Mark E Robson

List of Publications by Citations

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256 24,035 151 75 h-index g-index citations papers 281 6.3 30,032 9.1 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
256	Mutational landscape of metastatic cancer revealed from prospective clinical sequencing of 10,000 patients. <i>Nature Medicine</i> , 2017 , 23, 703-713	50.5	1638
255	Olaparib for Metastatic Breast Cancer in Patients with a Germline BRCA Mutation. <i>New England Journal of Medicine</i> , 2017 , 377, 523-533	59.2	1405
254	Oral poly(ADP-ribose) polymerase inhibitor olaparib in patients with BRCA1 or BRCA2 mutations and advanced breast cancer: a proof-of-concept trial. <i>Lancet, The,</i> 2010 , 376, 235-44	40	1395
253	Risk-reducing salpingo-oophorectomy in women with a BRCA1 or BRCA2 mutation. <i>New England Journal of Medicine</i> , 2002 , 346, 1609-15	59.2	1198
252	Inherited DNA-Repair Gene Mutations in Men with Metastatic Prostate Cancer. <i>New England Journal of Medicine</i> , 2016 , 375, 443-53	59.2	791
251	Gene-panel sequencing and the prediction of breast-cancer risk. <i>New England Journal of Medicine</i> , 2015 , 372, 2243-57	59.2	587
250	Tamoxifen and risk of contralateral breast cancer in BRCA1 and BRCA2 mutation carriers: a case-control study. Hereditary Breast Cancer Clinical Study Group. <i>Lancet, The</i> , 2000 , 356, 1876-81	40	473
249	Risk-reducing salpingo-oophorectomy for the prevention of BRCA1- and BRCA2-associated breast and gynecologic cancer: a multicenter, prospective study. <i>Journal of Clinical Oncology</i> , 2008 , 26, 1331-7	2.2	465
248	Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. <i>Nature Genetics</i> , 2013 , 45, 371-84, 384e1-2	36.3	422
247	Pathology of breast and ovarian cancers among BRCA1 and BRCA2 mutation carriers: results from the Consortium of Investigators of Modifiers of BRCA1/2 (CIMBA). <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2012 , 21, 134-47	4	411
246	Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. <i>American Journal of Human Genetics</i> , 2019 , 104, 21-34	11	363
245	American Society of Clinical Oncology Policy Statement Update: Genetic and Genomic Testing for Cancer Susceptibility. <i>Journal of Clinical Oncology</i> , 2015 , 33, 3660-7	2.2	360
244	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	2.2	349
243	Therapy-Related Clonal Hematopoiesis in Patients with Non-hematologic Cancers Is Common and Associated with Adverse Clinical Outcomes. <i>Cell Stem Cell</i> , 2017 , 21, 374-382.e4	18	339
242	The Genomic Landscape of Endocrine-Resistant Advanced Breast Cancers. Cancer Cell, 2018, 34, 427-43	82 9 63	339
241	Prediction of breast cancer risk based on profiling with common genetic variants. <i>Journal of the National Cancer Institute</i> , 2015 , 107,	9.7	324
240	Impairment of BRCA1-related DNA double-strand break repair leads to ovarian aging in mice and humans. <i>Science Translational Medicine</i> , 2013 , 5, 172ra21	17.5	290

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239	Association of type and location of BRCA1 and BRCA2 mutations with risk of breast and ovarian cancer. <i>JAMA - Journal of the American Medical Association</i> , 2015 , 313, 1347-61	27.4	286
238	Oral contraceptives and the risk of breast cancer in BRCA1 and BRCA2 mutation carriers. <i>Journal of the National Cancer Institute</i> , 2002 , 94, 1773-9	9.7	266
237	High-intensity sequencing reveals the sources of plasma circulating cell-free DNA variants. <i>Nature Medicine</i> , 2019 , 25, 1928-1937	50.5	263
