

## ORIGINAL ARTICLE

# A Population-Based Study of Genes Previously Implicated in Breast Cancer

C. Hu, S.N. Hart, R. Gnanaolivu, H. Huang, K.Y. Lee, J. Na, C. Gao, J. Lilyquist, S. Yadav, N.J. Boddicker, R. Samara, J. Klebba, C.B. Ambrosone, H. Anton-Culver, P. Auer, E.V. Bandera, L. Bernstein, K.A. Bertrand, E.S. Burnside, B.D. Carter, H. Eliassen, S.M. Gapstur, M. Gaudet, C. Haiman, J.M. Hodge, D.J. Hunter, E.J. Jacobs, E.M. John, C. Kooperberg, A.W. Kurian, L. Le Marchand, S. Lindstrom, T. Lindstrom, H. Ma, S. Neuhausen, P.A. Newcomb, K.M. O'Brien, J.E. Olson, I.M. Ong, T. Pal, J.R. Palmer, A.V. Patel, S. Reid, L. Rosenberg, D.P. Sandler, C. Scott, R. Tamimi, J.A. Taylor, A. Trentham-Dietz, C.M. Vachon, C. Weinberg, S. Yao, A. Ziogas, J.N. Weitzel, D.E. Goldgar, S.M. Domchek, K.L. Nathanson, P. Kraft, E.C. Polley, and F.J. Couch

## ABSTRACT

**BACKGROUND**

Population-based estimates of the risk of breast cancer associated with germline pathogenic variants in cancer-predisposition genes are critically needed for risk assessment and management in women with inherited pathogenic variants.

**METHODS**

In a population-based case-control study, we performed sequencing using a custom multigene amplicon-based panel to identify germline pathogenic variants in 28 cancer-predisposition genes among 32,247 women with breast cancer (case patients) and 32,544 unaffected women (controls) from population-based studies in the Cancer Risk Estimates Related to Susceptibility (CARRIERS) consortium. Associations between pathogenic variants in each gene and the risk of breast cancer were assessed.

**RESULTS**

Pathogenic variants in 12 established breast cancer-predisposition genes were detected in 5.03% of case patients and in 1.63% of controls. Pathogenic variants in *BRCA1* and *BRCA2* were associated with a high risk of breast cancer, with odds ratios of 7.62 (95% confidence interval [CI], 5.33 to 11.27) and 5.23 (95% CI, 4.09 to 6.77), respectively. Pathogenic variants in *PALB2* were associated with a moderate risk (odds ratio, 3.83; 95% CI, 2.68 to 5.63). Pathogenic variants in *BARD1*, *RAD51C*, and *RAD51D* were associated with increased risks of estrogen receptor-negative breast cancer and triple-negative breast cancer, whereas pathogenic variants in *ATM*, *CDH1*, and *CHEK2* were associated with an increased risk of estrogen receptor-positive breast cancer. Pathogenic variants in 16 candidate breast cancer-predisposition genes, including the c.657\_661del5 founder pathogenic variant in *NBN*, were not associated with an increased risk of breast cancer.

**CONCLUSIONS**

This study provides estimates of the prevalence and risk of breast cancer associated with pathogenic variants in known breast cancer-predisposition genes in the U.S. population. These estimates can inform cancer testing and screening and improve clinical management strategies for women in the general population with inherited pathogenic variants in these genes. (Funded by the National Institutes of Health and the Breast Cancer Research Foundation.)

The authors' full names, academic degrees, and affiliations are listed in the Appendix. Address reprint requests to Dr. Couch at the Department of Laboratory Medicine and Pathology, Mayo Clinic, Stable 2-42, 200 First St. SW, Rochester, MN 55905, or at couch.fergus@mayo.edu.

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**G**ERMLINE PATHOGENIC VARIANTS IN cancer-predisposition genes included in hereditary cancer multigene testing panels have been associated with an increased risk of breast cancer.<sup>1-4</sup> Identification of pathogenic variants in predisposition genes has provided benefit through improving access to risk-reducing prophylactic surgery and targeted therapies among carriers of pathogenic variants in *BRCA1* and *BRCA2* and access to enhanced mammography and magnetic resonance imaging (MRI)-based screening among carriers of pathogenic variants in several established breast cancer-predisposition genes.<sup>5,6</sup> The aggregate prevalence of pathogenic variants in these genes has been estimated at 7 to 10% among women with breast cancer.<sup>1,7-10</sup> However, these prevalences and associated risks of breast cancer are based on high-risk populations enriched with women who had a family history of breast and ovarian cancers, received a breast cancer diagnosis at a young age, had estrogen receptor (ER)-negative tumors, or had founder mutations. Only studies of limited size have evaluated pathogenic variants in multigene panels in women with breast cancer unselected for family history or age at diagnosis.<sup>11,12</sup> Thus, current risk estimates of breast cancer with respect to predisposition genes have uncertain application to the general population.<sup>13</sup>

Genetic testing recommendations have been developed to provide guidance on the selection of women for multigene panel testing. The U.S. Preventive Services Task Force has suggested that the selection of unaffected women for testing be based on risk stratification.<sup>14</sup> The National Comprehensive Cancer Network has also suggested that risk stratification be used in the selection of unaffected and affected women for testing.<sup>15</sup> In contrast, the American Society of Breast Surgeons has recommended that germline genetic testing for hereditary cancer be performed in all women with breast cancer. Separately, population-based screening for *BRCA1* and *BRCA2* in all women older than 30 years of age has been proposed.<sup>16</sup> However, large-scale population-based studies that provide estimates of the prevalence of pathogenic variants in predisposition genes in the general population are lacking.

