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Enrollment and biospecimen collection in a multiethnic family cohort: The Northern California site of the Breast Cancer Family Registry

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Abstract

Purpose: Racial/ethnic minorities are often assumed to be less willing to participate in and provide biospecimens for biomedical research. We examined racial/ethnic differences in enrollment of women with breast cancer (probands) and their first-degree relatives in the Northern California site of the Breast Cancer Family Registry from 1996-2011.

Methods: We evaluated participation in several study components, including biospecimen collection, for probands and relatives by race/ethnicity, cancer history, and other factors.

Results: Of 4,780 eligible probands, 76% enrolled in the family registry by completing the family history and risk factor questionnaires and 68% also provided a blood or mouthwash sample. Enrollment was highest (81%) for non-Hispanic whites (NHWs) and intermediate (73%-76%) for

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EMJ acquired the data, conceived of and designed the analysis, interpreted the data, and drafted and revised the manuscript. MS contributed to the literature review and the writing. JK and MS performed data management and the statistical analysis, interpreted the data, and participated in the revision of the manuscript. ASW and DWW acquired the data, interpreted the data, and participated in the revision of the manuscript. All authors approved the final manuscript.

Conflict of interest:

The authors declare they have no conflicts of interest.

Ethical approval:

All procedures performed in this study involving human participants were in accordance with the ethical standards of the Institutional Review Board at the Cancer Prevention Institute of California and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Informed consent:

Informed consent was obtained from all individual participants included in this study.

Hispanics, African Americans, and all Asian American subgroups, except Filipina women (66%). Of 4,279 eligible relatives, 77% enrolled in the family registry, and 65% also provided a biospecimen sample. Enrollment was highest for NHWs (87%) and lowest for Chinese (68%) and Filipinas (67%). Among those enrolled, biospecimen collection rates were similar for NHW, Hispanic, and African American women, both for probands (92%-95%) and relatives (82%-87%), but lower for some Asian-American subgroups (probands: 72%-88%; relatives: 71%-88%), foreign-born Asian Americans, and probands those who were more recent immigrants or had low English language proficiency.

Conclusions: These results show that racial/ethnic minority populations are willing to provide biospecimen samples for research, although some Asian American subgroups in particular may need more directed recruitment methods. To address long-standing and well-documented cancer health disparities, minority populations need equal opportunities to contribute to biomedical research.

Keywords

Biospecimen collection; Study participation; Race/ethnicity; Breast cancer; Epidemiology; Hispanics; African Americans; Asian Americans

Introduction

Breast cancer disparities by race/ethnicity span the continuum from etiology, prevention, early detection, diagnosis, treatment, and survivorship [1–3]. In order to understand the complex interactions between biologic, lifestyle, environmental, social, cultural, and community-level factors that underlie the disparities, research in diverse populations is critical [4,5]. Racial/ethnic minorities are under-represented in observational studies [5,6], intervention and clinical trials [7,8], and biorepositories [9,10], and it is often assumed that minorities are less willing to participate in biomedical research and provide biospecimens [11]. Multiple barriers precluding minority participation in biospecimen collection and genomics research have been identified [12–15]. Under-representation of racial/ethnic minorities may also be due to failures in recruitment methodology or lack of opportunities to engage in research rather than an inherent unwillingness to participate in biomedical research [16,17,14,18].

We report on the enrollment experience from the Northern California site of the Breast Cancer Family Registry [19,20]. Family studies are a powerful study design to investigate gene-environment interactions [21], but they also present unique challenges, as they depend on participants' willingness to grant permission to contact family members. We evaluated participation in multiple study components, including biospecimen collection, by race/ethnicity and other factors.

Materials and Methods

Study sample

The Northern California family registry site recruited female probands ages 18-64 years newly diagnosed with breast cancer through population-based cancer registries that are part

of the National Cancer Institute's Surveillance, Epidemiology, and End Results (SEER) program and the California Cancer Registry. The San Francisco Bay Area-based recruitment included invasive or *in situ* breast cancer cases of any race/ethnicity diagnosed between 1/1/1995 and 9/30/1998 (Phase I); Hispanic, African American, Chinese, Filipina, and Japanese invasive cases diagnosed between 10/1/1998 and 4/30/2003 (Phase II); Hispanic and African American invasive cases diagnosed between 5/1/2003 and 8/31/2009 (Phase III); and triple negative cases (estrogen receptor negative, progesterone receptor negative, human epidermal growth factor receptor 2 negative) diagnosed between 1/1/2007 and 6/30/2009 (Phase IV). Additionally, Hispanic and African American invasive cases from the Sacramento area diagnosed between 1/1/2005 and 12/31/2006 were identified through the Sacramento and Sierra Cancer Registries.

Of 34,517 ascertained cases, 1,235 (4%) were deceased, 270 (0.8%) had no physician approval to be contacted, and 3,092 (9.1%) had outdated addresses. Except for 361 Phase I cases (diagnosed before age 35 years, with a prior ovarian or childhood cancer, or bilateral breast cancer with a first diagnosis before age 50 years) which were enrolled without screening, the remaining 29,559 cases were screened by telephone to determine self-identified race/ethnicity and study eligibility. Cases with characteristics suggestive of inherited breast cancer (i.e., diagnosis before age 35 years, prior ovarian or childhood cancer, bilateral breast cancer with a first diagnosis before age 50 years, and first-degree family history of breast, ovarian, or childhood cancer) were invited to enroll in the family registry. Cases diagnosed at ages 35-64 years not meeting these criteria were randomly sampled; racial/ethnic minorities at 33% and non-Hispanic whites (NHWs) at 2.5%, given the high volume of NHW cases.

We also enrolled the probands' adult relatives living in North America, primarily first-degree relatives. The present analysis was limited to parents, full and half-sisters, and adult daughters or sons with a prior diagnosis of breast, ovarian, or childhood cancer.

Data collection

For probands we collected a family history questionnaire by telephone. Probands and relatives completed a risk factor questionnaire (by home visit if residing in the Bay area or by telephone if residing elsewhere) and a mailed food frequency questionnaire. Probands and relatives with a prior breast cancer completed a treatment questionnaire and a signed medical release to collect the pathology report and tumor tissue. Living parents of probands also completed the risk factor questionnaire, except for parents of probands diagnosed from 1995-1998, for whom we collected the risk factor questionnaire only if they had a prior breast or ovarian cancer. The risk factor questionnaire included questions about race/ethnicity, education, country of birth, year of migration to the U.S., years of residence in the U.S. if foreign-born, and first language learned. If English was not the participant's first language, English language proficiency was assessed in the questionnaire by asking "Which of these choices best describes how well you speak English?", with response options of "well", "medium", "little", or "not at all". Data on age and stage at diagnosis and neighborhood socioeconomic status (SES) based on U.S. census data were obtained from the cancer registries.

Biospecimen collection

All probands and relatives who enrolled in the family registry study by completing the risk factor questionnaire were invited to provide a biospecimen sample. For parents of probands diagnosed from 1995-1998, we collected a biospecimen sample only if they had a prior breast or ovarian cancer. For local participants (Bay Area residents; Alameda, Contra Costa, Marina, Monterey, Santa Clara, Santa Cruz, San Francisco and San Mateo counties), interviewers/phlebotomists collected the blood sample at the home visit after they administered the risk factor questionnaire; for non-local participants, we mailed a blood collection kit with prepaid postage for return if they were willing to have their blood drawn at their doctor's office. Participants who declined a blood draw were invited to provide a mouthwash sample using a mailed mouthwash collection kit.

