Karin Kast

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Risks of Breast, Ovarian, and Contralateral Breast Cancer for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. JAMA - Journal of the American Medical Association, 2017, 317, 2402.	7.4	1,898
2	Germline mutations in breast and ovarian cancer pedigrees establish RAD51C as a human cancer susceptibility gene. Nature Genetics, 2010, 42, 410-414.	21.4	638
3	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</i> /i>2 (CIMBA). Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 134-147.	2.5	513
4	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
5	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691.	21.4	356
6	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778.	21.4	289
7	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
8	Prevalence of <i>BRCA1/2</i> germline mutations in 21â€401 families with breast and ovarian cancer. Journal of Medical Genetics, 2016, 53, 465-471.	3.2	179
9	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
10	Impact of breast cancer subtypes and patterns of metastasis on outcome. Breast Cancer Research and Treatment, 2015, 150, 621-629.	2.5	157
11	Prediction of Breast and Prostate Cancer Risks in Male <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers Using Polygenic Risk Scores. Journal of Clinical Oncology, 2017, 35, 2240-2250.	1.6	152
12	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
13	RPA and Rad51 constitute a cell intrinsic mechanism to protect the cytosol from self DNA. Nature Communications, 2016, 7, 11752.	12.8	127
14	Ovarian and Breast Cancer Risks Associated With Pathogenic Variants in <i>RAD51C</i> and <i>RAD51D</i> . Journal of the National Cancer Institute, 2020, 112, 1242-1250.	6.3	106
15	Prevalence of deleterious germline variants in risk genes including BRCA1/2 in consecutive ovarian cancer patients (AGO-TR-1). PLoS ONE, 2017, 12, e0186043.	2.5	105
16	High-risk breast cancer surveillance with MRI: 10-year experience from the German consortium for hereditary breast and ovarian cancer. Breast Cancer Research and Treatment, 2019, 175, 217-228.	2.5	94
17	Male breast cancer in BRCA1 and BRCA2 mutation carriers: pathology data from the Consortium of Investigators of Modifiers of BRCA1/2. Breast Cancer Research, 2016, 18, 15.	5.0	88
18	Prevalence of pathogenic BRCA1/2 germline mutations among 802 women with unilateral triple-negative breast cancer without family cancer history. BMC Cancer, 2018, 18, 265.	2.6	84

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19	Intense dose-dense epirubicin, paclitaxel, cyclophosphamideÂversus weekly paclitaxel, liposomal doxorubicin (plus carboplatin in triple-negative breast cancer) for neoadjuvant treatment of high-risk early breast cancer (GeparOcto—GBG 84): A randomised phase III trial. European Journal of Cancer, 2019, 106, 181-192.	2.8	84
20	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
21	MLPA screening in the <i>BRCA1</i> gene from 1,506 German hereditary breast cancer cases: novel deletions, frequent involvement of exon 17, and occurrence in single early-onset cases. Human Mutation, 2008, 29, 948-958.	2.5	81
22	DNA Glycosylases Involved in Base Excision Repair May Be Associated with Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. PLoS Genetics, 2014, 10, e1004256.	3.5	47
23	Risk-reducing salpingo-oophorectomy, natural menopause, and breast cancer risk: an international prospective cohort of BRCA1 and BRCA2 mutation carriers. Breast Cancer Research, 2020, 22, 8.	5.0	41
24	Association of Genomic Domains in <i>BRCA1</i> and <i>BRCA2</i> with Prostate Cancer Risk and Aggressiveness. Cancer Research, 2020, 80, 624-638.	0.9	39
25	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
26	Deleterious somatic variants in 473 consecutive individuals with ovarian cancer: results of the observational AGO-TR1 study (NCT02222883). Journal of Medical Genetics, 2019, 56, 574-580.	3.2	34
27	Oral contraceptive use and ovarian cancer risk for BRCA1/2 mutation carriers: an international cohort study. American Journal of Obstetrics and Gynecology, 2021, 225, 51.e1-51.e17.	1.3	34
28	Oral Contraceptive Use and Breast Cancer Risk: Retrospective and Prospective Analyses From a BRCA1 and BRCA2 Mutation Carrier Cohort Study. JNCI Cancer Spectrum, 2018, 2, pky023.	2.9	33
29	Breast cancer risk in <i>BRCA1/2</i> mutation carriers and noncarriers under prospective intensified surveillance. International Journal of Cancer, 2020, 146, 999-1009.	5.1	32
30	Height and Body Mass Index as Modifiers of Breast Cancer Risk in <i>BRCA1</i> / <i>2</i> Mutation Carriers: A Mendelian Randomization Study. Journal of the National Cancer Institute, 2019, 111, 350-364.	6.3	30
31	Validation of the Manchester scoring system for predicting <i>BRCA1/2</i> mutations in 9,390 families suspected of having hereditary breast and ovarian cancer. International Journal of Cancer, 2014, 135, 2352-2361.	5.1	29
32	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
33	Benefits and risks of a percutaneous endoscopic gastrostomy (PEG) for decompression in patients with malignant gastrointestinal obstruction. Supportive Care in Cancer, 2017, 25, 2849-2856.	2.2	24
34	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
35	Trastuzumab and survival of patients with metastatic breast cancer. Archives of Gynecology and Obstetrics, 2017, 296, 303-312.	1.7	20
36	Cancer surveillance and distress among adult pathogenic <i>TP53</i> germline variant carriers in Germany: A multicenter feasibility and acceptance survey. Cancer, 2020, 126, 4032-4041.	4.1	20