236	OlympiAD final overall survival and tolerability results: Olaparib versus chemotherapy treatment of physician@ choice in patients with a germline BRCA mutation and HER2-negative metastatic breast cancer. <i>Annals of Oncology</i> , 2019 , 30, 558-566	10.3	240
235	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, 825-835	27.4	235
234	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> , 2004 , 6, R8-R17	8.3	225
233	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	6	209
232	Genome doubling shapes the evolution and prognosis of advanced cancers. <i>Nature Genetics</i> , 2018 , 50, 1189-1195	36.3	208
231	Microsatellite Instability Is Associated With the Presence of Lynch Syndrome Pan-Cancer. <i>Journal of Clinical Oncology</i> , 2019 , 37, 286-295	2.2	203
230	Counselling framework for moderate-penetrance cancer-susceptibility mutations. <i>Nature Reviews Clinical Oncology</i> , 2016 , 13, 581-8	19.4	200
229	Germline Variants in Targeted Tumor Sequencing Using Matched Normal DNA. <i>JAMA Oncology</i> , 2016 , 2, 104-11	13.4	198
228	Germline BRCA mutations denote a clinicopathologic subset of prostate cancer. <i>Clinical Cancer Research</i> , 2010 , 16, 2115-21	12.9	196
227	Clinical practice. Management of an inherited predisposition to breast cancer. <i>New England Journal of Medicine</i> , 2007 , 357, 154-62	59.2	194
226	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691	36.3	190
225	An emerging entity: pancreatic adenocarcinoma associated with a known BRCA mutation: clinical descriptors, treatment implications, and future directions. <i>Oncologist</i> , 2011 , 16, 1397-402	5.7	190
224	Multiplex genetic testing for cancer susceptibility: out on the high wire without a net?. <i>Journal of Clinical Oncology</i> , 2013 , 31, 1267-70	2.2	184
223	Tumour lineage shapes BRCA-mediated phenotypes. <i>Nature</i> , 2019 , 571, 576-579	50.4	170
222	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, 2141-7	2.2	170

221	Fallopian tube and primary peritoneal carcinomas associated with BRCA mutations. <i>Journal of Clinical Oncology</i> , 2003 , 21, 4222-7	2.2	169
220	BRCA germline mutations in Jewish patients with pancreatic adenocarcinoma. <i>Journal of Clinical Oncology</i> , 2009 , 27, 433-8	2.2	160
219	Shared genetic susceptibility to breast cancer, brain tumors, and Fanconi anemia. <i>Journal of the National Cancer Institute</i> , 2003 , 95, 1548-51	9.7	160
218	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,	9.7	153
217	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,	3.6	151
216	Uterine Cancer After Risk-Reducing Salpingo-oophorectomy Without Hysterectomy in Women With BRCA Mutations. <i>JAMA Oncology</i> , 2016 , 2, 1434-1440	13.4	151
215	BRCA mutations and risk of prostate cancer in Ashkenazi Jews. Clinical Cancer Research, 2004, 10, 2918-	2 12.9	139
214	Mutational spectrum in a worldwide study of 29,700 families with BRCA1 or BRCA2 mutations. <i>Human Mutation</i> , 2018 , 39, 593-620	4.7	138
213	Risk of endometrial carcinoma associated with BRCA mutation. <i>Gynecologic Oncology</i> , 2001 , 80, 395-8	4.9	135
212	Cancer Risks Associated With Germline Pathogenic Variants: An International Study of 524 Families. Journal of Clinical Oncology, 2020 , 38, 674-685	2.2	133
211	Diverse and Reversion Mutations in Circulating Cell-Free DNA of Therapy-Resistant Breast or Ovarian Cancer. <i>Clinical Cancer Research</i> , 2017 , 23, 6708-6720	12.9	132
210	Genome-wide association studies of cancer. <i>Journal of Clinical Oncology</i> , 2010 , 28, 4255-67	2.2	127