Cancer Risk Estimates Related to Susceptibility (CARRIERS) is a United States-based consortium consisting of population-based and family-based studies of breast cancer. Here, we used the

population-based studies in the CARRIERS consortium to estimate the prevalence and risk of breast cancer associated with pathogenic variants in breast cancer-predisposition genes in the U.S. general population.

## METHODS

### STUDY POPULATION

The CARRIERS consortium includes 17 studies — 7 nested case-control studies in prospective cohorts, 2 case-cohort studies in prospective cohorts, 3 case-control studies, and 5 case-control or case-cohort studies enriched with women with early-onset disease or a family history of breast cancer (Table S1 in the Supplementary Appendix, available with the full text of this article at NEJM.org). The characteristics of the 39,553 women with breast cancer (case patients) and 35,867 study-matched unaffected women (controls) are provided in Table S2. Population-based estimates were derived from 32,247 case patients and 32,544 controls from the 12 studies in the CARRIERS consortium that were not enriched with patients with a family history or early onset of disease (the Black Women's Health Study, the Cancer Prevention Study II, the Cancer Prevention Study 3, the California Teachers' Study, the Mayo Clinic Breast Cancer Study, the Multiethnic Cohort Study, the Mayo Mammography Health Study, the Nurses' Health Study, the Nurses' Health Study II, the Women's Circle of Health Study, the Women's Health Initiative, and the Wisconsin Women's Health Study). All participants provided informed consent for research. The CARRIERS study was approved by the institutional review board at the Mayo Clinic.

### DNA SEQUENCING AND BIOINFORMATICS ANALYSIS

Germline DNA samples were subjected to dual bar-coded QIAseq (Qiagen) multiplex amplicon-based analysis of 746 target regions in 37 cancer-predisposition genes.<sup>17</sup> Libraries from 768 samples were pooled and sequenced in each lane of a HiSeq 4000 system (Illumina). Genetic variants were identified with the use of the Genome Analysis Toolkit (GATK) Haplotype Caller tool and VarDict variant caller tool. High-quality sequence data (read depth of >20 times) were obtained for 99.3% of the target regions. Twenty-eight cancer predisposition genes including 12 established breast cancer-predisposition genes (*ATM*, *BARD1*,

*BRCA1*, *BRCA2*, *CDH1*, *CHEK2*, *NF1*, *PALB2*, *PTEN*, *RAD51C*, *RAD51D*, and *TP53*) and 16 candidate predisposition genes were evaluated (Table S3).<sup>18–</sup>

<sup>30</sup> Loss-of-function variants and variants identified as “pathogenic” or “likely pathogenic” in the ClinVar database were classified as pathogenic variants (see the Supplementary Appendix).<sup>17</sup> Pathogenic variants in *NF1* and *TP53* were restricted to those with an alternate allele fraction (calculated as the number of alternate allele reads divided by the total number of reads at a specific genomic position) between 0.3 and 0.7 in an effort to exclude potential clonal hematopoiesis variants.<sup>31</sup>

#### STATISTICAL ANALYSIS

Prevalences of pathogenic variants and variants of uncertain significance in each gene were tabulated for the case patients and controls in the population-based CARRIERS analysis, and 95% confidence intervals were estimated with the use of the Wilson score method without continuity correction. A generalized additive model for pathogenic-variant status and a smoothing spline function for age<sup>32</sup> were used to estimate the relationship between the prevalence of a pathogenic variant and age. Associations between pathogenic variants in each gene and the risk of breast cancer were assessed by means of logistic regression, with adjustment for study, age, first-degree family history of breast cancer, and race or ethnic group. Sensitivity analyses were conducted to assess the effect of each of the 12 studies with the use of a leave-one-study-out cross-validation approach. Comparisons between unaffected controls and women with ER-positive cancer, women with ER-negative cancer, and women with triple-negative breast cancer (ER-negative, progesterone receptor [PR]–negative, and human epidermal growth factor receptor type 2 [HER2]–negative) were conducted with the use of binomial logistic-regression models. All analyses were performed with R software (version 3.5.2), and all tests were two-sided. Lifetime absolute risk of breast cancer to age 85 years was estimated for pathogenic-variant carriers by combining age-specific odds ratio estimates with age-specific breast cancer incidence rates from the Surveillance, Epidemiology, and End Results (SEER) Program of the National Cancer Institute (Supplementary Appendix).

## RESULTS

### PARTICIPANT CHARACTERISTICS

The distributions of age at diagnosis for 39,553 case patients and age at the time of selection into the study for 35,867 controls from the 17 studies in the CARRIERS consortium are shown in Figure S1. In the 12 population-based studies, the mean age at the time of breast cancer diagnosis among 32,247 case patients was 62.1 years, and the mean age at the time of enrollment among 32,544 controls was 61.2 years, ages that are similar to those derived from the SEER 18 registries (Table 1 and Table S4). A family history of breast cancer was reported in 20.4% of case patients and 14.3% of controls (Table 1). Among the case patients with available data on tumor biomarkers, 82.9% had ER-positive breast cancer and 11.3% had triple-negative breast cancer (Table 1), prevalences that are consistent with those derived from the SEER 18 registries. Data on HER2 status were available for only 41.1% of tumors.

