Siddhartha P Kar

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3503434/publications.pdf

Version: 2024-02-01

66 papers

4,537 citations

28 h-index 61 g-index

78 all docs

78 does citations

times ranked

78

7398 citing authors

#	Article	IF	CITATIONS
1	Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94.	27.8	1,099
2	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691.	21.4	356
3	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778.	21.4	289
4	Intravenous autologous bone marrow mononuclear cells for ischemic stroke. Annals of Neurology, 2011, 70, 59-69.	5.3	259
5	Genetic predisposition to mosaic Y chromosome loss in blood. Nature, 2019, 575, 652-657.	27.8	198
6	Genome-Wide Meta-Analyses of Breast, Ovarian, and Prostate Cancer Association Studies Identify Multiple New Susceptibility Loci Shared by at Least Two Cancer Types. Cancer Discovery, 2016, 6, 1052-1067.	9.4	157
7	Breast cancer risk variants at 6q25 display different phenotype associations and regulate ESR1, RMND1 and CCDC170. Nature Genetics, 2016, 48, 374-386.	21.4	125
8	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. Nature Genetics, 2020, 52, 56-73.	21.4	120
9	Genome-wide analyses of 200,453 individuals yield new insights into the causes and consequences of clonal hematopoiesis. Nature Genetics, 2022, 54, 1155-1166.	21.4	109
10	Evidence that breast cancer risk at the 2q35 locus is mediated through IGFBP5 regulation. Nature Communications, 2014, 5, 4999.	12.8	105
11	Smoking, alcohol consumption, and cancer: A mendelian randomisation study in UK Biobank and international genetic consortia participants. PLoS Medicine, 2020, 17, e1003178.	8.4	103
12	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. Nature Communications, 2016, 7, 11375.	12.8	93
13	A transcriptome-wide association study of high-grade serous epithelial ovarian cancer identifies new susceptibility genes and splice variants. Nature Genetics, 2019, 51, 815-823.	21.4	89
14	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breast–ovarian cancer susceptibility locus. Nature Communications, 2016, 7, 12675.	12.8	78
15	Fine-Scale Mapping of the 5q11.2 Breast Cancer Locus Reveals at Least Three Independent Risk Variants Regulating MAP3K1. American Journal of Human Genetics, 2015, 96, 5-20.	6.2	76
16	Is Type 2 Diabetes Causally Associated With Cancer Risk? Evidence From a Two-Sample Mendelian Randomization Study. Diabetes, 2020, 69, 1588-1596.	0.6	75
17	Effects of tumour necrosis factor on cardiovascular disease and cancer: A two-sample Mendelian randomization study. EBioMedicine, 2020, 59, 102956.	6.1	74
18	Expression QTL-based analyses reveal candidate causal genes and loci across five tumor types. Human Molecular Genetics, 2014, 23, 5294-5302.	2.9	71

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19	Cis-eQTL analysis and functional validation of candidate susceptibility genes for high-grade serous ovarian cancer. Nature Communications, 2015, 6, 8234.	12.8	63
20	Identification of Novel Genetic Markers of Breast Cancer Survival. Journal of the National Cancer Institute, 2015, 107, .	6.3	56
21	Fineâ€scale mapping of 8q24 locus identifies multiple independent risk variants for breast cancer. International Journal of Cancer, 2016, 139, 1303-1317.	5.1	51
22	Genetic Data from Nearly 63,000 Women of European Descent Predicts DNA Methylation Biomarkers and Epithelial Ovarian Cancer Risk. Cancer Research, 2019, 79, 505-517.	0.9	49
23	Causal associations of thyroid function and dysfunction with overall, breast and thyroid cancer: A twoâ€sample Mendelian randomization study. International Journal of Cancer, 2020, 147, 1895-1903.	5.1	45
24	Germline whole exome sequencing and large-scale replication identifies FANCM as a likely high grade serous ovarian cancer susceptibility gene. Oncotarget, 2017, 8, 50930-50940.	1.8	43
25	Breast cancer risk factors and their effects on survival: a Mendelian randomisation study. BMC Medicine, 2020, 18, 327.	5.5	40
26	Polymorphisms in a Putative Enhancer at the 10q21.2 Breast Cancer Risk Locus Regulate NRBF2 Expression. American Journal of Human Genetics, 2015, 97, 22-34.	6.2	37
27	Insulinâ€like growth factorâ€1 and siteâ€specific cancers: A Mendelian randomization study. Cancer Medicine, 2020, 9, 6836-6842.	2.8	36
28	Body size and composition and risk of site-specific cancers in the UK Biobank and large international consortia: A mendelian randomisation study. PLoS Medicine, 2021, 18, e1003706.	8.4	35
29	Genome-wide Analysis Identifies Novel Loci Associated with Ovarian Cancer Outcomes: Findings from the Ovarian Cancer Association Consortium. Clinical Cancer Research, 2015, 21, 5264-5276.	7.0	33
30	Common Genetic Variation and Susceptibility to Ovarian Cancer: Current Insights and Future Directions. Cancer Epidemiology Biomarkers and Prevention, 2018, 27, 395-404.	2.5	33
31	Germline and Somatic Genetic Variants in the p53 Pathway Interact to Affect Cancer Risk, Progression, and Drug Response. Cancer Research, 2021, 81, 1667-1680.	0.9	32
32	Identification of independent association signals and putative functional variants for breast cancer risk through fine-scale mapping of the 12p11 locus. Breast Cancer Research, 2016, 18, 64.	5.0	31
33	eQTL Colocalization Analyses Identify NTN4 as a Candidate Breast Cancer Risk Gene. American Journal of Human Genetics, 2020, 107, 778-787.	6.2	29
34	Germline polymorphisms in an enhancer of <i>PSIP1</i> are associated with progression-free survival in epithelial ovarian cancer. Oncotarget, 2016, 7, 6353-6368.	1.8	29
35	Network-Based Integration of GWAS and Gene Expression Identifies a <i>HOX</i> -Centric Network Associated with Serous Ovarian Cancer Risk. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 1574-1584.	2.5	28
36	Genome-wide association studies identify susceptibility loci for epithelial ovarian cancer in east Asian women. Gynecologic Oncology, 2019, 153, 343-355.	1.4	28