Data and biospecimen collection procedures

We used several strategies to maximize enrollment and biospecimen collection. Trained professional study interviewers and phlebotomists made multiple attempts by phone to reach study participants or by mail to obtain updated telephone numbers. They made up to 10 attempts to conduct the telephone screening or to schedule a home visit or telephone interview. All study materials, except the diet questionnaire, were translated into Spanish and Chinese, and data and biospecimens were collected by bi-cultural and bi-lingual interviewers and phlebotomists. We matched participants and interviewers/phlebotomists on language and cultural background, when possible. To reduce participant burden, we collected all questionnaire data by home visit or telephone interview at a time that was convenient to the participant, including evenings and weekends, and re-scheduled cancelled appointments. Participants received \$25 for completing the risk factor questionnaire and \$25 for providing a biospecimen. Through the consent form participants were informed that de-identified data and biospecimens would be stored for future research by approved investigators. The study protocol was approved by the Institutional Review Board of the Cancer Prevention Institute of California and participants provided written informed consent.

Analytic variables

Analyses of case participation in telephone screening relied on race/ethnicity from the cancer registries, whereas analyses of proband and relative participation in data and biospecimen collection relied on self-reported race/ethnicity. For eligible relatives who did not enroll, we used the proband's race/ethnicity. We classified race/ethnicity as NHW, Hispanic, African American, Chinese, Filipina, Japanese, other Asian American/Pacific Islander, or other (Native American, mixed race/ethnicity). Stage at diagnosis was based on SEER summary stage (in situ, localized, regional, distant); neighborhood SES is a composite measure of seven SES indicators from 2000 census data [22] and was categorized according to the quintile distribution of all ascertained breast cancer cases. Cancer family history was defined as breast, ovarian, or childhood cancer in first-degree relatives, and personal cancer history was defined as a prior diagnosis of breast, ovarian or childhood cancer.

Statistical analysis

We evaluated racial/ethnic differences in study participation, defined as completion of telephone screening interview (breast cancer cases); family history and risk factor questionnaires (probands); risk factor questionnaire (relatives); and biospecimen collection (probands and relatives); and calculated participation rates as the number of subjects who completed the study component divided by the number of eligible subjects. For both enrollment and biospecimen collection, we examined differences in proband participation by race/ethnicity, age at diagnosis, stage, cancer family history, and neighborhood SES, and differences in relative participation by race/ethnicity, proband's age at diagnosis and personal cancer history. To evaluate differences in biospecimen collection by other characteristics collected in the risk factor questionnaire (i.e., education, country of birth, age at migration to the U.S., years of residence in the U.S., and English language proficiency), we restricted the analyses to enrolled probands and relatives. For single predictors of study participation, we assessed the statistical significance of differences using chi-square tests. To assess differences in study participation adjusting for multiple predictors, we used multivariable models to calculate odds ratios (OR) and 95% confidence intervals (CI). The ORs represent odds ratios for participation. For probands, we used unconditional logistic regression, whereas for relatives we used the generalized estimating equations (GEE) method on logistic models to account for correlation among relatives from the same family. We used the Wald test to test for significant differences in study participation. Two-sided $P < 0.05$ were considered statistically significant and all analyses were performed using SAS Version 9.4 (SAS Institute, Cary, NC).

Of the 4,841 eligible probands who were alive and selected to enroll in the family registry, 61 were from multiple proband families. For this analysis, secondary or tertiary probands were classified as relatives.

Results

Case screening interview

Of 29,559 incident female breast cancer cases contacted, 25,183 (85%) completed the telephone screening interview (Table 1). Participation differed by race/ethnicity ($P < 0.01$), with higher rates for African Americans, NHWs and Hispanics (87%-88%) than for Asian American subgroups (76%-81%). Participation also differed by age (lowest for ages 18-34 years) and stage at diagnosis (lowest for distant stage), but not by neighborhood SES. Lower participation in screening by Asian Americans was also seen in multivariable adjusted models.

Proband participation

Family history and risk factor questionnaires.—Of 4,780 breast cancer cases selected as probands, 3,620 (76%) enrolled in the family registry study by completing the family history and risk factor questionnaires (Table 2). Characteristics of enrolled probands are shown in Supplemental Table 1. Significant differences by race/ethnicity were found for age at diagnosis, stage, cancer family history, education and country of birth, but not for neighborhood SES. Enrollment varied by race/ethnicity ($P < 0.01$) and was highest for

NHWs (81%), intermediate (74%-76%) for all other groups except Filipinas (66%). Enrollment did not differ by age at diagnosis ($P = 0.55$) or neighborhood SES ($P = 0.45$), but was higher for those with a cancer family history ($P = 0.01$) and lower for those with missing stage ($P = 0.01$). In multivariable adjusted models, compared to NHWs, enrollment was similar for Hispanics and African Americans, but significantly lower for Asian Americans (OR=0.68, 95% CI=0.55-0.83).

Biospecimen collection.—Of eligible probands, 3,244 (68%) provided a biospecimen sample. Participation differed by race/ethnicity, ranging from 76% among NHW women to 50% among Filipinas, and was higher for family history positive vs. negative probands (71% vs. 66%, $P < 0.01$). Among enrolled probands who completed the risk factor questionnaire, biospecimen collection differed by race/ethnicity ($P < 0.01$), with similarly high participation by African Americans, NHWs and Hispanics (92%-95%), but notably lower participation by some Asian American subgroups (72%-88%; Table 2). Participation in biospecimen collection also differed by cancer family history ($P < 0.01$), education ($P < 0.01$), and country of birth ($P < 0.01$). In multivariable adjusted models, participation in biospecimen collection was 70% lower for Asian Americans compared to NHWs, but not significantly different for Hispanics and African Americans, and 34% lower for foreign-born vs. U.S.-born probands ($P = 0.01$). Overall, 92% of biospecimens collected were blood samples and 8% were mouthwash samples, with some variation by race/ethnicity ($P = 0.01$). The proportion of blood vs. mouthwash samples was highest for Hispanics (97%), followed by Japanese (94%), NHWs (93%), African Americans (90%), Filipinas (89%), Chinese (86%) and other Asians/others (86%) (data not shown in tables).

Relative participation

Of 3,620 enrolled probands, only 62% had eligible first-degree relatives that could be contacted (Supplemental Table 2). Seven percent had no living first-degree relatives, ranging from 3% for Filipinas to 14% for NHWs; 9% had no first-degree relatives living in North America, ranging from 1% for African Americans to 25% for other Asian Americans/others; and 21% did not give permission to contact their relatives, ranging from 11% for NHWs to 35% for Chinese. Overall, 38% of probands did not have a first-degree relative we could contact for enrollment, ranging from 28% for NHWs to 58% for Chinese.

Risk factor questionnaire.—Of 4,279 eligible first-degree relatives, 3,306 (77%) enrolled in the family registry study by completing the risk factor questionnaire (Table 3). Characteristics of enrolled relatives are shown in Supplemental Table 3. Statistically significant differences by race/ethnicity were found for age at interview, personal cancer history, education, and country of birth. Relative enrollment differed by race/ethnicity ($P < 0.01$), and was highest for NHWs (87%), intermediate (74%-80%) for African Americans, Hispanics, Japanese and other Asian Americans/others, and lowest (67%) for Chinese and Filipinas. Enrollment was similar for relatives with or without a personal cancer history (78% vs. 77%) and was higher for relatives of younger probands than those of older probands (84% vs. 76%, $P < 0.01$). In multivariable adjusted models, race/ethnicity and proband's age at diagnosis were significant predictors of enrollment.

