## Gadi Rennert

## List of Publications by Year in descending order

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9786 14208 20,490 287 73 128 citations h-index g-index papers 306 306 306 25152 times ranked docs citations citing authors all docs

#	Article	IF	CITATIONS
1	Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94.	27.8	1,099
2	Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. American Journal of Human Genetics, 2019, 104, 21-34.	6.2	711
3	Statins and the Risk of Colorectal Cancer. New England Journal of Medicine, 2005, 352, 2184-2192.	27.0	706
4	Network modeling links breast cancer susceptibility and centrosome dysfunction. Nature Genetics, 2007, 39, 1338-1349.	21.4	602
5	Genome-wide association scan identifies a colorectal cancer susceptibility locus on 11q23 and replicates risk loci at 8q24 and 18q21. Nature Genetics, 2008, 40, 631-637.	21.4	542
6	Large-scale association analysis identifies new lung cancer susceptibility loci and heterogeneity in genetic susceptibility across histological subtypes. Nature Genetics, 2017, 49, 1126-1132.	21.4	472
7	Breast Cancer Risk Following Bilateral Oophorectomy in BRCA1 and BRCA2 Mutation Carriers: An International Case-Control Study. Journal of Clinical Oncology, 2005, 23, 7491-7496.	1.6	408
8	Discovery of common and rare genetic risk variants for colorectal cancer. Nature Genetics, 2019, 51, 76-87.	21.4	377
9	European guidelines for quality assurance in colorectal cancer screening and diagnosis: Overview and introduction to the full Supplement publication. Endoscopy, 2012, 45, 51-59.	1.8	356
10	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691.	21.4	356
11	Oral Contraceptives and the Risk of Breast Cancer in BRCA1 and BRCA2 Mutation Carriers. Journal of the National Cancer Institute, 2002, 94, 1773-1779.	6.3	318
12	A locus on 19p13 modifies risk of breast cancer in BRCA1 mutation carriers and is associated with hormone receptor–negative breast cancer in the general population. Nature Genetics, 2010, 42, 885-892.	21.4	309
13	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778.	21.4	289
14	Clinical Outcomes of Breast Cancer in Carriers of <i>BRCA1 </i> BRCA2 Mutations. New England Journal of Medicine, 2007, 357, 115-123.	27.0	268
15	Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses. Nature Genetics, 2020, 52, 572-581.	21.4	265
16	Common Breast Cancer-Predisposition Alleles Are Associated with Breast Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. American Journal of Human Genetics, 2008, 82, 937-948.	6.2	257
17	Genome-Wide Association Study in BRCA1 Mutation Carriers Identifies Novel Loci Associated with Breast and Ovarian Cancer Risk. PLoS Genetics, 2013, 9, e1003212.	3.5	244
18	Determining Risk of Colorectal Cancer and Starting Age of Screening Based on Lifestyle, Environmental, and Genetic Factors. Gastroenterology, 2018, 154, 2152-2164.e19.	1.3	226

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19	Phenotype of Microsatellite Unstable Colorectal Carcinomas: Well-Differentiated and Focally Mucinous Tumors and the Absence of Dirty Necrosis Correlate With Microsatellite Instability. American Journal of Surgical Pathology, 2003, 27, 563-570.	3.7	225
20	Pathologic Predictors of Microsatellite Instability in Colorectal Cancer. American Journal of Surgical Pathology, 2009, 33, 126-133.	3.7	222
21	Identification of six new susceptibility loci for invasive epithelial ovarian cancer. Nature Genetics, 2015, 47, 164-171.	21.4	221
22	RAD51 135Gâ†'C Modifies Breast Cancer Risk among BRCA2 Mutation Carriers: Results from a Combined Analysis of 19 Studies. American Journal of Human Genetics, 2007, 81, 1186-1200.	6.2	217
23	Targeted Prostate Cancer Screening in BRCA1 and BRCA2 Mutation Carriers: Results from the Initial Screening Round of the IMPACT Study. European Urology, 2014, 66, 489-499.	1.9	195
24	Physical activity and risks of breast and colorectal cancer: a Mendelian randomisation analysis. Nature Communications, 2020, 11, 597.	12.8	193