209	Appropriateness of breast-conserving treatment of breast carcinoma in women with germline mutations in BRCA1 or BRCA2: a clinic-based series. <i>Cancer</i> , 2005 , 103, 44-51	6.4	123
208	Breast cancer phenotype in women with TP53 germline mutations: a Li-Fraumeni syndrome consortium effort. <i>Breast Cancer Research and Treatment</i> , 2012 , 133, 1125-30	4.4	120
207	Identification of germline genetic mutations in patients with pancreatic cancer. <i>Cancer</i> , 2015 , 121, 4382	2 -6 .4	117
206	Quality of life in women at risk for ovarian cancer who have undergone risk-reducing oophorectomy. <i>Gynecologic Oncology</i> , 2003 , 89, 281-7	4.9	111
205	Hereditary breast cancer. Current Problems in Surgery, 2001, 38, 387-480	2.8	111
204	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, 4071-4	4 07 8	110

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203	Age- and Tumor Subtype-Specific Breast Cancer Risk Estimates for CHEK2*1100delC Carriers. Journal of Clinical Oncology, 2016 , 34, 2750-60	2.2	107
202	Baseline Surveillance in Li-Fraumeni Syndrome Using Whole-Body Magnetic Resonance Imaging: A Meta-analysis. <i>JAMA Oncology</i> , 2017 , 3, 1634-1639	13.4	107
201	Role of Genetic Testing for Inherited Prostate Cancer Risk: Philadelphia Prostate Cancer Consensus Conference 2017. <i>Journal of Clinical Oncology</i> , 2018 , 36, 414-424	2.2	107
200	Prospective Evaluation of Germline Alterations in Patients With Exocrine Pancreatic Neoplasms. <i>Journal of the National Cancer Institute</i> , 2018 , 110, 1067-1074	9.7	103
199	Cancer therapy shapes the fitness landscape of clonal hematopoiesis. <i>Nature Genetics</i> , 2020 , 52, 1219-7	123263	103
198	Prediction of Breast and Prostate Cancer Risks in Male BRCA1 and BRCA2 Mutation Carriers Using Polygenic Risk Scores. <i>Journal of Clinical Oncology</i> , 2017 , 35, 2240-2250	2.2	101
197	Cancer genomics and inherited risk. Journal of Clinical Oncology, 2014, 32, 687-98	2.2	100
196	Management of Hereditary Breast Cancer: American Society of Clinical Oncology, American Society for Radiation Oncology, and Society of Surgical Oncology Guideline. <i>Journal of Clinical Oncology</i> , 2020 , 38, 2080-2106	2.2	95
195	Breast-conserving therapy achieves locoregional outcomes comparable to mastectomy in women with T1-2N0 triple-negative breast cancer. <i>Annals of Surgical Oncology</i> , 2013 , 20, 3469-76	3.1	95
194	Estimated risk of radiation-induced breast cancer from mammographic screening for young BRCA mutation carriers. <i>Journal of the National Cancer Institute</i> , 2009 , 101, 205-9	9.7	94
193	Screening for germline EGFR T790M mutations through lung cancer genotyping. <i>Journal of Thoracic Oncology</i> , 2012 , 7, 1049-52	8.9	92
192	TBCRC 048: Phase II Study of Olaparib for Metastatic Breast Cancer and Mutations in Homologous Recombination-Related Genes. <i>Journal of Clinical Oncology</i> , 2020 , 38, 4274-4282	2.2	92
191	Pleomorphic characteristics of a germ-line KIT mutation in a large kindred with gastrointestinal stromal tumors, hyperpigmentation, and dysphagia. <i>Clinical Cancer Research</i> , 2004 , 10, 1250-4	12.9	91
190	Frequency of CHEK2*1100delC in New York breast cancer cases and controls. <i>BMC Medical Genetics</i> , 2003 , 4, 1	2.1	91
189	Identification of a BRCA2-specific modifier locus at 6p24 related to breast cancer risk. <i>PLoS Genetics</i> , 2013 , 9, e1003173	6	90
188	Absence of premalignant histologic, molecular, or cell biologic alterations in prophylactic oophorectomy specimens from BRCA1 heterozygotes. <i>Cancer</i> , 2000 , 89, 383-90	6.4	90
187	BRCA Challenge: BRCA Exchange as a global resource for variants in BRCA1 and BRCA2. <i>PLoS Genetics</i> , 2018 , 14, e1007752	6	90
