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Impact of a Randomized Controlled Educational Trial to Improve Physician Practice Behaviors Around Screening for Inherited Breast Cancer

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BACKGROUND: Many primary care physicians (PCPs) are ill-equipped to provide screening and counseling for inherited breast cancer.

OBJECTIVE: To evaluate the outcomes of an interactive web-based genetics curriculum versus text curriculum for primary care physicians.

DESIGN: Randomized two-group design.

PARTICIPANTS: 121 California and Pennsylvania community physicians.

INTERVENTION: Web-based interactive genetics curriculum, evaluated against a control group of physicians who studied genetics review articles. After education, physicians interacted with an announced standardized patient (SP) at risk for inherited breast cancer.

MAIN MEASURES: Transcripts of visit discussions were coded for presence or absence of 69 topics relevant to inherited breast cancer.

KEY RESULTS: Across all physicians, history-taking, discussions of test result implications, and exploration of ethical and legal issues were incomplete. Approximately half of physicians offered a genetic counseling referral (54.6 %), and fewer (43.8 %) recommended testing. Intervention physicians were more likely than controls to explore genetic counseling benefits (78.3 % versus 60.7 %, $P=0.048$), encourage genetic counseling before testing (38.3 % versus 21.3 %, $P=0.048$), ask about a family history of prostate cancer (25.0 % versus 6.6 %, $P=0.006$), and report that a positive result indicated an increased risk of prostate cancer for male relatives (20.0 % versus 1.6 %, $P=0.001$). Intervention-group physicians were less likely than controls to ask about Ashkenazi heritage (13.3 % versus 34.4 %, $P=0.01$) or to reply that they would get tested when asked, "What would you do?" (33.3 % versus 54.1 %, $P=0.03$).

CONCLUSIONS: Physicians infrequently performed key counseling behaviors, and this was true regardless of whether they had completed the web-based interactive training or read clinical reviews.

KEY WORDS: inherited breast cancer; physician training; BRCA; genetic counseling; genetic testing.

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INTRODUCTION

A BRCA mutation is present in about 1 in 400 women in the general population¹; for these women, the average cumulative cancer risk by age 70 ranges from 45 % to 65 % for breast cancer and from 11 % to 39 % for ovarian cancer. Those with BRCA mutations account for 7 % of all breast cancers and 10 % to 15 % of ovarian cancers.^{2,3} The discovery of links between BRCA mutations and breast cancer creates an opportunity to identify individuals at risk and to intervene early in the disease process.⁴

Knowing how to screen appropriately for genetic risk factors for breast cancer and when to offer genetic testing are core skills for primary care physicians (PCPs), especially given the relative scarcity of genetic counselors.^{5,6} Identification of women with genetic mutations allows for aggressive surveillance, risk-reduction interventions, and identification of family members who may also be at risk.^{7,8} Many PCPs have an inadequate understanding of hereditary breast cancer, risk assessment, shared decision-making, and legal and ethical issues,^{9,10} and this results in inappropriate testing: underuse of testing and genetic counseling referral for patients at high risk^{9,11–18} and overuse for those at low risk.¹⁸

Various interventions aimed at increasing patient knowledge have been shown to be effective.^{19–25} That said, there are few educational interventions focusing on hereditary breast cancer that target physicians or trainees.²⁶ Here, we report an evaluation of an interactive web-based genetics curriculum for PCPs using announced standardized patient (SP) visits. The SP approach provides insights into clinical behaviors not provided by physician and patient surveys.²⁷ Physicians were not told that the SP visit would focus on breast cancer. Our primary research objective was to determine whether the educational intervention was more effective than a control

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curriculum at eliciting: 1) appropriate physician behaviors with regard to assessments of the patient’s risk for hereditary breast cancer; 2) discussion of genetic testing and counseling; and 3) exploration of ethical, legal, and social issues (ELSI) surrounding genetic testing.

