Study protocol for developing #CuttingCRC: a barbershop-based trial on masculinity barriers to care and colorectal cancer screening uptake among African-American men using an exploratory sequential mixed-methods design

Charles R Rogers,¹ Kola Okuyemi,¹ Electra D Paskett,² Roland J Thorpe,³ Tiana N Rogers,⁴ Man Hung,⁵ Susan Zickmund,⁶ Colin Riley,¹ Michael D Fetters⁷

ABSTRACT

Introduction Colorectal cancer (CRC) is preventable, as screening leads to the identification and removal of precancerous polyps. African-American men consistently have the highest CRC mortality rates, and their CRC-screening uptake remains low for complex reasons. Culture-specific masculinity barriers to care may contribute to the low uptake among African-American men. Examining these barriers to care is vital as CRC screening may challenge cultural role expectations of African-American men, whose tendency is to delay help-seeking medical care. Barbershops provide a pathway for reaching African-American men with masculinity barriers to care who are not regularly receiving healthcare services and CRC screening. This study aims to develop and pilot test a theory-driven, culture-specific, barbershop-based intervention targeting masculinity barriers to care and CRC-screening uptake among African-American men ages 45–75.

Methods and analysis Guided by the theory of planned behaviour and the behaviour change wheel, we will use a multistage mixed-methods study design, beginning with an exploratory sequential approach to validate items for subsequent use in a pilot mixed-methods intervention. First, we will collect and analyse qualitative data from focus groups, cognitive interviews and expert item review to validate and test a culture-specific Masculinity Barriers to Care Scale (MBCS) among African-American men. Next, we will administer the MBCS to our target population as an online quantitative survey and evaluate the association between scores and CRC-screening uptake. Then, we will consider existing evidence-based approaches, our integrated results (qualitative + quantitative), and community input to design a culture-specific Masculinity Barriers to Care Scale (MBCS) among African-American men. Next, we will administer the MBCS to our target population as an online quantitative survey and evaluate the association between scores and CRC-screening uptake. Then, we will consider existing evidence-based approaches, our integrated results (qualitative + quantitative), and community input to design a culture-specific Masculinity Barriers to Care Scale (MBCS) among African-American men. Next, we will administer the MBCS to our target population as an online quantitative survey and evaluate the association between scores and CRC-screening uptake. Then, we will consider existing evidence-based approaches, our integrated results (qualitative + quantitative), and community input to design a culture-specific Masculinity Barriers to Care Scale (MBCS) among African-American men. We will test the peer intervention in a pilot study with a two-arm cluster randomised design (six barbershops, randomised by site) to reduce contamination and account for barbershop culture differences. Our primary outcomes for the pilot are recruitment, sample size estimation, preliminary efficacy and acceptability.

Strengths and limitations of this study

► By drawing on constructs of the theory of planned behaviour and the behaviour change wheel, our study will be among the first to offer a structured approach to designing a behaviour change focused, culture-specific arm for our pilot intervention, while considering a range of psychosocial factors associated with colorectal cancer (CRC) screening among African-American men.

► Our study proposes a new, culture-specific Masculinity Barriers to Care Scale for understanding and reducing CRC-screening disparities among African-American men.

► Given the rising CRC burden among young adults, our study engages African-American men starting at age 45 years.

► Though self-report questionnaires are a common behaviour science methodology, social desirability and non-response bias are potential concerns that we will offset by testing the reliability and validity of the data, while collecting it electronically and securely.

► Additional research will be needed to ascertain the generalisability of the findings to other settings, since this study limits involvement to African-American men from two metropolitan areas in Utah and Minnesota.

© Author(s) (or their employer(s)) 2019. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to Dr Charles R Rogers; charles.rogers@utah.edu
through publications in peer-reviewed journals, community dialogue sessions, and presentations at conferences.