### PREVALENCE OF PATHOGENIC VARIANTS IN PREDISPOSITION GENES

In the overall CARRIERS analysis that included data from all 17 studies, the prevalence of pathogenic variants in 12 established breast cancer–predisposition genes (*ATM*, *BARD1*, *BRCA1*, *BRCA2*, *CDH1*, *CHEK2*, *NF1*, *PALB2*, *PTEN*, *RAD51C*, *RAD51D*, and *TP53*) was 5.67% (95% confidence interval [CI], 5.44 to 5.90) among case patients and 1.73% (95% CI, 1.60 to 1.87) among controls (Tables S5 and S6). However, in the population-based CARRIERS analysis, the prevalence was 5.03% (95% CI, 4.79 to 5.27) among cases patients and 1.63% (95% CI, 1.50 to 1.78) among controls (Table 2). The prevalence of pathogenic variants was similar among non-Hispanic White, non-Hispanic Black, and Hispanic case patients and controls (Table S7). A lower overall prevalence of pathogenic variants was detected among Asian American case patients (1.64%; 95% CI, 1.07 to 2.49) (Table S7). Among the case patients, the highest prevalence of pathogenic variants was observed for *BRCA2* (1.29%; 95% CI, 1.18 to 1.42), *CHEK2* (1.08%; 95% CI, 0.98 to 1.20), and *BRCA1* (0.85%; 95% CI, 0.76 to 0.96) (Table 2). *BRCA1* carriers had a relatively young mean ( $\pm$ SD) age at diagnosis (50.9 $\pm$ 13.3 years among those

with ER-positive status and 50.3±12.4 among those with ER-negative status), whereas *BRCA2* carriers had a slightly older mean age at diagnosis (55.4±12.8 years among those with ER-positive status and 58.6±12.2 among those with ER-negative status) (Table S8). The prevalence of variants of uncertain significance in the 12 established breast cancer genes was 18.9% (95% CI, 18.5 to 19.4) among case patients and 18.5% (95% CI, 18.1 to 19.0) among controls (Table S9).

#### **PATHOGENIC VARIANTS IN PREDISPOSITION GENES AND BREAST CANCER RISK**

Case-control association analyses, with adjustment for study, age, family history of breast cancer, and race or ethnic group, were performed with data from all 17 studies in the CARRIERS consortium and with data from the 12 population-based studies in the CARRIERS consortium (Table 2 and Table S5 and S6). In the population-based studies, *BRCA1* and *BRCA2* were associated with a high risk of breast cancer, with odds ratios of 7.62 (95% CI, 5.33 to 11.27) and 5.23 (95% CI, 4.09 to 6.77), respectively (Table 2 and Table S10). Pathogenic variants in *PALB2* and *CHEK2* were associated with a moderate risk, with odds ratios of 3.83 (95% CI, 2.68 to 5.63) and 2.47 (95% CI, 2.02 to 3.05), respectively. The common *CHEK2* pathogenic variants p.Ile157Thr and p.Ser428Phe had limited clinical importance (i.e., odds ratio, <1.5), with odds ratios of 1.30 (95% CI, 1.06 to 1.59; *P*=0.01) and 1.26 (95% CI, 0.76 to 2.12; *P*=0.37), respectively, and were excluded from the analyses. Pathogenic variants in *ATM* and *NF1* were associated with an increased risk of breast cancer, with odds ratios of 1.82 (95% CI, 1.46 to 2.27) and 1.93 (95% CI, 0.91 to 4.31), respectively. Pathogenic variants in *BARD1*, *RAD51C*, and *RAD51D* were associated with a moderate risk of ER-negative breast cancer and triple-negative breast cancer but not ER-positive breast cancer, whereas pathogenic variants in *ATM*, *CDH1*, and *CHEK2* were associated only with ER-positive breast cancer (Table 3). Limited numbers of women with pathogenic variants in *PTEN* and *TP53* did not allow us to assess associations with breast cancer (Tables 2 and 3). Sensitivity analyses verified that individual studies did not influence associations with breast cancer risk (Fig. S2).

None of the 16 candidate predisposition genes,

including the mismatch repair genes (*MLH1*, *MSH2*, and *MSH6*), were significantly associated with increased risk of breast cancer in analyses involving the participants overall (Table S10) or the participants stratified according to ER status (Table S11) (*P*>0.05 for all). An increased risk of breast cancer was not observed among the participants with any pathogenic variant in *NBN* (odds ratio, 1.05; 95% CI, 0.71 to 1.56) or among those with the *NBN* pathogenic variant c.657\_661del5 (odds ratio, 0.93; 95% CI, 0.52 to 1.68), which was previously associated with breast cancer, or those with c.657\_661del5 who were homozygous for the GG allele of the c.553 (Table S10).<sup>33,34</sup>

To investigate the influence of a family history of breast cancer on associations with breast cancer risk in the general population, analyses were conducted separately for the pathogenic-variant carriers with (20.4%) or without (79.6%) a first-degree relative with breast cancer. Among the participants with a family history, pathogenic variants in *BRCA1*, *BRCA2*, *CDH1*, and *PALB2* were associated with a high risk of breast cancer (odds ratio, >4) and pathogenic variants in *ATM* and *RAD51D* were associated with a moderate risk (odds ratio, >2) (Table S12). The influence of age at breast cancer diagnosis on associations was also evaluated.<sup>35</sup> With respect to the pathogenic variants in the 12 established predisposition genes, the associations with breast cancer risk were unchanged among the participants who received a breast cancer diagnosis at an age of 50 years or younger, except among those with pathogenic variants in *BRCA1* or *BRCA2*, who were at an increased risk (Table S13). Among the participants who received a diagnosis of breast cancer at an age of more than 50 years, the associations with breast cancer observed for pathogenic variants in most genes were similar to those among the participants overall (Table S14). To assess the influence of older age at enrollment, associations with breast cancer were evaluated in the studies in which women younger than 45 years of age were eligible for enrollment; the results did not differ from those of the population-based CARRIERS analysis (Table S15).