Biospecimen collection.—Of eligible relatives, 2,774 (65%) provided a biospecimen sample. Participation differed by race/ethnicity ($P < 0.01$), with participation ranging from 76% among NHWs to 53% among Filipinas. Participation was higher for relatives of younger probands ($P = <0.01$), but did not differ by the relative's personal cancer history. Of 3,306 enrolled relatives, 2,774 (84%) provided a blood or mouthwash sample, with significant differences by race/ethnicity ($P < 0.01$), ranging from 82%-87% for African Americans, Hispanics, and NHWs (Table 3). For Asian Americans, biospecimen collection was 78% overall, and ranged from 71%-88% for specific subgroups. Biospecimen collection was higher for younger than older relatives ($P < 0.01$), but did not differ by education ($P = 0.65$) or personal history of cancer ($P = 0.69$). In multivariable models, ORs for biospecimen collection were 1.00 (95% CI=0.71-1.42) for Hispanics, 0.70 (95% CI=0.51-0.97) for African Americans, and 0.59 (95% CI=0.42-0.82) for Asian Americans, compared to NHWs.

Proband and relative biospecimen collection by migration history

Among enrolled Hispanic probands, biospecimen collection (95% participation overall) did not differ by education, country of birth, age at migration to the U.S., duration of residence in the U.S., or English proficiency (Table 4). In multivariable adjusted models, none of the differences in participation were statistically significant. In contrast, among Asian Americans, biospecimen collection rates were higher among more educated probands, and those who were U.S.-bom, migrated to the U.S. before age 20 years, lived in the U.S. for 40 years, or spoke English well or English only (all P values < 0.01). In multivariable models, education, country of birth, years of residence in the U.S., and English language proficiency remained significant predictors of biospecimen collection.

Among enrolled Hispanic relatives, biospecimen collection was lower compared to Hispanic probands (87% vs. 95%), and higher for less educated vs. more educated relatives ($P = 0.02$), foreign-born vs. U.S.-bom relatives (90% vs. 84%; $P = 0.01$) and those who lived in the U.S. 40 years vs. > 40 years (89%-96% vs. 85%; $P < 0.01$) (Table 4). In multivariable models, country of birth and years of residence in the U.S. remained significant predictors of biospecimen collection. In contrast, among enrolled Asian Americans, biospecimen collection was similar for relatives and probands (78% vs. 77%), lowest for those with low education ($P < 0.01$), higher for U.S.-bom than foreign-bom relatives (87% vs. 73%; $P < 0.01$) and differed by migration history, with the highest participation for relatives who migrated to the U.S. before age 30 years (79%-81%), lived in the U.S. for > 40 years (87%), and spoke English well or English only (82%). In multivariable models, only education remained a significant predictor of biospecimen collection; country of birth was a predictor of borderline significance.

Discussion

In this population-based family cohort, study participation was generally high, with some variation by race/ethnicity. Participation in telephone screening was similar for female Hispanic, African American, and NHW breast cancer cases, but lower for Asian American subgroups. Proband enrollment was highest for NHWs, and intermediate for all other

groups, except Filipinas. A similar enrollment pattern by race/ethnicity was seen for first-degree relatives. Biospecimen collection rates both for probands and relatives were lowest for Asian Americans, with considerable variation across Asian American subgroups.

Our enrollment rates for Hispanic, African American, and NHW probands are comparable to the participation rates in the San Francisco Bay Area Breast Cancer Study (SFBCS), a population-based case-control study [23]. Both studies used similar recruitment methods and collected interview data and biospecimens through home visits. Proband enrollment was somewhat lower than case participation in SFBCS (Hispanics: 75% vs. 89%; African Americans: 76% vs. 87%; NHWs: 81% vs. 86%), possibly due to long-term follow-up and involvement of family members.

One of our key findings is the high biospecimen collection rate for enrolled Hispanic and African American probands, consistent with the high rates for SFBCS cases who completed the interview [24] (Hispanics: 95% vs. 88%; African Americans: 92% vs. 85%; NHWs: 94% vs. 90%). Furthermore, most participants provided a blood vs. mouthwash sample, demonstrating that California Hispanic and African American women with breast cancer are as willing as their NHW counterparts to donate blood for biomedical research.

Reports on biospecimen collection rates in racial/ethnic minorities are sparse. The Southern Community Cohort Study obtained blood or buccal samples for 96% (half were blood samples) of African Americans recruited from community health centers [25]. Blood or saliva collection by home visits was also high for African Americans in the North Carolina Colorectal Cancer Study (94% overall, 79% for blood) [26]. The Black Women's Health Study obtained mailed buccal samples for 51% [27], and a blood sample collected at a nearby clinical center for 35% of 1,500 pilot study participants [28]. These data suggest that high biospecimen collection rates are more difficult to attain in studies that increase participant burden (i.e., return of biospecimens by mail, clinic visits). Studies, such as ours, that reduce participant burden through home visits may be more effective in achieving high biospecimen collection. Home visits are more costly, particularly for geographically dispersed participants, but large-scale repeated mailings of biospecimen collection kits that are not returned also come at a considerable cost [29].

Our high biospecimen collection rate for Hispanics is consistent with the Mano a Mano cohort of Mexican Americans [30] that obtained biospecimens (blood, cheek cell, or urine samples) for 94% (collection rates for specific biospecimens were not provided), with similar collection rates for U.S.-born and foreign-born Hispanics, consistent with our findings for enrolled probands, whereas foreign-born relatives were twice as likely to participate in biospecimen collection. Consent to give blood and urine samples for future research did not differ between Mexican Americans and NHWs who participated in the 2011-2012 National Health and Nutrition Examination Survey [31]. Focus groups and surveys assessing knowledge and beliefs about biospecimens among Hispanics have also found high willingness to provide biospecimens for biomedical research [32–36].

For Asian Americans, proband and relative enrollment and biospecimen collection were considerably lower, as reported by others [37,31] and differed between Asian American

subgroups. Associations with migration-related variables differed from those for Hispanic women. Among both enrolled Asian American probands and relatives, biospecimen collection was lower for foreign-born than U.S.-born Asian Americans, whereas among enrolled Hispanics, biospecimen collection was higher for foreign-born relatives. Duration of residence in the U.S. and English language proficiency were significant predictors for enrolled Asian American probands only. The lack of interviewers who spoke a Filipino language may have contributed to the lower enrollment of Filipina cases in our study. Greater reluctance to participate in biospecimen-based research by foreign-born Asian Americans and more recent immigrants may be related to cultural beliefs and lack of knowledge about cancer, biospecimens, and biobanking [38]. Culturally relevant educational programs for Chinese Americans have been successful at increasing knowledge about biospecimens, addressing informed consent procedures and privacy concerns, and generally encouraging participation in biomedical research [39,40,14,41].

Enrollment of family members presented several challenges. The willingness to grant access to first-degree relatives varied by race/ethnicity, with about a third of Chinese probands not granting permission to contact relatives. Family size and immigrant background also affected the availability of relatives for enrollment; 14% of NHW probands did not have any first-degree relatives who were alive, and 19% of Chinese probands did not have any first-degree relatives who lived in North America. Overall, only 62% of probands had relatives whom we could approach for enrollment, with the lowest percentage for Chinese Americans (42%). Therefore, the enrolled relatives may not be representative of all eligible relatives.

