

Kenneth Offit

List of Publications by Year in descending order

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Version: 2024-02-01

381
papers

39,831
citations

1704

104
h-index

3486

182
g-index

395
all docs

395
docs citations

395
times ranked

35172
citing authors

#	ARTICLE	IF	CITATIONS
1	Risk-Reducing Salpingo-oophorectomy in Women with a <i>BRCA1</i> or <i>BRCA2</i> Mutation. New England Journal of Medicine, 2002, 346, 1609-1615.	27.0	1,363
2	Inherited DNA-Repair Gene Mutations in Men with Metastatic Prostate Cancer. New England Journal of Medicine, 2016, 375, 443-453.	27.0	1,205
3	Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. American Journal of Human Genetics, 2019, 104, 21-34.	6.2	711
4	Integrative clinical genomics of metastatic cancer. Nature, 2017, 548, 297-303.	27.8	685
5	Germline mutations in BAP1 predispose to melanocytic tumors. Nature Genetics, 2011, 43, 1018-1021.	21.4	662
6	Familial colorectal cancer in Ashkenazim due to a hypermutable tract in APC. Nature Genetics, 1997, 17, 79-83.	21.4	630
7	Network modeling links breast cancer susceptibility and centrosome dysfunction. Nature Genetics, 2007, 39, 1338-1349.	21.4	602
8	Hereditary Cancer Predisposition Syndromes. Journal of Clinical Oncology, 2005, 23, 276-292.	1.6	534
9	Risk-Reducing Salpingo-Oophorectomy for the Prevention of BRCA1- and BRCA2-Associated Breast and Gynecologic Cancer: A Multicenter, Prospective Study. Journal of Clinical Oncology, 2008, 26, 1331-1337.	1.6	522
10	Pathology of Breast and Ovarian Cancers among <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Results from the Consortium of Investigators of Modifiers of <i>BRCA1/2</i> (CIMBA). Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 134-147.	2.5	513
11	Genetic/Familial High-Risk Assessment: Breast, Ovarian, and Pancreatic, Version 2.2021, NCCN Clinical Practice Guidelines in Oncology. Journal of the National Comprehensive Cancer Network: JNCCN, 2021, 19, 77-102.	4.9	498
12	Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. Nature Genetics, 2013, 45, 371-384.	21.4	493
13	Prediction of Breast Cancer Risk Based on Profiling With Common Genetic Variants. Journal of the National Cancer Institute, 2015, 107, .	6.3	428
14	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
15	NCCN Guidelines Insights: Genetic/Familial High-Risk Assessment: Breast and Ovarian, Version 2.2017. Journal of the National Comprehensive Cancer Network: JNCCN, 2017, 15, 9-20.	4.9	408
16	Microsatellite Instability Is Associated With the Presence of Lynch Syndrome Pan-Cancer. Journal of Clinical Oncology, 2019, 37, 286-295.	1.6	397
17	Outcome of Preventive Surgery and Screening for Breast and Ovarian Cancer in <i>BRCA</i> Mutation Carriers. Journal of Clinical Oncology, 2002, 20, 1260-1268.	1.6	395
18	Association of Type and Location of <i>BRCA1</i> and <i>BRCA2</i> Mutations With Risk of Breast and Ovarian Cancer. JAMA - Journal of the American Medical Association, 2015, 313, 1347.	7.4	390

