

# Ruth McPherson

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6744082/publications.pdf>

Version: 2024-02-01

89  
papers

16,702  
citations

101543

36  
h-index

49909

87  
g-index

91  
all docs

91  
docs citations

91  
times ranked

24345  
citing authors

#	ARTICLE	IF	CITATIONS
1	Rare coding variants in 35 genes associate with circulating lipid levelsâ€”A multi-ancestry analysis of 170,000 exomes. <i>American Journal of Human Genetics</i> , 2022, 109, 81-96.	6.2	24
2	Common Polymorphism That Protects From Cardiovascular Disease Increases Fibronectin Processing and Secretion. <i>Circulation Genomic and Precision Medicine</i> , 2022, 15, CIRCGEN121003428.	3.6	5
3	Association of muscle fiber type with measures of obesity: A systematic review. <i>Obesity Reviews</i> , 2022, 23, e13444.	6.5	10
4	Genetically Determined Reproductive Aging and Coronary Heart Disease: A Bidirectional 2-sample Mendelian Randomization. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e2952-e2961.	3.6	13
5	A novel anti-inflammatory role links the CARS2 locus to protection from coronary artery disease. <i>Atherosclerosis</i> , 2022, 348, 8-15.	0.8	3
6	Convergence of biomarkers and risk factor trait loci of coronary artery disease at 3p21.31 and HLA region. <i>Npj Genomic Medicine</i> , 2021, 6, 12.	3.8	6
7	Common Polymorphism in the FADS1 Locus Links miR1908 to Low-Density Lipoprotein Cholesterol Through BMP-1. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2021, 41, 2252-2262.	2.4	4
8	Epigenome-Wide Study Identified Methylation Sites Associated with the Risk of Obesity. <i>Nutrients</i> , 2021, 13, 1984.	4.1	8
9	Interindividual variability in weight loss in the treatment of obesity. <i>American Journal of Clinical Nutrition</i> , 2021, 114, 824-825.	4.7	3
10	Understanding the Function of a Locus Using the Knowledge Available at Single-Nucleotide Polymorphisms. <i>Neurology International</i> , 2021, 11, 255-262.	0.5	2
11	miR1908-5p regulates energy homeostasis in hepatocyte models. <i>Scientific Reports</i> , 2021, 11, 23748.	3.3	2
12	Multomics Screening Identifies Molecular Biomarkers Causally Associated With the Risk of Coronary Artery Disease. <i>Circulation Genomic and Precision Medicine</i> , 2020, 13, e002876.	3.6	9
13	RIPK1 gene variants associate with obesity in humans and can be therapeutically silenced to reduce obesity in mice. <i>Nature Metabolism</i> , 2020, 2, 1113-1125.	11.9	34
14	Factors affecting weight loss variability in obesity. <i>Metabolism: Clinical and Experimental</i> , 2020, 113, 154388.	3.4	50
15	Association of Factor V Leiden With Subsequent Atherothrombotic Events. <i>Circulation</i> , 2020, 142, 546-555.	1.6	11
16	Heterozygous <i>ABCG5</i> Gene Deficiency and Risk of Coronary Artery Disease. <i>Circulation Genomic and Precision Medicine</i> , 2020, 13, 417-423.	3.6	45
17	Molecular mechanism linking a novel PCSK9 copy number variant to severe hypercholesterolemia. <i>Atherosclerosis</i> , 2020, 304, 39-43.	0.8	3
18	<i>SGCG</i> rs679482 Associates With Weight Loss Success in Response to an Intensively Supervised Outpatient Program. <i>Diabetes</i> , 2020, 69, 2017-2026.	0.6	8