186	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	11	86

185	Refined histopathological predictors of BRCA1 and BRCA2 mutation status: a large-scale analysis of breast cancer characteristics from the BCAC, CIMBA, and ENIGMA consortia. <i>Breast Cancer Research</i> , 2014 , 16, 3419	8.3	82
184	Epithelial lesions in prophylactic mastectomy specimens from women with BRCA mutations. <i>Cancer</i> , 2003 , 97, 1601-8	6.4	79
183	The contribution of pathogenic variants in breast cancer susceptibility genes to familial breast cancer risk. <i>Npj Breast Cancer</i> , 2017 , 3, 22	7.8	78
182	A prospective, longitudinal study of the functional status and quality of life of older patients with breast cancer receiving adjuvant chemotherapy. <i>Journal of the American Geriatrics Society</i> , 2006 , 54, 11	1 5 :24	76
181	A phase II open-label study of ganetespib, a novel heat shock protein 90 inhibitor for patients with metastatic breast cancer. <i>Clinical Breast Cancer</i> , 2014 , 14, 154-60	3	75
180	Genetic analysis of the early natural history of epithelial ovarian carcinoma. <i>PLoS ONE</i> , 2010 , 5, e10358	3.7	74
179	Heterogenic loss of the wild-type BRCA allele in human breast tumorigenesis. <i>Annals of Surgical Oncology</i> , 2007 , 14, 2510-8	3.1	72
178	Prevalence of BRCA1 and BRCA2 mutations in Ashkenazi Jewish families with breast and pancreatic cancer. <i>Cancer</i> , 2012 , 118, 493-9	6.4	71
177	Effect of adjuvant breast cancer chemotherapy on cognitive function from the older patient@perspective. <i>Breast Cancer Research and Treatment</i> , 2006 , 98, 343-8	4.4	71
176	Risk of ovarian cancer in BRCA1 and BRCA2 mutation-negative hereditary breast cancer families. Journal of the National Cancer Institute, 2005, 97, 1382-4	9.7	70
175	Analysis of genetic variants in never-smokers with lung cancer facilitated by an Internet-based blood collection protocol: a preliminary report. <i>Clinical Cancer Research</i> , 2010 , 16, 755-63	12.9	69
174	Comparison of screening CEDM and MRI for women at increased risk for breast cancer: A pilot study. <i>European Journal of Radiology</i> , 2017 , 97, 37-43	4.7	66
173	Associations of common variants at 1p11.2 and 14q24.1 (RAD51L1) with breast cancer risk and heterogeneity by tumor subtype: findings from the Breast Cancer Association Consortium. <i>Human Molecular Genetics</i> , 2011 , 20, 4693-706	5.6	66
172	Prevalence of Germline Mutations in Cancer Susceptibility Genes in Patients With Advanced Renal Cell Carcinoma. <i>JAMA Oncology</i> , 2018 , 4, 1228-1235	13.4	66
171	The Landscape of Somatic Genetic Alterations in Breast Cancers From ATM Germline Mutation Carriers. <i>Journal of the National Cancer Institute</i> , 2018 , 110, 1030-1034	9.7	65
170	BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. Journal of the National Cancer Institute, 2016 , 108,	9.7	65
169	A Phase II Study of Talazoparib after Platinum or Cytotoxic Nonplatinum Regimens in Patients with Advanced Breast Cancer and Germline Mutations (ABRAZO). <i>Clinical Cancer Research</i> , 2019 , 25, 2717-27	² 24.9	65
168	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. <i>Nature Communications</i> , 2016 , 7, 11375	17.4	64

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167	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	3.7	64	
166	Population Frequency of Germline BRCA1/2 Mutations. <i>Journal of Clinical Oncology</i> , 2016 , 34, 4183-418	352.2	64	