METHODS

Setting and Participants

The study took place in California (Los Angeles and the Greater Sacramento areas) and rural Pennsylvania. PCPs were eligible if they were an MD or DO, English-speaking, and had Internet and e-mail access. In California, a list of PCPs was compiled through an internet search. Physicians were then sent information about the study through faxes and flyers. Colleagues at clinics in two large health systems made recruitment appeals on our behalf. In Pennsylvania, PCPs were identified via the Pennsylvania Area Health Education Center, which sent personalized letters of invitation,

recruitment flyers, and business reply postcards to prospective participants. The Pennsylvania State University (PSU) team sent recruitment materials to PCPs around the state. Across all sites, 121 PCPs were recruited between September 2011 and April 2013 (Fig. 1). Participants were offered six units of continuing medical education (CME) credit and a payment of \$250 after study completion; credit and payment were not affected by performance during the SP visit.

Design and Procedure Overview

Study procedures were IRB-approved in California and Pennsylvania. Each physician was randomly assigned to an intervention group ($n=60$) or control group ($n=61$). Upon enrollment, physicians participated in a pre-intervention objective structured video exercise (OSVE), responded to an online pre-visit questionnaire, completed the educational curriculum, took an online post-visit survey, and finally, completed a post-intervention OSVE. OSVEs provided no instruction on inherited breast cancer. After the curriculum had been completed, a time was scheduled for the SP visit. PCPs were

