Human Molecular Genetics, 2021, 30, 2190-2204.	2.9	8
320	Sugar-Sweetened Beverage Consumption May Modify Associations Between Genetic Variants in the CHREBP (Carbohydrate Responsive Element Binding Protein) Locus and HDL-C (High-Density Lipoprotein) Tj ETQq e003288.	0,0,0 rgBT	/Overlock 1
321	Demographic and sociocultural risk factors for adulthood weight gain in Hispanic/Latinos: results from the Hispanic Community Health Study / Study of Latinos (HCHS/SOL). BMC Public Health, 2021, 21, 2064.	2.9	8
322	Smoking and Selected DNA Repair Gene Polymorphisms in Controls: Systematic Review and Meta-Analysis. Cancer Epidemiology Biomarkers and Prevention, 2010, 19, 3055-3086.	2.5	7
323	No evidence of interaction between known lipid-associated genetic variants and smoking in the multi-ethnic PAGE population. Human Genetics, 2013, 132, 1427-1431.	3.8	7
324	Discovery and replication of SNP-SNP interactions for quantitative lipid traits in over 60,000 individuals. BioData Mining, 2017, 10, 25.	4.0	7

#	Article	IF	CITATIONS
325	Genome-wide Association Study of Susceptibility to Particulate Matter–Associated QT Prolongation. Environmental Health Perspectives, 2017, 125, 067002.	6.0	7
326	Arsenic-gene interactions and beta-cell function in the Strong Heart Family Study. Toxicology and Applied Pharmacology, 2018, 348, 123-129.	2.8	7
327	The Value of Rare Genetic Variation in the Prediction of Common Obesity in European Ancestry Populations. Frontiers in Endocrinology, 2022, 13, 863893.	3.5	7
328	Gaseous air pollutants and DNA methylation in a methylome-wide association study of an ethnically and environmentally diverse population of U.S. adults. Environmental Research, 2022, 212, 113360.	7. 5	7
329	Sex-specific effects of ACE I/D and AGT-M235T on pulse pressure: the HyperGEN Study. Human Genetics, 2007, 122, 33-40.	3.8	6
330	Variation in the checkpoint kinase 2 gene is associated with type 2 diabetes in multiple populations. Acta Diabetologica, 2010, 47, 199-207.	2.5	6
331	Complex patterns of direct and indirect association between the transcription Factor-7 like 2 gene, body mass index and type 2 diabetes diagnosis in adulthood in the Hispanic Community Health Study/Study of Latinos. BMC Obesity, 2018, 5, 26.	3.1	6
332	Methylome-wide association study of central adiposity implicates genes involved in immune and endocrine systems. Epigenomics, 2020, 12, 1483-1499.	2.1	6
333	Genetically Determined Height and Risk of Non-hodgkin Lymphoma. Frontiers in Oncology, 2019, 9, 1539.	2.8	6
334	Multi-ethnic genome-wide association analyses of white blood cell and platelet traits in the Population Architecture using Genomics and Epidemiology (PAGE) study. BMC Genomics, 2021, 22, 432.	2.8	6
335	Genetic Risk Factors for BMI and Obesity in an Ethnically Diverse Population: Results From the Population Architecture Using Genomics and Epidemiology (PAGE) Study. Obesity, 0, , .	3.0	6
336	Enrichment analyses identify shared associations for 25 quantitative traits in over 600,000 individuals from seven diverse ancestries. American Journal of Human Genetics, 2022, 109, 871-884.	6.2	6
337	Findings from the Hispanic Community Health Study/Study of Latinos on the Importance of Sociocultural Environmental Interactors: Polygenic Risk Score-by-Immigration and Dietary Interactions. Frontiers in Genetics, 2021, 12, 720750.	2.3	6
338	Interactions Between Obesity, Parental History of Hypertension, and Age on Prevalent Hypertension. Asia-Pacific Journal of Public Health, 2012, 24, 970-980.	1.0	5