165	Common breast cancer susceptibility alleles are associated with tumour subtypes in BRCA1 and BRCA2 mutation carriers: results from the Consortium of Investigators of Modifiers of BRCA1/2. Breast Cancer Research, 2011 , 13, R110	8.3	62	
164	ESMO / ASCO Recommendations for a Global Curriculum in Medical Oncology Edition 2016. <i>ESMO Open</i> , 2016 , 1, e000097	6	59	
163	Variation in anastrozole metabolism and pharmacodynamics in women with early breast cancer. <i>Cancer Research</i> , 2010 , 70, 3278-86	10.1	58	
162	Health literacy, numeracy, and interpretation of graphical breast cancer risk estimates. <i>Patient Education and Counseling</i> , 2011 , 83, 92-8	3.1	58	
161	Genomic Methods Identify Homologous Recombination Deficiency in Pancreas Adenocarcinoma and Optimize Treatment Selection. <i>Clinical Cancer Research</i> , 2020 , 26, 3239-3247	12.9	58	
160	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	8.3	58	
159	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. <i>Nature Genetics</i> , 2020 , 52, 56-73	36.3	56	
158	Breast cancer detection and tumor characteristics in BRCA1 and BRCA2 mutation carriers. <i>Breast Cancer Research and Treatment</i> , 2017 , 163, 565-571	4.4	53	
157	Ovarian and Breast Cancer Risks Associated With Pathogenic Variants in RAD51C and RAD51D. Journal of the National Cancer Institute, 2020 , 112, 1242-1250	9.7	51	
156	Revealing the incidentalome when targeting the tumor genome. <i>JAMA - Journal of the American Medical Association</i> , 2013 , 310, 795-6	27.4	51	
155	Contralateral breast cancer after radiotherapy among BRCA1 and BRCA2 mutation carriers: a WECARE study report. <i>European Journal of Cancer</i> , 2013 , 49, 2979-85	7.5	51	
154	The safety of dose-dense doxorubicin and cyclophosphamide followed by paclitaxel with trastuzumab in HER-2/neu overexpressed/amplified breast cancer. <i>Journal of Clinical Oncology</i> , 2008 , 26, 1216-22	2.2	51	
153	Effect of mammography on breast cancer risk in women with mutations in BRCA1 or BRCA2. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2006 , 15, 2311-3	4	51	
152	American Society of Clinical Oncology policy statement: the role of the oncologist in cancer prevention and risk assessment. <i>Journal of Clinical Oncology</i> , 2009 , 27, 986-93	2.2	50	
151	Alterations in and promote clinical resistance to alpelisib plus aromatase inhibitors. <i>Nature Cancer</i> , 2020 , 1, 382-393	15.4	49	
150	Favorable prognosis in patients with T1a/T1bN0 triple-negative breast cancers treated with multimodality therapy. <i>Cancer</i> , 2012 , 118, 4944-52	6.4	49	

149	Comparison of 6q25 breast cancer hits from Asian and European Genome Wide Association Studies in the Breast Cancer Association Consortium (BCAC). <i>PLoS ONE</i> , 2012 , 7, e42380	3.7	49
148	Germline EGFR T790M mutation found in multiple members of a familial cohort. <i>Journal of Thoracic Oncology</i> , 2014 , 9, 554-8	8.9	47
147	Associations of common breast cancer susceptibility alleles with risk of breast cancer subtypes in BRCA1 and BRCA2 mutation carriers. <i>Breast Cancer Research</i> , 2014 , 16, 3416	8.3	46
146	TSPYL5 SNPs: association with plasma estradiol concentrations and aromatase expression. <i>Molecular Endocrinology</i> , 2013 , 27, 657-70		42
145	Increased progesterone receptor expression in benign epithelium of BRCA1-related breast cancers. <i>Cancer Research</i> , 2004 , 64, 5051-3	10.1	42
144	A comparison of bilateral breast cancers in BRCA carriers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2005 , 14, 1534-8	4	42
143	Feasibility trial of letrozole in combination with bevacizumab in patients with metastatic breast cancer. <i>Journal of Clinical Oncology</i> , 2010 , 28, 628-33	2.2	41