#### **LIFETIME ABSOLUTE RISK OF BREAST CANCER**

The prevalences of pathogenic variants in the commonly mutated genes *ATM*, *BRCA1*, *BRCA2*, *CHEK2*, and *PALB2* were assessed among the case

**Table 1. Demographic and Clinical Characteristics of the Participants in the Population-Based CARRIERS Analysis.\***

Variable	Case Patients (N = 32,247)	Controls (N = 32,544)
<b>Demographic characteristic</b>		
Age†		
Mean — yr	62.07±11.44	61.22±11.82
Range — yr	21.00–94.00	21.80–94.30
Distribution — no./total no. (%)		
≤40 yr	1,099/31,708 (3.5)	1,599/32,542 (4.9)
41–50 yr	4,197/31,708 (13.2)	4,443/32,542 (13.7)
51–60 yr	7,999/31,708 (25.2)	8,580/32,542 (26.4)
61–70 yr	10,357/31,708 (32.7)	10,095/32,542 (31.0)
>70 yr	8,056/31,708 (25.4)	7,825/32,542 (24.0)
Race or ethnic group — no./total no. (%)‡		
Asian	1,282/32,068 (4.0)	1,269/32,498 (3.9)
Non-Hispanic Black	3,946/32,068 (12.3)	4,954/32,498 (15.2)
Hispanic	1,019/32,068 (3.2)	998/32,498 (3.1)
Non-Hispanic White	25,287/32,068 (78.9)	24,770/32,498 (76.2)
Other	534/32,068 (1.7)	507/32,498 (1.6)
Participants in included study — no./total no. (%)		
BWHS	1,437/32,247 (4.5)	2,896/32,544 (8.9)
CPSII	4,037/32,247 (12.5)	3,935/32,544 (12.1)
CPS3	1,537/32,247 (4.8)	1,729/32,544 (5.3)
CTS	2,226/32,247 (6.9)	2,134/32,544 (6.6)
MCBCS	4,517/32,247 (14.0)	3,249/32,544 (10.0)
MEC	3,641/32,247 (11.3)	3,689/32,544 (11.3)
MMHS	291/32,247 (0.9)	1,257/32,544 (3.9)
NHS	2,088/32,247 (6.5)	2,420/32,544 (7.4)
NHSII	935/32,247 (2.9)	1,391/32,544 (4.3)
WCHS	2,215/32,247 (6.9)	1,705/32,544 (5.2)
WHI	4,994/32,247 (15.5)	4,535/32,544 (13.9)
WWHS	4,329/32,247 (13.4)	3,604/32,544 (11.1)
Family history of breast cancer — no./total no. (%)§		
No	24,873/31,234 (79.6)	27,016/31,527 (85.7)
Yes	6,361/31,234 (20.4)	4,511/31,527 (14.3)
Family history of ovarian cancer — no./total no. (%)§		
No	27,494/28,534 (96.4)	28,002/28,949 (96.7)
Yes	1,040/28,534 (3.6)	947/28,949 (3.3)
<b>Clinical characteristic</b>		
Tumor behavior — no./total no. (%)		
In situ	4,446/31,221 (14.2)	NA
Invasive	26,775/31,221 (85.8)	NA

**Table 1. (Continued.)**

Variable	Case Patients (N = 32,247)	Controls (N = 32,544)
Estrogen-receptor status — no./total no. (%)		
Negative	3,805/22,233 (17.1)	NA
Positive	18,428/22,233 (82.9)	NA
Progesterone-receptor status — no./total no. (%)		
Negative	6,186/21,643 (28.6)	NA
Positive	15,457/21,643 (71.4)	NA
HER2 status — no./total no. (%)		
Negative	11,077/13,252 (83.6)	NA
Positive	2,175/13,252 (16.4)	NA
Triple-negative breast cancer — no./total no. (%)		
No	11,452/12,915 (88.7)	NA
Yes	1,463/12,915 (11.3)	NA

\* Plus–minus values are means  $\pm$ SD. Percentages may not total 100 because of rounding. The population-based Cancer Risk Estimates Related to Susceptibility (CARRIERS) analysis included data from the Black Women's Health Study (BWHS), the Cancer Prevention Study II (CPSII), the Cancer Prevention Study 3 (CPS3), the California Teachers' Study (CTS), the Mayo Clinic Breast Cancer Study (MCBCS), the Multiethnic Cohort Study (MEC), the Mayo Mammography Health Study (MMHS), the Nurses' Health Study (NHS), the Nurses' Health Study II (NHSII), the Women's Circle of Health Study (WCHS), the Women's Health Initiative (WHI), and the Wisconsin Women's Health Study (WWHS). NA denotes not applicable and HER2 human epidermal growth factor receptor 2.

† Age at the time of breast cancer diagnosis was used for case patients, and age at the time of selection into the study was used for controls. Data on age were missing for 539 case patients and for 2 controls.

‡ Race or ethnic group was reported by the participant.

§ Family history was restricted to first-degree relatives.

patients according to age at diagnosis and among the controls according to age at the time of selection into the study (Fig. S3). The prevalence of pathogenic variants in *BRCA1* and *BRCA2* among case patients decreased rapidly after age 40 years, whereas a constant and limited decline in the prevalence of pathogenic variants in *ATM*, *CHEK2*, and *PALB2* was observed among the case patients 40 to 85 years of age (Fig. S3). Pathogenic variants in *ATM*, *BRCA1*, *BRCA2*, *CHEK2*, and *PALB2* were associated with lifetime absolute risk of breast cancer of greater than 20% by age 85 years among non-Hispanic Whites; pathogenic variants in *BRCA1* or *BRCA2* yielded a lifetime risk of approximately 50%, and in *PALB2*, a lifetime risk of approximately 32% (Fig. 1).