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19	American Society of Clinical Oncology Policy Statement Update: Genetic and Genomic Testing for Cancer Susceptibility. <i>Journal of Clinical Oncology</i> , 2010, 28, 893-901.	1.6	389
20	Discovery of common and rare genetic risk variants for colorectal cancer. <i>Nature Genetics</i> , 2019, 51, 76-87.	21.4	377
21	Rearrangement of the bcl-6 Gene as a Prognostic Marker in Diffuse Large-Cell Lymphoma. <i>New England Journal of Medicine</i> , 1994, 331, 74-80.	27.0	375
22	The carrier frequency of the BRCA2 6174delT mutation among Ashkenazi Jewish individuals is approximately 1%. <i>Nature Genetics</i> , 1996, 14, 188-190.	21.4	375
23	Cancer therapy shapes the fitness landscape of clonal hematopoiesis. <i>Nature Genetics</i> , 2020, 52, 1219-1226.	21.4	367
24	Mutation Detection in Patients With Advanced Cancer by Universal Sequencing of Cancer-Related Genes in Tumor and Normal DNA vs Guideline-Based Germline Testing. <i>JAMA - Journal of the American Medical Association</i> , 2017, 318, 825.	7.4	366
25	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. <i>Nature Genetics</i> , 2017, 49, 680-691.	21.4	356
26	Prediction of Germline Mutations and Cancer Risk in the Lynch Syndrome. <i>JAMA - Journal of the American Medical Association</i> , 2006, 296, 1479.	7.4	328
27	A germline JAK2 SNP is associated with predisposition to the development of JAK2V617F-positive myeloproliferative neoplasms. <i>Nature Genetics</i> , 2009, 41, 455-459.	21.4	322
28	Oral Contraceptives and the Risk of Breast Cancer in BRCA1 and BRCA2 Mutation Carriers. <i>Journal of the National Cancer Institute</i> , 2002, 94, 1773-1779.	6.3	318
29	NCCN Guidelines Insights: Genetic/Familial High-Risk Assessment: Breast, Ovarian, and Pancreatic, Version 1.2020. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2020, 18, 380-391.	4.9	314
30	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. <i>Nature Genetics</i> , 2010, 42, 885-892.	21.4	309
31	Chromosomal and Gene Amplification in Diffuse Large B-Cell Lymphoma. <i>Blood</i> , 1998, 92, 234-240.	1.4	308
32	Two Decades After <i>BRCA</i> Setting Paradigms in Personalized Cancer Care and Prevention. <i>Science</i> , 2014, 343, 1466-1470.	12.6	300
33	Cancer Survivorship Genetic Susceptibility and Second Primary Cancers: Research Strategies and Recommendations. <i>Journal of the National Cancer Institute</i> , 2006, 98, 15-25.	6.3	295
34	Tumour lineage shapes BRCA-mediated phenotypes. <i>Nature</i> , 2019, 571, 576-579.	27.8	295
35	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. <i>Nature Genetics</i> , 2017, 49, 1767-1778.	21.4	289
36	Prospective Genomic Profiling of Prostate Cancer Across Disease States Reveals Germline and Somatic Alterations That May Affect Clinical Decision Making. <i>JCO Precision Oncology</i> , 2017, 2017, 1-16.	3.0	286