**Trial registration number** ClinicalTrials.gov identifier: NCT03733197 (Pre-results);https://clinicaltrials.gov/ct2/show/NCT03733197

**INTRODUCTION**

Colorectal cancer (CRC) is one of the most treatable and preventable cancers. Despite CRC screening’s (CRCS) life-saving potential, however, nearly 28% of Americans aged 50–75 years have not received timely screening.1 Across all gender and racial/ethnic groups, African-American men have the highest CRC mortality and shortest survival.2 In 2010, national CRCS uptake rates among African Americans (56%) were significantly lower than among non-Hispanic whites (62%).3–5 CRC incidence and mortality rates are 27% and 52% higher, respectively, among African-American men than among non-Hispanic white men.4,6

**Recommendations for CRCS**

The US Preventive Services Task Force endorses a CRCS-screening age range of 50–75 years for average-risk men and screening initiation at age 40 years for those with a family history of CRC.7 Because African-American men are more likely than non-Hispanic white men to be diagnosed at both a younger age and a more advanced disease stage,3 the American College of Gastroenterology has lowered its recommended age of screening initiation to 45 years for African-American men.2,3,8 The proportion of CRC cases diagnosed in individuals aged under 55 years has doubled in the past two decades, and CRC incidence among younger adults (aged 35–49 years), including African-American men, is predicted to increase 28%–46% by 2030.9

**Masculinity and psychosocial factors may contribute to low CRCS**

Masculinity is an important aspect of gendered and cultural identity for men10–12 and plays a critical role in African-American men’s healthcare use, health behaviours and mortality.13–17 Because CRCS challenges some cultural role expectations of African-American men, who tend to delay seeking medical care, examination of masculinity barriers to care is perilous. However, the specific influence of cultural masculinity perceptions on African-American men’s CRCS rates is not well studied. An unacknowledged sense of vulnerability that conflicts with culturally accepted gender norms is also often inherent in men’s experience of CRCS. Previous research suggests that inadequate existing validated measures and biases towards Western culture (norms, values, customs, etc, associated with Europe and European descent) may explain the absence of a significant association between masculinity and CRCS attitudes among African-American men.18–25 In a systematic review of the literature examining connections between masculinity, racism, social support and CRCS uptake among African-American men,7 few studies have examined how masculinity relates to poor CRCS uptake and, of these, none used validated measures. Further, no validated masculinity measures have been developed for African-American men in the context of CRCS uptake or medical care.18–25

Consideration of how psychosocial factors relate to CRCS uptake is also critical. Previous research18–22 with African-American men has documented the influence of factors such as attitudes, knowledge, racism and perceived barriers (eg, embarrassment, fear) on CRCS. Medical mistrust is another widely cited attitudinal barrier to CRCS and treatment seeking21,26–27 and is related to the low health services utilisation among African-American men.21 yet it is unclear whether trust-related barriers are related to CRCS.18–20 Since each of the aforementioned factors represents deeply intricate aspects of the social milieu in which African-American men make health decisions, the first author leads the creation and psychometric evaluation of the reliability and validity of his Male Role Norms, Knowledge, Attitudes, and Perceptions associated with CRCS (MKAP-CRCS) survey.21 On average, our sample of young adult African-American men (ages 19–45) disagreed with traditional masculinity ideology—as measured by the 21-item Male Role Norms Inventory-Short Form (MRNI-SF) scale.32 Our principal component analysis revealed the MKAP-CRCS measure was psychometrically sound, but some participants may have withdrawn prematurely from the MRNI-SF portion as they found the measures’ norms offensive/taboo, or felt awkward sharing beliefs about the roles expected of men. We concluded that research focused on developing a scale explicitly considering African-American men’s masculinity beliefs in the medical care context and with more rigorous psychometric assessments (eg, exploratory factor analysis) is needed.

**Barbershops as a site for interventions to improve CRCS**

Barbershops serving African-American men are favourable settings for reaching our target population.23 Previous multicomponent, barbershop-based trials have been conducted with African-American men on HIV risk reduction, prostate cancer education, heart disease control and hypertension detection.34–36 Few trials of CRCS uptake among African-American men have found significant results. The MISTER B study, the first and to date only barbershop-based CRCS trial, tested a phone-based patient-navigation intervention to urge CRCS among older (mean age 57 years), low-income African-American men with uncontrolled hypertension.37 Intervention completion was associated with a 16-fold increase in the odds of CRCS uptake by 6 months; however, although nearly 70% of participants voiced the intent to obtain colonoscopy screening in the next 6 months, only 17% in the intervention groups and 8% in the control group did so. Our study will help fill this gap between uptake and intention by creating a new, culture-specific intervention that directly addresses masculinity barriers to care, psychosocial factors and CRCS uptake among African-American men beginning at age 45, then test its
feasibility and acceptability in a cluster randomised pilot intervention (at the barbershop level).