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37	Late onset Li-Fraumeni Syndrome with bilateral breast cancer and other malignancies: case report and review of the literature. BMC Cancer, 2012, 12, 217.	2.6	19
38	Spectrum of genetic variants of BRCA1 and BRCA2 in a German single center study. Archives of Gynecology and Obstetrics, 2017, 295, 1227-1238.	1.7	18
39	Germline truncating-mutations in BRCA1 and MSH6 in a patient with early onset endometrial cancer. BMC Cancer, 2012, 12, 531.	2.6	12
40	Prostate-specific antigen velocity in a prospective prostate cancer screening study of men with genetic predisposition. British Journal of Cancer, 2018, 118, 266-276.	6.4	12
41	Survival analysis of the randomised phase III GeparOcto trial comparing neoadjuvant chemotherapy of intense dose-dense epirubicin, paclitaxel, cyclophosphamide versus weekly paclitaxel, liposomal doxorubicin (plus carboplatin in triple-negative breast cancer) for patients with high-risk early breast cancer. Furopean Journal of Cancer. 2022, 160, 100-111.	2.8	12
42	Pathological Response in the Breast and Axillary Lymph Nodes after Neoadjuvant Systemic Treatment in Patients with Initially Node-Positive Breast Cancer Correlates with Disease Free Survival: An Exploratory Analysis of the GeparOcto Trial. Cancers, 2022, 14, 521.	3.7	12
43	Fine-Scale Mapping at 9p22.2 Identifies Candidate Causal Variants That Modify Ovarian Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. PLoS ONE, 2016, 11, e0158801.	2.5	10
44	Changes in classification of genetic variants in BRCA1 and BRCA2. Archives of Gynecology and Obstetrics, 2018, 297, 279-280.	1.7	8
45	BRCA1/2 missense mutations and the value of in-silico analyses. European Journal of Medical Genetics, 2017, 60, 572-577.	1.3	7
46	Patient-Reported Satisfaction after Prophylactic Operations of the Breast. Breast Care, 2019, 14, 217-223.	1.4	7
47	Breast cancer characteristics and surgery among women with Liâ€Fraumeni syndrome in Germany—A retrospective cohort study. Cancer Medicine, 2021, 10, 7747-7758.	2.8	7
48	Oral Contraceptive Use in <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Absolute Cancer Risks and Benefits. Journal of the National Cancer Institute, 2022, 114, 540-552.	6.3	7
49	Familial Breast Cancer - Targeted Therapy in Secondary and Tertiary Prevention. Breast Care, 2015, 10, 27-31.	1.4	5
50	Prevalence of Lynch syndrome in unselected patients with endometrial or ovarian cancer. Archives of Gynecology and Obstetrics, 2016, 294, 1299-1303.	1.7	4
51	Sensitivity and specificity of loss of heterozygosity analysis for the classification of rare germline variants in BRCA1/2: results of the observational AGO-TR1 study (NCT02222883). Journal of Medical Genetics, 2020, , jmedgenet-2020-107353.	3.2	3
52	Recommendation and Acceptance of Counselling for Familial Cancer Risk in Newly Diagnosed Breast Cancer Cases. Breast Care, 2022, 17, 153-158.	1.4	1
53	LIFESTYLE und erblicher Brustkrebs. Medizinische Genetik, 2015, 27, 237-243.	0.2	0