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37	Recurrent BRCA2 6174delT mutations in Ashkenazi Jewish women affected by breast cancer. <i>Nature Genetics</i> , 1996, 13, 126-128.	21.4	282
38	The OncoArray Consortium: A Network for Understanding the Genetic Architecture of Common Cancers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2017, 26, 126-135.	2.5	278
39	MDM2 SNP309 Accelerates Tumor Formation in a Gender-Specific and Hormone-Dependent Manner. <i>Cancer Research</i> , 2006, 66, 5104-5110.	0.9	277
40	Genome-wide association study provides evidence for a breast cancer risk locus at 6q22.33. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 4340-4345.	7.1	274
41	A recurrent germline PAX5 mutation confers susceptibility to pre-B cell acute lymphoblastic leukemia. <i>Nature Genetics</i> , 2013, 45, 1226-1231.	21.4	270
42	Germline Variants in Targeted Tumor Sequencing Using Matched Normal DNA. <i>JAMA Oncology</i> , 2016, 2, 104.	7.1	270
43	Cancer Risks Associated With Germline <i>PALB2</i> Pathogenic Variants: An International Study of 524 Families. <i>Journal of Clinical Oncology</i> , 2020, 38, 674-685.	1.6	270
44	Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses. <i>Nature Genetics</i> , 2020, 52, 572-581.	21.4	265
45	Germline <i>BRCA</i> Mutations Denote a Clinicopathologic Subset of Prostate Cancer. <i>Clinical Cancer Research</i> , 2010, 16, 2115-2121.	7.0	263
46	A combined analysis of outcome following breast cancer: differences in survival based on BRCA1/BRCA2 mutation status and administration of adjuvant treatment. <i>Breast Cancer Research</i> , 2003, 6, R8-R17.	5.0	262
47	Counselling framework for moderate-penetrance cancer-susceptibility mutations. <i>Nature Reviews Clinical Oncology</i> , 2016, 13, 581-588.	27.6	258
48	Genetic/Familial High-Risk Assessment: Breast and Ovarian. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2010, 8, 562-594.	4.9	253
49	Tamoxifen and contralateral breast cancer in BRCA1 and BRCA2 carriers: An update. <i>International Journal of Cancer</i> , 2006, 118, 2281-2284.	5.1	246
50	Genome-Wide Association Study in BRCA1 Mutation Carriers Identifies Novel Loci Associated with Breast and Ovarian Cancer Risk. <i>PLoS Genetics</i> , 2013, 9, e1003212.	3.5	244
51	Evaluation of Polygenic Risk Scores for Breast and Ovarian Cancer Risk Prediction in BRCA1 and BRCA2 Mutation Carriers. <i>Journal of the National Cancer Institute</i> , 2017, 109, .	6.3	242
52	The "Duty to Warn" a Patient's Family Members About Hereditary Disease Risks. <i>JAMA - Journal of the American Medical Association</i> , 2004, 292, 1469.	7.4	228
53	Mutational spectrum in a worldwide study of 29,700 families with <i>BRCA1</i> or <i>BRCA2</i> mutations. <i>Human Mutation</i> , 2018, 39, 593-620.	2.5	224
54	Management of an Inherited Predisposition to Breast Cancer. <i>New England Journal of Medicine</i> , 2007, 357, 154-162.	27.0	222

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55	Identification of six new susceptibility loci for invasive epithelial ovarian cancer. <i>Nature Genetics</i> , 2015, 47, 164-171.	21.4	221
56	Reproductive risk factors for ovarian cancer in carriers of BRCA1 or BRCA2 mutations: a case-control study. <i>Lancet Oncology</i> , The, 2007, 8, 26-34.	10.7	220
57	RAD51 135Gâ†C Modifies Breast Cancer Risk among BRCA2 Mutation Carriers: Results from a Combined Analysis of 19 Studies. <i>American Journal of Human Genetics</i> , 2007, 81, 1186-1200.	6.2	217
58	Multiplex Genetic Testing for Cancer Susceptibility: Out on the High Wire Without a Net?. <i>Journal of Clinical Oncology</i> , 2013, 31, 1267-1270.	1.6	217
59	ASCO/SSO Review of Current Role of Risk-Reducing Surgery in Common Hereditary Cancer Syndromes. <i>Journal of Clinical Oncology</i> , 2006, 24, 4642-4660.	1.6	214
60	The Scientific Foundation for Personal Genomics: Recommendations from a National Institutes of Healthâ€Centers for Disease Control and Prevention Multidisciplinary Workshop. <i>Genetics in Medicine</i> , 2009, 11, 559-567.	2.4	207
61	Reliable Detection of Mismatch Repair Deficiency in Colorectal Cancers Using Mutational Load in Next-Generation Sequencing Panels. <i>Journal of Clinical Oncology</i> , 2016, 34, 2141-2147.	1.6	204
62	Fallopian Tube and Primary Peritoneal Carcinomas Associated With BRCA Mutations. <i>Journal of Clinical Oncology</i> , 2003, 21, 4222-4227.	1.6	199
63	<i>BRCA</i> Germline Mutations in Jewish Patients With Pancreatic Adenocarcinoma. <i>Journal of Clinical Oncology</i> , 2009, 27, 433-438.	1.6	194
64	Physical activity and risks of breast and colorectal cancer: a Mendelian randomisation analysis. <i>Nature Communications</i> , 2020, 11, 597.	12.8	193
65	Uterine Cancer After Risk-Reducing Salpingo-oophorectomy Without Hysterectomy in Women With <i>BRCA</i> Mutations. <i>JAMA Oncology</i> , 2016, 2, 1434.	7.1	189
66	Shared Genetic Susceptibility to Breast Cancer, Brain Tumors, and Fanconi Anemia. <i>Journal of the National Cancer Institute</i> , 2003, 95, 1548-1551.	6.3	183
67	Genome-wide association study identifies multiple risk loci for chronic lymphocytic leukemia. <i>Nature Genetics</i> , 2013, 45, 868-876.	21.4	179
68	BLM Heterozygosity and the Risk of Colorectal Cancer. <i>Science</i> , 2002, 297, 2013-2013.	12.6	174
69	Personalized medicine: new genomics, old lessons. <i>Human Genetics</i> , 2011, 130, 3-14.	3.8	173
70	Genetics, genomics, and cancer risk assessment. <i>Ca-A Cancer Journal for Clinicians</i> , 2011, 61, 327-359.	329.8	172
71	Prospective Evaluation of Germline Alterations in Patients With Exocrine Pancreatic Neoplasms. <i>Journal of the National Cancer Institute</i> , 2018, 110, 1067-1074.	6.3	170
72	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. <i>Cancer Research</i> , 2010, 70, 9742-9754.	0.9	169