**Study objectives**

Disparities associated with CRCS uptake for African-American men, the failure of previous interventions to significantly increase screening rates, and the novel idea of using the barbershop as an intervention setting led to the current study, with the following objectives: (1) validate and test a culture-specific Masculinity Barriers To Care Scale (MBCS) relative to psychosocial factors and CRCS uptake among African-American men and (2) develop and pilot test a theory-driven, culture-specific peer intervention that targets masculinity barriers to care, psychosocial factors and uptake of CRCS (specifically, of the fecal immunochemical test (FIT)) among African-American men. Culture-specific refers to the embodiment of ‘an (African-American male’s) real-life experiences within a given cultural context (eg, neighbourhood) and his understanding of those experiences.’

**METHODS AND ANALYSIS**

**Overall study design**

The Standard Protocol Items: Recommendations for Interventional Trials checklist was used while developing this manuscript. For the proposed study, a multistage mixed-methods design that is shown in figure 1 will be employed. We will begin with an exploratory sequential approach intended to validate items for subsequent use in a pilot mixed-methods intervention. For Objectives 1A and 1B (years 1–2), we will collect and analyze qualitative data from focus groups, cognitive interviews and expert item review to validate and test a culture-specific MBCS among African-American men. Questions for the MBCS will stem from modifications to the (1) Barriers to Help-Seeking Scale developed by Mansfield et al, (2) the Group-Based Medical Mistrust Scale developed by Thompson et al, (3) Mincey et al Masculinity Inventory Scale, (4) the MRNI-SF by Levant et al, Bowleg et al Black Men’s Experiences Scale, and the Masculinity Salience scale developed by Hammond et al. Six factors are expected for the underlying structure of the 21 items in the MBCS: (1) need for control and self-reliance, (2) minimising health problems and resignation, (3) medical mistrust, (4) privacy, (5) emotional control and (6) black masculinity. For all factors, individual items will be assessed on a Likert-type scale. Higher scores will indicate a greater degree of endorsement of masculinity barriers to care. Next, we will administer the MBCS as an online quantitative survey of our target population to evaluate the association between scale scores and CRCS uptake.

For objective 2 (years 3–5), we will consider existing evidence-based approaches (eg, motivational interviewing, MI), our integrated results (qualitative + quantitative) from objectives 1A and 1B regarding masculinity barriers to care, and community input to design a novel, culture-specific, behavioural intervention that is (1) aimed at increasing CRCS uptake (via FIT) among African-American men and (2) feasible for delivery in barbershops. To reduce contamination and account for differences in barbershop culture, we will pilot test the peer intervention in a two-arm cluster randomised intervention (six barbershops, with participants randomised by site). Our primary outcomes for the pilot are recruitment, sample size estimation, preliminary efficacy and acceptability. We will also conduct postintervention interviews with participants from both arms to evaluate acceptability (ie, why and how each arm was or was not successful). To ensure confidentiality, data dispersed to project team members will be blinded of any identifying participant information.

**Patient and public involvement**

Neither patients nor the public were involved in the design of the study.

**Theoretical foundation**

A conceptual framework integrating constructs of the theory of planned behaviour (TPB) will guide our work. The TPB posits that behaviour is a function of intention, which is influenced by attitudes and beliefs.
illustrates how masculinity barriers to care and other psychosocial factors may influence CRCS intention and uptake among African-American men. We will also assess demographic characteristics (eg, age, marital status and health insurance status) that are known to influence African-American men’s masculinity, and CRCS perceptions and behaviours.