Race/ethnicity was the only statistically significant predictor consistently associated with enrollment and biospecimen collection, both among probands and relatives, with generally similar participation for Hispanics, African Americans and NHWs, but lower participation in some Asian American subgroups. It is reassuring that case screening, proband enrollment and biospecimen collection varied little by neighborhood SES. For enrolled probands and relatives, biospecimen collection varied by education, though only among Asian Americans (data not shown for NHWs and African Americans). Similarly, other studies found only small differences in biospecimen collection by education [42] and among Hispanics and African Americans specifically [9,27].

We employed several strategies to maximize enrollment and biospecimen collection, including data collection by home visits and telephone interviews which helps build rapport and trust with participants, allows for participants' concerns to be addressed and resolved in a timely manner, and overcomes literacy issues. Home visits also reduce participant burden and help mitigate barriers such as lack of transportation or interference with family or work responsibilities [43]. Bilingual research staff and multi-lingual and culturally sensitive study materials are essential in multiethnic and immigrant study populations, and concordance in language and culture between study participants and interviewers has been shown to increase participation [44]. Community-based participatory research approaches and community outreach have also been shown to be effective in the recruitment of minority populations [45–48].

Our data and those from other epidemiologic studies demonstrate that minorities are willing to donate biospecimens for biomedical research [24,26,25,30]. However, given numerous barriers that may hinder participation in research [49,13,12,50], special efforts should be directed towards giving minorities opportunities to participate in research studies and facilitating their participation by reducing barriers. Increasing knowledge about biospecimens through educational programs and greater transparency by study investigators also help overcome issues of distrust and reluctance to participate in biospecimen collection [51,14]. It is important that populations from diverse racial/ethnic and socioeconomic backgrounds are given the opportunity to participate in biomedical research [5] because research that is inclusive of all populations is critical to inform targeted cancer control and prevention strategies, personalized medicine, and health policy.

Our study has several important strengths, including the population-based design for the recruitment of probands, purposeful oversampling of racial/ethnic minorities, and comprehensive collection of breast cancer risk factors by questionnaire and clinical and tumor characteristics from cancer registry records. Recruitment shortly after diagnosis and during treatment may be challenging, but we were successful at contacting cases at least 6 months after breast cancer diagnosis as soon as cases from the cancer registries became available. Women who died soon after diagnosis did not have the opportunity to enroll in the study, but given the high survival rate, the proportion of women who had died before being contacted for eligibility screening was small (3%). Other limitations include inability to assess study eligibility for all breast cancer cases due to language barriers at the screening level, and the less than optimal participation by Filipina and Chinese women. It is also possible that study participation in other U.S. regions differs for minority populations with different cultural backgrounds, countries of origin or sociodemographic characteristics compared to the urban populations of the San Francisco Bay Area.

Conclusions

Our results show that racial/ethnic minority populations are willing to participate in research and provide biospecimen samples, although recent immigrants may need more directed recruitment methods. Future studies should prioritize culturally sensitive approaches in their design in order to maximize recruitment and biospecimen collection, especially among Asian Americans.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Abbreviations:

CI	confidence interval
NHW	non-Hispanic white
OR	odds ratio
SES	socioeconomic status
SFBCS	San Francisco Bay Area Breast Cancer Study
U.S.	United States

References

1. Kohler BA, Sherman RL, Howlader N, Jemal A, Ryerson AB, Henry KA, Boscoe FP, Cronin KA, Lake A, Noone AM, Henley SJ, Ehemann CR, Anderson RN, Penberthy L (2015) Annual Report to the Nation on the Status of Cancer, 1975-2011, Featuring Incidence of Breast Cancer Subtypes by Race/Ethnicity, Poverty, and State. *J Natl Cancer Inst* 107 (6):djv048. [PubMed: 25825511]
2. Hunt BR, Hurlbert MS (2016) Black:white disparities in breast cancer mortality in the 50 largest cities in the United States, 2005-2014. *Cancer Epidemiol* 45:169–73. [PubMed: 27720130]
3. Wheeler SB, Reeder-Hayes KE, Carey LA (2013) Disparities in breast cancer treatment and outcomes: biological, social, and health system determinants and opportunities for research. *Oncologist* 18 (9):986–993. [PubMed: 23939284]
4. Warnecke RB, Oh A, Breen N, Gehlert S, Paskett E, Tucker KL, Lurie N, Rebbeck T, Goodwin J, Flack J, Srinivasan S, Kerner J, Heurtin-Roberts S, Abeles R, Tyson FL, Patmios G, Hiatt RA (2008) Approaching health disparities from a population perspective: the National Institutes of Health Centers for Population Health and Health Disparities. *Am J Public Health* 98 (9):1608–1615. [PubMed: 18633099]
5. Martin DN, Lam TK, Brignole K, Ashing KT, Blot WJ, Burhansstipanov L, Chen JT, Dignan M, Gomez SL, Martinez ME, Matthews A, Palmer JR, Perez-Stable EJ, Schootman M, Vilchis H, Vu A, Srinivasan S (2016) Recommendations for Cancer Epidemiologic Research in Understudied Populations and Implications for Future Needs. *Cancer Epidemiol Biomarkers Prev* 25 (4):573–580. [PubMed: 27196089]
6. Aziz NM, Rowland JH (2002) Cancer survivorship research among ethnic minority and medically underserved groups. *Oncol Nurs Forum* 29 (5):789–801. [PubMed: 12058154]
7. Chen MS Jr, Lara PN, Dang JH, Paterniti DA, Kelly K (2014) Twenty years post-NIH Revitalization Act: enhancing minority participation in clinical trials (EMPaCT): laying the groundwork for improving minority clinical trial accrual: renewing the case for enhancing minority participation in cancer clinical trials. *Cancer* 120 Suppl 7:1091–1096. [PubMed: 24643646]
8. Nicholson LM, Schwirian PM, Groner JA (2015) Recruitment and retention strategies in clinical studies with low-income and minority populations: Progress from 2004-2014. *Contemp Clin Trials* 45 (Pt A):34–40. [PubMed: 26188163]
9. Simon MA, de la Riva EE, Bergan R, Norbeck C, McKoy JM, Kulesza P, Dong X, Schink J, Fleisher L (2014) Improving diversity in cancer research trials: the story of the Cancer Disparities Research Network. *J Cancer Educ* 29 (2):366–374. [PubMed: 24519744]