#	ARTICLE	IF	CITATIONS
19	A missense variant in Mitochondrial Amidoxime Reducing Component 1 gene and protection against liver disease. <i>PLoS Genetics</i> , 2020, 16, e1008629.	3.5	101
20	2018 George Lyman Duff Memorial Lecture. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2019, 39, 1925-1937.	2.4	6
21	Genome-wide identification of circulating-miRNA expression quantitative trait loci reveals the role of several miRNAs in the regulation of cardiometabolic phenotypes. <i>Cardiovascular Research</i> , 2019, 115, 1629-1645.	3.8	55
22	The selective peroxisome proliferator-activated receptor alpha modulator (SPPARM $\alpha$ ) paradigm: conceptual framework and therapeutic potential. <i>Cardiovascular Diabetology</i> , 2019, 18, 71.	6.8	104
23	Subsequent Event Risk in Individuals With Established Coronary Heart Disease. <i>Circulation Genomic and Precision Medicine</i> , 2019, 12, e002470.	3.6	17
24	Regulation of MFGE8 by the intergenic coronary artery disease locus on 15q26.1. <i>Atherosclerosis</i> , 2019, 284, 11-17.	0.8	26
25	Off-target effects of CRISPRa on interleukin-6 expression. , 2019, 14, e0224113.		0
26	Off-target effects of CRISPRa on interleukin-6 expression. , 2019, 14, e0224113.		0
27	Off-target effects of CRISPRa on interleukin-6 expression. , 2019, 14, e0224113.		0
28	Off-target effects of CRISPRa on interleukin-6 expression. , 2019, 14, e0224113.		0
29	Partitioning the Pleiotropy Between Coronary Artery Disease and Body Mass Index Reveals the Importance of Low Frequency Variants and Central Nervous System-Specific Functional Elements. <i>Circulation Genomic and Precision Medicine</i> , 2018, 11, e002050.	3.6	16
30	Analysis of predicted loss-of-function variants in UK Biobank identifies variants protective for disease. <i>Nature Communications</i> , 2018, 9, 1613.	12.8	78
31	Phenotypic Consequences of a Genetic Predisposition to Enhanced Nitric Oxide Signaling. <i>Circulation</i> , 2018, 137, 222-232.	1.6	87
32	Is Type 2 Diabetes in Adults Associated With Impaired Capacity for Weight Loss?. <i>Canadian Journal of Diabetes</i> , 2018, 42, 313-316.e1.	0.8	7
33	Obesity shows preserved plasma proteome in large independent clinical cohorts. <i>Scientific Reports</i> , 2018, 8, 16981.	3.3	45
34	The Cardiovascular Burden of Undiagnosed Familial Hypercholesterolemia: Need to Modify Guidelines to Encourage Earlier Diagnosis and Therapy. <i>Canadian Journal of Cardiology</i> , 2018, 34, 1112-1113.	1.7	4
35	Systematic Evaluation of Pleiotropy Identifies 6 Further Loci Associated With Coronary Artery Disease. <i>Journal of the American College of Cardiology</i> , 2017, 69, 823-836.	2.8	214
36	Relations between lipoprotein(a) concentrations, LPA genetic variants, and the risk of mortality in patients with established coronary heart disease: a molecular and genetic association study. <i>Lancet Diabetes and Endocrinology</i> , 2017, 5, 534-543.	11.4	84