339	Chocolate consumption and prevalence of metabolic syndrome in the NHLBI Family Heart Study. E-SPEN Journal, 2012, 7, e139-e143.	0.5	5
340	Post-Genome-Wide Association Study Challenges for Lipid Traits: Describing Age as a Modifier of Gene-Lipid Associations in the Population Architecture Using Genomics and Epidemiology (PAGE) Study. Annals of Human Genetics, 2013, 77, 416-425.	0.8	5
341	Lack of association of apolipoprotein E (Apo E) polymorphism with the prevalence of metabolic syndrome: the National Heart, Lung and Blood Institute Family Heart Study. Diabetes/Metabolism Research and Reviews, 2015, 31, 582-587.	4.0	5
342	Genetic variation near <scp><i>IRS</i></scp> <i>1</i> <ii>is associated with adiposity and a favorable metabolic profile in <scp>U</scp>.<scp>S.</scp><scp>H</scp>ispanics/<scp>L</scp>atinos. Obesity, 2016, 24, 2407-2413.</ii>	3.0	5

#	Article	IF	Citations
343	Rare variant associations with waist-to-hip ratio in European-American and African-American women from the NHLBI-Exome Sequencing Project. European Journal of Human Genetics, 2016, 24, 1181-1187.	2.8	5
344	Gene-by-Psychosocial Factor Interactions Influence Diastolic Blood Pressure in European and African Ancestry Populations: Meta-Analysis of Four Cohort Studies. International Journal of Environmental Research and Public Health, 2017, 14, 1596.	2.6	5
345	Direct and indirect genetic effects on triglycerides through omics and correlated phenotypes. BMC Proceedings, 2018, 12, 22.	1.6	5
346	Generalization and fine mapping of red blood cell trait genetic associations to multiâ€ethnic populations: The PAGE study. American Journal of Hematology, 2018, 93, 1061-1073.	4.1	5
347	Genetic analysis of hsCRP in American Indians: The Strong Heart Family Study. PLoS ONE, 2019, 14, e0223574.	2.5	5
348	Whole Genome Sequencing Identifies CRISPLD2 as a Lung Function Gene in Children With Asthma. Chest, 2019, 156, 1068-1079.	0.8	5
349	Sociodemographic predictors of early postnatal growth: evidence from a Chilean infancy cohort. BMJ Open, 2020, 10, e033695.	1.9	5
350	The case-only independence assumption: associations between genetic polymorphisms and smoking among controls in two population-based studies. International Journal of Molecular Epidemiology and Genetics, 2012, 3, 333-60.	0.4	5
351	Predicted gene expression in ancestrally diverse populations leads to discovery of susceptibility loci for lifestyle and cardiometabolic traits. American Journal of Human Genetics, 2022, 109, 669-679.	6.2	5
352	Strengthening Causal Inference in Exposomics Research: Application of Genetic Data and Methods. Environmental Health Perspectives, 2022, 130, 55001.	6.0	5
353	Admixture mapping of pelvic organ prolapse in African Americans from the Women's Health Initiative Hormone Therapy trial. PLoS ONE, 2017, 12, e0178839.	2.5	4
354	The perils of standardizing infant weight to assess weight change differences across exposure groups. Annals of Epidemiology, 2018, 28, 515-520.	1.9	4
355	Genetic Variants Related to Cardiometabolic Traits Are Associated to B Cell Function, Insulin Resistance, and Diabetes Among AmeriCan Indians: The Strong Heart Family Study. Frontiers in Genetics, 2018, 9, 466.	2.3	4
356	Maternal diet as a risk factor for primary congenital glaucoma and defects of the anterior segment of the eye in the <scp>National Birth Defects Prevention Study</scp> . Birth Defects Research, 2020, 112, 503-514.	1.5	4