Evidence-based cultural grounding to facilitate understanding of African-American men’s culture and engage community stakeholders as trial-development partners is the best achieved by an iterative, participatory and reflexive research process.44 45 Hence, we will use the behaviour change wheel (BCW) as the conceptual framework driving our study’s intervention-development phase. Developed from 19 behavioural change frameworks, the BCW offers a structured approach to inclusively analysing available intervention options and designing behavioural change interventions.46

Setting
We will conduct this research in the Salt Lake City, Utah and Minneapolis–St. Paul (Twin Cities), Minnesota, metro areas, regions notable for having the largest populations of African Americans in their respective states.47 48 Moreover, in both states, CRCS rates among African Americans are well below statewide averages (53% vs 72% for all ethnic and racial groups combined in Utah; 57% vs 73% for both non-Hispanic whites and all ethnic and racial groups combined in Minnesota).32-44 Nationally, African-American men exhibit a lower screening likelihood than African-American women.35 49-51

Focus groups
Participants and procedures
To inform MBCS development, we will conduct twelve 2-hour focus groups (a sufficient number to reach saturation),32 33 each involving eight men who (1) self-identify as non-Hispanic black/African American; (2) were born in the USA; (3) are aged 45–75 years; (4) have a working telephone; (5) speak English and (6) reside in the Salt Lake City or Twin Cities metro area. Six focus groups will be conducted in each metro area. Because participants may be more comfortable with other African-American men of similar age who either have or have not completed CRCS, each group will be clustered by age and CRCS status (table 1). Men aged 45–49 years will be included because African-American men are diagnosed with CRC at both an earlier age and a more advanced disease stage.3 7

We will use culture-specific marketing materials to promote the study through existing social networks, including newspaper advertisements, social media, predominantly African-American churches, air time on two radio stations (one in Minneapolis, one in Salt Lake City) with a predominantly African-American male audience and African-American male serving barbershops. The principal investigator (PI), CRR, has a record of success in recruiting African-American men using these strategies.19 21 22

Potential participants will be encouraged to visit www.cuttingCRC.com to express interest in focus group participation. Basic demographic information will be collected (and kept confidential) to enable research team members to contact participants by phone to confirm eligibility and discuss participation arrangements. Food and drink will be provided during each session. Each participant will receive a US$20 target gift card and participants may choose to be entered into a random drawing to win one of three incentives: (1) a US$100 target gift card, (2) two tickets to a Utah Jazz or Minnesota Timberwolves

<table>
<thead>
<tr>
<th>Groups</th>
<th>Age range</th>
<th>Colorectal cancer screening status</th>
</tr>
</thead>
<tbody>
<tr>
<td>1–2</td>
<td>45–49</td>
<td>Never completed</td>
</tr>
<tr>
<td>3–4</td>
<td>50–65</td>
<td>Not current</td>
</tr>
<tr>
<td>5–6</td>
<td>66–75</td>
<td>Not current</td>
</tr>
<tr>
<td>7–8</td>
<td>45–49</td>
<td>Completed/current</td>
</tr>
<tr>
<td>9–10</td>
<td>50–65</td>
<td>Completed/current</td>
</tr>
<tr>
<td>10–12</td>
<td>66–75</td>
<td>Completed/current</td>
</tr>
</tbody>
</table>
basketball game in 2019-20 season (regardless of their home state), or (3) a Samsung 55’ 4K UHDTV.

**Data collection and analyses**
CRR will facilitate the focus groups, using an interview guide stemming from modifications to existing measures that examine masculinity as well as attitudes and practices precluding men from seeking healthcare access. Another team member will assist with consenting and note taking. The 2-hour sessions will be audio-recorded with two voice recorders, transcribed, and checked for accuracy. De-identified transcripts will be imported into NVivo V.11 software (QSR International, Melbourne, Australia). Our NVivo-proficient coding team (TNR, CRR, and the research assistant) will use constant comparative and content analysis methods to independently code transcripts for themes. After identifying themes relevant to our research questions from a sample reading and initial coding, we will automate term searches, code all documents and run reports to ascertain the code text related to study themes. NVivo can organise data by participant characteristics, allowing us to compare the responses of participants who have or have not undergone CRCS. To interpret and discuss findings and develop a codebook depicting how our codes inter-relate, we will track coding decisions in NVivo and adjudicate them at team meetings. MDF and/or SZ will re-code transcripts, as needed. Key themes will be identified and incorporated into the new MBCS.