#	ARTICLE	IF	CITATIONS
37	TRIB1 is a positive regulator of hepatocyte nuclear factor 4-alpha. <i>Scientific Reports</i> , 2017, 7, 5574.	3.3	26
38	Association analyses based on false discovery rate implicate new loci for coronary artery disease. <i>Nature Genetics</i> , 2017, 49, 1385-1391.	21.4	571
39	Partitioning the heritability of coronary artery disease highlights the importance of immune-mediated processes and epigenetic sites associated with transcriptional activity. <i>Cardiovascular Research</i> , 2017, 113, 973-983.	3.8	31
40	Can response to dietary restriction predict weight loss after Roux-Y gastroplasty?. <i>Obesity</i> , 2016, 24, 805-811.	3.0	7
41	Diagnostic Yield and Clinical Utility of Sequencing Familial Hypercholesterolemia Genes in Patients With Severe Hypercholesterolemia. <i>Journal of the American College of Cardiology</i> , 2016, 67, 2578-2589.	2.8	723
42	Coding Variation in <i>ANGPTL4</i> , <i>LPL</i> and <i>SVEP1</i> and the Risk of Coronary Disease. <i>New England Journal of Medicine</i> , 2016, 374, 1134-1144.	27.0	427
43	Adverse Effects of $\beta$ -Blocker Therapy on Weight Loss in Response to a Controlled Dietary Regimen. <i>Canadian Journal of Cardiology</i> , 2016, 32, 1246.e21-1246.e26.	1.7	5
44	The Sum of Its Parts: The Polygenic Basis of Coronary Artery Disease. <i>Canadian Journal of Cardiology</i> , 2016, 32, 1372-1374.	1.7	1
45	Role of Tribbles Pseudokinase 1 (TRIB1) in human hepatocyte metabolism. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2016, 1862, 223-232.	3.8	16
46	Functional Analysis of a Novel Genome-Wide Association Study Signal in <i>SMAD3</i> That Confers Protection From Coronary Artery Disease. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2016, 36, 972-983.	2.4	48
47	Genome-wide association study and targeted metabolomics identifies sex-specific association of CPS1 with coronary artery disease. <i>Nature Communications</i> , 2016, 7, 10558.	12.8	108
48	Genetics of Coronary Artery Disease. <i>Circulation Research</i> , 2016, 118, 564-578.	4.5	288
49	Association of exome sequences with plasma C-reactive protein levels in >9000 participants. <i>Human Molecular Genetics</i> , 2015, 24, 559-571.	2.9	36
50	New genetic loci link adipose and insulin biology to body fat distribution. <i>Nature</i> , 2015, 518, 187-196.	27.8	1,328
51	Genetic variants primarily associated with type 2 diabetes are related to coronary artery disease risk. <i>Atherosclerosis</i> , 2015, 241, 419-426.	0.8	26
52	Runs of Homozygosity: Association with Coronary Artery Disease and Gene Expression in Monocytes and Macrophages. <i>American Journal of Human Genetics</i> , 2015, 97, 228-237.	6.2	37
53	A pharmacodynamic comparison of a personalized strategy for anti-platelet therapy versus ticagrelor in achieving a therapeutic window. <i>International Journal of Cardiology</i> , 2015, 197, 318-325.	1.7	15
54	IRF2BP2 Reduces Macrophage Inflammation and Susceptibility to Atherosclerosis. <i>Circulation Research</i> , 2015, 117, 671-683.	4.5	64

#	ARTICLE	IF	CITATIONS
55	Prognostic and Therapeutic Implications of Statin and Aspirin Therapy in Individuals With Nonobstructive Coronary Artery Disease. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2015, 35, 981-989.	2.4	147
56	Ezetimibe. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2015, 35, e13-5.	2.4	15
57	Obesity and Ischemic Heart Disease. <i>Circulation Research</i> , 2015, 116, 570-571.	4.5	7
58	Nonstatin Low-Density Lipoproteinâ€“Lowering Therapy and Cardiovascular Risk Reductionâ€“Statement From <i>ATVB</i> Council. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2015, 35, 2269-2280.	2.4	58
59	Functional interaction between COL4A1/COL4A2 and SMAD3 risk loci for coronary artery disease. <i>Atherosclerosis</i> , 2015, 242, 543-552.	0.8	55
60	<i>PHACTR1</i>. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2015, 35, 1293-1295.	2.4	2
61	A comprehensive 1000 Genomesâ€“based genome-wide association meta-analysis of coronary artery disease. <i>Nature Genetics</i> , 2015, 47, 1121-1130.	21.4	2,054
62	Exome sequencing identifies rare LDLR and APOA5 alleles conferring risk for myocardial infarction. <i>Nature</i> , 2015, 518, 102-106.	27.8	581
63	Acylcarnitines: potential implications for skeletal muscle insulin resistance. <i>FASEB Journal</i> , 2015, 29, 336-345.	0.5	191
64	Lower Mitochondrial Proton Leak and Decreased Glutathione Redox in Primary Muscle Cells of Obese Diet-Resistant Versus Diet-Sensitive Humans. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, 4223-4230.	3.6	17
65	Distribution and Medical Impact of Loss-of-Function Variants in the Finnish Founder Population. <i>PLoS Genetics</i> , 2014, 10, e1004494.	3.5	351
66	Adiposity significantly modifies genetic risk for dyslipidemia. <i>Journal of Lipid Research</i> , 2014, 55, 2416-2422.	4.2	33
67	A Lowâ€“Frequency Variant in MAPK14 Provides Mechanistic Evidence of a Link With Myeloperoxidase: A Prognostic Cardiovascular Risk Marker. <i>Journal of the American Heart Association</i> , 2014, 3, .	3.7	7
68	Functional Analysis of the TRIB <i>1</i> Associated Locus Linked to Plasma Triglycerides and Coronary Artery Disease. <i>Journal of the American Heart Association</i> , 2014, 3, e000884.	3.7	42
69	Risk Stratification and Selection for Statin Therapy: Going Beyond Framingham. <i>Canadian Journal of Cardiology</i> , 2014, 30, 667-670.	1.7	7
70	Genome-Wide Association Studies of Cardiovascular Disease in European and Non-European Populations. <i>Current Genetic Medicine Reports</i> , 2014, 2, 1-12.	1.9	16
71	Association of Low-Frequency and Rare Coding-Sequence Variants with Blood Lipids and Coronary Heart Disease in 56,000 Whites and Blacks. <i>American Journal of Human Genetics</i> , 2014, 94, 223-232.	6.2	287
72	Low copy number of the salivary amylase gene predisposes to obesity. <i>Nature Genetics</i> , 2014, 46, 492-497.	21.4	214