25	A transcriptome-wide association study of 229,000 women identifies new candidate susceptibility genes for breast cancer. Nature Genetics, 2018, 50, 968-978.	21.4	184
26	Meta-analysis of new genome-wide association studies of colorectal cancer risk. Human Genetics, 2012, 131, 217-234.	3.8	183
27	Genetic insights into biological mechanisms governing human ovarian ageing. Nature, 2021, 596, 393-397.	27.8	183
28	Organ-Specific Molecular Classification of Primary Lung, Colon, and Ovarian Adenocarcinomas Using Gene Expression Profiles. American Journal of Pathology, 2001, 159, 1231-1238.	3.8	180
29	Hormone Therapy and the Risk of Breast Cancer in BRCA1 Mutation Carriers. Journal of the National Cancer Institute, 2008, 100, 1361-1367.	6.3	179
30	BLM Heterozygosity and the Risk of Colorectal Cancer. Science, 2002, 297, 2013-2013.	12.6	174
31	Clinical Phenotype of Families with Longevity. Journal of the American Geriatrics Society, 2004, 52, 274-277.	2.6	174
32	Replication of Lung Cancer Susceptibility Loci at Chromosomes 15q25, 5p15, and 6p21: A Pooled Analysis From the International Lung Cancer Consortium. Journal of the National Cancer Institute, 2010, 102, 959-971.	6.3	174
33	Use of Bisphosphonates and Risk of Postmenopausal Breast Cancer. Journal of Clinical Oncology, 2010, 28, 3577-3581.	1.6	172
34	Common Breast Cancer Susceptibility Alleles and the Risk of Breast Cancer for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Implications for Risk Prediction. Cancer Research, 2010, 70, 9742-9754.	0.9	169
35	Tumor-Infiltrating Lymphocytes, Crohn's-Like Lymphoid Reaction, and Survival From Colorectal Cancer. Journal of the National Cancer Institute, 2016, 108, .	6.3	162
36	Effect of pregnancy as a risk factor for breast cancer in <i>BRCA1</i> li>BRCA2mutation carriers. International Journal of Cancer, 2005, 117, 988-991.	5.1	152

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37	Beyond aspirinâ€"cancer prevention with statins, metformin and bisphosphonates. Nature Reviews Clinical Oncology, 2013, 10, 625-642.	27.6	150
38	Factor XI deficiency is associated with lower risk for cardiovascular and venous thromboembolism events. Blood, 2017, 129, 1210-1215.	1.4	149
39	Interim Results from the IMPACT Study: Evidence for Prostate-specific Antigen Screening in BRCA2 Mutation Carriers. European Urology, 2019, 76, 831-842.	1.9	148
40	The underrecognized progressive nature of N370S Gaucher disease and assessment of cancer risk in 403 patients. American Journal of Hematology, 2009, 84, 208-214.	4.1	146
41	Global implementation of genomic medicine: We are not alone. Science Translational Medicine, 2015, 7, 290ps13.	12.4	146
42	Increased risk of lung cancer in individuals with a family history of the disease: A pooled analysis from the International Lung Cancer Consortium. European Journal of Cancer, 2012, 48, 1957-1968.	2.8	143
43	<i>MRE11</i> Deficiency Increases Sensitivity to Poly(ADP-ribose) Polymerase Inhibition in Microsatellite Unstable Colorectal Cancers. Cancer Research, 2011, 71, 2632-2642.	0.9	140
44	Genome-wide association study of colorectal cancer identifies six new susceptibility loci. Nature Communications, 2015, 6, 7138.	12.8	138
45	Novel Common Genetic Susceptibility Loci for Colorectal Cancer. Journal of the National Cancer Institute, 2019, 111, 146-157.	6.3	129
46	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
47	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. Nature Genetics, 2020, 52, 56-73.	21.4	120
48	A Novel Founder Mutation in the RNASEL Gene, 471delAAAG, Is Associated with Prostate Cancer in Ashkenazi Jews. American Journal of Human Genetics, 2002, 71, 981-984.	6.2	113
49	Usefulness of CHADS2 and CHA2DS2-VASc Scores in the Prediction of New-Onset Atrial Fibrillation: A Population-Based Study. American Journal of Medicine, 2016, 129, 843-849.	1.5	111