**Cognitive interviews**

**Participants and procedures**
We will pilot test the MBCS with 10 CRC advocates and survivors from across the USA (men and women who speak English, have a working telephone and are aged 18–75 years), using 1-hour cognitive interviews (conducted in person or by phone) to elicit input as participants respond to the survey in real time. Interviews will probe (1) how participants understand each question and response option; (2) whether the questions are likely to elicit an honest response; (3) the clarity of question wording; (4) the user-friendliness of the online survey set-up and (5) the questions’ cultural specificity. Participants will engage in a thinking-aloud process with follow-up probes such as ‘How did you arrive at that answer?’ These approaches will improve feasibility, reduce response error and enhance face validity (an estimate of the degree to which the scale is clearly tapping the desired construct we aim to assess, that is, culture-specific masculinity barriers to care). Cognitive interviews also allow us to assess participants’ comfort with online survey completion via PsychData (PsychData, State College, Pennsylvania, USA), a secure, web-based application that supports data capture for research objective 1B. Our MBCS will be modified as a result of the cognitive interviews when necessary. Interviewees will receive a US$20 Amazon gift card. RJT and two additional leaders in African-American men’s health will provide expert item review of the final MBCS using a questionnaire appraisal system.

**Online survey**

**Participants and procedures**
During year 2, we will recruit 400 African-American men to complete an online survey, administered via smartphone, to test the relationship between masculinity barriers to care and CRCS. Eligible respondents are men who (1) self-describe as non-Hispanic black/African American; (2) were born in the USA; (3) are aged 45–75 years; (4) reside in the Salt Lake City or Twin Cities metro area; (5) have a telephone with internet access and (6) speak English. With the aid of barbers and culture-specific marketing materials, we will recruit participants from African-American male serving barbershops. Survey participants will have the opportunity to participate in drawings for one of five incentives: (1) a US$50 visa gift card, (2) a US$75 gas station gift card, (3) a US$100 grocery store gift card, (4) an Apple iPad or (5) Samsung 43’ 4K UHDTV.

Barbershops are cultural hubs of trust essential in the growth and development of African-American men. Men usually spend at least 30 min waiting for or getting a haircut or chatting with others in the barbershop. Using PsychData, participants will be able to complete our survey within 15 min on their smartphones while waiting for or getting a haircut in participating barbershops. PsychData prevents survey alterations and eliminates transcription errors. The PI has a successful record of recruiting African-American men to complete surveys using mobile technology, and African Americans outpace all groups for smartphone use. For men who want to complete the survey but do not own a smartphone, each participating barbershop will be provided one smartphone courtesy of the study.

**Dependent variables**
We will use two Behavioral Risk Factor Surveillance System (BRFSS) questions to assess CRCS uptake: (1) ‘A blood stool test is a test that may use a special kit at home to determine whether the stool contains blood. Have you ever had this test using a home kit?’ and (2) ‘Sigmoidoscopy and colonoscopy are examinations in which a tube is inserted in the rectum to view the colon for signs of cancer or other health problems. Have you ever had either of these examinations?’

**Independent variables**
In accordance with our conceptual model illustrated in figure 2 and the PI’s MKAP-CRCS tool, our independent variables will be masculinity barriers to care from our new scale and five factors known to influence CRCS uptake among African-American men: knowledge, social support, beliefs and attitudes towards CRCS and two CRCS exams (FIT, colonoscopy).

