Is Discussion of Colorectal Cancer Screening Options Associated with Heightened Patient Confusion?

Resa M. Jones1,3, Sally W. Vernon5, and Steven H. Woolf1,2,4

Abstract

Background: Clinical guidelines recommend offering patients options for colorectal cancer (CRC) screening, but the modalities vary by frequency, accuracy, preparations, discomfort, and cost, which may cause confusion and reduce screening rates. We examined whether patients reported confusion about the options and whether confusion was associated with socio-demographic characteristics, number of options discussed, and adherence.

Methods: Patients ages 50 to 75 years who had visited a clinician within 2 years were randomly selected for a cross-sectional study (n = 6,100). A questionnaire mailed in 2007 asked the following: whether a clinician had ever discussed CRC screening options; which of four recommended tests (i.e., fecal occult blood testing, flexible sigmoidoscopy, colonoscopy, and barium enema) were presented; and whether the options were confusing. Analyses were restricted to respondents who reported discussing one or more screening options (n = 1,707). Weighted frequencies were calculated and multivariate logistic regression was done.

Results: The sample was 55.5% female, 15.6% African American, and 83.2% adherent to screening recommendations, and 56.0% had discussed two or more screening options. In adjusted analyses, nonadherent patients reported greater confusion than adherent patients (P < 0.01). Adults who discussed two or more options were 1.6 times more likely to be confused than those who discussed one option [95% confidence interval (CI), 1.08-2.26]. Patients who reported being confused were 1.8 times more likely to be nonadherent to screening than those who did not (95% CI, 1.14-2.75).

Conclusions: Our study provides the first empirical evidence linking multiple options with confusion and confusion with screening adherence.

Impact: Confusion may act as a barrier to screening and should be considered in public health messages and interventions. Cancer Epidemiol Biomarkers Prev; 19(11); 2821–5. ©2010 AACR.

Introduction

Colorectal cancer (CRC) is the second leading cause of cancer death (1). Although efficacious early detection screening methods exist (2), known barriers to screening contribute to whether people undergo recommended CRC screening modalities (e.g., refs. 3-5). Public education efforts have raised awareness about the importance of CRC screening, and several studies have evaluated interventions to increase screening uptake (6, 7). Despite these efforts, screening rates are lower than those for breast and cervical cancer (8).

Expert panels and professional organizations recommend several options for CRC screening [e.g., fecal occult blood testing (FOBT), flexible sigmoidoscopy, and colonoscopy]. They further recommend that clinicians offer and discuss the options with patients and let patients choose the one they prefer (2, 9). This approach was adopted because there was inadequate evidence that any one test offered substantially greater clinical effectiveness and cost-effectiveness over another and because personal preferences of patients vary (10, 11). Some argue that giving patients a choice may enhance uptake (12), whereas others argue that offering too many choices may cause confusion and may partially explain low rates of CRC screening (13). Data from the 2005 Health Information National Trends Survey found that 49% of adults who had not had CRC agreed that the number of options for CRC screening made it difficult to know which one to
choose (16). Lafata et al. (17) found that patients offered multiple screening options were less likely to be screened compared with those offered one choice [odds ratio (OR), 0.57; 95% confidence interval (CI), 0.37-0.86]. Similarly, based on tape-recorded patient-physician encounters, McQueen and colleagues (18) hypothesized that patients’ confusion may contribute to low CRC screening rates.

We conducted a secondary analysis of data collected for a study of patient barriers to CRC screening (5) to explore whether patients who discussed CRC screening with their clinician discussed multiple screening options, which options they discussed, and whether they reported being confused about the options. We also examined whether confusion was associated with socio-demographic characteristics, the number of options discussed, and self-reported adherence.