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73	Cytogenetic analysis of 434 consecutively ascertained specimens of non-Hodgkin's lymphoma: Correlations between recurrent aberrations, histology, and exposure to cytotoxic treatment. <i>Genes Chromosomes and Cancer</i> , 1991, 3, 189-201.	2.8	167
74	Breast Conservation Therapy for Invasive Breast Cancer in Ashkenazi Women With BRCA Gene Founder Mutations. <i>Journal of the National Cancer Institute</i> , 1999, 91, 2112-2117.	6.3	167
75	Identification of germline genetic mutations in patients with pancreatic cancer. <i>Cancer</i> , 2015, 121, 4382-4388.	4.1	167
76	Low incidence of BRCA2 mutations in breast carcinoma and other cancers. <i>Nature Genetics</i> , 1996, 13, 241-244.	21.4	162
77	Genome-Wide Association Studies of Cancer. <i>Journal of Clinical Oncology</i> , 2010, 28, 4255-4267.	1.6	159
78	BRCA Mutations and Risk of Prostate Cancer in Ashkenazi Jews. <i>Clinical Cancer Research</i> , 2004, 10, 2918-2921.	7.0	156
79	Sequencing an Ashkenazi reference panel supports population-targeted personal genomics and illuminates Jewish and European origins. <i>Nature Communications</i> , 2014, 5, 4835.	12.8	156
80	Immunohistochemistry as First-line Screening for Detecting Colorectal Cancer Patients at Risk for Hereditary Nonpolyposis Colorectal Cancer Syndrome. <i>American Journal of Surgical Pathology</i> , 2009, 33, 1639-1645.	3.7	155
81	Genetic/Familial High-Risk Assessment: Breast and Ovarian, Version 2.2015. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2016, 14, 153-162.	4.9	153
82	Effect of pregnancy as a risk factor for breast cancer in <i>BRCA1</i> and <i>BRCA2</i> mutation carriers. <i>International Journal of Cancer</i> , 2005, 117, 988-991.	5.1	152
83	Prevention and Management of Hereditary Breast Cancer. <i>Journal of Clinical Oncology</i> , 2005, 23, 1656-1663.	1.6	152
84	Analysis of Heritability and Shared Heritability Based on Genome-Wide Association Studies for Thirteen Cancer Types. <i>Journal of the National Cancer Institute</i> , 2015, 107, djv279.	6.3	152
85	Age- and Tumor Subtype-Specific Breast Cancer Risk Estimates for <i>CH</i> and <i>EK</i> <i>2</i> *1100delC Carriers. <i>Journal of Clinical Oncology</i> , 2016, 34, 2750-2760.	1.6	152
86	Prediction of Breast and Prostate Cancer Risks in Male <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers Using Polygenic Risk Scores. <i>Journal of Clinical Oncology</i> , 2017, 35, 2240-2250.	1.6	152
87	Haplotype and Phenotype Analysis of Nine Recurrent BRCA2 Mutations in 111 Families: Results of an International Study. <i>American Journal of Human Genetics</i> , 1998, 62, 1381-1388.	6.2	150
88	BRCA Challenge: BRCA Exchange as a global resource for variants in BRCA1 and BRCA2. <i>PLoS Genetics</i> , 2018, 14, e1007752.	3.5	148
89	Risk of Endometrial Carcinoma Associated with BRCA Mutation. <i>Gynecologic Oncology</i> , 2001, 80, 395-398.	1.4	147
90	Genome-wide association study identifies multiple susceptibility loci for diffuse large B cell lymphoma. <i>Nature Genetics</i> , 2014, 46, 1233-1238.	21.4	147