50	Large-Scale Genome-Wide Association Study of East Asians Identifies Loci Associated With Risk for Colorectal Cancer. Gastroenterology, 2019, 156, 1455-1466.	1.3	111
51	Cumulative Burden of Colorectal Cancer–Associated Genetic Variants Is More Strongly Associated With Early-Onset vs Late-Onset Cancer. Gastroenterology, 2020, 158, 1274-1286.e12.	1.3	110
52	Higher risk of venous thrombosis associated with drospirenone-containing oral contraceptives: a population-based cohort study. Cmaj, 2011, 183, E1319-E1325.	2.0	108
53	The Relationship Between Serum 25(OH)D and Parathyroid Hormone Levels. American Journal of Medicine, 2011, 124, 1165-1170.	1.5	105
54	Identification of a BRCA2-Specific Modifier Locus at 6p24 Related to Breast Cancer Risk. PLoS Genetics, 2013, 9, e1003173.	3.5	105

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55	Accurate classification of <i>MLH1/MSH2</i> missense variants with multivariate analysis of protein polymorphisms-mismatch repair (MAPP-MMR). Human Mutation, 2008, 29, 852-860.	2.5	101
56	Common variants in LSP1, 2q35 and 8q24 and breast cancer risk for BRCA1 and BRCA2 mutation carriers. Human Molecular Genetics, 2009, 18, 4442-4456.	2.9	99
57	BRCA germline mutations in Jewish women with uterine serous papillary carcinoma. Gynecologic Oncology, 2004, 92, 521-524.	1.4	98
58	Use of Bisphosphonates and Reduced Risk of Colorectal Cancer. Journal of Clinical Oncology, 2011, 29, 1146-1150.	1.6	96
59	Hypothyroidism Is a Risk Factor for New-Onset Diabetes: A Cohort Study. Diabetes Care, 2015, 38, 1657-1664.	8.6	93
60	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. Nature Communications, 2016, 7, 11375.	12.8	93
61	The MLH1 D132H variant is associated with susceptibility to sporadic colorectal cancer. Nature Genetics, 2004, 36, 694-699.	21.4	92
62	Interplay between BRCA1 and RHAMM Regulates Epithelial Apicobasal Polarization and May Influence Risk of Breast Cancer. PLoS Biology, 2011, 9, e1001199.	5.6	91
63	Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. Nature Communications, 2019, 10, 1741.	12.8	90
64	Circulating Levels of Insulin-like Growth Factor 1 and Insulin-like Growth Factor Binding Protein 3 Associate With Risk of Colorectal Cancer Based on Serologic and Mendelian Randomization Analyses. Gastroenterology, 2020, 158, 1300-1312.e20.	1.3	90
65	Shared heritability and functional enrichment across six solid cancers. Nature Communications, 2019, 10, 431.	12.8	88
66	The effect of statins on risk and survival of gynecological malignancies. Gynecologic Oncology, 2013, 130, 615-619.	1.4	87
67	From the Bench to Public Health. American Journal of Preventive Medicine, 2014, 46, 273-280.	3.0	87
68	Cancer-driving H3G34V/R/D mutations block H3K36 methylation and H3K36me3–MutSα interaction. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 9598-9603.	7.1	87
69	Common Genetic Variants and Modification of Penetrance of BRCA2-Associated Breast Cancer. PLoS Genetics, 2010, 6, e1001183.	3.5	85
70	BRCA1 and BRCA2 Founder Mutations and the Risk of Colorectal Cancer. Journal of the National Cancer Institute, 2004, 96, 15-21.	6.3	83
71	Use of Hormone Replacement Therapy and the Risk of Colorectal Cancer. Journal of Clinical Oncology, 2009, 27, 4542-4547.	1.6	83
72	Polygenic risk scores and breast and epithelial ovarian cancer risks for carriers of BRCA1 and BRCA2 pathogenic variants. Genetics in Medicine, 2020, 22, 1653-1666.	2.4	82

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73	Associations of obesity and circulating insulin and glucose with breast cancer risk: a Mendelian randomization analysis. International Journal of Epidemiology, 2019, 48, 795-806.	1.9	81
74	Obesity, metabolic factors and risk of different histological types of lung cancer: A Mendelian randomization study. PLoS ONE, 2017, 12, e0177875.	2.5	79
75	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breast–ovarian cancer susceptibility locus. Nature Communications, 2016, 7, 12675.	12.8	78