10. Tham HM, Hohl S, Copeland W, Briant KJ, Marquez-Magana L, Thompson B (2017) Enhancing Biospecimen Knowledge Among Health Care Providers and Representatives From Community Organizations. *Health Promot Pract* 18 (5):715–725. [PubMed: 27118784]
11. Wendler D, Kington R, Madans J, Wye GV, Christ-Schmidt H, Pratt LA, Brawley OW, Gross CP, Emanuel E (2006) Are Racial and Ethnic Minorities Less Willing to Participate in Health Research? *PLoS Med* 3 (2):e19 [PubMed: 16318411]
12. Ford JG, Howerton MW, Lai GY, Gary TL, Bolen S, Gibbons MC, Tilburt J, Baffi C, Tanpitukpongse TP, Wilson RF, Powe NR, Bass EB (2008) Barriers to recruiting underrepresented populations to cancer clinical trials: a systematic review. *Cancer* 112 (2):228–242. [PubMed: 18008363]
13. James RD, Yu JH, Henrikson NB, Bowen DJ, Fullerton SM (2008) Strategies and stakeholders: minority recruitment in cancer genetics research. *Community Genet* 11 (4):241–249. [PubMed: 18417972]
14. Dang JH, Rodriguez EM, Luque JS, Erwin DO, Meade CD, Chen MS Jr. (2014) Engaging diverse populations about biospecimen donation for cancer research. *J Community Genet* 5 (4):313–327. [PubMed: 24664489]
15. Hann KEJ, Freeman M, Fraser L, Waller J, Sanderson SC, Rahman B, Side L, Gessler S, Lanceley A, team Ps (2017) Awareness, knowledge, perceptions, and attitudes towards genetic testing for cancer risk among ethnic minority groups: a systematic review. *BMC Public Health* 17 (1):503. [PubMed: 28545429]
16. Thompson B, Hebert JR (2014) Involving disparate populations in clinical trials and biobanking protocols: experiences from the community network program centers. *Cancer Epidemiol Biomarkers Prev* 23 (3):370–373. [PubMed: 24609845]
17. Partridge EE (2014) Yes, minority and underserved populations will participate in biospecimen collection. *Cancer Epidemiol Biomarkers Prev* 23 (6):895–897. [PubMed: 24895442]
18. Hagiwara N, Berry-Bobovski L, Francis C, Ramsey L, Chapman RA, Albrecht TL (2014) Unexpected findings in the exploration of African American underrepresentation in biospecimen collection and biobanks. *J Cancer Educ* 29 (3):580–587. [PubMed: 24243440]
19. John EM, Hopper JL, Beck JC, Knight JA, Neuhausen SL, Senie RT, Ziogas A, Andrulis IL, Anton-Culver H, Boyd N, Buys SS, Daly MB, O'Malley FP, Santella RM, Southey MC, Venne VL, Venter DJ, West DW, Whittemore AS, Seminars D (2004) The Breast Cancer Family Registry: an infrastructure for cooperative multinational, interdisciplinary and translational studies of the genetic epidemiology of breast cancer. *Breast Cancer Res* 6 (4):R375–389 [PubMed: 15217505]
20. Terry MB, Phillips KA, Daly MB, John EM, Andrulis IL, Buys SS, Goldgar DE, Knight JA, Whittemore AS, Chung WK, Apicella C, Hopper JL (2016) Cohort Profile: The Breast Cancer Prospective Family Study Cohort (ProF-SC). *Int J Epidemiol* 45 (3):683–692. [PubMed: 26174520]
21. Hopper JL, Bishop DT, Easton DF (2005) Population-based family studies in genetic epidemiology. *Lancet* 366 (9494):1397–1406. [PubMed: 16226618]
22. Yost K, Perkins C, Cohen R, Morris C, Wright W (2001) Socioeconomic status and breast cancer incidence in California for different race/ethnic groups. *Cancer Causes Control* 12 (8):703–711. [PubMed: 11562110]
23. John EM, Phipps AI, Davis A, Koo J (2005) Migration history, acculturation, and breast cancer risk in Hispanic women. *Cancer Epidemiol Biomarkers Prev* 14 (12):2905–2913 [PubMed: 16365008]
24. John EM, Schwartz GG, Koo J, Wang W, Ingles SA (2007) Sun exposure, vitamin D receptor gene polymorphisms and breast cancer risk in a multiethnic population. *Am J Epidemiol* 166 (12):1409–1419. [PubMed: 17934201]
25. Signorello LB, Hargreaves MK, Blot WJ (2010) The Southern Community Cohort Study: investigating health disparities. *J Health Care Poor Underserved* 21 (1 Suppl):26–37.
26. Bussey-Jones J, Garrett J, Henderson G, Moloney M, Blumenthal C, Corbie-Smith G (2010) The role of race and trust in tissue/blood donation for genetic research. *Genet Med* 12 (2):116–121. [PubMed: 20098329]

27. Adams-Campbell LL, Dash C, Palmer JR, Wiedemeier MV, Russell CW, Rosenberg L, Cozier YC (2016) Predictors of biospecimen donation in the Black Women's Health Study. *Cancer Causes Control* 27 (6):797–803. [PubMed: 27106577]
28. Cozier YC, Albert MA, Castro-Webb N, Coogan PF, Ridker P, Kaufman HW, Palmer JR, Rosenberg L (2016) Neighborhood socioeconomic status in relation to serum biomarkers in the Black Women's Health Study. *J Urban Health* 93 (2):279–291. [PubMed: 27000125]
29. Lacey JV Jr., Savage KE (2016) 50 % Response rates: half-empty, or half-full? *Cancer Causes Control* 27 (6):805–808. [PubMed: 27100357]
30. Lopez DS, Fernandez ME, Cano MA, Mendez C, Tsai CL, Wetter DW, Strom SS (2014) Association of acculturation, nativity, and years living in the United States with biobanking among individuals of Mexican descent. *Cancer Epidemiol Biomarkers Prev* 23 (3):402–408. [PubMed: 24609849]
31. Gabriel A, Cohen CC, Sun C (2014) Consent to specimen storage and continuing studies by race and ethnicity: a large dataset analysis using the 2011-2012 National Health and Nutrition Examination Survey. *ScientificWorldJournal* 2014:120891. [PubMed: 25485292]
32. Loffredo CA, Luta G, Wallington S, Makgoeng SB, Selsky C, Mandelblatt JS, Adams-Campbell LL, Region 1 Bio-specimen Management of Cancer Health Disparities P (2013) Knowledge and willingness to provide research biospecimens among foreign-born Latinos using safety-net clinics. *J Community Health* 38 (4):652–659. [PubMed: 23543371]
33. Hohl SD, Gonzalez C, Carosso E, Ibarra G, Thompson B (2014) "I did it for us and I would do it again": perspectives of rural latinos on providing biospecimens for research. *Am J Public Health* 104 (5):911–916. [PubMed: 24625153]
34. Ceballos RM, Knerr S, Scott MA, Hohl SD, Malen RC, Vilchis H, Thompson B (2014) Latino beliefs about biomedical research participation: a qualitative study on the U.S.-Mexico border. *J Empir Res Hum Res Ethics* 9 (4):10–21. [PubMed: 25747293]
35. Rodriguez EM, Saad-Harfouche FG, Miller A, Mahoney MC, Ambrosone CB, Morrison CD, Underwood WR, Erwin DO (2016) Engaging diverse populations in biospecimen donation: results from the Hoy y Manana study. *J Community Genet* 7 (4):271–277. [PubMed: 27488840]
36. Nodora JN, Komenaka IK, Bouton ME, Ohno-Machado L, Schwab R, Kim HE, Farcas C, Perez G, Elena Martinez M (2017) Biospecimen Sharing Among Hispanic Women in a Safety-Net Clinic: Implications for the Precision Medicine Initiative. *J Natl Cancer Inst* 109 (2).
37. Wenzel L, Bowen D, Habbal R, Leighton N, Vu T, Anton-Culver H (2008) Testing targeted approaches to enhance Cancer Genetics Network minority recruitment within Asian populations. *Community Genet* 11 (4):234–240. [PubMed: 18417971]
38. Gao W, Ma GX, Tan Y, Fang C, Weaver J, Jin M, Lai P (2014) Factors associated with willingness to participate in biospecimen research among Chinese Americans. *Biopreserv Biobank* 12 (2):131–138. [PubMed: 24749880]
39. Tong EK, Fung LC, Stewart SL, Paterniti DA, Dang JH, Chen MS Jr. (2014) Impact of a biospecimen collection seminar on willingness to donate biospecimens among Chinese Americans: results from a randomized, controlled community-based trial. *Cancer Epidemiol Biomarkers Prev* 23 (3):392–401. [PubMed: 24609848]
40. Gao W, Ma GX, Tan Y, Fang C, Weaver J, Jin M, Lai P, Godwin AK (2014) Culturally appropriate education intervention on biospecimen research participation among Chinese Americans. *Cancer Epidemiol Biomarkers Prev* 23 (3):383–391. [PubMed: 24609847]
41. Ma GX, Tan Y, Blakeney NC, Seals BF, Ma XS, Zhai S, Liu A, Tai Y, Michaels M (2014) The impact of a community-based clinical trial educational intervention among underrepresented Chinese Americans. *Cancer Epidemiol Biomarkers Prev* 23 (3):424–432. [PubMed: 24092627]
42. McQuillan GM, Pan Q, Porter KS (2006) Consent for genetic research in a general population: an update on the National Health and Nutrition Examination Survey experience. *Genet Med* 8 (6): 354–360. [PubMed: 16778597]
43. Brown DR, Fouad MN, Basen-Engquist K, Tortolero-Luna G (2000) Recruitment and retention of minority women in cancer screening, prevention, and treatment trials. *Ann Epidemiol* 10 (8 Suppl):S13–21. [PubMed: 11189088]

