# Mary B Daly

List of Publications by Year
in descending order



| | | | |
|---|---|---|---|
| 101 papers | 8,162 citations | 109137 35 h-index | 56606 83 g-index |
| 104 all docs | 104 docs citations | 104 times ranked | 10916 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Maternal and prenatal factors and age at thelarche in the LEGACY Girls Study cohort: implications for breast cancer risk. International Journal of Epidemiology, 2023, 52, 272-283. | 0.9 | 1 |
| 2 | Cancer Risks Associated With BRCA1 and BRCA2 Pathogenic Variants. Journal of Clinical Oncology, 2022, 40, 1529-1541. | 0.8 | 90 |
| 3 | Polygenic risk modeling for prediction of epithelial ovarian cancer risk. European Journal of Human Genetics, 2022, 30, 349-362. | 1.4 | 23 |
| 4 | Common variants in breast cancer risk loci predispose to distinct tumor subtypes. Breast Cancer Research, 2022, 24, 2. | 2.2 | 15 |
| 5 | OUP accepted manuscript. International Journal of Epidemiology, 2022, , . | 0.9 | 0 |
| 6 | Therapeutic implications of germline vulnerabilities in DNA repair for precision oncology. Cancer Treatment Reviews, 2022, 104, 102337. | 3.4 | 6 |
| 7 | Investigating the impact of the COVID-19 pandemic on breast cancer cliniciansâ€™ communication about sexual health. Supportive Care in Cancer, 2022, 30, 5801-5810. | 1.0 | 4 |
| 8 | Coping With Changes to Sex and Intimacy After a Diagnosis of Metastatic Breast Cancer: Results From a Qualitative Investigation With Patients and Partners. Frontiers in Psychology, 2022, 13, 864893. | 1.1 | 4 |
| 9 | Weight is More Informative than Body Mass Index for Predicting Postmenopausal Breast Cancer Risk: Prospective Family Study Cohort (ProF-SC). Cancer Prevention Research, 2022, 15, 185-191. | 0.7 | 4 |
| 10 | Comparing 5-Year and Lifetime Risks of Breast CancerÂusing the Prospective Family Study Cohort. Journal of the National Cancer Institute, 2021, 113, 785-791. | 3.0 | 13 |
| 11 | Association of Risk-Reducing Salpingo-Oophorectomy With Breast Cancer Risk in Women With BRCA1 and BRCA2 Pathogenic Variants. JAMA Oncology, 2021, 7, 585-592. | 3.4 | 16 |
| 12 | The predictive ability of the 313 variantâ€“based polygenic risk score for contralateral breast cancer risk prediction in women of European ancestry with a heterozygous BRCA1 or BRCA2 pathogenic variant. Genetics in Medicine, 2021, 23, 1726-1737. | 1.1 | 16 |
| 13 | Functional annotation of the 2q35 breast cancer risk locus implicates a structural variant in influencing activity of a long-range enhancer element. American Journal of Human Genetics, 2021, 108, 1190-1203. | 2.6 | 6 |
| 14 | Genetic risk assessment for hereditary renal cell carcinoma: Clinical consensus statement. Cancer, 2021, 127, 3957-3966. | 2.0 | 11 |
| 15 | Cascade Genetic Testing for Hereditary Cancer Risk: An Underutilized Tool for Cancer Prevention. JCO Precision Oncology, 2021, 5, 1387-1396. | 1.5 | 23 |
| 16 | Association of germline genetic variants with breast cancer-specific survival in patient subgroups defined by clinic-pathological variables related to tumor biology and type of systemic treatment. Breast Cancer Research, 2021, 23, 86. | 2.2 | 7 |
| 17 | Mendelian randomisation study of smoking exposure in relation to breast cancer risk. British Journal of Cancer, 2021, 125, 1135-1145. | 2.9 | 9 |