76	BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. Journal of the National Cancer Institute, 2016, 108, djv315.	<b>6.</b> 3	77
77	β2â€adrenoceptor agonists and antagonists and risk of Parkinson's disease. Movement Disorders, 2018, 33, 1465-1471.	3.9	76
78	Adiposity, metabolites, and colorectal cancer risk: Mendelian randomization study. BMC Medicine, 2020, 18, 396.	5.5	76
79	Catheter ablation of atrial fibrillation is associated with reduced risk of stroke and mortality: A propensity score–matched analysis. Heart Rhythm, 2017, 14, 635-642.	0.7	74
80	Causal relationships between body mass index, smoking and lung cancer: Univariable and multivariable Mendelian randomization. International Journal of Cancer, 2021, 148, 1077-1086.	5.1	73
81	Clinical implications of <i>UGT1A1</i> *28 genotype testing in colorectal cancer patients. Cancer, 2011, 117, 3156-3162.	4.1	72
82	Genetic variation in 8q24 associated with risk of colorectal cancer. Cancer Biology and Therapy, 2007, 6, 1143-1147.	3.4	70
83	Gene Expression Differences between Colon and Rectum Tumors. Clinical Cancer Research, 2011, 17, 7303-7312.	7.0	69
84	Coffee Consumption and the Risk of Colorectal Cancer. Cancer Epidemiology Biomarkers and Prevention, 2016, 25, 634-639.	2.5	68
85	Genetic Variation in 3-Hydroxy-3-Methylglutaryl CoA Reductase Modifies the Chemopreventive Activity of Statins for Colorectal Cancer. Cancer Prevention Research, 2010, 3, 597-603.	1.5	66
86	Accurate Molecular Classification of Human Cancers Based on Gene Expression Using a Simple Classifier with a Pathological Tree-Based Framework. American Journal of Pathology, 2003, 163, 1985-1995.	3.8	64
87	Smoking, Gender, and Ethnicity Predict Somatic <i>BRAF</i> Mutations in Colorectal Cancer. Cancer Epidemiology Biomarkers and Prevention, 2010, 19, 838-843.	2.5	64
88	<i>MutYH</i> mutation carriers have increased breast cancer risk. Cancer, 2012, 118, 1989-1993.	4.1	63
89	Survival of first and second primary breast cancer. Cancer, 1993, 71, 172-176.	4.1	62
90	Genetic Anthropology of the Colorectal Cancer–Susceptibility Allele APC I1307K: Evidence of Genetic Drift within the Ashkenazim. American Journal of Human Genetics, 2003, 73, 1250-1260.	6.2	61

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91	Human papillomavirus is not associated with colorectal cancer in a large international study. Cancer Causes and Control, 2010, 21, 737-743.	1.8	60
92	Identification of susceptibility pathways for the role of chromosome 15q25.1 in modifying lung cancer risk. Nature Communications, 2018, 9, 3221.	12.8	60
93	Calcium supplementation provides an extended window of opportunity for bone mass accretion after menarche. American Journal of Clinical Nutrition, 2003, 78, 993-998.	4.7	58
94	Gene Expression Patterns in Mismatch Repair-Deficient Colorectal Cancers Highlight the Potential Therapeutic Role of Inhibitors of the Phosphatidylinositol 3-Kinase-AKT-Mammalian Target of Rapamycin Pathway. Clinical Cancer Research, 2009, 15, 2829-2839.	<b>7.</b> O	57
95	Associations of common breast cancer susceptibility alleles with risk of breast cancer subtypes in BRCA1 and BRCA2 mutation carriers. Breast Cancer Research, 2014, 16, 3416.	5.0	57
96	Factors influencing ovulation and the risk of ovarian cancer in <scp><i>BRCA1</i></scp> and <scp><i>BRCA2</i></scp> mutation carriers. International Journal of Cancer, 2015, 137, 1136-1146.	5.1	56
97	A Transcriptome-Wide Association Study Among 97,898 Women to Identify Candidate Susceptibility Genes for Epithelial Ovarian Cancer Risk. Cancer Research, 2018, 78, 5419-5430.	0.9	54
98	N-Methylpurine DNA Glycosylase and OGG1 DNA Repair Activities: Opposite Associations With Lung Cancer Risk. Journal of the National Cancer Institute, 2012, 104, 1765-1769.	6.3	53
99	Genome-wide association study of germline variants and breast cancer-specific mortality. British Journal of Cancer, 2019, 120, 647-657.	6.4	52
100	A Case–Control Study of Levothyroxine and the Risk of Colorectal Cancer. Journal of the National Cancer Institute, 2010, 102, 568-572.	6.3	51