357	Multi-Ethnic Genome-Wide Association Study of Decomposed Cardioelectric Phenotypes Illustrates Strategies to Identify and Characterize Evidence of Shared Genetic Effects for Complex Traits. Circulation Genomic and Precision Medicine, 2020, 13, e002680.	3.6	4
358	Epigenetically mediated electrocardiographic manifestations of sub-chronic exposures to ambient particulate matter air pollution in the Women's Health Initiative and Atherosclerosis Risk in Communities Study. Environmental Research, 2021, 198, 111211.	7.5	4
359	Do adverse childhood experiences and genetic obesity risk interact in relation to body mass index in young adulthood? Findings from the National Longitudinal Study of Adolescent to Adult Health. Pediatric Obesity, 2022, 17, e12885.	2.8	4
360	Novel diabetes gene discovery through comprehensive characterization and integrative analysis of longitudinal gene expression changes. Human Molecular Genetics, 2022, 31, 3191-3205.	2.9	4

#	Article	IF	CITATIONS
361	The Utility of Bayesian Model Averaging for Detecting Known Oligogenic Effects. Genetic Epidemiology, 2001, 21, S789-93.	1.3	3
362	Mapping of a blood pressure QTL on chromosome 17 in American Indians of the strong heart family study. BMC Cardiovascular Disorders, 2014, 14, 158.	1.7	3
363	Evidence for Association between <i>SH2B1</i> Gene Variants and Glycated Hemoglobin in Nondiabetic European American Young Adults: The Add Health Study. Annals of Human Genetics, 2016, 80, 294-305.	0.8	3
364	Comparison of 2 models for gene–environment interactions: an example of simulated gene–medication interactions on systolic blood pressure in family-based data. BMC Proceedings, 2016, 10, 371-377.	1.6	3
365	Characterization of the contribution of shared environmental and genetic factors to metabolic syndrome methylation heritability and familial correlations. BMC Genetics, 2018, 19, 69.	2.7	3
366	Tracing the Distribution of European Lactase Persistence Genotypes Along the Americas. Frontiers in Genetics, 2021, 12, 671079.	2.3	3
367	Genomeâ€wide association study identifying novel variant for fasting insulin and allelic heterogeneity in known glycemic loci in Chilean adolescents: The Santiago Longitudinal Study. Pediatric Obesity, 2021, 16, e12765.	2.8	3
368	Dynamic relationships between body fat and circulating adipokine levels from adolescence to young adulthood: The Santiago Longitudinal Study. Nutrition, Metabolism and Cardiovascular Diseases, 2022, 32, 1055-1063.	2.6	3
369	Ancestral diversity improves discovery and fine-mapping of genetic loci for anthropometric traitsâ€"The Hispanic/Latino Anthropometry Consortium. Human Genetics and Genomics Advances, 2022, 3, 100099.	1.7	3
370	Extended Human Papillomavirus Genotyping to Predict Progression to High-Grade Cervical Precancer: A Prospective Cohort Study in the Southeastern United States. Cancer Epidemiology Biomarkers and Prevention, 2022, 31, 1564-1571.	2.5	3
371	Multi-ancestry genome-wide association study accounting for gene-psychosocial factor interactions identifies novel loci for blood pressure traits. Human Genetics and Genomics Advances, 2021, 2, 100013.	1.7	2
372	Full title: A largeâ€scale transcriptomeâ€wide association study (TWAS) of 10 blood cell phenotypes reveals complexities of TWAS fineâ€mapping. Genetic Epidemiology, 2021, , .	1.3	2
373	Rare coding variants in RCN3 are associated with blood pressure. BMC Genomics, 2022, 23, 148.	2.8	2
374	The Association of Cell Cycle Checkpoint 2 Variants and Kidney Function: Findings of the Family Blood Pressure Program and the Atherosclerosis Risk in Communities Study. American Journal of Hypertension, 2009, 22, 552-558.	2.0	1