| 18 | Mobile Technology-Based (mLearning) Intervention to Enhance Breast Cancer Clinicians' Communication About Sexual Health: A Pilot Trial. Journal of the National Comprehensive Cancer Network: JNCCN, 2021, , . | 2.3 | 7 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Germline variants and breast cancer survival in patients with distant metastases at primary breast cancer diagnosis. Scientific Reports, 2021, 11, 19787. | 1.6 | 2 |
| 20 | Recreational Physical Activity and Outcomes After Breast Cancer in Women at High Familial Risk. JNCI Cancer Spectrum, 2021, 5, pkab090. | 1.4 | 1 |
| 21 | Association of Genomic Domains in BRCA1 and BRCA2 with Prostate Cancer Risk and Aggressiveness. Cancer Research, 2020, 80, 624-638. | 0.4 | 39 |
| 22 | Considerations When Using Breast Cancer Risk Models for Women with Negative BRCA1/BRCA2 Mutation Results. Journal of the National Cancer Institute, 2020, 112, 418-422. | 3.0 | 1 |
| 23 | Recreational Physical Activity Is Associated with Reduced Breast Cancer Risk in Adult Women at High Risk for Breast Cancer: A Cohort Study of Women Selected for Familial and Genetic Risk. Cancer Research, 2020, 80, 116-125. | 0.4 | 37 |
| 24 | Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. Nature Genetics, 2020, 52, 56-73. | 9.4 | 120 |
| 25 | Points to consider: is there evidence to support BRCA1/2 and other inherited breast cancer genetic testing for all breast cancer patients? A statement of the American College of Medical Genetics and Genomics (ACMG). Genetics in Medicine, 2020, 22, 681-685. | 1.1 | 20 |
| 26 | 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. | 1.1 | 82 |
| 27 | Navigating the Intersection between Genomic Research and Clinical Practice. Cancer Prevention Research, 2020, 13, 219-222. | 0.7 | 4 |
| 28 | Breast Cancer Polygenic Risk Score and Contralateral Breast Cancer Risk. American Journal of Human Genetics, 2020, 107, 837-848. | 2.6 | 39 |
| 29 | Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses. Nature Genetics, 2020, 52, 572-581. | 9.4 | 265 |
| 30 | Longitudinal follow-up after telephone disclosure in the randomized COGENT study. Genetics in Medicine, 2020, 22, 1401-1406. | 1.1 | 4 |
| 31 | Germline HOXB13 mutations p.G84E and p.R217C do not confer an increased breast cancer risk. Scientific Reports, 2020, 10, 9688. | 1.6 | 2 |
| 32 | Transcriptomeâ€wide association study of breast cancer risk by estrogenâ€receptor status. Genetic Epidemiology, 2020, 44, 442-468. | 0.6 | 32 |
| 33 | Evaluating a couple-based intervention addressing sexual concerns for breast cancer survivors: study protocol for a randomized controlled trial. Trials, 2020, 21, 173. | 0.7 | 18 |
| 34 | Alcohol Consumption, Cigarette Smoking, and Risk of Breast Cancer for BRCA1 and BRCA2 Mutation Carriers: Results from The BRCA1 and BRCA2 Cohort Consortium. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 368-378. | 1.1 | 24 |
| 35 | A network analysis to identify mediators of germline-driven differences in breast cancer prognosis. Nature Communications, 2020, 11, 312. | 5.8 | 30 |
| 36 | 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. | 2.2 | 41 |

| # | Article | IF | Citations |
|---|---|---|---|
| 37 | Two truncating variants in FANCC and breast cancer risk. Scientific Reports, 2019, 9, 12524. | 1.6 | 5 |
| 38 | Breast Cancer Chemopreventionâ€"Can We Make a Case for Precision Medicine?. JAMA Oncology, 2019, 5, 1542. | 3.4 | 2 |
| 39 | Shared heritability and functional enrichment across six solid cancers. Nature Communications, 2019, 10, 431. | 5.8 | 88 |