101	Assessing Lung Cancer Absolute Risk Trajectory Based on a Polygenic Risk Model. Cancer Research, 2021, 81, 1607-1615.	0.9	50
102	Risk of colorectal cancer in selfâ€reported inflammatory bowel disease and modification of risk by statin and NSAID use. Cancer, 2011, 117, 1640-1648.	4.1	49
103	Associated Links Among Smoking, Chronic Obstructive Pulmonary Disease, and Small Cell Lung Cancer: A Pooled Analysis in the International Lung Cancer Consortium. EBioMedicine, 2015, 2, 1677-1685.	6.1	49
104	Oral Bisphosphonates and Improved Survival of Breast Cancer. Clinical Cancer Research, 2017, 23, 1684-1689.	7.0	48
105	A prospective prostate cancer screening programme for men with pathogenic variants in mismatch repair genes (IMPACT): initial results from an international prospective study. Lancet Oncology, The, 2021, 22, 1618-1631.	10.7	48
106	Aspirin and NSAID use and lung cancer risk: a pooled analysis in the International Lung Cancer Consortium (ILCCO). Cancer Causes and Control, 2011, 22, 1709-1720.	1.8	47
107	Age at first birth and the risk of breast cancer in BRCA1 and BRCA2 mutation carriers. Breast Cancer Research and Treatment, 2007, 105, 221-228.	2.5	45
108	Combined Associations of a Polygenic Risk Score and Classical Risk Factors With Breast Cancer Risk. Journal of the National Cancer Institute, 2021, 113, 329-337.	6.3	45

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109	The relationship between obesity and the increase in serum 25(OH)D levels in response to vitamin D supplementation. Osteoporosis International, 2013, 24, 1447-1454.	3.1	44
110	Genetic variant predictors of gene expression provide new insight into risk of colorectal cancer. Human Genetics, 2019, 138, 307-326.	3.8	44
111	Genetic architectures of proximal and distal colorectal cancer are partly distinct. Gut, 2021, 70, 1325-1334.	12.1	44
112	The Risk of All-Cause Mortality Is Inversely Related to Serum 25(OH)D Levels. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 2792-2798.	3.6	43
113	The Association Between Red Cell Distribution Width and Stroke in Patients with Atrial Fibrillation. American Journal of Medicine, 2015, 128, 192.e11-192.e18.	1.5	43
114	Fine mapping of MHC region in lung cancer highlights independent susceptibility loci by ethnicity. Nature Communications, 2018, 9, 3927.	12.8	43
115	Sustained effect of short-term calcium supplementation on bone mass in adolescent girls with low calcium intake. American Journal of Clinical Nutrition, 2005, 81, 168-174.	4.7	42
116	The association between obesity and urinary tract infection. European Journal of Internal Medicine, 2013, 24, 127-131.	2.2	42
117	Neutrophil to lymphocyte ratio and risk of a first episode of stroke in patients with atrial fibrillation: a cohort study. Journal of Thrombosis and Haemostasis, 2015, 13, 1971-1979.	3.8	42
118	A Recurrent <i>ERCC3</i> Truncating Mutation Confers Moderate Risk for Breast Cancer. Cancer Discovery, 2016, 6, 1267-1275.	9.4	41
119	A comparison of the clinical characteristics of first and second primary head and neck cancers. A population-based study. Cancer, 1991, 68, 189-192.	4.1	40
120	CHA2DS2-VASc Score Is Directly Associated with the Risk of Pulmonary Embolism in Patients with Atrial Fibrillation. American Journal of Medicine, 2014, 127, 45-52.	1.5	40
121	Glycated hemoglobin and risk of first episode stroke in diabetic patients with atrial fibrillation: A cohort study. Heart Rhythm, 2015, 12, 886-892.	0.7	40
122	Association of Diabetes and Glycated Hemoglobin With the Risk of Intracerebral Hemorrhage: A Population-Based Cohort Study. Diabetes Care, 2019, 42, 682-688.	8.6	39
123	Cancer Risk After Radioactive Iodine Treatment for Hyperthyroidism: A Cohort Study. Thyroid, 2020, 30, 243-250.	4.5	39
124	Breast Cancer Polygenic Risk Score and Contralateral Breast Cancer Risk. American Journal of Human Genetics, 2020, 107, 837-848.	6.2	39