**Demographic covariates**
Age, educational level, marital status, employment status and other covariates will be included as previous studies
by the PI and others have found these factors to be related to CRCS among African-American men.2 18–22 67 68

Sample size and power considerations
With a sample of 400 African-American men, we will have 80% power at the 0.05 level to detect a masculinity barriers to care effect on the odds of having had CRCS, assuming 35% of men have a masculinity barriers to care index equal to the mean have had CRCS compared with 25% of men with a masculinity barriers to care index 1 SD above the mean. We estimate that the average screening rate will be 35%, as the screening rate for African Americans is 53.1% in Utah and 52% in Minnesota and African-American men in both states tend to have lower CRCS rates than women.4 6 12 45 This assumes a moderately strong relationship between the masculinity barriers to care index and confounders (ie, R²=0.25 for the linear model that regresses the masculinity barriers to care index on the confounders). If the relationship between the confounders and the masculinity index is weaker, the power will be higher: 82% power with R²=0.2% and 87% power with R²=0.1. Power calculations were performed using PASS V.15.

Data collection and analyses
We will test for associations between masculinity barriers to care, psychosocial factors and CRCS uptake. Our central hypothesis is that masculinity barriers to care will be negatively associated with CRCS uptake. The masculinity barriers to care items emerging from objective 1 will be used to create a latent variable that represents the construct being measured. The CRCS uptake outcome will be a binary variable that indicates whether a participant self-reported CRCS uptake (ie, answered yes to either BRFSS dependent-variable question). We will fit a structural equation model with CRCS uptake as the outcome and masculinity barriers to care as the predictor. We will adjust for potential confounders (eg, age, educational level). We will also present the estimate and 95% CI for the OR comparing the odds of CRCS uptake between participants with a one-point difference in masculinity barriers to care scores. Descriptive statistics will summarise the participants’ characteristics.

Two-arm intervention
Integration
In an exploratory sequential designed study, a key step is to apply the qualitative data captured in objective 1A to assist with building objective 1B’s quantitative phase. As described by Fetters et al,69 we will ‘merge’ qualitative and quantitative data from objective 1 to identify content areas for contrasting, comparing and synthesising results. During the first 6 months of year 3, two team members (MDF and CRR) will determine to what degree and how the results from the combined qualitative and quantitative datasets yield a richer and more comprehensive understanding of the impact of masculinity barriers to care on CRCS uptake among African-American men. Through this process, we will apply what we learn about the role of our variable of interest on CRCS to develop a pilot intervention for overcoming these barriers.

Development
During the first 6 months of year 3, we will adopt the BCW approach, working with community advisory board members (2 hours small-group discussions via conference call and/or in person, three members per meeting) to develop the culture-specific intervention arm of our pilot intervention. We will use information from (1) our integrated objective 1e results, (2) existing CRCS intervention evidence and (3) study team expertise to apply the Acceptability, Practicability, Effectiveness/cost-effectiveness, Affordability, Safety/side effects, Equity (APEASE criteria) (table 2). Our hypothesis is that CRCS uptake will be higher in the culture-specific arm than in the control arm.

We anticipate that the culture-specific arm will include at least two core components: barbers as motivational interviewers and InSure FIT kits distributed by barbers. MI is ‘a collaborative conversation style for strengthening a person’s own motivation and commitment to change.’69 Telephone-based MI is an effective way to improve cancer screening among under-represented groups. Community-member–led MI has proved successful, but it is unknown if barbers as motivational interviewers can assist with reducing CRCS inequalities among African-American men.70–73 Also, randomised trials have shown that the

<table>
<thead>
<tr>
<th>Table 2</th>
<th>Behaviour change wheel (BCW) activities to drive development of culture-specific trial arm</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Behavioural diagnosis</td>
<td>Use the BCW to determine what needs to change for colorectal cancer (CRC) screening uptake to increase among African-American men.</td>
</tr>
<tr>
<td>2. Intervention strategy selection</td>
<td>Use1 to decide which intervention functions to apply (eg, education, persuasion, enablement).</td>
</tr>
<tr>
<td>3. Behaviour change technique Identification</td>
<td>Develop a detailed culture-specific arm plan by selecting from among a range of specific, evidence-based behaviour change techniques (eg, intervention components such as barbers as motivational interviewers plus barbers distributing fecal immunochemical test kits; info about health consequences related to negating CRC screening).</td>
</tr>
<tr>
<td>4. Draft full intervention specifications</td>
<td>Create the detailed intervention specifications covering all aspects of content and delivery of the intervention structured around.</td>
</tr>
</tbody>
</table>

FIT is the first-choice fecal occult blood test for CRCS, is less invasive, less costly and may be better accepted than other CRCS tests.74 If we choose this route for the culture-specific arm, preliminary data from our barbers suggest that the PI may teach the barbers the MI technique using content stems from objective 1 findings. Additional components for this arm may be developed during the APEASE process.

