

Cassandra N Spracklen

List of Publications by Year in descending order

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Version: 2024-02-01

34
papers

3,665
citations

430874

18
h-index

361022

35
g-index

45
all docs

45
docs citations

45
times ranked

6278
citing authors

#	ARTICLE	IF	CITATIONS
1	A catalog of genetic loci associated with kidney function from analyses of a million individuals. <i>Nature Genetics</i> , 2019, 51, 957-972.	21.4	549
2	The Polygenic and Monogenic Basis of Blood Traits and Diseases. <i>Cell</i> , 2020, 182, 1214-1231.e11.	28.9	388
3	Trans-ethnic and Ancestry-Specific Blood-Cell Genetics in 746,667 Individuals from 5 Global Populations. <i>Cell</i> , 2020, 182, 1198-1213.e14.	28.9	353
4	The power of genetic diversity in genome-wide association studies of lipids. <i>Nature</i> , 2021, 600, 675-679.	27.8	353
5	The trans-ancestral genomic architecture of glycemic traits. <i>Nature Genetics</i> , 2021, 53, 840-860.	21.4	341
6	Identification of type 2 diabetes loci in 433,540 East Asian individuals. <i>Nature</i> , 2020, 582, 240-245.	27.8	282
7	Target genes, variants, tissues and transcriptional pathways influencing human serum urate levels. <i>Nature Genetics</i> , 2019, 51, 1459-1474.	21.4	251
8	Multi-ancestry genetic study of type 2 diabetes highlights the power of diverse populations for discovery and translation. <i>Nature Genetics</i> , 2022, 54, 560-572.	21.4	250
9	Association analyses of East Asian individuals and trans-ancestry analyses with European individuals reveal new loci associated with cholesterol and triglyceride levels. <i>Human Molecular Genetics</i> , 2017, 26, 1770-1784.	2.9	135
10	Exome chip meta-analysis identifies novel loci and East Asian-specific coding variants that contribute to lipid levels and coronary artery disease. <i>Nature Genetics</i> , 2017, 49, 1722-1730.	21.4	129
11	Associations of autozygosity with a broad range of human phenotypes. <i>Nature Communications</i> , 2019, 10, 4957.	12.8	84
12	Interethnic analyses of blood pressure loci in populations of East Asian and European descent. <i>Nature Communications</i> , 2018, 9, 5052.	12.8	75
13	Birth weight and subsequent risk of cancer. <i>Cancer Epidemiology</i> , 2014, 38, 538-543.	1.9	57
14	Adipose Tissue Gene Expression Associations Reveal Hundreds of Candidate Genes for Cardiometabolic Traits. <i>American Journal of Human Genetics</i> , 2019, 105, 773-787.	6.2	45
15	Physical Activity During Pregnancy and Subsequent Risk of Preeclampsia and Gestational Hypertension: A Case Control Study. <i>Maternal and Child Health Journal</i> , 2016, 20, 1193-1202.	1.5	31
16	Identification and functional analysis of glycemic trait loci in the China Health and Nutrition Survey. <i>PLoS Genetics</i> , 2018, 14, e1007275.	3.5	30
17	Genetic Studies of Leptin Concentrations Implicate Leptin in the Regulation of Early Adiposity. <i>Diabetes</i> , 2020, 69, 2806-2818.	0.6	26
18	Multi-SNP mediation intersection-union test. <i>Bioinformatics</i> , 2019, 35, 4724-4729.	4.1	23

#	ARTICLE	IF	CITATIONS
19	Genetic discovery and risk characterization in type 2 diabetes across diverse populations. <i>Human Genetics and Genomics Advances</i> , 2021, 2, 100029.	1.7	23
20	Exome-Derived Adiponectin-Associated Variants Implicate Obesity and Lipid Biology. <i>American Journal of Human Genetics</i> , 2019, 105, 15-28.	6.2	21
21	Identification of genetic effects underlying type 2 diabetes in South Asian and European populations. <i>Communications Biology</i> , 2022, 5, 329.	4.4	21
22	Genetic Predisposition to Dyslipidemia and Risk of Preeclampsia. <i>American Journal of Hypertension</i> , 2015, 28, 915-923.	2.0	19
23	Genetic Risk Score for Essential Hypertension and Risk of Preeclampsia. <i>American Journal of Hypertension</i> , 2016, 29, 17-24.	2.0	19
24	Influence of a loop electrosurgical excision procedure (LEEP) on levels of cytokines in cervical secretions. <i>Journal of Reproductive Immunology</i> , 2015, 109, 74-83.	1.9	18
25	Differential and shared genetic effects on kidney function between diabetic and non-diabetic individuals. <i>Communications Biology</i> , 2022, 5, .	4.4	17
26	Genetic variants associated with patent ductus arteriosus in extremely preterm infants. <i>Journal of Perinatology</i> , 2019, 39, 401-408.	2.0	16
27	Adiponectin GWAS loci harboring extensive allelic heterogeneity exhibit distinct molecular consequences. <i>PLoS Genetics</i> , 2020, 16, e1009019.	3.5	11
28	Allelic Heterogeneity at the CRP Locus Identified by Whole-Genome Sequencing in Multi-ancestry Cohorts. <i>American Journal of Human Genetics</i> , 2020, 106, 112-120.	6.2	9
29	Validation of birth certificate and maternal recall of events in labor and delivery with medical records in the Iowa health in pregnancy study. <i>BMC Pregnancy and Childbirth</i> , 2022, 22, 232.	2.4	8
30	Inferring Regulatory Networks From Mixed Observational Data Using Directed Acyclic Graphs. <i>Frontiers in Genetics</i> , 2020, 11, 8.	2.3	7
31	Genetic variation in CYB5R3 is associated with methemoglobin levels in preterm infants receiving nitric oxide therapy. <i>Pediatric Research</i> , 2015, 77, 472-476.	2.3	5
32	Progress in Defining the Genetic Contribution to Type 2 Diabetes in Individuals of East Asian Ancestry. <i>Current Diabetes Reports</i> , 2021, 21, 17.	4.2	5
33	Low Birth Weight and Risk of Later-Life Physical Disability in Women. <i>Journals of Gerontology - Series A Biological Sciences and Medical Sciences</i> , 2016, 72, glw134.	3.6	3
34	Birthweight and subsequent risk for thyroid and autoimmune conditions in postmenopausal women. <i>Journal of Developmental Origins of Health and Disease</i> , 2021, , 1-8.	1.4	2