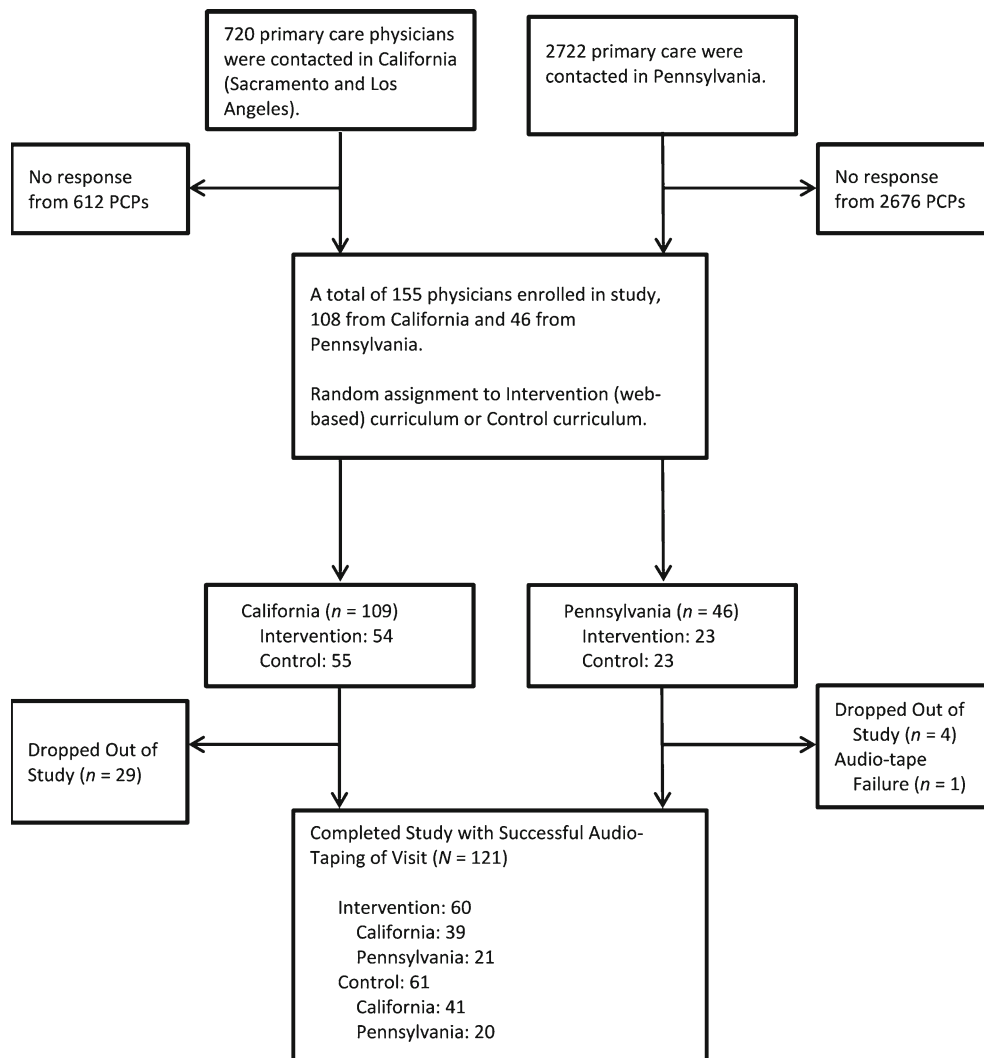


Fig. 1 Study flow diagram.

told only that the SP would be used to evaluate the curriculum; no mention was made of the clinical condition (breast cancer). Visits took place an average of one month after completion of the curriculum. These visits were audio-taped, transcribed, and then coded.

Intervention and Control Group Curricula

Intervention Group. Intervention-group physicians completed a six-hour interactive web-based curriculum covering information about genetic testing, risk assessment, practice behaviors, and communication skills. The curriculum featured four clinical patient cases, with tutorials developed around specific learning objectives to illustrate common genetic conditions likely to be encountered by PCPs—breast cancer, cystic fibrosis, Huntington’s Disease, and inherited thrombophilia. Each case included video vignettes that modeled physician communication, raised questions requiring application of principles, and provided hyperlinks to additional written and video materials. The tutorials included in-depth considerations of epidemiology and ELSI related to genetics. A more complete description of the technology underlying the intervention has been reported elsewhere.⁴ Essential competencies were derived from organizational recommendations,²⁸ published reports,²⁹ and the input of 20 expert consultants from six academic institutions.⁴ Because of resource limitations, just one of the four medical conditions, breast cancer, was addressed in the SP study described below.

Control Group. Control group participants read eight review articles from leading journals and an information sheet extracted from the National Cancer Institute website.^{30–38} These nine resources covered clinical genetics, ethical issues, doctor–patient issues, and clinical reasoning, and were topically similar to the material contained in the intervention group. Participant access and downloading of the articles was recorded electronically. It was estimated that physicians would need at least six hours to read all articles. The use of such a control group allowed us to examine the impact of our web-based curriculum relative to a more traditional learning format. Given the newness of the genetics information provided in the intervention, evaluating it against a control group of untaught physicians would have been a less meaningful comparison.

Standardized Patient Procedure

Each physician was visited by one of five actresses playing the role of “Catherine Douglas,” a trained and rehearsed SP portraying a woman at risk for inherited breast cancer. For logistic reasons, SP visits were announced to the clinics in advance. Catherine arrived at the physician office with documents, purportedly prepared by her last physician from another state, describing her as a 41-year old, divorced, college-educated

white woman with an unremarkable medical history other than intermittent headaches. She had a son (age 13) and daughter (age 17). She had just moved to the area to take care of her younger sister, who was recently diagnosed with breast cancer.

Catherine initiated the encounter with the following: “My sister was recently diagnosed with breast cancer. Although I am very concerned about her, I was wondering how much I have to worry about getting breast cancer myself.” Though her mammogram was normal, she had read a story about genetic testing for breast cancer and would like to know more. If the physician asked about her family history, Catherine reported that her mother had been diagnosed with breast cancer at age 50, had a mastectomy, but died from the disease at age 52. If asked about other cancers in the family, the doctor would learn that Catherine’s maternal aunt had died at age 40 of cancer—possibly ovarian cancer, but Catherine is uncertain. Answers to anticipated physician questions were scripted such that Catherine had a greater-than-average risk of breast cancer and would be a reasonable candidate for genetic counseling and testing.

Coding of Physician Behavior

One author (HD) coded transcripts of all visits, and a second (RB) independently coded 60 visits in order to assess coding reliability (average Cohen’s kappa, 0.91). Coding consisted of a determination of the presence or absence of 69 specific physician behaviors pertaining to the SP’s family history and personal health history, implications of genetic test results for the SP and her family, ELSI, genetic counseling, and genetic tests. These codes (described below) captured widely accepted core competencies.^{39,40}

Statistical Analysis

Data were analyzed using Stata (version 12.1). Descriptive statistics were used to describe characteristics of the sample. Cross-tabulations were used to compare intervention and control groups on dichotomous behavioral outcomes. Fisher’s exact test was used to test for statistical significance.