**Materials and Methods**

**Study population and data collection**

Twelve family medicine practices affiliated with the Virginia Ambulatory Care Outcomes Research Network, a practice-based research network, participated in a study to assess the importance of patient-reported barriers to CRC screening. The sampling frame for the parent study, data collection methods, and the relative importance of screening barriers for each of four CRC screening modalities are reported elsewhere (5).

Briefly, the sampling frame consisted of patients, age 50 to 75 years old, who had an office visit in the last 2 years. Beginning in February 2007, a questionnaire was distributed to 6,100 randomly selected adults using a modified Dillman sequential mailing protocol, with a $2 unconditional incentive enclosed (19, 20). A total of 3,357 patients completed the questionnaire. The subsample selected for this study was the group who reported discussing at least one option for CRC screening with a clinician (n = 1,707). The questionnaire, which is available online (21), included items asking participants (a) whether their healthcare provider ever discussed different choices or options for CRC screening, (b) which of four recommended CRC screening tests (i.e., FOBT, flexible sigmoidoscopy, colonoscopy, and barium enema) they had discussed as options, and (c) whether they would “strongly agree,” “somewhat agree,” “neither agree nor...
disagree,” “somewhat disagree,” or “strongly disagree” with the following statement: “The information about the different colon cancer screening options was confusing.” CRC screening history questions (22) and demographics were also assessed.

The study was approved by the institutional review boards of Virginia Commonwealth University and Riverside Medical Center.

Statistical analysis

All analyses were done using SAS/STAT software (version 9.2; SAS Institute, Inc.). Given the probabilistic sampling scheme, procedures SurveyFreq, SurveyReg, and SurveyLogistic were used to calculate weighted frequencies of the proportion of adults who talked to their clinician about screening options and which of the four modalities they discussed.

Responses to the question about confusion were rated on a five-point scale (strongly agree to strongly disagree), where higher scores reflect greater confusion. Due to a skewed distribution, the response categories “strongly agreed” and “somewhat agreed” were combined and compared with the other three categories.

Based on published guidelines in effect at the time (23), respondents were classified as nonadherent if they did not report a FOBT within the last year, flexible sigmoidoscopy or barium enema within the last 5 years, or colonoscopy within the last 10 years. Finite population adjustments were made to SEs using the sampling rates, and sampling weights were adjusted for nonresponse.

Multivariate logistic regression (OR and 95% CI) was done to determine whether confusion was independently associated with patient characteristics (i.e., gender, age, race, education, annual income, insurance coverage, and number of screening options discussed with the clinician) and with screening adherence.

Results

The sample was 55.5% female, 15.6% were African American, 26.9% were age 65 to 75, 22.8% had no more than a high school education, and 83.2% were adherent to screening recommendations. A total of 56.0% had discussed two or more screening options with their clinician. Of those who discussed options, the majority discussed colonoscopy (94.7%), followed by FOBT (50.3%), flexible sigmoidoscopy (31.8%), and barium enema (14.7%). Overall, 13.0% strongly agreed or somewhat agreed that the information about various screening options was confusing. Adults who discussed two or more screening options (weighted \( n = 7,556 \)) with their clinician were 1.57 times more likely to be confused than those who discussed one option (weighted \( n = 5,930 \); 95% CI, 1.08-2.26). The test for linear trend was statistically nonsignificant (\( P = 0.17 \)). However, whereas the ORs were similar when only one option was presented compared with two (OR, 1.59; 95% CI, 1.02-2.49; weighted \( n = 4,010 \)) or three (OR, 1.24; 95% CI, 0.73-2.11; weighted \( n = 2,313 \)) options, confusion was greater when four options were discussed (OR, 2.33; 95% CI, 1.14-4.77; weighted \( n = 1,233 \)).

Confusion about the options was reported by 17.8% of patients who were nonadherent to CRC screening recommendations and 12.0% of those who were adherent. People who reported being confused were 1.77 times more likely to be nonadherent to screening compared with people who were not confused (95% CI, 1.14-2.75).