| 40 | Mendelian randomisation study of height and body mass index as modifiers of ovarian cancer risk in 22,588 BRCA1 and BRCA2 mutation carriers. British Journal of Cancer, 2019, 121, 180-192. | 2.9 | 19 |
| 41 | BRCA1andBRCA2pathogenic sequence variants in women of African origin or ancestry. Human Mutation, 2019, 40, 1781-1796. | 1.1 | 26 |
| 42 | Regular use of aspirin and other non-steroidal anti-inflammatory drugs and breast cancer risk for women at familial or genetic risk: a cohort study. Breast Cancer Research, 2019, 21, 52. | 2.2 | 44 |
| 43 | Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. Nature Communications, 2019, 10, 1741. | 5.8 | 90 |
| 44 | Genome-wide association study of germline variants and breast cancer-specific mortality. British Journal of Cancer, 2019, 120, 647-657. | 2.9 | 52 |
| 45 | Association of Prepubertal and Adolescent Androgen Concentrations With Timing of Breast Development and Family History of Breast Cancer. JAMA Network Open, 2019, 2, e190083. | 2.8 | 7 |
| 46 | Benign breast disease increases breast cancer risk independent of underlying familial risk profile: Findings from a Prospective Family Study Cohort. International Journal of Cancer, 2019, 145, 370-379. | 2.3 | 9 |
| 47 | A brief intervention to enhance breast cancer clinicians' communication about sexual health: Feasibility, acceptability, and preliminary outcomes. Psycho-Oncology, 2019, 28, 872-879. | 1.0 | 21 |
| 48 | 10-year performance of four models of breast cancer risk: a validation study. Lancet Oncology, The, 2019, 20, 504-517. | 5.1 | 116 |
| 49 | Alcohol consumption, cigarette smoking, and familial breast cancer risk: findings from the Prospective Family Study Cohort (ProF-SC). Breast Cancer Research, 2019, 21, 128. | 2.2 | 27 |
| 50 | Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. American Journal of Human Genetics, 2019, 104, 21-34. | 2.6 | 711 |
| 51 | Preferences for inâ€person disclosure: Patients declining telephone disclosure characteristics and outcomes in the multicenter Communication Of GENetic Test Results by Telephone study. Clinical Genetics, 2019, 95, 293-301. | 1.0 | 16 |
| 52 | Patient-clinician communication about sexual health in breast cancer: A mixed-methods analysis of clinic dialogue. Patient Education and Counseling, 2019, 102, 436-442. | 1.0 | 38 |
| 53 | Height and Body Mass Index as Modifiers of Breast Cancer Risk in BRCA1/2 Mutation Carriers: A Mendelian Randomization Study. Journal of the National Cancer Institute, 2019, 111, 350-364. | 3.0 | 30 |
| 54 | Risk-Reducing Oophorectomy and Breast Cancer Risk Across the Spectrum of Familial Risk. Journal of the National Cancer Institute, 2019, 111, 331-334. | 3.0 | 31 |

| # | Article | IF | Citations |
|---|---|---|---|
| 55 | Prostate cancer genetic testing: NCCN familial high-risk assessment: breast/ovarian. Canadian Journal of Urology, 2019, 26, 29-30. | 0.0 | 2 |
| 56 | Randomized Noninferiority Trial of Telephone vs In-Person Disclosure of Germline Cancer Genetic Test Results. Journal of the National Cancer Institute, 2018, 110, 985-993. | 3.0 | 35 |
| 57 | Breast cancer family history and allele-specific DNA methylation in the legacy girls study. Epigenetics, 2018, 13, 240-250. | 1.3 | 10 |
| 58 | Comparison of methods to assess onset of breast development in the LEGACY Girls Study: methodological considerations for studies of breast cancer. Breast Cancer Research, 2018, 20, 33. | 2.2 | 9 |