## DISCUSSION

Here, we report the prevalence of pathogenic variants in breast cancer–predisposition genes among 32,247 women with breast cancer and

32,544 study-matched unaffected women from U.S. population-based studies in the CARRIERS consortium and provide estimates of breast cancer risk with respect to these pathogenic variants in the general population. On the basis of the American Cancer Society estimate of 276,000 new diagnoses of breast cancer in the United States in 2020, the population-based CARRIERS analysis suggests that at least 13,800 (approximately 5%) will occur in women with germline pathogenic variants in predisposition genes. However, many of these women are not known to have underlying genetic susceptibility to breast cancer.

Currently, there are differing recommendations for the selection of patients with breast cancer for clinical genetic testing, with considerable controversy regarding which patients to test and which genes to include in the testing process.<sup>15,36–38</sup> The National Comprehensive Cancer Network and the National Institute for Health and Care Excellence provide criteria for the selection of women with breast cancer, ovarian cancer,

**Table 2. Associations between Pathogenic Variants in Established Breast Cancer–Predisposition Genes and Risk of Breast Cancer.\***

Breast Cancer–Predisposition Gene <sup>1,2,7</sup>	Case Patients (N=32,247) no. with pathogenic variant (%)	Controls (N=32,544) no. with pathogenic variant (%)	Odds Ratio (95% CI) <sup>†</sup>	P Value
<i>ATM</i>	253 (0.78)	134 (0.41)	1.82 (1.46–2.27)	<0.001
<i>BARD1</i>	49 (0.15)	35 (0.11)	1.37 (0.87–2.16)	0.18
<i>BRCA1</i>	275 (0.85)	37 (0.11)	7.62 (5.33–11.27)	<0.001
<i>BRCA2</i>	417 (1.29)	78 (0.24)	5.23 (4.09–6.77)	<0.001
<i>CDH1</i>	17 (0.05)	6 (0.02)	2.50 (1.01–7.07)	0.06
<i>CHEK2</i>	349 (1.08)	138 (0.42)	2.47 (2.02–3.05)	<0.001
<i>NF1</i> <sup>‡</sup>	19 (0.06)	11 (0.03)	1.93 (0.91–4.31)	0.09
<i>PALB2</i>	148 (0.46)	38 (0.12)	3.83 (2.68–5.63)	<0.001
<i>PTEN</i>	8 (0.02)	3 (0.01)	NA	NA
<i>RAD51C</i>	41 (0.13)	35 (0.11)	1.20 (0.75–1.93)	0.44
<i>RAD51D</i>	26 (0.08)	14 (0.04)	1.72 (0.88–3.51)	0.12
<i>TP53</i> <sup>‡</sup>	19 (0.06)	2 (0.01)	NA	NA
Total	1621 (5.03)	531 (1.63)	—	—

\* The studies in the CARRIERS consortium that were included in this population-based analysis were BWHS, CPSII, CPS3, CTS, MCBCS, MEC, MMHS, NHS, NHSII, WCHS, WHI, and WWHS. NA denotes not applicable (too few events [ $<5$ ] to calculate a stable odds ratio).

<sup>†</sup> Odds ratio estimates for any breast cancer were adjusted for study, age, family history of breast cancer, and race or ethnic group.

<sup>‡</sup> Pathogenic variants in *NF1* and *TP53* were restricted to those with an alternate allele fraction (calculated as the number of alternate allele reads divided by the total number of reads at a specific genomic position) between 0.3 and 0.7.

or both for testing on the basis of personal and family history of these and other cancers.<sup>15,36–38</sup> In contrast, the American Society of Breast Surgeons has suggested offering testing to all patients with breast cancer, which increases the number of pathogenic-variant carriers by 30%.<sup>37</sup> Among the patients with breast cancer in the population-based CARRIERS analysis, 5.03% had pathogenic variants in the 12 established, clinically actionable predisposition genes, with 0.85% and 1.29% having pathogenic variants in *BRCA1* and *BRCA2*, respectively. These refined estimates of the prevalences of pathogenic variants among women with breast cancer in the overall population, as opposed to selected high-risk patients, may inform ongoing discussions regarding testing in patients with breast cancer. The risks of breast cancer associated with pathogenic variants in the genes evaluated in the population-based CARRIERS analysis also provide important information for risk assessment and counseling of women with breast cancer who do not meet high-risk selection criteria.

The population-based CARRIERS analysis also showed that certain subgroups of patients with breast cancer are at substantially increased risk of having pathogenic variants in high-penetrance, clinically actionable genes. For instance, pathogenic variants in *BRCA1*, *BRCA2*, and *PALB2* were observed in 8.13% of the patients with triple-negative breast cancer, as compared with 1.84% of the patients with ER-positive breast cancer. In addition, approximately 3.3% of women with ER-positive breast cancer without a family history had pathogenic variants in actionable breast cancer genes, with pathogenic variants in *ATM*, *CHEK2*, and *BRCA2* accounting for the majority. Furthermore, pathogenic variants in *BARD1*, *RAD51C*, and *RAD51D* showed weak associations with breast cancer risk among the participants overall but were associated with a moderate risk of ER-negative breast cancer (odds ratio,  $>2$ ). These findings were consistent with previously reported associations with ER-negative and triple-negative breast cancer among women who qualified for clinical genetic testing<sup>3,39</sup> and among non-