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91	Conflicting Interpretation of Genetic Variants and Cancer Risk by Commercial Laboratories as Assessed by the Prospective Registry of Multiplex Testing. <i>Journal of Clinical Oncology</i> , 2016, 34, 4071-4078.	1.6	147
92	Value of Immunohistochemical Detection of DNA Mismatch Repair Proteins in Predicting Germline Mutation in Hereditary Colorectal Neoplasms. <i>American Journal of Surgical Pathology</i> , 2005, 29, 96-104.	3.7	136
93	Evidence of Gene-Environment Interactions between Common Breast Cancer Susceptibility Loci and Established Environmental Risk Factors. <i>PLoS Genetics</i> , 2013, 9, e1003284.	3.5	136
94	Appropriateness of breast-conserving treatment of breast carcinoma in women with germline mutations in BRCA1 or BRCA2. <i>Cancer</i> , 2005, 103, 44-51.	4.1	132
95	Prevalence of Germline Mutations in Cancer Susceptibility Genes in Patients With Advanced Renal Cell Carcinoma. <i>JAMA Oncology</i> , 2018, 4, 1228.	7.1	132
96	Quality of life in women at risk for ovarian cancer who have undergone risk-reducing oophorectomy. <i>Gynecologic Oncology</i> , 2003, 89, 281-287.	1.4	130
97	Novel Common Genetic Susceptibility Loci for Colorectal Cancer. <i>Journal of the National Cancer Institute</i> , 2019, 111, 146-157.	6.3	129
98	Germline ETV6 Mutations Confer Susceptibility to Acute Lymphoblastic Leukemia and Thrombocytopenia. <i>PLoS Genetics</i> , 2015, 11, e1005262.	3.5	128
99	Variants of the Adiponectin (<i>ADIPOQ</i>) and Adiponectin Receptor 1 (<i>ADIPOR1</i>) Genes and Colorectal Cancer Risk. <i>JAMA - Journal of the American Medical Association</i> , 2008, 300, 1523.	7.4	127
100	Breast cancer risk variants at 6q25 display different phenotype associations and regulate <i>ESR1</i> , <i>RMND1</i> and <i>CCDC170</i> . <i>Nature Genetics</i> , 2016, 48, 374-386.	21.4	125
101	Genome-wide Modeling of Polygenic Risk Score in Colorectal Cancer Risk. <i>American Journal of Human Genetics</i> , 2020, 107, 432-444.	6.2	124
102	Cancer Genomics and Inherited Risk. <i>Journal of Clinical Oncology</i> , 2014, 32, 687-698.	1.6	121
103	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. <i>Nature Genetics</i> , 2020, 52, 56-73.	21.4	120
104	Genetic/Familial High-Risk Assessment: Breast and Ovarian, Version 1.2014. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2014, 12, 1326-1338.	4.9	119
105	Ovarian cancer risk in Ashkenazi Jewish carriers of BRCA1 and BRCA2 mutations. <i>Clinical Cancer Research</i> , 2002, 8, 3776-81.	7.0	116
106	Evaluation of ACMG-Guideline-Based Variant Classification of Cancer Susceptibility and Non-Cancer-Associated Genes in Families Affected by Breast Cancer. <i>American Journal of Human Genetics</i> , 2016, 98, 801-817.	6.2	113
107	Comprehensive detection of germline variants by MSK-IMPACT, a clinical diagnostic platform for solid tumor molecular oncology and concurrent cancer predisposition testing. <i>BMC Medical Genomics</i> , 2017, 10, 33.	1.5	111
108	Cumulative Burden of Colorectal Cancer-Associated Genetic Variants Is More Strongly Associated With Early-Onset vs Late-Onset Cancer. <i>Gastroenterology</i> , 2020, 158, 1274-1286.e12.	1.3	110