375	Fine mapping and identification of serum urate loci in American Indians: The Strong Heart Family Study. Scientific Reports, 2019, 9, 17899.	3.3	1
376	Genetic variants and physical activity interact to affect bone density in Hispanic children. BMC Pediatrics, 2021, 21, 79.	1.7	1
377	Discrimination and Leukocyte Telomere Length by Depressive Symptomatology: The Jackson Heart Study. Healthcare (Switzerland), 2021, 9, 639.	2.0	1
378	Nine-Year Ethanol Intake Trajectories and Their Association With 15-Year Cognitive Decline Among Black and White Adults. American Journal of Epidemiology, 2020, 189, 788-800.	3.4	1

#	Article	IF	Citations
379	OUP accepted manuscript. American Journal of Epidemiology, 2022, , .	3.4	1
380	Metabolomic Associations of Asthma in the Hispanic Community Health Study/Study of Latinos. Metabolites, 2022, 12, 359.	2.9	1
381	Reply to â∈̃Misestimation of heritability and prediction accuracy of male-pattern baldness'. Nature Communications, 2018, 9, 2538.	12.8	0
382	Genome-wide homozygosity and risk of four non-Hodgkin lymphoma subtypes., 2021, 5, 200-217.		0
383	Gene-Environment Joint Linkage and Association Analysis of Arsenic Exposure and Diabetes-Related Traits in the Strong Heart Family Study. ISEE Conference Abstracts, 2021, 2021, .	0.0	0
384	Genetic determinants of metabolic biomarkers and their associations with cardiometabolic traits in Hispanic/Latino adolescents. Pediatric Research, 2021, , .	2.3	0
385	Abstract 051: Trans-ethnic Metabochip Genotyping of Established Lipid Loci Identifies Low Frequency Susceptibility Variants and Additional Independent Signals in Known Loci. Circulation, 2012, 125, .	1.6	0
386	Abstract P223: Larger Effect Sizes of Established BMI Genetic Variants During Adolescence, a Vulnerable Period of Weight Gain. Circulation, 2012, 125, .	1.6	0
387	Abstract 050: Meta-analysis of Genetic Associations in up to 339,224 Individuals Identify 66 New Loci for Bmi, Confirming a Neuronal Contribution to Body Weight Regulation and Implicating Several Novel Pathways. Circulation, 2013, 127, .	1.6	0
388	Abstract P157: Does Physical Activity Modify the Association of 15 Well-established Obesity Loci with BMI: The ARIC Study. Circulation, 2013, 127, .	1.6	0
389	The interaction between physical activity and obesity gene variants in association with BMI: Does the obesogenic environment matter?. FASEB Journal, 2013, 27, 236.5.	0.5	0
390	Abstract 21: Accounting For Smoking Behavior In Genome-wide Analysis Of Obesity Phenotypes: The Giant (genetic Investigation Of Anthropometric Traits) Consortium. Circulation, 2014, 129, .	1.6	0
391	Abstract P261: Genome-wide Association Study of Susceptibility to Particulate Matter-associated Reduced Heart Rate Variability. Circulation, 2016, 133, .	1.6	0
392	Abstract 053: Genome-wide Tcea3 -SNP Interaction Study Identifies Novel QT Interval Loci. Circulation, 2018, 137, .	1.6	0
393	Abstract 002: Adherence to Ideal Life's Simple 7 Metrics is Associated With Epigenetic Biomarkers of Aging in African Americans: The Atherosclerosis Risk in Communities (ARIC) Study. Circulation, 2019, 139, .	1.6	0
394	The dynamic genetic architecture of early childhood BMI. Nature Metabolism, 2022, 4, 308-309.	11.9	0
395	Title is missing!. , 2020, 16, e1008684.		0
396	Title is missing!. , 2020, 16, e1008684.		0

#	Article	lF	CITATIONS
397	Title is missing!. , 2020, 16, e1008684.		0
398	Title is missing!. , 2020, 16, e1008684.		0
399	Title is missing!. , 2020, 16, e1008684.		0
400	Title is missing!. , 2020, 16, e1008684.		0