**Table 3. Associations between Pathogenic Variants in Established Breast Cancer–Predisposition Genes and Risk of Breast Cancer According to Estrogen-Receptor and Triple-Negative Breast Cancer Status of Tumors.\***

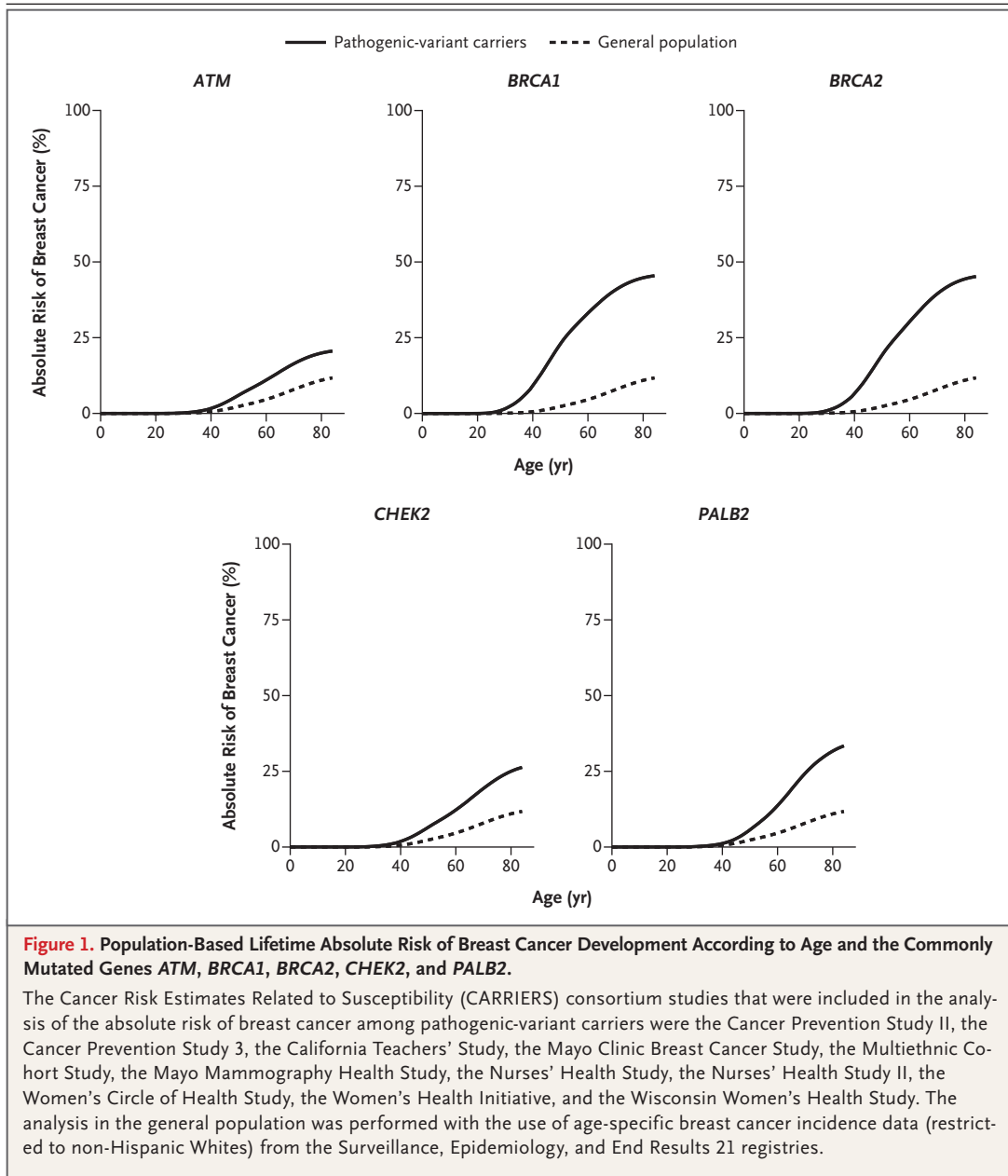
Breast Cancer–Predisposition Gene	ER-Positive Breast Cancer (N = 18,428)		ER-Negative Breast Cancer (N = 3805)		Triple-Negative Breast Cancer (N = 1463)	
	Participants with Pathogenic Variant no. (%)	Odds Ratio (95% CI)†	Participants with Pathogenic Variant no. (%)	Odds Ratio (95% CI)†	Participants with Pathogenic Variant no. (%)	Odds Ratio (95% CI)†
ATM	151 (0.82)	1.96 (1.52–2.53)	19 (0.50)	1.04 (0.59–1.72)	5 (0.34)	0.50 (0.12–1.36)
BARD1	20 (0.11)	0.91 (0.49–1.64)	11 (0.29)	2.52 (1.18–5.00)	6 (0.41)	3.18 (1.16–7.42)
BRCA1	73 (0.40)	3.39 (2.17–5.45)	114 (3.00)	26.33 (17.28–41.52)	65 (4.44)	42.88 (26.56–71.25)
BRCA2	201 (1.09)	4.66 (3.52–6.23)	82 (2.16)	8.89 (6.36–12.47)	30 (2.05)	9.70 (5.97–15.47)
CDH1	13 (0.07)	3.37 (1.24–10.72)	3 (0.08)	NA	1 (0.07)	NA
CHEK2	205 (1.11)	2.60 (2.05–3.31)	20 (0.53)	1.40 (0.83–2.25)	8 (0.55)	1.63 (0.72–3.20)
NF1‡	10 (0.05)	1.63 (0.65–4.03)	2 (0.05)	NA	1 (0.07)	NA
PALB2	64 (0.35)	3.13 (2.02–4.96)	42 (1.10)	9.22 (5.63–15.25)	24 (1.64)	13.03 (7.08–23.75)
PTEN	3 (0.02)	NA	0	NA	0	NA
RAD51C	16 (0.09)	0.83 (0.44–1.54)	9 (0.24)	2.19 (0.97–4.49)	4 (0.27)	NA
RAD51D	13 (0.07)	1.61 (0.71–3.70)	7 (0.18)	3.93 (1.40–10.29)	1 (0.07)	NA
TP53‡	9 (0.05)	NA	2 (0.05)	NA	2 (0.14)	NA