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109	Functional and genomic approaches reveal an ancient CHEK2 allele associated with breast cancer in the Ashkenazi Jewish population. <i>Human Molecular Genetics</i> , 2005, 14, 555-563.	2.9	109
110	The contribution of pathogenic variants in breast cancer susceptibility genes to familial breast cancer risk. <i>Npj Breast Cancer</i> , 2017, 3, 22.	5.2	108
111	Cancer Genetic Testing and Assisted Reproduction. <i>Journal of Clinical Oncology</i> , 2006, 24, 4775-4782.	1.6	107
112	Frequency of CHEK2*1100delC in New York breast cancer cases and controls. <i>BMC Medical Genetics</i> , 2003, 4, 1.	2.1	106
113	Identification of a BRCA2-Specific Modifier Locus at 6p24 Related to Breast Cancer Risk. <i>PLoS Genetics</i> , 2013, 9, e1003173.	3.5	105
114	<i>TGFBR1</i> *6A and Cancer Risk: A Meta-Analysis of Seven Case-Control Studies. <i>Journal of Clinical Oncology</i> , 2003, 21, 3236-3243.	1.6	104
115	Variants of the Adiponectin and Adiponectin Receptor 1 Genes and Breast Cancer Risk. <i>Cancer Research</i> , 2008, 68, 3178-3184.	0.9	104
116	Genomic Profiles for Disease Risk. <i>JAMA - Journal of the American Medical Association</i> , 2008, 299, 1353.	7.4	100
117	Common variants in LSP1, 2q35 and 8q24 and breast cancer risk for BRCA1 and BRCA2 mutation carriers. <i>Human Molecular Genetics</i> , 2009, 18, 4442-4456.	2.9	99
118	18q21 rearrangement in diffuse large cell lymphoma: incidence and clinical significance. <i>British Journal of Haematology</i> , 1989, 72, 178-183.	2.5	97
119	Absence of premalignant histologic, molecular, or cell biologic alterations in prophylactic oophorectomy specimens from BRCA1 heterozygotes. <i>Cancer</i> , 2000, 89, 383-390.	4.1	97
120	Genome-wide Association Study Identifies Five Susceptibility Loci for Follicular Lymphoma outside the HLA Region. <i>American Journal of Human Genetics</i> , 2014, 95, 462-471.	6.2	96
121	Meta-analysis of genome-wide association studies discovers multiple loci for chronic lymphocytic leukemia. <i>Nature Communications</i> , 2016, 7, 10933.	12.8	94
122	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. <i>Nature Communications</i> , 2016, 7, 11375.	12.8	93
123	Interplay between BRCA1 and RHAMM Regulates Epithelial Apicobasal Polarization and May Influence Risk of Breast Cancer. <i>PLoS Biology</i> , 2011, 9, e1001199.	5.6	91
124	<i>FANCM</i> c.5791C>T nonsense mutation (rs144567652) induces exon skipping, affects DNA repair activity and is a familial breast cancer risk factor. <i>Human Molecular Genetics</i> , 2015, 24, 5345-5355.	2.9	91
125	Epithelial lesions in prophylactic mastectomy specimens from women with BRCA mutations. <i>Cancer</i> , 2003, 97, 1601-1608.	4.1	90
126	Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. <i>Nature Communications</i> , 2019, 10, 1741.	12.8	90