RESULTS

Physician and Visit Characteristics

Physicians were primarily male, white, non-Hispanic, and middle-aged (Table 1). The SP visit occurred an average of one month after completion of learning activities. There were no significant differences between the intervention and control groups with regard to demographics, years of practice, or experience with inherited breast cancer. Additionally, there were no significant interactions between region (CA versus PA) and study variables, allowing for aggregation of data across states.

Table 1 Physician Characteristics by Study Group

Characteristic	Control (n=61)	Intervention (n=60)	Combined (n=121)	P*
Age, mean (SD)	48.7 (9.8)	49.1 (10.6)	48.9 (10.1)	0.81
Years since MD, mean (SD)	21.7 (9.7)	21.3 (12.3)	21.5 (11.0)	0.86
Female, %	34.4	46.7	40.5	0.20
White race, %	67.2	70.0	68.6	0.85
Hispanic, %	4.9	8.3	6.6	0.49
Experience, either in physician practice or personally, with inherited breast cancer, %	49.2	53.3	51.2	0.72

* The significance of mean differences between study groups were examined with *t* tests. Differences for the categorical demographic and experience variables were examined with Fisher's exact test. Nonsignificant differences are suggestive of successful randomization of participants to the two study arms.

Clinical Behaviors

History-Taking. The number (and percentage) of physicians who asked about each of 10 family issues is reported in the top section of Table 2. Physicians asked an average of 2.2 (SD=1.5) questions relating to the family history issues listed in the table. For only one topic (SP's mother's age at onset of breast cancer) did more than 50 % of physicians ascertain information. Specific questions about cancers in the family, including ovarian, breast, and prostate cancers, were not usually asked. Significant differences were found between physicians in the control and intervention groups on two family history variables. Intervention-group participants were

more likely than control physicians to ask about a history of prostate cancer among relatives, but were less likely to ask about Ashkenazi Jewish heritage.

Queries about the SP's personal history are reported in the bottom section of Table 2. Physicians asked an average of 2.0 (SD=1.5) of these 11 personal history questions. More than two-thirds asked the SP about her age and whether she underwent regular mammography screening. All other personal history queries were made in less than 16 % of visits. There were no significant differences between intervention and control-group learners on any of the personal history-taking queries.

Discussions About Implications of Test Results. Counseling on test result implications for the SP are reported in Table 3. Intervention and control-group physicians did not differ significantly on any of these counseling behaviors. Physicians discussed an average of 5.2 (SD=1.8) of the 13 topics reported in the table. A majority of physicians informed the SP that a positive test result points to an increased risk of developing breast or ovarian cancer. With regard to treatment implications of a positive result, most physicians emphasized that such results would lead to a consideration of surgical options. Other implications of treatment were discussed by about one-fourth or less of the physicians. The third section of the table reports that most physicians counseled the SP about the implications of a positive result regarding ongoing screening, but few discussed what such a test result would mean for self-monitoring with breast exams. In the fourth section of Table 3, it is shown that the limitations of current genetic testing received very little attention. The last section of the table

Table 2 Physician History-Taking Queries, Sorted by Combined Frequency of Occurrence*