125	Nongenetic Determinants of Risk forÂEarly-Onset Colorectal Cancer. JNCI Cancer Spectrum, 2021, 5, pkab029.	2.9	39
126	Treatment of infertility does not increase the risk of ovarian cancer among women with a BRCA1 or BRCA2 mutation. Fertility and Sterility, 2016, 105, 781-785.	1.0	38

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127	Fecal DNA Biomarkers for the Detection of Colorectal Neoplasia: Attractive, but Is It Feasible?. Journal of the National Cancer Institute, 2005, 97, 1107-1109.	6.3	37
128	Serum 25(OH)D concentrations in sunny Israel. Osteoporosis International, 2012, 23, 687-694.	3.1	36
129	Identifying Novel Susceptibility Genes for Colorectal Cancer Risk From a Transcriptome-Wide Association Study of 125,478 Subjects. Gastroenterology, 2021, 160, 1164-1178.e6.	1.3	36
130	Polymorphisms in Alcohol Metabolism Genes ADH1B and ALDH2, Alcohol Consumption and Colorectal Cancer. PLoS ONE, 2013, 8, e80158.	2.5	36
131	Alcohol and lung cancer risk among never smokers: A pooled analysis from the international lung cancer consortium and the SYNERGY study. International Journal of Cancer, 2017, 140, 1976-1984.	5.1	35
132	Mendelian randomization analysis of C-reactive protein on colorectal cancer risk. International Journal of Epidemiology, 2019, 48, 767-780.	1.9	35
133	Assessing Associations between the AURKA-HMMR-TPX2-TUBG1 Functional Module and Breast Cancer Risk in BRCA1/2 Mutation Carriers. PLoS ONE, 2015, 10, e0120020.	2.5	34
134	Genetic polymorphisms in fatty acid metabolism genes and colorectal cancer. Mutagenesis, 2012, 27, 169-176.	2.6	33
135	Association of statin use with spontaneous intracerebral hemorrhage. Neurology, 2018, 91, e400-e409.	1.1	33
136	Transcriptomeâ€wide association study reveals candidate causal genes for lung cancer. International Journal of Cancer, 2020, 146, 1862-1878.	5.1	33
137	Size of acute myocardial infarcts in patients with diabetes mellitus. American Journal of Cardiology, 1985, 55, 1629-1630.	1.6	32
138	The effect of bisphosphonates on the risk of endometrial and ovarian malignancies. Gynecologic Oncology, 2014, 133, 309-313.	1.4	32
139	Mendelian Randomization and mediation analysis of leukocyte telomere length and risk of lung and head and neck cancers. International Journal of Epidemiology, 2019, 48, 751-766.	1.9	32
140	Transcriptomeâ€wide association study of breast cancer risk by estrogenâ€receptor status. Genetic Epidemiology, 2020, 44, 442-468.	1.3	32
141	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
142	Protein-altering germline mutations implicate novel genes related to lung cancer development. Nature Communications, 2020, 11, 2220.	12.8	31
143	A novel colorectal cancer risk locus at 4q32.2 identified from an international genome-wide association study. Carcinogenesis, 2014, 35, 2512-2519.	2.8	30
144	A network analysis to identify mediators of germline-driven differences in breast cancer prognosis. Nature Communications, 2020, 11, 312.	12.8	30

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145	Association of atrial fibrillation and cancer: Analysis from two large population-based case-control studies. PLoS ONE, 2018, 13, e0190324.	2.5	30
146	CDX2 Polymorphisms, RNA Expression, and Risk of Colorectal Cancer. Cancer Research, 2005, 65, 5488-5492.	0.9	29
147	Genome-wide interaction study of smoking behavior and non-small cell lung cancer risk in Caucasian population. Carcinogenesis, 2018, 39, 336-346.	2.8	29
148	The gradient in mammography screening behavior: a lifestyle marker. Social Science and Medicine, 1999, 48, 1281-1290.	3.8	28
149	K-ras mutations in sporadic colorectal tumors in Israel: unusual high frequency of codon 13 mutations and evidence for nonhomogeneous representation of mutation subtypes. Digestive Diseases and Sciences, 2002, 47, 1073-1079.	2.3	28
150	Unraveling Seasonality in Population Averages: An Examination of Seasonal Variation in Glucose Levels in Diabetes Patients Using a Large Population-based Data Set. Chronobiology International, 2011, 28, 352-360.	2.0	28