44. Hughes C, Peterson SK, Ramirez A, Gallion KJ, McDonald PG, Skinner CS, Bowen D (2004) Minority recruitment in hereditary breast cancer research. *Cancer Epidemiol Biomarkers Prev* 13 (7):1146–1155. [PubMed: 15247125]
45. Brown BA, Long HL, Gould H, Weitz T, Milliken N (2000) A conceptual model for the recruitment of diverse women into research studies. *J Womens Health Gend Based Med* 9 (6):625–632. [PubMed: 10957751]
46. Paskett ED, Reeves KW, McLaughlin JM, Katz ML, McAlearney AS, Ruffin MT, Halbert CH, Merete C, Davis F, Gehlert S (2008) Recruitment of minority and underserved populations in the United States: the Centers for Population Health and Health Disparities experience. *Contemp Clin Trials* 29 (6):847–861. [PubMed: 18721901]
47. Greiner KA, Friedman DB, Adams SA, Gwede CK, Cupertino P, Engelman KK, Meade CD, Hebert JR (2014) Effective recruitment strategies and community-based participatory research: community networks program centers' recruitment in cancer prevention studies. *Cancer Epidemiol Biomarkers Prev* 23 (3):416–423. [PubMed: 24609851]
48. Wallington SF, Dash C, Sheppard VB, Goode TD, Opong BA, Dodson EE, Hamilton RN, Adams-Campbell LL (2016) Enrolling Minority and Underserved Populations in Cancer Clinical Research. *Am J Prev Med* 50 (1):111–117. [PubMed: 26470805]
49. Giuliano AR, Mokuau N, Hughes C, Tortolero-Luna G, Risendal B, Ho RCS, Prewitt TE, McCaskill-Stevens WJ (2000) Participation of minorities in cancer research: the influence of structural, cultural, and linguistic factors. *Ann Epidemiol* 10 (8 Suppl):S22–34. [PubMed: 11189089]
50. Salman A, Nguyen C, Lee YH, Cooksey-James T (2016) A Review of Barriers to Minorities' Participation in Cancer Clinical Trials: Implications for Future Cancer Research. *J Immigr Minor Health* 18 (2):447–453. doi:10.1007/s10903-015-0198-9. [PubMed: 25822567]
51. Dash C, Wallington SF, Muthra S, Dodson E, Mandelblatt J, Adams-Campbell LL (2014) Disparities in knowledge and willingness to donate research biospecimens: a mixed-methods study in an underserved urban community. *J Community Genet* 5 (4):329–336. [PubMed: 24771039]

Table 1

Participation in screening interview, by race/ethnicity and neighborhood socioeconomic status

	Female breast cancer cases diagnosed at ages 18-64 years ^{a,b} n = 29,559			
	n	%	OR ^c	95% CI
Participation in screening	25,183	85		
Race/ethnicity ^d				
Non-Hispanic whites	16,976	87	1.0	
Hispanics	2,259	88	1.11	0.97-1.26
African Americans	1,713	87	1.08	0.94-1.26
Asian Americans	3,650	77	0.52	0.48-0.56
Chinese	1,370	76	0.47	0.42-0.53
Japanese	345	78	0.51	0.41-0.65
Filipinas	1,031	81	0.64	0.55-0.74
Other Asian Americans/Pacific Islander	904	76	0.48	0.42-0.56
Others ^e	585	79	0.54	0.45-0.65
P value for differences by race/ethnicity ^f		<0.01	<0.01	
Age at diagnosis (years)				
18-34	505	83	1.05	0.84-1.31
35-49	9,365	86	1.21	1.13-1.30
50-64	15,313	85	1.0	
P value for differences by age ^f		<0.01	<0.01	
Stage at diagnosis ^g				
In situ	1,316	89	1.38	1.16-1.65
Localized	12,667	86	1.0	
Regional	6,733	87	1.06	0.98-1.15
Distant	421	80	0.63	0.51-0.79
Missing	4,046	79	0.61	0.56-0.66
P value for differences by stage ^f		0.01	<0.01	
Neighborhood socioeconomic status (quintiles) ^{h,i}				
1 (low)	5,026	84	1.0	
2	4,948	85	0.93	0.84-1.04
3	4,979	85	0.95	0.86-1.06
4	5,035	86	1.02	0.92-1.14
5 (high)	5,055	86	1.04	0.94-1.16
P value for differences by neighborhood SES ^f		0.10	0.18	

Abbreviations: *SES* socioeconomic status, *OR* odds ratio, *CI* confidence interval, *SEER* Surveillance, Epidemiology, and End Results Program^aIncludes incident cases diagnosed in Greater Bay Area (Alameda, Contra Costa, Marin, Monterey, San Francisco, San Mateo, Santa Clara, or Santa Cruz counties) from 1995-2009 and incident cases diagnosed in Sacramento or Solano counties from 2005-2006.^bExcludes 361 cases from Phase I recruitment who were enrolled in the family registry as probands without telephone screening for eligibility.

^cMutually adjusted for race/ethnicity, age at diagnosis, stage, and neighborhood SES.

^dBased on cancer registry records.

^eIncludes 28 Native American cases, 48 cases listed as other race/ethnicity, and 509 cases of unknown race/ethnicity.

^f*P* value from chi square test; in multivariable models, *P* value from Wald test.

^gSEER summary stage.

^hNeighborhood SES based on a composite measure of seven SES indicators from Census data at the level of block groups and categorized according to the quintile distribution of eligible breast cancer cases.

ⁱNeighborhood SES was missing for 151 cases.

Table 2

Proband enrollment and biospecimen collection, by race/ethnicity, age at diagnosis, stage, first-degree cancer family history, neighborhood socioeconomic status, education, and country of birth.