Topic	Control (n=61)	Intervention (n=60)	Combined (n=121)	P†
SP's family history‡				
Mother's age at onset of breast cancer	34 (55.7 %)	34 (56.8 %)	68 (56.2 %)	1.0
Other cancers in family	29 (47.5 %)	21 (35.0 %)	50 (41.3 %)	0.20
Any relative with ovarian cancer diagnosis	22 (36.1 %)	20 (33.3 %)	42 (34.7 %)	0.85
Second-degree relative w/ breast cancer	21 (34.4 %)	16 (26.7 %)	37 (30.6 %)	0.43
Ashkenazi Jewish heritage	21 (34.4 %)	8 (13.3 %)	29 (24.0 %)	0.01
Any male relative w/ prostate cancer	4 (6.6 %)	15 (25.0 %)	19 (15.7 %)	0.006
Was mother's breast cancer bilateral	4 (6.6 %)	4 (6.7 %)	8 (6.6 %)	1.0
Was sister's breast cancer bilateral	3 (4.9 %)	4 (6.7 %)	7 (5.8 %)	0.72
Third-degree relative with breast cancer	5 (8.2 %)	1 (1.7 %)	6 (5.0 %)	0.21
Any male relative w/ breast cancer	3 (4.9 %)	2 (3.3 %)	5 (4.1 %)	1.0
SP's personal history				
Regular mammography	45 (73.8 %)	46 (76.7 %)	91 (75.2 %)	0.83
Current age	43 (70.5 %)	40 (66.7 %)	83 (68.6 %)	0.70
Current or past use of oral contraceptive pills	11 (18.0 %)	8 (13.3 %)	19 (15.7 %)	0.62
Lifetime no. of pregnancies	9 (14.8 %)	8 (13.3 %)	17 (14.1 %)	1.0
Regular intake of alcohol	7 (11.5 %)	5 (8.3 %)	12 (9.9 %)	0.76
Breastfeeding of children	4 (6.6 %)	4 (6.7 %)	8 (6.6 %)	1.0
Age at first live birth	3 (4.9 %)	2 (3.3 %)	5 (4.1 %)	1.0
Weight/BMI	2 (3.3 %)	1 (1.7 %)	3 (2.5 %)	1.0
Past breast biopsy	2 (3.3 %)	1 (1.7 %)	3 (2.5 %)	1.0
Age at first menstrual period	2 (3.3 %)	0 (0.0 %)	2 (1.7 %)	0.50
Current or past hormone replacement therapy	2 (3.3 %)	0 (0.0 %)	2 (1.7 %)	0.50

*Values reported are the numbers (percentages) of physicians who made the query during the visit

†Probability values are based on Fisher's exact test

‡Queries about the cancer history of the SP's mother and sister were not coded because this information was volunteered by the SP as part of the standardized script

Table 3 Physician Counseling About Test Result Implications for the SP, Sorted by Combined Frequency of Occurrence*

Topic	Control (n=61)	Intervention (n=60)	Combined (n=121)	P†
Cancer risks				
Positive result means increased risk of breast cancer	45 (73.8 %)	43 (71.7 %)	88 (72.3 %)	0.84
Positive result means increased risk of ovarian cancer	38 (62.3 %)	32 (53.3 %)	70 (57.9 %)	0.36
Treatment implications				
Positive result introduces option of surgical treatment	57 (93.4 %)	55 (91.7 %)	112 (92.6 %)	0.74
Positive result has fertility implications due to effects of prophylactic treatments (oophorectomy)	13 (21.3 %)	19 (31.7 %)	32 (26.5 %)	0.22
Positive result introduces option of prophylaxis with medications	19 (31.2 %)	11 (18.3 %)	30 (24.8 %)	0.14
Screening implications				
Positive result means more ongoing testing/screening	49 (80.3 %)	52 (86.7 %)	101 (83.5 %)	0.46
Positive result underscores the importance of self-monitoring with breast exams	6 (9.8 %)	5 (8.3 %)	11 (9.1 %)	1.0
Test limitations				
Even with negative result, SP (and/or sister) could still get breast cancer	21 (34.4 %)	17 (28.3 %)	38 (31.4 %)	0.56
Tests only look for known mutations	15 (24.6 %)	17 (28.3 %)	32 (26.5 %)	0.68
Genetic testing is not 100 % accurate	8 (13.1 %)	4 (6.7 %)	12 (9.9 %)	0.36
Emotional issues				
Physician discusses emotional repercussions of testing for SP	27 (44.3 %)	30 (50.0 %)	57 (47.1 %)	0.59
Physician discusses how a negative test result could be beneficial to the SP (e.g., reassurance)	23 (37.7 %)	19 (31.7 %)	42 (34.7 %)	0.57
Physician asks if SP has support system (in relation to BRCA testing)	2 (3.3 %)	3 (5.0 %)	5 (4.1 %)	0.68

*Values reported are the numbers (percentages) of physicians who raised the topic during the visit.