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127	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. <i>Gastroenterology</i> , 2020, 158, 1300-1312.e20.	1.3	90
128	Male breast cancer in BRCA1 and BRCA2 mutation carriers: pathology data from the Consortium of Investigators of Modifiers of BRCA1/2. <i>Breast Cancer Research</i> , 2016, 18, 15.	5.0	88
129	Shared heritability and functional enrichment across six solid cancers. <i>Nature Communications</i> , 2019, 10, 431.	12.8	88
130	TGFBR1*6A and Cancer: A Meta-Analysis of 12 Case-Control Studies. <i>Journal of Clinical Oncology</i> , 2004, 22, 756-758.	1.6	85
131	Common Genetic Variants and Modification of Penetrance of BRCA2-Associated Breast Cancer. <i>PLoS Genetics</i> , 2010, 6, e1001183.	3.5	85
132	Combined Genetic Assessment of Transforming Growth Factor- β Signaling Pathway Variants May Predict Breast Cancer Risk. <i>Cancer Research</i> , 2005, 65, 3454-3461.	0.9	83
133	Prevalence of <i>BRCA1</i> and <i>BRCA2</i> mutations in Ashkenazi Jewish families with breast and pancreatic cancer. <i>Cancer</i> , 2012, 118, 493-499.	4.1	83
134	Therapeutic Implications of Germline Testing in Patients With Advanced Cancers. <i>Journal of Clinical Oncology</i> , 2021, 39, 2698-2709.	1.6	83
135	Heterogenic Loss of the Wild-Type BRCA Allele in Human Breast Tumorigenesis. <i>Annals of Surgical Oncology</i> , 2007, 14, 2510-2518.	1.5	82
136	Polygenic risk scores and breast and epithelial ovarian cancer risks for carriers of BRCA1 and BRCA2 pathogenic variants. <i>Genetics in Medicine</i> , 2020, 22, 1653-1666.	2.4	82
137	Clonal hematopoiesis is associated with risk of severe Covid-19. <i>Nature Communications</i> , 2021, 12, 5975.	12.8	81
138	Risk of Ovarian Cancer in BRCA1 and BRCA2 Mutation-Negative Hereditary Breast Cancer Families. <i>Journal of the National Cancer Institute</i> , 2005, 97, 1382-1384.	6.3	80
139	Familial Kidney Cancer: Implications of New Syndromes and Molecular Insights. <i>European Urology</i> , 2019, 76, 754-764.	1.9	80
140	Common variants at 12p11, 12q24, 9p21, 9q31.2 and in ZNF365 are associated with breast cancer risk for BRCA1 and/or BRCA2 mutation carriers. <i>Breast Cancer Research</i> , 2012, 14, R33.	5.0	78
141	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breast-ovarian cancer susceptibility locus. <i>Nature Communications</i> , 2016, 7, 12675.	12.8	78
142	Preimplantation Genetic Diagnosis for Cancer Syndromes. <i>JAMA - Journal of the American Medical Association</i> , 2006, 296, 2727.	7.4	77
143	BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. <i>Journal of the National Cancer Institute</i> , 2016, 108, djv315.	6.3	77
144	Adiposity, metabolites, and colorectal cancer risk: Mendelian randomization study. <i>BMC Medicine</i> , 2020, 18, 396.	5.5	76

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145	Genome-wide association analysis implicates dysregulation of immunity genes in chronic lymphocytic leukaemia. <i>Nature Communications</i> , 2017, 8, 14175.	12.8	75
146	Susceptibility Loci Associated with Prostate Cancer Progression and Mortality. <i>Clinical Cancer Research</i> , 2010, 16, 2819-2832.	7.0	74
147	Prospective pan-cancer germline testing using MSK-IMPACT informs clinical translation in 751 patients with pediatric solid tumors. <i>Nature Cancer</i> , 2021, 2, 357-365.	13.2	74
148	Psychosocial predictors of BRCA counseling and testing decisions among urban African-American women. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2002, 11, 1579-85.	2.5	74
149	Improved survival for BRCA2-associated serous ovarian cancer compared with both BRCA-negative and BRCA1-associated serous ovarian cancer. <i>Cancer</i> , 2012, 118, 3703-3709.	4.1	72
150	Frequency of BRCA1 and BRCA2 Mutations in Unselected Ashkenazi Jewish Patients With Colorectal Cancer. <i>Journal of the National Cancer Institute</i> , 2004, 96, 68-70.	6.3	71
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