	Breast cancer cases eligible as probands <i>n</i> = 4,780						Enrolled probands <i>n</i> = 3,620 ^{<i>a</i>}							
	Enrollment ^{<i>a</i>}			Biospecimen collection ^{<i>b</i>}			Enrollment ^{<i>a</i>}			Biospecimen collection ^{<i>c</i>}				
	<i>n</i>	%	OR ^{<i>d</i>}	95% CI	<i>n</i>	%	<i>n</i>	%	OR ^{<i>e</i>}	95% CI	<i>n</i>	%	OR ^{<i>e</i>}	95% CI
Overall participation	3,620	76			3,244	68					3,244	90		
Race/ethnicity ^{<i>f</i>}														
Non-Hispanic whites	866	81	1.0		816	76					816	94	1.0	
Hispanics	1,072	75	0.85	0.69-1.04	1,013	71					1,013	95	1.55	0.99-2.41
African Americans	831	76	0.88	0.71-1.10	762	69					762	92	0.83	0.56-1.25
Asian Americans	851	72	0.68	0.55-0.83	653	55					653	76	0.30	0.20-0.45
Chinese	387	74	0.80	0.62-1.04	279	54					279	72	0.23	0.15-0.36
Japanese	94	75	0.77	0.50-1.20	83	66					83	88	0.51	0.25-1.02
Filipinas	267	66	0.51	0.39-0.66	201	50					201	75	0.27	0.16-0.44
Others ^{<i>g</i>}	103	75	0.79	0.52-1.21	90	65					90	87	0.50	0.25-1.00
<i>P</i> value for differences by race/ethnicity ^{<i>h</i>}		<0.01	<0.01			<0.01						<0.01		<0.01
Age at diagnosis (years)														
18-34	371	76	1.0		343	70					343	92	1.0	
35-49	1,395	75	0.93	0.73-1.18	1,252	67					1,252	90	1.57	1.01-2.43
50-64	1,854	76	0.93	0.80-1.07	1,649	68					1,649	89	1.12	0.88-1.42
<i>P</i> value for differences by age ^{<i>h</i>}		0.55	0.56			0.40						0.13	0.12	
Stage at diagnosis ^{<i>i</i>}														
In situ	299	79	1.01	0.77-1.33	277	73					277	93	1.32	0.82-2.14
Localized	1,915	78	1.0		1,745	71					1,745	91	1.0	
Regional	1,087	79	1.06	0.90-1.24	965	70					965	89	0.77	0.59-1.00
Distant	70	76	0.88	0.54-1.44	55	60					55	79	0.36	0.19-0.68
Missing	249	52	0.31	0.26-0.39	202	42					202	42	0.57	0.39-0.82

	Breast cancer cases eligible as probands <i>n</i> = 4,780						Enrolled probands <i>n</i> = 3,620 ^a					
	Enrollment ^d			Biospecimen collection ^b			Enrollment ^d			Biospecimen collection ^c		
	<i>n</i>	%	OR ^d	95% CI	<i>n</i>	%	OR ^e	95% CI	<i>n</i>	%	OR ^e	95% CI
<i>P</i> value for differences by stage ^h		0.01	<0.01			0.01				<0.01		<0.01
Cancer family history ^j												
Yes	1,388	78	1.06	0.91-1.23	1,269	71			1,269	91	1.10	0.85-1.42
No	2,232	75	1.0		1,975	66			1,975	88	1.0	
<i>P</i> value for differences by family history ^h		0.01	0.46			<0.01				<0.01		0.45
Neighborhood socioeconomic status ^k												
1 (low)	680	78	1.0		609	70			609	90	1.0	
2	718	76	0.89	0.71-1.11	646	68			646	90	1.03	0.72-1.48
3	690	75	0.84	0.67-1.04	623	67			623	90	1.08	0.75-1.57
4	745	75	0.86	0.69-1.07	661	66			661	89	0.91	0.64-1.30
5 (high)	787	75	0.88	0.71-1.09	705	68			705	90	1.00	0.70-1.43
<i>P</i> value for differences by neighborhood SES ^h		0.45	0.57			0.58				0.90		0.92
Education ^l												
Some high school or less									469	91	1.0	
High school graduate									476	87	0.60	0.39-0.93
Some college or other ^m									1,129	92	1.20	0.79-1.82
College graduate									1,164	88	0.98	0.66-1.48
<i>P</i> value for differences by education ^h										<0.01		<0.01
Country of birth ⁿ												
U.S.-born									2,128	93	1.0	
Foreign-born									1,115	84	0.66	0.49-0.90
<i>P</i> value for differences by country of birth ^h										<0.01		0.01

Abbreviations: *SES* socioeconomic status, *OR* odds ratio, *CI* confidence interval

^aCompleted the family history and risk factor questionnaires.

- ^q Completed the family history and risk factor questionnaires and provided a blood or mouthwash sample.
- ^r Enrolled probands who provided a blood or mouthwash sample.
- ^s Odds ratio of participation and 95% confidence interval, mutually adjusted for race/ethnicity, age at diagnosis, neighborhood SES.
- ^t Odds ratio of participation and 95% confidence interval, mutually adjusted for race/ethnicity, age at diagnosis, neighborhood SES, education, and country of birth, stage, cancer family history, and stage, cancer family history.
- ^u Based on self-report.
- ^v Includes other Asian Americans, Pacific Islanders, Native Americans, and mixed race/ethnicity.
- ^w *P* value from chi square test; in multivariable models, *P* value from Wald test.
- ^x SEER summary stage.
- ^y Family history of breast, ovarian or childhood cancer in first-degree relatives.
- ^z Neighborhood SES based on a composite measure of seven SES indicators from Census data at the level of block groups and categorized according to the quintile distribution of eligible breast cancer cases.
- ^{aa} Education was missing for 14 probands.
- ^{ab} Technical or vocational school.
- ^{ac} Country of birth was missing for 4 probands.

Table 3

First-degree relative enrollment and biospecimen collection, by race/ethnicity, age, personal cancer history, education, and country of birth

	Eligible relatives ^a n = 4,279				Enrolled relatives ^b n = 3,306					
	Enrollment ^b		Biospecimen collection ^c		Biospecimen collection ^c		Biospecimen collection ^d			
	n	%	OR ^e	95% CI	n	%	n	%	OR ^f	95% CI
Overall participation	3,306	77			2,774	65	2,774	84		
Race/ethnicity ^g										
Non-Hispanic whites	872	87	1.0		755	76	755	87	1.0	
Hispanics	1,045	77	0.47	0.37-0.61	907	67	907	87	1.00	0.71-1.42
African Americans	807	74	0.44	0.34-0.57	660	60	660	82	0.70	0.51-0.97
Asian Americans	582	70	0.35	0.26-0.45	452	54	452	78	0.59	0.42-0.82
Chinese	204	67	0.29	0.20-0.43	144	48	144	71	0.40	0.25-0.62
Japanese	78	80	0.58	0.33-1.01	69	70	69	88	1.27	0.62-2.60
Filipinas	226	67	0.32	0.22-0.45	178	53	178	79	0.61	0.37-1.02
Others ^h	74	76	0.46	0.24-0.87	61	63	61	82	0.58	0.32-1.06
P value for differences by race/ethnicity ⁱ		<0.01	<0.01			<0.01		<0.01	<0.01	
Proband's age at diagnosis (years)										
18-34	396	84	1.0		357	75				
35-49	1,355	77	0.75	0.55-1.02	1,135	64				
50-64	1,555	76	0.68	0.50-0.92	1,282	63				
P value for differences by age ⁱ		<0.01	0.03			<0.01				
Age at interview (years)										
18-34							233	91	1.0	
35-49							906	85	0.58	0.37-0.90
50-64							1,106	86	0.68	0.44-1.05
65							529	76	0.47	0.30-0.75
P value for differences by age ⁱ								<0.01	<0.01	
Personal cancer history ^j										