142	Olaparib for Metastatic Germline BRCA-Mutated Breast Cancer. <i>New England Journal of Medicine</i> , 2017 , 377, 1792-3	59.2	39
141	A phase IIA trial of acupuncture to reduce chemotherapy-induced peripheral neuropathy severity during neoadjuvant or adjuvant weekly paclitaxel chemotherapy in breast cancer patients. <i>European Journal of Cancer</i> , 2018 , 101, 12-19	7.5	39
140	Susceptibility loci associated with specific and shared subtypes of lymphoid malignancies. <i>PLoS Genetics</i> , 2013 , 9, e1003220	6	38
139	Poly(ADP-ribose) polymerase inhibitors in triple-negative breast cancer. <i>Cancer Journal (Sudbury, Mass)</i> , 2010 , 16, 48-52	2.2	38
138	Association of a Polygenic Risk Score With Breast Cancer Among Women Carriers of High- and Moderate-Risk Breast Cancer Genes. <i>JAMA Network Open</i> , 2020 , 3, e208501	10.4	38
137	Risk of metachronous breast cancer after BRCA mutation-associated ovarian cancer. <i>Cancer</i> , 2013 , 119, 1344-8	6.4	37
136	Inherited predisposition to gastrointestinal stromal tumor. <i>Hematology/Oncology Clinics of North America</i> , 2009 , 23, 1-13, vii	3.1	36
135	Patient-reported outcomes in patients with a germline BRCA mutation and HER2-negative metastatic breast cancer receiving olaparib versus chemotherapy in the OlympiAD trial. <i>European Journal of Cancer</i> , 2019 , 120, 20-30	7.5	35
134	Pre- and postmenopausal high-risk women undergoing screening for ovarian cancer: anxiety, risk perceptions, and quality of life. <i>Gynecologic Oncology</i> , 2003 , 89, 440-6	4.9	35
133	Ductal lavage in patients undergoing mastectomy for mammary carcinoma: a correlative study. <i>Cancer</i> , 2003 , 98, 2170-6	6.4	35
132	Increased CpG methylation of the estrogen receptor gene in BRCA1-linked estrogen receptor-negative breast cancers. <i>Oncogene</i> , 2002 , 21, 7034-41	9.2	34

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131	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	8.1	34	
130	DNA glycosylases involved in base excision repair may be associated with cancer risk in BRCA1 and BRCA2 mutation carriers. <i>PLoS Genetics</i> , 2014 , 10, e1004256	6	33	
129	Genome-wide association studies of cancer predisposition. <i>Hematology/Oncology Clinics of North America</i> , 2010 , 24, 973-96	3.1	33	
128	Breast MRI for women with hereditary cancer risk. <i>JAMA - Journal of the American Medical Association</i> , 2004 , 292, 1368-70	27.4	33	
127	Clinical considerations in the management of individuals at risk for hereditary breast and ovarian cancer. <i>Cancer Control</i> , 2002 , 9, 457-65	2.2	33	
126	Estrogens and their precursors in postmenopausal women with early breast cancer receiving anastrozole. <i>Steroids</i> , 2015 , 99, 32-8	2.8	32	
125	Evaluation of germline PTEN mutations in endometrial cancer patients. <i>Gynecologic Oncology</i> , 2005 , 96, 21-4	4.9	32	
124	Germline BRCA mutation does not prevent response to taxane-based therapy for the treatment of castration-resistant prostate cancer. <i>BJU International</i> , 2012 , 109, 713-9	5.6	31	
123	Hereditary ovarian cancer in Ashkenazi Jews. Familial Cancer, 2004, 3, 259-64	3	31	
122	Next generation sequencing and tumor mutation profiling: are we ready for routine use in the oncology clinic?. <i>BMC Medicine</i> , 2014 , 12, 140	11.4	30	
121	COMPLEXO: identifying the missing heritability of breast cancer via next generation collaboration. <i>Breast Cancer Research</i> , 2013 , 15, 402	8.3	30	
120	A randomized Phase II study of veliparib with temozolomide or carboplatin/paclitaxel versus placebo with carboplatin/paclitaxel in BRCA1/2 metastatic breast cancer: design and rationale. <i>Future Oncology</i> , 2017 , 13, 307-320	3.6	30	