† Probability values are based on Fisher's exact test.

reports that the emotional repercussions of genetic testing for the SP were explored by nearly half of physicians. The potential benefits of receiving a negative test result were discussed by approximately one-third of physicians. Rarely did the physician ask about the SP's support system.

Implications of Testing for the SP's Family. Genetic test results have implications for a patient's relatives. Physician counseling on such implications are reported in Table 4. On average, physicians covered 1.9 (SD=1.6) of the eight topics reported in the table. Only one significant difference between groups was found. Intervention-group physicians were more likely than control physicians to tell the SP that if she tested positive, it would indicate that her male relatives were at greater risk of prostate cancer. With regard to relatives' cancer risks, a majority of physicians discussed the daughter's elevated risk of cancer. Physician counseling on what a positive

test result would mean for the genetic testing of other family members was rarely offered. Physicians rarely mentioned that a positive test result could necessitate breast cancer screening and possibly surgery or prophylaxis with medications for other family members. The emotional impact on the family of a positive test result for the SP was seldom discussed.

Discussions About Genetic Counseling. The frequency of discussions about counseling and testing are reported in Table 5. Most physicians discussed the benefits of genetic counseling, but intervention-group physicians were significantly more likely to do this than control-group physicians. A majority of physicians offered a referral to a genetic counselor. Cost issues related to testing were not typically discussed with the SP, and did not differ by study group. Genetic testing recommendations varied. The most common recommendation was that the SP should delay her testing

Table 4 Physician Counseling About Test Result Implications for the SP's Family Members, Sorted by Combined Frequency of Occurrence*

Topic	Control (n=61)	Intervention (n=60)	Combined (n=121)	P†
Family cancer risks				
Positive result means daughter is at increased risk of cancer	35 (57.4 %)	33 (55.0 %)	68 (56.2 %)	0.86
Positive result means son is at increased risk of cancer	14 (23.0 %)	21 (35.0 %)	35 (28.9 %)	0.16
Positive result means increased risk of prostate cancer for male relatives	1 (1.6 %)	12 (20.0 %)	13 (10.7 %)	.001
Genetic testing				
Positive result means female family members (in addition to the sister) may need genetic testing	28 (49.5 %)	24 (40.0 %)	52 (43.0 %)	0.58
Positive result means male family members may need genetic testing	7 (11.5 %)	6 (10.0 %)	13 (10.7 %)	1.0
Screening and treatment				
Positive result means family members may need additional breast cancer screening	12 (19.7 %)	9 (15.0 %)	21 (17.4 %)	0.63
Positive result means family members may need treatment (including surgery and prophylaxis with medications)	7 (11.5 %)	3 (5.0 %)	10 (8.3 %)	0.32
Emotional issues				
Testing creates potential for emotional repercussions for the family	12 (19.7 %)	9 (15.0 %)	21 (17.4 %)	0.63

* Values reported are the numbers (percentages) of physicians who raised the topic during the visit.

† Probability values are based on Fisher's exact test.

Table 5 Physician Discussion and Recommendations Regarding Genetic Counseling, Genetic Tests, and ELSI, Sorted by Combined Frequency of Occurrence*