Based on the PI’s research and evidence-based strategies,72 the control arm will include an informational CRCS brochure developed by the American Cancer Society76 plus an FIT kit distributed by the barbers. Since the FIT kits will be free and the study will cover postage and processing fees, participants will be able to complete screening regardless of whether they have health insurance. Participants will mail the completed FIT kits to our local laboratory for processing. We will refer participants with positive FIT results to Huntsman Cancer Institute for a colonoscopy.

Participants and procedures

Intervention participants will be non-Hispanic black/African-American men (n=60) who (1) have never completed CRCS; (2) are aged 45–75 years; (3) were born in the USA; (4) reside in the Salt Lake City metro area; (5) have a telephone with internet access and (6) speak English. As a feasibility intervention, sample size is not based on the power to detect a certain effect size.77 Rather, n=60 (30 per arm) was chosen based on practical considerations (eg, cost, recruitment).

The intervention will comprise a recruitment phase and an implementation phase. The recruitment phase will occur during the last 6 months of year 3. With the assistance of barbers and culture-specific marketing materials, we will enrol 10 eligible African-American men at each of 6 barbershops. At baseline, participants will complete the demographic portion of our online survey. Once total enrolment is reached and consent obtained, the six barbershops will be randomised to the culture-specific or control arm using a permuted block size of 6. Then the implementation phase will begin. These distinct phases provide advantages in a cluster randomised design. First, we eliminate recruitment bias as we blind participants to the intervention at enrolment.77 Second, the distinct phases allow each man to be exposed to the intervention for the same amount of time. We foresee the recruitment phase lasting 3 months and the implementation phase 7 months, resulting in a 10-month intervention, allowing ample time for participants to obtain CRCS.

During the last 6 months of year 4, our coding team will conduct exit interviews. Prior literature documents that 6–12 individual interviews per homogeneous group are sufficient to reach data saturation.7879 Thus, 18 in-depth, 60 min participant interviews (2 participants from each of the 6 barbershops plus 3 barbers from each arm) will permit us to obtain rigorous outcomes data as well as participant accounts of what worked well and what did not for our two-arm intervention’s implementation.

Analyses

Descriptive statistics will summarise participants’ baseline characteristics. Continuous variables will be summarised by mean (SD) or median (IQR), and categorical variables in contingency tables. Feasibility of the study protocol will be evaluated as follows:

Recruitment

We will calculate the number of days needed to reach full enrolment at each barbershop, the percentage of men meeting eligibility criteria and, of those, the percentage who chose to enrol.

Sample size estimation

The intraclass correlation coefficient will be estimated from our study data and inflated (due to the expected downward bias) to estimate the necessary sample size for the full trial.80

Preliminary efficacy

By treatment arm, we will calculate the percentage for whom we can ascertain FIT uptake. We will assess intervention adherence 7 months after the recruitment phase by FIT kits returned to our laboratory for processing. Because we will have only three barbershops per arm, no formal statistical analysis of FIT uptake will be performed. Instead, percentages for these outcomes will be calculated by barbershop. We will perform this as intention to treat, with men included based in the arm to which their shop was randomised. A per-protocol analysis will also be performed. Many more barbershops are needed to accurately account for the correlation of African-American men within the same barbershop and achieve a cluster-level confounding balance.81 In our pilot trial, a logistic mixed-effects model with a random intercept for each barbershop will be used to estimate the ORs comparing CRCS uptake between our control and culture-specific arms.