| 59 | Use and Patient-Reported Outcomes of Clinical Multigene Panel Testing for Cancer Susceptibility in the Multicenter Communication of Genetic Test Results by Telephone Study. JCO Precision Oncology, 2018, 2, 1-12. | 1.5 | 10 |
| 60 | Age-specific breast cancer risk by body mass index and familial risk: prospective family study cohort (ProF-SC). Breast Cancer Research, 2018, 20, 132. | 2.2 | 51 |
| 61 | 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. | 0.4 | 54 |
| 62 | A transcriptome-wide association study of 229,000 women identifies new candidate susceptibility genes for breast cancer. Nature Genetics, 2018, 50, 968-978. | 9.4 | 184 |
| 63 | Breast cancer risk prediction using a polygenic risk score in the familial setting: a prospective study from the Breast Cancer Family Registry and kConFab. Genetics in Medicine, 2017, 19, 30-35. | 1.1 | 53 |
| 64 | Dietary isoflavone intake and allâ€cause mortality in breast cancer survivors: The Breast Cancer Family Registry. Cancer, 2017, 123, 2070-2079. | 2.0 | 67 |
| 65 | Multigene Panel Testing and Breast Cancer Risk. JAMA Oncology, 2017, 3, 1176. | 3.4 | 8 |
| 66 | Effective patient-provider communication about sexual concerns in breast cancer: a qualitative study. Supportive Care in Cancer, 2017, 25, 3199-3207. | 1.0 | 63 |
| 67 | Risks of Breast, Ovarian, and Contralateral Breast Cancer for BRCA1 and BRCA2 Mutation Carriers. JAMA - Journal of the American Medical Association, 2017, 317, 2402. | 3.8 | 1,898 |
| 68 | LincIN, a novel NF90-binding long non-coding RNA, is overexpressed in advanced breast tumors and involved in metastasis. Breast Cancer Research, 2017, 19, 62. | 2.2 | 36 |
| 69 | Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691. | 9.4 | 356 |
| 70 | Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94. | 13.7 | 1,099 |
| 71 | Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778. | 9.4 | 289 |
| 72 | The Protein Encoded by the CCDC170 Breast Cancer Gene Functions to Organize the Golgi-Microtubule Network. EBioMedicine, 2017, 22, 28-43. | 2.7 | 26 |

| # | Article | IF | Citations |
|---|---|---|---|
| 73 | Talking about women's sexual health after cancer: Why is it so hard to move the needle?. Cancer, 2017, 123, 4757-4763. | 2.0 | 34 |
| 74 | Non-invasive optical spectroscopic monitoring of breast development during puberty. Breast Cancer Research, 2017, 19, 12. | 2.2 | 14 |
| 75 | Pubertal development in girls by breast cancer family history: the LEGACY girls cohort. Breast Cancer Research, 2017, 19, 69. | 2.2 | 18 |
| 76 | Patient-provider communication about sexual concerns in cancer: a systematic review. Journal of Cancer Survivorship, 2017, 11, 175-188. | 1.5 | 123 |
| 77 | Inherited Mutations in Men Undergoing Multigene Panel Testing for Prostate Cancer: Emerging Implications for Personalized Prostate Cancer Genetic Evaluation. JCO Precision Oncology, 2017, 1, 1-17. | 1.5 | 27 |
| 78 | Uterine Cancer After Risk-Reducing Salpingo-oophorectomy Without Hysterectomy in Women With BRCA Mutations. JAMA Oncology, 2016, 2, 1434. | 3.4 | 189 |
| 79 | 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. | 2.2 | 31 |
| 80 | 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. | 2.2 | 88 |
| 81 | Psychosocial Adjustment and Perceived Risk Among Adolescent Girls From Families With BRCA1/2 or Breast Cancer History. Journal of Clinical Oncology, 2016, 34, 3409-3416. | 0.8 | 16 |
