## Colleen M Sitlani

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1272003/publications.pdf

Version: 2024-02-01

62 papers

2,988 citations

<sup>331538</sup>
21
h-index

50 g-index

62 all docs

62 docs citations

times ranked

62

8286 citing authors

#	Article	IF	CITATIONS
1	New genetic loci link adipose and insulin biology to body fat distribution. Nature, 2015, 518, 187-196.	13.7	1,328
2	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.	2.6	123
3	Circulating Sphingolipids, Insulin, HOMA-IR, and HOMA-B: The Strong Heart Family Study. Diabetes, 2018, 67, 1663-1672.	0.3	120
4	Global Electric Heterogeneity Risk Score for Prediction of Sudden Cardiac Death in the General Population. Circulation, 2016, 133, 2222-2234.	1.6	118
5	Plasma phospholipid very-long-chain saturated fatty acids and incident diabetes in older adults: the Cardiovascular Health Study. American Journal of Clinical Nutrition, 2015, 101, 1047-1054.	2.2	97
6	Novel genetic associations for blood pressure identified via gene-alcohol interaction in up to 570K individuals across multiple ancestries. PLoS ONE, 2018, 13, e0198166.	1.1	94
7	Analysis commons, a team approach to discovery in a big-data environment for genetic epidemiology. Nature Genetics, 2017, 49, 1560-1563.	9.4	93
8	Multiethnic meta-analysis identifies ancestry-specific and cross-ancestry loci for pulmonary function. Nature Communications, 2018, 9, 2976.	5.8	85
9	Plasma Phospholipid Saturated Fatty Acids and Incident Atrial Fibrillation: The Cardiovascular Health Study. Journal of the American Heart Association, 2014, 3, e000889.	1.6	71
10	Circulating sphingolipids, fasting glucose, and impaired fasting glucose: The Strong Heart Family Study. EBioMedicine, 2019, 41, 44-49.	2.7	48
11	Circulating Very Longâ€Chain Saturated Fatty Acids and Heart Failure: The Cardiovascular Health Study. Journal of the American Heart Association, 2018, 7, e010019.	1.6	45
12	Whole Blood DNA Methylation Signatures of Diet Are Associated With Cardiovascular Disease Risk Factors and All-Cause Mortality. Circulation Genomic and Precision Medicine, 2020, 13, e002766.	1.6	42
13	Parental smoking during pregnancy and offspring cardio-metabolic risk factors at ages 17 and 32. Atherosclerosis, 2014, 235, 430-437.	0.4	39
14	Plasma Ceramide Species Are Associated with Diabetes Risk in Participants of the Strong Heart Study. Journal of Nutrition, 2020, 150, 1214-1222.	1.3	38
15	Generalized estimating equations for genomeâ€wide association studies using longitudinal phenotype data. Statistics in Medicine, 2015, 34, 118-130.	0.8	37
16	Epigenetic Age and the Risk of Incident Atrial Fibrillation. Circulation, 2021, 144, 1899-1911.	1.6	35
17	Genomeâ€Wide Associations of Global Electrical Heterogeneity ECG Phenotype: The ARIC (Atherosclerosis Risk in Communities) Study and CHS (Cardiovascular Health Study). Journal of the American Heart Association, 2018, 7, .	1.6	31
18	Plasma Ceramides and Sphingomyelins in Relation to Atrial Fibrillation Risk: The Cardiovascular Health Study. Journal of the American Heart Association, 2020, 9, e012853.	1.6	31

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19	Associations of Plasma Phospholipid SFAs with Total and Cause-Specific Mortality in Older Adults Differ According to SFA Chain Length. Journal of Nutrition, 2016, 146, 298-305.	1.3	29
20	Innate and adaptive immune cell subsets as risk factors for coronary heart disease in two population-based cohorts. Atherosclerosis, 2020, 300, 47-53.	0.4	28
21	Variation in resting heart rate over 4â€years and the risks of myocardial infarction and death among older adults. Heart, 2015, 101, 132-138.	1.2	27
22	Genetic Studies of Leptin Concentrations Implicate Leptin in the Regulation of Early Adiposity. Diabetes, 2020, 69, 2806-2818.	0.3	26
23	Exome-Derived Adiponectin-Associated Variants Implicate Obesity and Lipid Biology. American Journal of Human Genetics, 2019, 105, 15-28.	2.6	21
24	Circulating Ceramides and Sphingomyelins and Risk of Mortality: The Cardiovascular Health Study. Clinical Chemistry, 2021, 67, 1650-1659.	1.5	21
25	Association of Brain Volumes and White Matter Injury With Race, Ethnicity, and Cardiovascular Risk Factors: The Multiâ€Ethnic Study of Atherosclerosis. Journal of the American Heart Association, 2022, 11, e023159.	1.6	21
26	Incident Atrial Fibrillation and Disabilityâ€Free Survival in the Cardiovascular Health Study. Journal of the American Geriatrics Society, 2016, 64, 838-843.	1.3	20
27	Plasma ceramides containing saturated fatty acids are associated with risk of type 2 diabetes. Journal of Lipid Research, 2021, 62, 100119.	2.0	19
28	Meta-analysis of exome array data identifies six novel genetic loci for lung function. Wellcome Open Research, 2018, 3, 4.	0.9	19
29	Pharmacogenomics of statin-related myopathy: Meta-analysis of rare variants from whole-exome sequencing. PLoS ONE, 2019, 14, e0218115.	1.1	18
30	Longitudinal structural mixed models for the analysis of surgical trials with noncompliance. Statistics in Medicine, 2012, 31, 1738-1760.	0.8	17
31	Genome-Wide Association Study of Apparent Treatment-Resistant Hypertension in the CHARGE Consortium: The CHARGE Pharmacogenetics Working Group. American Journal of Hypertension, 2019, 32, 1146-1153.	1.0	17
32	Common variation in fatty acid metabolic genes and risk of incident sudden cardiac arrest. Heart Rhythm, 2014, 11, 471-477.	0.3	16
33	Parent-of-Origin Effects of the APOB Gene on Adiposity in Young Adults. PLoS Genetics, 2015, 11, e1005573.	1.5	16
34	Drug-Gene Interactions of Antihypertensive Medications and Risk of Incident Cardiovascular Disease: A Pharmacogenomics Study from the CHARGE Consortium. PLoS ONE, 2015, 10, e0140496.	1.1	15
35	Identifying genetic loci associated with antidepressant drug response with drug–gene interaction models in a population-based study. Journal of Psychiatric Research, 2015, 62, 31-37.	1.5	13
36	The Challenges of Genome-Wide Interaction Studies: Lessons to Learn from the Analysis of HDL Blood Levels. PLoS ONE, 2014, 9, e109290.	1.1	13

