Dieter Niederacher

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/589559/publications.pdf

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24 papers 2,334 citations

471509 17 h-index 677142 22 g-index

24 all docs

24 docs citations

times ranked

24

5243 citing authors

#	Article	IF	CITATIONS
1	Association of Type and Location of <i>BRCA1 </i> li>and <i>BRCA2 </i> li>Mutations With Risk of Breast and Ovarian Cancer. JAMA - Journal of the American Medical Association, 2015, 313, 1347.	7.4	390
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	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
5	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
6	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
7	Cancer Risks Associated With <i>BRCA1</i> and <i>BRCA2</i> Pathogenic Variants. Journal of Clinical Oncology, 2022, 40, 1529-1541.	1.6	90
8	Challenges for CTC-based liquid biopsies: low CTC frequency and diagnostic leukapheresis as a potential solution. Expert Review of Molecular Diagnostics, 2016, 16, 147-164.	3.1	89
9	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
10	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
11	BRIP1 loss-of-function mutations confer high risk for familial ovarian cancer, but not familial breast cancer. Breast Cancer Research, 2018, 20, 7.	5.0	78
12	Association Between Loss-of-Function Mutations Within the <i>FANCM</i> Gene and Early-Onset Familial Breast Cancer. JAMA Oncology, 2017, 3, 1245.	7.1	74
13	AluY-mediated germline deletion, duplication and somatic stem cell reversion in <i>UBE2T</i> defines a new subtype of Fanconi anemia. Human Molecular Genetics, 2015, 24, 5093-5108.	2.9	62
14	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
15	A Novel Workflow to Enrich and Isolate Patient-Matched EpCAMhigh and EpCAMlow/negative CTCs Enables the Comparative Characterization of the PIK3CA Status in Metastatic Breast Cancer. International Journal of Molecular Sciences, 2017, 18, 1885.	4.1	37
16	Frequent allele loss on 9p21–22 defines a smallest common region in the vicinity of theCDKN2 gene in sporadic breast cancer. , 1996, 17, 14-20.		34
17	Resistance to CD95-mediated apoptosis in breast cancer is not due to somatic mutation of the CD95 gene. International Journal of Cancer, 2001, 92, 309-310.	5.1	25
18	Breast and Prostate Cancer Risks for Male <i>BRCA1</i> BRCA1li>BRCA2Pathogenic Variant Carriers Using Polygenic Risk Scores. Journal of the National Cancer Institute, 2022, 114, 109-122.	6.3	19

#	Article	IF	CITATION
19	EZH2 Loss Drives Resistance to Carboplatin and Paclitaxel in Serous Ovarian Cancers Expressing ATM. Molecular Cancer Research, 2020, 18, 278-286.	3.4	12
20	Prevalence of Cancer Predisposition Germline Variants in Male Breast Cancer Patients: Results of the German Consortium for Hereditary Breast and Ovarian Cancer. Cancers, 2022, 14, 3292.	3.7	11
21	Evaluation of HER2 expression in urothelial carcinoma cells as a biomarker for circulating tumor cells. Cytometry Part B - Clinical Cytometry, 2020, 98, 355-367.	1.5	10
22	Predictors of Impaired Postpartum Renal Function in Women after Preeclampsia: Results of a Prospective Single Center Study. Disease Markers, 2016, 2016, 1-8.	1.3	8
23	The <i>GPRC5A</i> frameshift variant c.183del is not associated with increased breast cancer risk in <i>BRCA1</i> mutation carriers. International Journal of Cancer, 2019, 144, 1761-1763.	5.1	2
24	Implementing microwell slides for detection and isolation of single circulating tumor cells from complex cell suspensions. Cytometry Part A: the Journal of the International Society for Analytical Cytology, 2022, 101, 1057-1067.	1.5	1