Topic	Control (n=61)	Intervention (n=60)	Combined (n=121)	P†
Benefits and referral				
Benefits of genetic counseling	37 (60.7 %)	47 (78.3 %)	84 (69.4 %)	0.05
Offers genetic counseling referral	32 (52.5 %)	34 (56.7 %)	66 (54.6 %)	0.72
Cost discussions				
Insurance coverage for testing	22 (36.1 %)	18 (30.0 %)	40 (33.1 %)	0.56
Cost of office genetic test	16 (26.2 %)	19 (31.7 %)	35 (28.9 %)	0.55
Cost of genetic counseling (including insurance coverage)	3 (4.9 %)	5 (8.3 %)	8 (6.6 %)	0.49
Testing recommendations				
Sister should be tested first before SP makes a testing decision	23 (37.7 %)	33 (55.0 %)	56 (46.3 %)	0.07
SP should “think about it” and then revisit the physician	22 (36.1 %)	16 (26.7 %)	38 (31.4 %)	0.33
SP should make testing decision after genetic counseling visit	13 (21.3 %)	23 (38.3 %)	36 (29.8 %)	0.05
Testing expectations and concerns				
Asks what SP would do if she received a positive BRCA test result	27 (44.3 %)	32 (53.3 %)	59 (48.8 %)	0.37
Asks the SP for her expectations for screening and/or today’s visit	9 (14.8 %)	14 (23.3 %)	23 (19.0 %)	0.26
Asks how SP would feel if she received a positive BRCA test result	4 (6.6 %)	10 (16.7 %)	14 (11.6 %)	0.10
Response to, “what would you do?” ‡				
I would get tested	33 (54.1 %)	20 (33.3 %)	53 (43.8 %)	0.03
It’s up to you. / Only you can decide.	14 (23.0 %)	24 (40.0 %)	38 (31.4 %)	0.052
I don’t know	2 (3.3 %)	5 (8.3 %)	7 (5.8 %)	0.27
I would not get tested	0 (0.0 %)	3 (5.0 %)	3 (2.5 %)	0.12
ELSI for the SP				
Confidentiality of the medical record	31 (50.9 %)	34 (56.7 %)	65 (53.7 %)	0.59
Health insurance discrimination	32 (52.5 %)	32 (53.3 %)	64 (52.9 %)	1.0
Federal legal protections (GINA)	35 (57.4 %)	26 (43.3 %)	61 (50.4 %)	0.15
Life insurance discrimination	30 (49.2 %)	30 (50.0 %)	60 (49.6 %)	1.0
Employment discrimination	11 (18.0 %)	9 (15.0 %)	20 (16.5 %)	0.82
Stigma as a social issue	0 (0.0 %)	0 (0.0 %)	0 (0.0 %)	—
ELSI for family members				
Life insurance discrimination	1 (1.6 %)	1 (1.7 %)	2 (1.7 %)	1.0
Health insurance discrimination	1 (1.6 %)	0 (0.0 %)	1 (0.8 %)	1.0
Confidentiality of the medical record	1 (1.6 %)	0 (0.0 %)	1 (0.8 %)	1.0
Employment discrimination	0 (0.0 %)	0 (0.0 %)	0 (0.0 %)	—
Federal legal protections (GINA)	0 (0.0 %)	0 (0.0 %)	0 (0.0 %)	—
Stigma as a social issue	0 (0.0 %)	0 (0.0 %)	0 (0.0 %)	—

ELSI ethical, legal, and social issues

* Values reported are the numbers (percentages) of physicians who raised the topic or made the specific statement during the visit.

† Probability values are based on Fisher’s exact test.

‡ Numbers for this grouping need not sum to 100 %. A physician could ignore the query or respond in a manner not represented by our codes.

decision until after her sister had been tested. Intervention physicians were significantly more likely than control physicians to advise making a testing decision after counseling. Fewer than one in five physicians asked the SP about her expectations for cancer screening and for the visit. Nearly half of physicians asked the SP what she would *do* if she received a positive test result, but the SP was asked how she would *feel* in such an event in only one-tenth of visits.

As part of the standardized script, physicians were asked if they would get the test “if you were me.” Intervention-group physicians were less likely than control-group physicians to respond that they would be tested, and were more inclined to tell the SP that she needed to decide what was best for her. This difference narrowly failed to reach conventional standards of statistical significance. Physicians rarely said that they did not know what they would do if placed in the SP’s position or that they would not get tested.

Ethical, Legal, and Social Issues. Coded topics pertaining to the relevance of ethical, legal, and social issues for the patient are shown in the sixth section of Table 5; there was no

difference between intervention and control groups. The most frequently discussed ELSI issue was medical record confidentiality, followed by health insurance discrimination issues, Genetic Information Nondiscrimination Act (GINA) legislation, and life insurance discrimination. Employment discrimination was rarely discussed, and the potential for social stigma stemming from a positive test result was not considered in any visit. The last section of the table reports how often these same issues were discussed with regard to the impact of the SP’s test result on other family members. Such discussions were absent in virtually all visits.