As shown in Table 1, women reported greater confusion about screening options than men (OR, 1.53; 95% CI, 1.05-2.22). Respondents with education beyond high school or higher annual household incomes (≥$20,000) were more likely to report confusion than those with no more than a high school education or lower income. However, the uninsured were six times more likely to express confusion than those with insurance coverage (95% CI, 1.96-17.6). Although statistically significant, the CI for that estimate was very wide because few members of the sample population were uninsured.

Discussion

Although the majority of patients did not report being confused about CRC screening options, our findings are consistent with the results of qualitative studies that suggest that presenting multiple options for CRC screening may increase patient confusion (3, 18). In our sample, respondents who reported discussing only one screening option were significantly less likely to report confusion than those who discussed two to four options, as recent guidelines recommend (24). Our findings also are consistent with those of Lafta et al. (17) who reported that offering multiple screening options reduced the likelihood of screening adherence. To our knowledge, however, our study provides the first empirical evidence linking multiple options with confusion and linking confusion with screening adherence.

Our findings are consistent with the view expressed by Schwartz (25) who described a “paradox of choice” in which people desire choices but become overwhelmed and “paralyzed” by the number of options during the decision-making process. This observation supports Neugut and Lebwohl recommendation that clinicians recommend only one test (i.e., colonoscopy) to patients and that the review of a menu of options be eliminated from guidelines (26). Another argument for promoting a single test is that some screening options are not being used or are not widely available (e.g., flexible sigmoidoscopy or double-contrast barium enema). In general, we observed a threshold effect when more than one option was presented. We saw no evidence of a linear dose-response association between confusion and the number of options presented; however, discussing four options resulted in much greater confusion than one option. Our analyses did not take into account which combination of two or three screening modalities were actually discussed. Further, perhaps clinicians discussed screening options without necessarily
giving equal weight to each modality. Thus, although our findings suggest that presenting one screening option may reduce confusion, our data do not indicate which modality should be recommended.

Women were more likely to report confusion than men as were the uninsured, and respondents who reported higher household incomes and more education also were more likely to report being confused. Further research is needed to understand these patterns and to make sense of potentially inconsistent findings (e.g., the uninsured are more likely to have low incomes). Confusion has cognitive and affective components and as such likely encompasses much more than simple knowledge transfer. Psychological and sociological research on the causes and determinants of confusion is important to understand how gender, socioeconomic status, and related covariates (e.g., health literacy) might influence one’s desire and ability to obtain information about options and the risk of becoming confused or overwhelmed by the volume or complexity of information. In the context of a choice presented by a clinician, as in the case of CRC screening, a related question is whether patients’ gender, socioeconomic status, or related covariates influence practitioner behavior in ways that contribute to confusion. In short, although explanations of the complexity of screening need to be improved, it is crucial to better understand the various components of confusion to ultimately aid in patient-physician communication.

The medical literature provides few insights on how screening advice may foment confusion. The most pertinent studies have focused on gaps in information provided during such counseling. In a study of informed decision-making elements covered with men during a primary care visit for CRC screening, Ling et al. found that >50% of visits included no informed decision elements. In <6% of visits was there discussion of uncertainties, assessment of patient understanding, probing about input from trusted people, and discussion of the patient’s role in the decision making (10). McQueen et al. (18) reported that patients’ confusion persisted even after discussion with their physicians because questions and issues remained unaddressed.