| 82 | Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. Nature Communications, 2016, 7, 11375. | 5.8 | 93 |
| 83 | Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breastâ€“ovarian cancer susceptibility locus. Nature Communications, 2016, 7, 12675. | 5.8 | 78 |
| 84 | Communicating genetic test results within the family: Is it lost in translation? A survey of relatives in the randomized six-step study. Familial Cancer, 2016, 15, 697-706. | 0.9 | 68 |
| 85 | Cohort Profile: The Breast Cancer Prospective Family Study Cohort (ProF-SC). International Journal of Epidemiology, 2016, 45, 683-692. | 0.9 | 48 |
| 86 | Family Communication of Genetic Risk: A Personalized Approach. Current Genetic Medicine Reports, 2016, 4, 35-40. | 1.9 | 5 |
| 87 | Comparison of Clinical, Maternal, and Self Pubertal Assessments: Implications for Health Studies. Pediatrics, 2016, 138, . | 1.0 | 36 |
| 88 | Breast cancer risk variants at 6q25 display different phenotype associations and regulate ESR1, RMND1 and CCDC170. Nature Genetics, 2016, 48, 374-386. | 9.4 | 125 |
| 89 | Utilizing Remote Real-Time Videoconferencing to Expand Access to Cancer Genetic Services in Community Practices: A Multicenter Feasibility Study. Journal of Medical Internet Research, 2016, 18, e23. | 2.1 | 79 |
| 90 | Assessing Associations between the AURKA-HMMR-TPX2-TUBG1 Functional Module and Breast Cancer Risk in BRCA1/2 Mutation Carriers. PLoS ONE, 2015, 10, e0120020. | 1.1 | 34 |

| # | Article | IF | Citations |
|---|---|---|---|
| 91 | Psychosocial Adjustment in School-age Girls With a Family History of Breast Cancer. Pediatrics, 2015, 136, 927-937. | 1.0 | 13 |
| 92 | Salpingectomy as a Means to Reduce Ovarian Cancer Risk. Cancer Prevention Research, 2015, 8, 342-348. | 0.7 | 54 |
| 93 | Germline Mutation in BRCA1 or BRCA2 and Ten-Year Survival for Women Diagnosed with Epithelial Ovarian Cancer. Clinical Cancer Research, 2015, 21, 652-657. | 3.2 | 138 |
| 94 | Development of a tiered and binned genetic counseling model for informed consent in the era of multiplex testing for cancer susceptibility. Genetics in Medicine, 2015, 17, 485-492. | 1.1 | 79 |
| 95 | Correlation of DNA methylation levels in blood and saliva DNA in young girls of the LEGACY Girls study. Epigenetics, 2014, 9, 929-933. | 1.3 | 32 |
| 96 | Human Subjects Protection: An Event Monitoring Committee for Research Studies of Girls From Breast Cancer Families. Journal of Adolescent Health, 2014, 55, 352-357. | 1.2 | 5 |
| 97 | Identifying a Highly-Aggressive DCIS Subgroup by Studying Intra-Individual DCIS Heterogeneity among Invasive Breast Cancer Patients. PLoS ONE, 2014, 9, e100488. | 1.1 | 21 |
| 98 | Development of a Communication Protocol for Telephone Disclosure of Genetic Test Results for Cancer Predisposition. JMIR Research Protocols, 2014, 3, e49. | 0.5 | 26 |
| 99 | Implementation and outcomes of telephone disclosure of clinical BRCA1/2 test results. Patient Education and Counseling, 2013, 93, 413-419. | 1.0 | 26 |
| 100 | Delivery of Internet-based cancer genetic counselling services to patients' homes: a feasibility study. Journal of Telemedicine and Telecare, 2011, 17, 36-40. | 1.4 | 47 |
| 101 | Women's thoughts on receiving and sharing genetic information: Considerations for genetic counseling. Journal of Genetic Counseling, 0, , . | 0.9 | 1 |