\* The studies in the CARRIERS consortium that were included in this population-based analysis were BWHS, CPSII, CPS3, CTS, MCBCS, MEC, MMHS, NHS, NHSII, WCHS, WHI, and WWHS. ER denotes estrogen-receptor.

† Odds ratio estimates for any breast cancer were adjusted for study, age, family history of breast cancer, and race or ethnic group.

‡ Pathogenic variants in *NF1* and *TP53* were restricted to those with an alternate allele fraction between 0.3 and 0.7.





Hispanic Black women with breast cancer, a relatively high proportion of whom have ER-negative disease.<sup>4</sup> Thus, risk stratification of women with breast cancer in the general population based on features such as tumor markers is an important method for identifying women at the highest risk of having a mutation, especially in underserved, minority populations.

There is also increasing discussion regarding screening for pathogenic variants in *BRCA1* and

*BRCA2* in the unaffected population.<sup>40</sup> Such testing for the Ashkenazi Jewish founder mutations in *BRCA1* and *BRCA2* is currently available in Israel.<sup>41</sup> However, beyond founder mutations, estimates of the prevalence of pathogenic variants in *BRCA1* and *BRCA2* or other breast cancer–predisposition genes are not well established in the general population. Here, we provide prevalence estimates for the 12 predisposition genes, showing that pathogenic variants in *CHEK2* and *ATM*

are the most common, and we note that pathogenic variants in *BRCA1* and *BRCA2* were found in 0.35% of the participants (or, 1 in 280). These estimates may inform the debate about population-based testing.

Most commercial genetic testing for hereditary cancer is based on multigene panels. However, many genes included on commercially available panels were not associated with an increased risk of breast cancer in the population-based CARRIERS analysis. Furthermore, several genes previously associated with an increased risk of breast cancer, including *NBN*, *BRIP1*, and *RECQL*, showed no associations in this population-based study. In particular, the finding that the *NBN* c.657\_661del5 Slavic founder mutation was not associated with an increased risk of breast cancer suggests that the recommendation by management guidelines<sup>15,36-38</sup> to increase screening among women with *NBN* pathogenic variants may need to be reevaluated. Among the established breast cancer–predisposition genes, *ATM* yielded an odds ratio of 1.82 (95% CI, 1.46 to 2.27) among all the women in the population-based CARRIERS analysis, an odds ratio of 1.72 (95% CI, 1.37 to 2.16) among women with no family history of breast cancer, and an odds ratio of 1.68 (95% CI, 1.31 to 2.17) among those who received a diagnosis of breast cancer at an age of more than 50 years. These findings suggest that carriers of pathogenic variants in *ATM* in the general population may have a substantially lower risk than what is often communicated to carriers of pathogenic variants who are identified through clinical testing (i.e., a risk that is said to be 2.5 times as high as that among noncarriers).<sup>1,42</sup> However, the estimated lifetime risk of breast cancer by age 85 years among carriers of pathogenic variants in *ATM* was still over the 20% threshold used clinically for enhanced screening. Pathogenic variants in *PALB2* were associated with a moderate risk of breast cancer in the population-based CARRIERS analysis (odds ratio, 3.83; 95% CI, 2.68 to 5.63), a finding that is similar to that in a study of two *PALB2* founder mutations in a prospective cohort study in Poland (odds ratio, 4.39; 95% CI, 2.30 to 8.37).<sup>43</sup> However, *PALB2* was identified as a high-risk gene (odds ratio, 8.04; 95% CI, 5.33 to 12.29) among case patients with a family history of breast cancer in the population-based CARRIERS analysis

(Table S12), a finding consistent with the results of a study of 524 families with *PALB2* mutations (relative risk, 7.18; 95% CI, 5.82 to 8.85).<sup>44</sup> These findings confirmed the effect of family history on breast cancer risk and identified family history as a critical factor for risk stratification of patients.

This study has some limitations. Enrollment was restricted to women 50 years of age or older in certain population-based studies in the CARRIERS consortium, which had the potential to influence the generalizability of the aggregate estimates of the prevalence of pathogenic variants in *BRCA1* and *BRCA2* to younger women; however, sensitivity analyses that excluded the Women's Health Initiative and the Cancer Prevention Study II (studies that involved women at an older age at enrollment) did not substantially influence the findings. In addition, the statistical model for penetrance estimation was based on the assumptions that the underlying population-based SEER rates, prevalence of pathogenic variants, and age-specific estimates of odds ratios reflect those in the general population. Future studies are needed to evaluate the calibration of the probability estimates. Another potential limitation is that the sequencing was conducted in a research laboratory rather than a commercial genetic-testing facility. However, the custom QIAseq panel was shown to have high sensitivity and specificity for pathogenic variants in predisposition genes.<sup>17</sup> Furthermore, because all samples were sequenced in a single center and variants were called through a single pipeline, issues with bioinformatics and batch effects were minimized. In addition, it was not possible to study the effects of individual pathogenic variants on breast cancer risk because of the rarity of the variants.