DISCUSSION

An increasing number of genetic tests are available that can identify carriers of mutations that elevate cancer risk, such as BRCA.¹⁰ Many patients with questions about their risks seek answers from their PCPs,⁴¹ underscoring the need for clinical genetics education in primary care. This study evaluated the

effectiveness of one such training program—an interactive web-based curriculum delivered as CME. The intervention's impact on practice behaviors was minimal, with a few notable exceptions.

Intervention-group physicians were more likely to inform patients that a test showing a BRCA mutation could indicate an increased risk for prostate cancer among relatives. However, they were less likely than controls to ask about Ashkenazi heritage, even though the link between Ashkenazi heritage and breast cancer has been known and taught for many years. Intervention-group physicians were also more likely than controls to explore the benefits of genetic counseling and to encourage a deferral of the testing decision until after genetic counseling. They were not more likely than control physicians, however, to offer genetic counseling to the SP.

In an era of increasingly complex and numerous genetic tests, PCPs may lack the time and expertise to provide the specialized counseling that a trained genetic counselor can offer. Modifying the online curriculum to facilitate appropriate referrals may help to ensure that patients receive the information they need to make informed choices. When asked what they would do if personally faced with a similar situation, intervention-group physicians were more inclined than control-group physicians to engage in a nuanced discussion around values and goals. This greater appreciation of contingent decision-making and respect for the patient's values may flow from the intervention curriculum's modeling of shared decision-making.

Regardless of the curriculum completed, participants were generally unprepared to provide robust counseling to an SP for whom genetic counseling and genetic testing was reasonable. The average physician in this study asked just 20 % of the family and personal history questions that would be appropriate in this setting, a finding consistent with other studies.^{11,27} Physicians also fell short in their counseling about the implications of test results, focusing heavily on surgical options and neglecting to explore familial implications, emotional impacts, or social support, though each of these is an essential component of the genetic counseling process.⁴²⁻⁴⁵

Furthermore, though considerable attention was devoted to ELSI issues in the web-based and control curriculum materials, only half of the participants talked with the SP about medical record confidentiality, insurance discrimination, and federal legal protections. Even fewer examined employment discrimination concerns, and none explored the important issue of social stigma.⁴³ These ELSI discussions rarely examined the potential for discrimination against family members that could result from a positive BRCA test result or legal protections for the patient or their family.

In light of these findings, clinical genetics training may need to be longer and periodically reinforced. Alternatively, it may be unrealistic to expect PCPs to provide necessary information to patients presenting with concerns about their inherited risk for cancer. Genetic counselors are specialty-trained to have these conversations and typically have more time and resources to discuss these matters than do PCPs.

Though computer-based curricula aimed at educating patients have been shown to be effective at increasing knowledge about genetic testing for inherited breast cancer,^{46,47} educating time-constrained physicians is challenging, and perhaps our goals should shift toward increasing availability of and referrals to genetic counselors.

Limitations

A large number of physicians invited to participate did not respond to the invitation, leading to the possibility of a selection bias wherein motivated and interested physicians were overrepresented. The effect of such a selection bias would be to overstate the true level of provider ability in the domain studied. The SPs were announced, and although physicians had no advanced knowledge of the clinical presentation, they did know that they were being evaluated. They may, therefore, have exhibited a higher level of clinical skills than would have been observed had the SPs been unannounced.⁴⁸ Further, we examined just one genetic condition; findings may not generalize to other conditions. Finally, the study was carried out in only two states.

CONCLUSIONS

Physicians infrequently performed key counseling behaviors surrounding inherited breast cancer, and this was true regardless of whether they had completed the web-based interactive training or read clinical reviews. Ultimately, we may conclude that the promise of the clinical genetics revolution can be realized only by preparing current and future primary care physicians to identify patients at risk for inherited conditions and to refer those patients for appropriate counseling and testing.

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