119	A Recurrent ERCC3 Truncating Mutation Confers Moderate Risk for Breast Cancer. <i>Cancer Discovery</i> , 2016 , 6, 1267-1275	24.4	30	
118	Phase II study of feasibility of dose-dense FEC followed by alternating weekly taxanes in high-risk, four or more node-positive breast cancer. <i>Clinical Cancer Research</i> , 2004 , 10, 5754-61	12.9	29	
117	Twenty-one-gene recurrence score assay in BRCA-associated versus sporadic breast cancers: Differences based on germline mutation status. <i>Cancer</i> , 2016 , 122, 1178-84	6.4	29	
116	Breast Cancer Family History and Contralateral Breast Cancer Risk in Young Women: An Update From the Women@ Environmental Cancer and Radiation Epidemiology Study. <i>Journal of Clinical Oncology</i> , 2018 , 36, 1513-1520	2.2	29	
115	A feasibility study of bevacizumab plus dose-dense doxorubicin-cyclophosphamide (AC) followed by nanoparticle albumin-bound paclitaxel in early-stage breast cancer. <i>Clinical Cancer Research</i> , 2011 , 17, 3398-407	12.9	28	
114	The TP53 mutational spectrum and frequency of CHEK2*1100delC in Li-Fraumeni-like kindreds. <i>Familial Cancer</i> , 2005 , 4, 177-81	3	28	

113	Adjuvant treatment recommendations in older women with breast cancer: a survey of oncologists. <i>Critical Reviews in Oncology/Hematology</i> , 2007 , 61, 255-60	7	27
112	Assessing associations between the AURKA-HMMR-TPX2-TUBG1 functional module and breast cancer risk in BRCA1/2 mutation carriers. <i>PLoS ONE</i> , 2015 , 10, e0120020	3.7	26
111	BRCA mutations in women with ductal carcinoma in situ. Clinical Cancer Research, 2007, 13, 4306-10	12.9	26
110	Genetic Testing Awareness and Attitudes among Latinos: Exploring Shared Perceptions and Gender-Based Differences. <i>Public Health Genomics</i> , 2016 , 19, 34-46	1.9	26
109	Should all BRCA1 mutation carriers with stage I breast cancer receive chemotherapy?. <i>Breast Cancer Research and Treatment</i> , 2013 , 138, 273-9	4.4	25
108	Assessment of SLX4 Mutations in Hereditary Breast Cancers. <i>PLoS ONE</i> , 2013 , 8, e66961	3.7	24
107	Germline PALB2 mutation analysis in breast-pancreas cancer families. <i>Journal of Medical Genetics</i> , 2011 , 48, 523-5	5.8	24
106	Functional redundancy of exon 12 of BRCA2 revealed by a comprehensive analysis of the c.6853A>G (p.I2285V) variant. <i>Human Mutation</i> , 2009 , 30, 1543-50	4.7	24
105	Pathologic complete response rate according to HER2 detection methods in HER2-positive breast cancer treated with neoadjuvant systemic therapy. <i>Breast Cancer Research and Treatment</i> , 2019 , 177, 61-66	4.4	23
104	Smoking and the risk of breast cancer in BRCA1 and BRCA2 carriers: an update. <i>Breast Cancer Research and Treatment</i> , 2009 , 114, 127-35	4.4	23
103	Breast cancer surveillance in women with hereditary risk due to BRCA1 or BRCA2 mutations. <i>Clinical Breast Cancer</i> , 2004 , 5, 260-8; discussion 269-71	3	23
102	Ovarian carcinoma screening in women at intermediate risk: impact on quality of life and need for invasive follow-up. <i>Cancer</i> , 2005 , 104, 314-20	6.4	23
101	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	15.4	23
100	Association of Common Genetic Variants With Contralateral Breast Cancer Risk in the WECARE Study. <i>Journal of the National Cancer Institute</i> , 2017 , 109,	9.7	22
99	Association of Genomic Domains in and with Prostate Cancer Risk and Aggressiveness. <i>Cancer Research</i> , 2020 , 80, 624-638	10.1	22
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