Our findings should be considered in the context of the following limitations. First, generalizability may be limited because our sample was drawn from primary care practices. Patients must have had an office visit within the past 2 years to be included, and the sample of uninsured patients was very small. People with more limited access to care may have different experiences with how and from whom they receive information about CRC screening options. Moreover, the practices involved in the study and the geographic areas they represented may differ from other settings. Second, although a slightly larger proportion of nonadherent persons reported confusion than those who were adherent, confusion was reported by only 13.0% of patients in this sample. Further, our sample only included patients who reported discussing options or choices for CRC screening and a large proportion of those patients was screened per recommendations (83.2%), which is much higher than national CRC screening rate (60.8%; ref. 27). Experiences about confusion could have differed in the 49.2% of the parent study sample who did not report discussing CRC screening with their clinician and were not included in this study (only 59.0% of whom were screened per recommendations). Third, this study reports secondary analysis of data originally collected for another purpose—to assess the relative importance of patient-reported barriers to CRC screening (3, 4). Confusion was measured with only one item. To limit respondent burden, we did not include questions to ask about the determinants or potential domains of confusion. Fourth, this study includes no comparison group because respondents were only asked about their confusion if they reported discussing screening options with their clinician. Fifth, cross-sectional data do not permit the assessment of temporal or causality in the association between confusion and CRC screening adherence. Finally, this study was completed when greater consistency existed among CRC screening guidelines. Now, the recommendations of the American Cancer Society–U.S. Multi-Society Task Force differ from those of the U.S. Preventive Services Task Force in that the former recommends more screening options, a situation that could exacerbate confusion for patients.

Given that current guidelines are inconsistent and recommend an even larger number of screening options, such as fecal DNA testing and virtual colonoscopy, confusion likely exists about the recommendations themselves, as well as the different modalities and what they entail (e.g., risks and benefits). One area for future research is to explore the sources and domains of confusion about CRC screening and to develop ways to measure them. Comprehensive measures of confusion about screening guidelines in general and about currently recommended modalities in particular are needed. Given that a cross-sectional association between confusion and screening rates does not prove causality, a prospective randomized trial would be of public health importance to compare screening rates among patients offered one versus multiple tests. Such findings would help determine whether guidelines, and clinicians who counsel patients, should recommend only one test or multiple options to accommodate heterogeneous preferences.

**Disclosure of Potential Conflicts of Interest**

No potential conflicts of interest were disclosed.

**Acknowledgments**

We thank the staff at the Virginia Commonwealth University Department of Family Medicine who were responsible for assembling the questionnaire packets and for data management, the Survey Evaluation and Research Laboratory at Virginia Commonwealth
University for doing survey data entry, Robert E. Johnson, Ph.D. for statistical consultation, and the 12 family medicine practices in the Virginia Ambulatory Care Outcomes Research Network that participated in the study: Nelson Clinic, Richmond, VA; Ambulatory Care Clinic, Richmond, VA; Town Center Family Medicine, Reston, VA; Vienna Family Medicine, Vienna, VA; Prince William Family Medicine, Manassas, VA; Front Royal Family Practice, Front Royal, VA; Riverside Family Medicine, Newport News, VA; Riverside Suburban Family Practice, Newport News, VA; Bruton Avenue Family Practice, Bruton, VA; Mathews Medical Center, Mathews, VA; and Williamsburg Medical Arts-Family Practice, Williamsburg, VA.

References

Is Discussion of Colorectal Cancer Screening Options Associated with Heightened Patient Confusion?

Resa M. Jones, Sally W. Vernon and Steven H. Woolf

Cancer Epidemiol Biomarkers Prev 2010;19:2821-2825. Published OnlineFirst September 17, 2010.

Updated version
Access the most recent version of this article at:
doi:10.1158/1055-9965.EPI-10-0695

Cited articles
This article cites 19 articles, 2 of which you can access for free at:
http://cebp.aacrjournals.org/content/19/11/2821.full.html#ref-list-1

Citing articles
This article has been cited by 4 HighWire-hosted articles. Access the articles at:
/content/19/11/2821.full.html#related-urls

E-mail alerts
Sign up to receive free email-alerts related to this article or journal.

Reprints and Subscriptions
To order reprints of this article or to subscribe to the journal, contact the AACR Publications Department at pubs@aacr.org.

Permissions
To request permission to re-use all or part of this article, contact the AACR Publications Department at permissions@aacr.org.