	Eligible relatives ^a n = 4,279				Enrolled relatives ^b n = 3,306			
	Enrollment ^b		Biospecimen collection ^c		Biospecimen collection ^d		Biospecimen collection ^d	
	n	%	n	%	n	%	OR ^f	95% CI
No	2,766	77	2,324	65	2,324	84	1.0	
Yes	540	78	450	65	450	83	1.20	0.91-1.59
<i>P</i> value for differences by personal cancer history ⁱ								
Education ^k								
Some high school or less		0.67		0.90		0.69		0.19
High school graduate								
Some college or other ^l								
College graduate								
<i>P</i> value for differences by education ⁱ								
Country of birth								
U.S.-born								0.65
Foreign-born								<0.01
<i>P</i> value for differences by country of birth ⁱ								

^aAdult first-degree relatives: mothers, fathers, full and half-sisters, and adult daughters and sons with a prior diagnosis of breast, ovarian or childhood cancer.

^bCompleted risk factor questionnaire.

^cCompleted risk factor questionnaire and provided a blood or mouthwash sample.

^dEnrolled relatives who provided a blood or mouthwash sample.

^eOdds ratio of participation and 95% confidence interval, mutually adjusted for race/ethnicity, proband's age at diagnosis, and personal cancer history.

^fOdds ratio of participation and 95% confidence interval, mutually adjusted for race/ethnicity, age at interview, personal cancer history, education, and country of birth.

^gFor enrolled relatives, based on self-report; for eligible relatives who did not enroll, based on proband's self-reported race/ethnicity.

^hIncludes other Asian Americans, Pacific Islanders, Native Americans, and mixed race/ethnicity.

ⁱ*P* value from chi square test; in multivariable models, *P* value from Wald test.

^jPersonal history of breast, ovarian or childhood cancer.

Education was unknown for 18 relatives.
Technical or vocational school.

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Table 4 Biospecimen collection for enrolled Hispanic and Asian American probands and first-degree relatives, by migration history

	Hispanics						Asian Americans							
	<i>n</i>	%	OR ^c	95% CI	Enrolled relatives ^b <i>n</i> = 1,045	Enrolled probands ^a <i>n</i> = 1,072	<i>n</i>	%	OR ^c	95% CI	Enrolled relatives ^b <i>n</i> = 851	Enrolled probands ^a <i>n</i> = 582		
Overall biospecimen collection ^e	1013	95			907	87	653	77			452	78		
Education ^f														
Some high school or less	347	96	1.0		354	90	39	62	1.0		28	55	1.0	
High school graduate	183	93	0.46	0.19-1.11	211	89	0.95	0.51-1.76	0.52	0.24-1.13	58	81	2.63	1.20-5.79
Some college or other ^g	301	95	0.57	0.24-1.39	245	86	0.73	0.42-1.28	1.45	0.64-3.28	134	88	3.62	1.69-7.79
College graduate	181	94	0.42	0.16-1.13	93	82	0.58	0.29-1.13	1.11	0.50-2.46	229	81	2.65	1.31-5.39
<i>P</i> value for differences by education ^h		0.43	0.29			0.02	0.18			0.01	<0.01	0.04		
Country of birth														
U.S.-born	474	94	1.0		483	84	1.0		1.0		178	87	1.0	
Foreign-born	539	95	1.35	0.59-0.9	424	90	2.04	1.04-3.99	0.56	0.35-0.91	273	73	0.62	0.37-1.04
<i>P</i> value for differences by country of birth ^h		0.83	0.48			0.01	0.03			0.02	<0.01	0.08		
Age at migration to the U.S. (years)														
U.S.-born	474	94	1.0		483	84	1.0		1.0		178	87	1.0	
<20	156	93	0.97	0.42-2.26	174	91	1.94	1.00-3.77	0.60	0.32-1.13	59	79	0.65	0.33-1.27
20-29	221	97	3.65	1.06-12.6	134	93	3.38	1.24-9.23	0.51	0.30-0.86	106	81	0.63	0.34-1.17
30-39	104	93	1.50	0.42-5.31	60	87	2.16	0.71-6.55	0.59	0.32-1.07	53	74	0.62	0.29-1.31
40	52	95	2.44	0.49-12.2	48	92	3.89	0.83-18.3	0.70	0.35-1.40	43	61	0.45	0.20-1.03
<i>P</i> value for differences by age at migration ^h		0.54	0.18			0.05	0.09			0.16	<0.01	0.38		
Years of residence in the U.S.														
U.S.-born	474	94	1.0		483	84	1.0		1.0		178	87	1.0	
40	86	92	0.96	0.36-2.53	73	85	1.19	0.59-2.41	1.03	0.37-2.93	39	87	1.17	0.46-2.97
30-39	106	97	2.70	0.67-10.9	82	96	5.92	1.76-19.9	0.42	0.24-0.74	55	75	0.57	0.25-1.27
20-29	132	93	1.11	0.37-3.30	96	89	2.46	1.02-5.97	0.61	0.35-1.08	82	71	0.41	0.22-0.77
<20	209	96	2.37	0.70-8.10	163	93	3.90	1.30-11.7	0.62	0.35-1.10	86	75	0.66	0.32-1.33

	Hispanics						Asian Americans					
	Enrolled probands ^a n = 1,072		Enrolled relatives ^b n = 1,045		Enrolled probands ^a n = 851		Enrolled relatives ^b n = 582					
	n	%	OR ^c	95% CI	n	%	OR ^d	95% CI	n	%	OR ^d	95% CI
<i>P</i> value for differences by years of residence in the U.S. ^h		0.61	0.35		<0.01	0.01		<0.01	0.04		<0.01	0.10
English proficiency ^f												
Well or English only	621	95	1.0		595	86	1.0		487	80	1.0	
Medium	58	97	0.91	0.18-4.59	40	89	0.67	0.25-1.78	89	74	0.83	0.50-1.38
Little	125	95	0.41	0.12-1.38	99	93	0.75	0.28-1.99	49	70	0.84	0.41-1.74
Not at all	207	94	0.31	0.09-1.07	172	88	0.39	0.14-1.08	12	35	0.21	0.08-0.56
<i>P</i> value for differences by English proficiency ^h		0.86	0.27		0.21	0.39		<0.01	0.01		<0.01	0.94

^aCompleted family history and risk factor questionnaires.

^bAdult first-degree relatives (mothers, fathers, full and half-sisters, and adult daughters and sons with a personal history of breast, ovarian or childhood cancer) who completed risk factor questionnaire.

^cOdds ratio of participation and 95% confidence interval, adjusted for age at diagnosis, stage (in situ, localized, regional, distant or missing), cancer family history, neighborhood SES, education, English language proficiency.

^dOdds ratio of participation and 95% confidence interval, adjusted for age at interview, personal cancer history, education, English language proficiency.

^eBlood or mouthwash sample.

^fAlso adjusted for age at migration.

^gTechnical or vocational school.

^h*P* value from chi square test; in multivariable models, *P* value from Wald test.