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37	Sleep duration and risk of overall and 22 siteâ€specific cancers: A Mendelian randomization study. International Journal of Cancer, 2021, 148, 914-920.	5.1	28
38	A multi-level investigation of the genetic relationship between endometriosis and ovarian cancer histotypes. Cell Reports Medicine, 2022, 3, 100542.	6.5	26
39	Fine-Scale Mapping of the 4q24 Locus Identifies Two Independent Loci Associated with Breast Cancer Risk. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 1680-1691.	2.5	24
40	Enrichment of putative PAX8 target genes at serous epithelial ovarian cancer susceptibility loci. British Journal of Cancer, 2017, 116, 524-535.	6.4	23
41	Predicting the effect of statins on cancer risk using genetic variants from a Mendelian randomization study in the UK Biobank. ELife, 2020, 9, .	6.0	23
42	Addressing underlying causes of violence against doctors in India. Lancet, The, 2017, 389, 1979-1980.	13.7	21
43	Iron Status and Cancer Risk in UK Biobank: A Two-Sample Mendelian Randomization Study. Nutrients, 2020, 12, 526.	4.1	21
44	Fine scale mapping of the 17q22 breast cancer locus using dense SNPs, genotyped within the Collaborative Oncological Gene-Environment Study (COGs). Scientific Reports, 2016, 6, 32512.	3.3	19
45	Genetically proxied milk consumption and risk of colorectal, bladder, breast, and prostate cancer: a two-sample Mendelian randomization study. BMC Medicine, 2020, 18, 370.	5.5	19
46	Assessing the role of cortisol in cancer: a wide-ranged Mendelian randomisation study. British Journal of Cancer, 2021, 125, 1025-1029.	6.4	17
47	Selenium and cancer risk: Wideâ€angled Mendelian randomization analysis. International Journal of Cancer, 2022, 150, 1134-1140.	5.1	17
48	The association between weight at birth and breast cancer risk revisited using Mendelian randomisation. European Journal of Epidemiology, 2019, 34, 591-600.	5.7	16
49	The Relationship between Common Genetic Markers of Breast Cancer Risk and Chemotherapy-Induced Toxicity: A Case-Control Study. PLoS ONE, 2016, 11, e0158984.	2.5	15
50	Do sex hormones confound or mediate the effect of chronotype on breast and prostate cancer? A Mendelian randomization study. PLoS Genetics, 2022, 18, e1009887.	3.5	14
51	Serum Estradiol and 20 Site-Specific Cancers in Women: Mendelian Randomization Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e467-e474.	3.6	13
52	Genetically predicted circulating protein biomarkers and ovarian cancer risk. Gynecologic Oncology, 2021, 160, 506-513.	1.4	12
53	Genetically predicted circulating B vitamins in relation to digestive system cancers. British Journal of Cancer, 2021, 124, 1997-2003.	6.4	8
54	Genetic Analysis of Lung Cancer and the Germline Impact on Somatic Mutation Burden. Journal of the National Cancer Institute, 2022, 114, 1159-1166.	6.3	8

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55	Pleiotropy-guided transcriptome imputation from normal and tumor tissues identifies candidate susceptibility genes for breast and ovarian cancer. Human Genetics and Genomics Advances, 2021, 2, 100042.	1.7	6
56	Large-scale cross-cancer fine-mapping of the 5p15.33 region reveals multiple independent signals. Human Genetics and Genomics Advances, 2021, 2, 100041.	1.7	6
57	Personalized therapy for pancreatic cancer: Myth or reality in 2010?. Journal of Gastrointestinal Oncology, 2010, 1, 24-33.	1.4	4
58	Rho GTPase gene expression and breast cancer risk: a Mendelian randomization analysis. Scientific Reports, 2022, 12, 1463.	3.3	4
59	Body mass index and the association between low-density lipoprotein cholesterol as predicted by HMGCR genetic variants and breast cancer risk. International Journal of Epidemiology, 2019, 48, 1727-1730.	1.9	3
60	Assessing the protective role of allergic disease in gastrointestinal tract cancers using Mendelian randomization analysis. Allergy: European Journal of Allergy and Clinical Immunology, 2021, 76, 1559-1562.	5.7	1
61	chromMAGMA: regulatory element-centric interrogation of risk variants. Life Science Alliance, 2022, 5, e202201446.	2.8	1
62	Title is missing!. , 2020, 17, e1003178.		0
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