151	A genome-wide association study for colorectal cancer identifies a risk locus in 14q23.1. Human Genetics, 2015, 134, 1249-1262.	3.8	28
152	Tyrosine kinase-targeting drugs-associated heart failure. British Journal of Cancer, 2017, 116, 1366-1373.	6.4	28
153	Circulating bilirubin levels and risk of colorectal cancer: serological and Mendelian randomization analyses. BMC Medicine, 2020, 18, 229.	5.5	28
154	Association of PHB 1630 C>T and MTHFR 677 C>T polymorphisms with breast and ovarian cancer risk in BRCA1/2 mutation carriers: results from a multicenter study. British Journal of Cancer, 2012, 106, 2016-2024.	6.4	27
155	Genetic modifiers of radon-induced lung cancer risk: a genome-wide interaction study in former uranium miners. International Archives of Occupational and Environmental Health, 2018, 91, 937-950.	2.3	27
156	Lung Cancer Risk in Never-Smokers of European Descent is Associated With Genetic Variation in the 5p15.33 TERT-CLPTM1Ll Region. Journal of Thoracic Oncology, 2019, 14, 1360-1369.	1.1	27
157	Genetically predicted circulating concentrations of micronutrients and risk of colorectal cancer among individuals of European descent: a Mendelian randomization study. American Journal of Clinical Nutrition, $2021, 113, 1490-1502$ .	4.7	27
158	The public prefers fecal occult blood test over colonoscopy for colorectal cancer screening. European Journal of Cancer Prevention, 2008, 17, 430-437.	1.3	26
159	FOXP3 germline polymorphisms are not associated with risk of breast cancer. Cancer Genetics and Cytogenetics, 2009, 190, 40-42.	1.0	26
160	Low Integrated DNA Repair Score and Lung Cancer Risk. Cancer Prevention Research, 2014, 7, 398-406.	1.5	26
161	An original phylogenetic approach identified mitochondrial haplogroup T1a1 as inversely associated with breast cancer risk in BRCA2 mutation carriers. Breast Cancer Research, 2015, 17, 61.	5.0	26
162	Mendelian Randomization of Circulating Polyunsaturated Fatty Acids and Colorectal Cancer Risk. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 860-870.	2.5	26

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163	Genetic interaction analysis among oncogenesis-related genes revealed novel genes and networks in lung cancer development. Oncotarget, 2019, 10, 1760-1774.	1.8	25
164	Lymphocytic infiltration in stage II microsatellite stable colorectal tumors: A retrospective prognosis biomarker analysis. PLoS Medicine, 2020, 17, e1003292.	8.4	25
165	Implications of Russian Immigration on Mortality Patterns in Israel. International Journal of Epidemiology, 1994, 23, 751-756.	1.9	24
166	Quality Control in a National Program for the Early Detection of Breast Cancer. Women's Health Issues, 2008, 18, 110-117.	2.0	24
167	Development of APE1 enzymatic DNA repair assays: low APE1 activity is associated with increase lung cancer risk. Carcinogenesis, 2015, 36, 982-991.	2.8	24
168	Inherited Rare, Deleterious Variants in ATM Increase Lung Adenocarcinoma Risk. Journal of Thoracic Oncology, 2020, 15, 1871-1879.	1.1	24
169	BRCA2 germline mutation in a woman with uterine serous papillary carcinoma—Case report. Gynecologic Oncology, 2005, 99, 486-488.	1.4	23
170	The role of CHADS2 and CHA2DS2â€VASc scores in the prediction of stroke in individuals without atrial fibrillation: a populationâ€based study. Journal of Thrombosis and Haemostasis, 2016, 14, 1155-1162.	3.8	23
171	Immune-mediated genetic pathways resulting in pulmonary function impairment increase lung cancer susceptibility. Nature Communications, 2020, $11$ , $27$ .	12.8	23
172	Candidate Genetic Modifiers for Breast and Ovarian Cancer Risk in <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 308-316.	2.5	22
173	Alcohol consumption and lung cancer risk: A pooled analysis from the International Lung Cancer Consortium and the SYNERGY study. Cancer Epidemiology, 2019, 58, 25-32.	1.9	22
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