Acceptability

After the 7-month intervention phase, we will conduct 18 postintervention interviews to obtain rigorous outcomes data. The audio-recorded and transcribed postintervention interviews will be analysed by our coding team using NVivo64 and Creswell’s methods.82 To increase our findings’ internal validity, data will be triangulated or compared from the perspectives of the two study arms.83

CONCLUSION

African-American men have the highest CRC mortality across all gender and racial/ethnic groups. Moreover, national findings predict a 28%–46% increase in CRC incidence among adults ages 35–49 years, including African-American men, by 2030.83 Our study aims to aid in reducing CRCS inequities among African-American men by creating a new, culture-specific intervention that directly addresses masculinity barriers to care, psychosocial factors, and CRCS uptake among African-American
men beginning at age 45 years. Subsequently, we will test its feasibility and acceptability in a cluster randomised pilot intervention.

Completing our objective will provide the preliminary data needed for an R01 application to test the new intervention’s efficacy in a large-scale, well-powered, cluster randomised controlled trial. More broadly, this research will demonstrate that decisions regarding CRCS uptake are not detached from cultural and other influences. We will use the culture-specific survey instrument we create to more rigorously assess the association between masculinity barriers to care and CRCS uptake. This will strengthen our scale’s predictive utility while endorsing optimal health for African-American men as warranted by Healthy People 2020. Additionally, our scale could be adapted for use in research on other types of cancer (eg, prostate cancer) that disproportionately affect African-American and other under-represented men.

Overall, our efforts will serve as a model for more culture-specific tailored approaches to prevention, diagnosis and treatment, a goal aligned with the National Cancer Institute’s Cancer Moonshot initiative.

**Author affiliations**

1. Family & Preventive Medicine, University of Utah School of Medicine, Salt Lake City, Utah, USA
2. Internal Medicine, College of Medicine, Ohio State University, Columbus, Ohio, USA
3. Program for Research on Men’s Health, Johns Hopkins University Bloomberg School of Public Health, Baltimore, Maryland, USA
4. Sorenson Impact Center, University of Utah Eccles School of Business, Salt Lake City, Utah, USA
5. College of Dental Medicine, Roseman University of Health Sciences, South Jordan, Utah, USA
6. Internal Medicine, University of Utah, Salt Lake City, Utah, USA
7. Department of Family Medicine, University of Michigan Medical School, Ann Arbor, Michigan, USA

**Acknowledgements**
The research team extends gratitude to Eleanor Mayfield for editorial assistance.

**Contributors**
KO, EDP, MH, SZ, RJT and MDF serve as data monitoring committee (DMC) members for this study, while additional data monitoring, harms and auditing logistics are available from the University of Utah Institutional Review Board. Only the DMC will have access to the full pilot trial dataset in order to ensure that the overall results are not disclosed by an individual study site prior to the main publication. CRR is the study PI, wrote the first draft of the study protocol and edited every draft thereafter. KO is a primary mentor for the study and edited the study protocol with the PI. EDP is a secondary mentor of the study and edited the study protocol with the PI. Study commentors MH, SZ and RJT edited the study protocol with the PI. CRR and TNR are co-investigators of the study and edited the study protocol with the PI. In addition to being a mixed-methods expert, MDF is a co-mentor on the study who edited the study protocol extensively with the PI as a co-mentor. All authors agreed to be accountable for all aspects of the study in ensuring that questions related to the accuracy or integrity of any part of the study are appropriately investigated and resolved. Lastly, all authors read and approved the final manuscript.

**Funding**
This work was supported by the National Cancer Institute of the National Institutes of Health (NIH) under Award Number K01CA234319. RJT was supported by the National Institute on Aging (K02AG059140) and the National Institute on Minority Health and Health Disparities (U54MD000214-6867).

**Disclaimer**
The content is solely the responsibility of the authors and does not necessarily represent the official views of the NIH.

**Competing interests**
None declared.

**Patient consent for publication**
Not required.

**Ethics approval**
This study protocol has received ethics approval from the University of Utah Institutional Review Board (00113679), who will also be responsible for receiving communication updates regarding important protocol modifications.

**Provenance and peer review**
Not commissioned; externally peer reviewed.

**Open access**
This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/.

**REFERENCES**