#	ARTICLE	IF	CITATIONS
73	Netrin-1 promotes adipose tissue macrophage retention and insulin resistance in obesity. <i>Nature Medicine</i> , 2014, 20, 377-384.	30.7	213
74	From Genome-Wide Association Studies to Functional Genomics: New Insights Into Cardiovascular Disease. <i>Canadian Journal of Cardiology</i> , 2013, 29, 23-29.	1.7	16
75	ERK1/2 regulates hepatocyte Trib1 in response to mitochondrial dysfunction. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2013, 1833, 3405-3414.	4.1	15
76	Chromosome 9p21.3 Locus for Coronary Artery Disease. <i>Journal of the American College of Cardiology</i> , 2013, 62, 1382-1383.	2.8	14
77	A 680 kb duplication at the FTO locus in a kindred with obesity and a distinct body fat distribution. <i>European Journal of Human Genetics</i> , 2013, 21, 1417-1422.	2.8	10
78	Remnant Cholesterol. <i>Journal of the American College of Cardiology</i> , 2013, 61, 437-439.	2.8	26
79	Inflammation and Coronary Artery Disease: Insights From Genetic Studies. <i>Canadian Journal of Cardiology</i> , 2012, 28, 662-666.	1.7	45
80	Blood Gene Expression Reveal Pathway Differences Between Diet-Sensitive and Resistant Obese Subjects Prior to Caloric Restriction. <i>Obesity</i> , 2011, 19, 457-463.	3.0	23
81	Biological, clinical and population relevance of 95 loci for blood lipids. <i>Nature</i> , 2010, 466, 707-713.	27.8	3,249
82	Distinct skeletal muscle fiber characteristics and gene expression in diet-sensitive versus diet-resistant obesity. <i>Journal of Lipid Research</i> , 2010, 51, 2394-2404.	4.2	52
83	New genetic loci implicated in fasting glucose homeostasis and their impact on type 2 diabetes risk. <i>Nature Genetics</i> , 2010, 42, 105-116.	21.4	1,982
84	Genetic contributors to obesity. <i>Canadian Journal of Cardiology</i> , 2007, 23, 23A-27A.	1.7	30
85	A Common Allele on Chromosome 9 Associated with Coronary Heart Disease. <i>Science</i> , 2007, 316, 1488-1491.	12.6	1,591
86	Canadian Cardiovascular Society position statement "Recommendations for the diagnosis and treatment of dyslipidemia and prevention of cardiovascular disease. <i>Canadian Journal of Cardiology</i> , 2006, 22, 913-927.	1.7	271
87	Molecular regulation of SREBP function: the Insig-SCAP connection and isoform-specific modulation of lipid synthesis. <i>Biochemistry and Cell Biology</i> , 2004, 82, 201-211.	2.0	98
88	Decreased Mitochondrial Proton Leak and Reduced Expression of Uncoupling Protein 3 in Skeletal Muscle of Obese Diet-Resistant Women. <i>Diabetes</i> , 2002, 51, 2459-2466.	0.6	113
89	Efficacy of atorvastatin in achieving National Cholesterol Education Program low-density lipoprotein targets in women with severe dyslipidemia and cardiovascular disease or risk factors for cardiovascular disease: The Women's Atorvastatin Trial on Cholesterol (WATCH). <i>American Heart Journal</i> , 2001, 141, 949-956.	2.7	21