Overall, the results of the population-based CARRIERS analysis showed that pathogenic variants in the predisposition genes *ATM*, *BRCA1*, *BRCA2*, *CHEK2*, and *PALB2* were associated with increased risks of breast cancer and that pathogenic variants in *BARD1*, *RAD51C*, and *RAD51D* were associated with increased risks of ER-negative breast cancer in the general population. To date, the management recommendations for women with pathogenic variants in these genes have been based on risk estimates from studies involving women at high risk. We anticipate that the estimates from the population-based CARRIERS

analysis will inform cancer screening and other risk-management strategies for women with pathogenic variants in cancer-predisposition genes in the general population.

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#### APPENDIX

The authors' full names and academic degrees are as follows: Chunling Hu, M.D., Ph.D., Steven N. Hart, Ph.D., Rohan Gnanaolivu, M.S., Hongyan Huang, Ph.D., Kun Y. Lee, Ph.D., Jie Na, M.S., Chi Gao, M.Sc., Jenna Lilyquist, Ph.D., Siddhartha Yadav, M.D., Nicholas J. Boddicker, Ph.D., Raed Samara, Ph.D., Josh Klebba, B.S., Christine B. Ambrosone, Ph.D., Hoda Anton-Culver, Ph.D., Paul Auer, Ph.D., Elisa V. Bandera, Ph.D., Leslie Bernstein, Ph.D., Kimberly A. Bertrand, Sc.D., Elizabeth S. Burnside, M.D., M.P.H., Brian D. Carter, M.P.H., Heather Eliassen, Sc.D., Susan M. Gapstur, Ph.D., Mia Gaudet, Ph.D., Christopher Haiman, Sc.D., James M. Hodge, M.P.H., J.D., David J. Hunter, Sc.D., Eric J. Jacobs, Ph.D., Esther M. John, Ph.D., Charles Kooperberg, Ph.D., Allison W. Kurian, M.D., Loïc Le Marchand, M.D., Ph.D., Sara Lindstroem, Ph.D., Tricia Lindstrom, B.S., Huiyan Ma, Ph.D., Susan Neuhausen, Ph.D., Polly A. Newcomb, Ph.D., M.P.H., Katie M. O'Brien, Ph.D., Janet E. Olson, Ph.D., Irene M. Ong, Ph.D., Tuya Pal, M.D., Julie R. Palmer, Sc.D., Alpa V. Patel, Ph.D., Sonya Reid, M.D., Lynn Rosenberg, Sc.D., Dale P. Sandler, Ph.D., Christopher Scott, M.S., Rulla Tamimi, Sc.D., Jack A. Taylor, M.D., Ph.D., Amy Trentham-Dietz, Ph.D., Celine M. Vachon, Ph.D., Clarice Weinberg, Ph.D., Song Yao, Ph.D., Argyrios Zogas, Ph.D., Jeffrey N. Weitzel, M.D., David E. Goldgar, Ph.D., Susan M. Domchek, M.D., Katherine L. Nathanson, M.D., Peter Kraft, Sc.D., Eric C. Polley, Ph.D., and Prof. Fergus J. Couch, Ph.D.

The authors' affiliations are as follows: Mayo Clinic, Rochester, MN (C. Hu, S.N.H., R.G., K.Y.L., J.N., J.L., S. Yadav, N.J.B., T.L., J.E.O., C.S., C.M.V., E.C.P., F.J.C.); Harvard University T.H. Chan School of Public Health (H.H., C.G., D.J.H., P.K.), Slone Epidemiology Center at Boston University (K.A.B., J.R.P., L.R.), and Brigham and Women's Hospital (H.E.) — all in Boston; Qiagen, Hilden, Germany (R.S., J.K.); Roswell Park Comprehensive Cancer Center, Buffalo (C.B.A., S. Yao), and Weill Cornell Medicine, New York (R.T.) — both in New York; the University of California, Irvine (H.A.-C., A.Z.), Beckman Research Institute of City of Hope, Duarte (L.B., H.M., S.N., J.N.W.), Keck School of Medicine, University of Southern California, Los Angeles (C. Haiman), and Stanford University School of Medicine, Stanford (E.M.J., A.W.K.) — all in California; the University of Wisconsin–Milwaukee Joseph J. Zilber School of Public Health, Milwaukee (P.A.), and the University of Wisconsin–Madison, Madison (E.S.B., I.M.O., A.T.-D.); the Cancer Prevention and Control Program, Rutgers Cancer Institute of New Jersey, State University of New Jersey, New Brunswick (E.V.B.); the Behavioral and Epidemiology Research Group, American Cancer Society, Atlanta (B.D.C., S.M.G., M.G., J.M.H., E.J.J., A.V.P.); the University of Oxford, Oxford, United Kingdom (D.J.H.); the Fred Hutchinson Cancer Research Center (C.K., P.A.N.) and the Department of Epidemiology, University of Washington (S.L.) — both in Seattle; the Epidemiology Program, University of Hawaii Cancer Center, Honolulu (L.L.M.); the National Institute of Environmental Health Sciences, Durham, NC (K.M.O., D.P.S., J.A.T., C.W.); Vanderbilt University, Nashville (T.P., S.R.); the University of Utah, Salt Lake City (D.E.G.); and the Department of Medicine and the Basser Center for BRCA, Abramson Cancer Center, Perelman School of Medicine at the University of Pennsylvania, Philadelphia (S.M.D., K.L.N.).

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