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37	Multiâ€phenotype analyses of hemostatic traits with cardiovascular events reveal novel genetic associations. Journal of Thrombosis and Haemostasis, 2022, 20, 1331-1349.	1.9	12
38	Role of Rare and Low-Frequency Variants in Gene-Alcohol Interactions on Plasma Lipid Levels. Circulation Genomic and Precision Medicine, 2020, 13, e002772.	1.6	11
39	A systematic analysis of protein-altering exonic variants in chronic obstructive pulmonary disease. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2021, 321, L130-L143.	1.3	11
40	Meta-analysis of exome array data identifies six novel genetic loci for lung function. Wellcome Open Research, 0, 3, 4.	0.9	11
41	Associations of Innate and Adaptive Immune Cell Subsets With Incident Type 2 Diabetes Risk: The MESA Study. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e848-e857.	1.8	10
42	Nonclassical Monocytes (CD14dimCD16+) Are Associated With Carotid Intima-Media Thickness Progression for Men but Not Women. Arteriosclerosis, Thrombosis, and Vascular Biology, 2021, 41, 1810-1817.	1.1	10
43	Natural killer cells, gamma delta T cells and classical monocytes are associated with systolic blood pressure in the multi-ethnic study of atherosclerosis (MESA). BMC Cardiovascular Disorders, 2021, 21, 45.	0.7	10
44	Obesity Partially Mediates the Diabetogenic Effect of Lowering LDL Cholesterol. Diabetes Care, 2022, 45, 232-240.	4.3	10
45	Maternal Genetic Variation Accounts in Part for the Associations of Maternal Size during Pregnancy with Offspring Cardiometabolic Risk in Adulthood. PLoS ONE, 2014, 9, e91835.	1.1	9
46	A genome-wide interaction analysis of tricyclic/tetracyclic antidepressants and RR and QT intervals: a pharmacogenomics study from the Cohorts for Heart and Aging Research in Genomic Epidemiology (CHARGE) consortium. Journal of Medical Genetics, 2017, 54, 313-323.	1.5	9
47	Associations of Early and Late Gestational Weight Gain with Infant Birth Size. Maternal and Child Health Journal, 2015, 19, 2462-2469.	0.7	8
48	Integrative analysis of clinical and epigenetic biomarkers of mortality. Aging Cell, 2022, 21, e13608.	3.0	8
49	Genome-wide association study and meta-analysis identify loci associated with ventricular and supraventricular ectopy. Scientific Reports, 2018, 8, 5675.	1.6	4
50	Genomeâ€wide metaâ€analysis of SNP and antihypertensive medication interactions on left ventricular traits in African Americans. Molecular Genetics & Enomic Medicine, 2019, 7, e00788.	0.6	4
51	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.	1.6	4
52	Plasma epoxyeicosatrienoic acids and dihydroxyeicosatrieonic acids, insulin, glucose and risk of diabetes: The strong heart study. EBioMedicine, 2021, 66, 103279.	2.7	4
53	Association of immune cell subsets with cardiac mechanics in the Multi-Ethnic Study of Atherosclerosis. JCI Insight, 2021, 6, .	2.3	4
54	Monocyte subsets, T cell activation profiles, and stroke in men and women: The Multi-Ethnic Study of Atherosclerosis and Cardiovascular Health Study. Atherosclerosis, 2022, 351, 18-25.	0.4	4

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55	Genome-wide meta-analysis of SNP-by9-ACEI/ARB and SNP-by-thiazide diuretic and effect on serum potassium in cohorts of European and African ancestry. Pharmacogenomics Journal, 2019, 19, 97-108.	0.9	3
56	Incorporating sampling weights into robust estimation of Cox proportional hazards regression model, with illustration in the Multi-Ethnic Study of Atherosclerosis. BMC Medical Research Methodology, 2020, 20, 62.	1.4	3
57	Analyzing longitudinal data to characterize the accuracy of markers used to select treatment. Statistics in Medicine, 2014, 33, 2881-2896.	0.8	2
58	Rare and low-frequency exonic variants and gene-by-smoking interactions in pulmonary function. Scientific Reports, $2021, 11, 19365$ .	1.6	2
59	Genome-wide gene–environment interactions on quantitative traits using family data. European Journal of Human Genetics, 2016, 24, 1022-1028.	1.4	1
60	Comparison of adaptive multiple phenotype association tests using summary statistics in genome-wide association studies. Human Molecular Genetics, 2021, 30, 1371-1383.	1.4	1
61	Meta-analysis of exome array data identifies six novel genetic loci for lung function. Wellcome Open Research, 0, 3, 4.	0.9	1
62	GWAS of Variant-by-Thiazide Interaction on Lipids Identifies a Novel Low-Density Lipoprotein Cholesterol Locus. Circulation Research, 0, , .	2.0	1