

BMJ Open Quality Faecal immunochemical testing implementation to increase colorectal cancer screening in primary care

Smita Bakhai,¹ Gaurav Ahluwalia,¹ Naren Nallapeta,¹ Amanpreet Mangat,¹ Jessica L Reynolds²

To cite: Bakhai S, Ahluwalia G, Nallapeta N, *et al*. Faecal immunochemical testing implementation to increase colorectal cancer screening in primary care. *BMJ Open Quality* 2018;7:e000400. doi:10.1136/bmjopen-2018-000400

Received 6 April 2018

Revised 14 August 2018

Accepted 22 September 2018

ABSTRACT

Colorectal cancer (CRC) is the second leading cause of cancer death in USA, and CRC screening remains suboptimal. The aim of this quality improvement was to increase CRC screening in the internal medicine clinic (IMC) patients, between the ages of 50–75 years, from a baseline rate of 50%–70% over 12 months with the introduction of faecal immunochemical test (FIT) testing. We used the Plan–Do–Study–Act (PDSA) method and performed a root cause analysis to identify barriers to acceptance of CRC screening. The quality improvement team created a driver diagram to identify and prioritise change ideas. We developed a process flow map to optimise opportunities to improve CRC screening. We performed eight PDSA cycles. The major components of interventions included: (1) leveraging health information technology; (2) optimising team work, (3) education to patient, physicians and IMC staff, (4) use of patient navigator for tracking FIT completion and (5) interactive workshops for the staff and physicians to learn motivational interview techniques. The outcome measure included CRC screening rates with either FIT or colonoscopy. The process measures included FIT order and completion rates. Data were analysed using a statistical process control and run charts. Four hundred and seven patients visiting the IMC were offered FIT, and 252 (62%) completed the test. Twenty-two (8.7%) of patients were FIT positive, 14 of those (63.6%) underwent a subsequent diagnostic colonoscopy. We achieved 75% CRC screening with FIT or colonoscopy within 12 months and exceeded our goal. Successful strategies included engaging the leadership, the front-line staff and a highly effective multidisciplinary team. For average-risk patients, FIT was the preferred method of screening. We were able to sustain a CRC screening rate of 75% during the 6-month postproject period. Sustainable annual FIT is required for successful CRC screening.

INTRODUCTION

An estimated 137 000 new cases of colorectal cancer (CRC) and 50 260 related deaths occurred in 2017.¹ CRC screening significantly reduces the incidence and mortality of this disease^{2–5}; however, it remains suboptimal, particularly among the underserved population.^{6–8} In the academic, safety-net Internal Medicine Clinic (IMC) at Erie County Medical Center (ECMC), less than 50% of active, eligible patients were screened for CRC by December of 2016. In March 2014, the American Cancer

Society (ACS), the Centers for Disease Control and Prevention and the National Colorectal Cancer Roundtable (NCCRT) proposed The ‘80% by 2018’ initiative with a goal of implementing CRC screening for 80% of adults between the ages of 50 years and 75 years by 2018.^{9 10} The ECMC leadership pledged 80% CRC screening by 2018; therefore, we designed this quality improvement (QI) to increase the CRC screening in the IMC population.

Preliminary studies suggest that biennial screening instead of annual screening may be effective.^{11 12} However, at this time, the US Preventive Service Task Force screening recommendations include colonoscopy (every 10 years) or home-based faecal testing (every year) for average-risk adults.^{13–17} Many individuals are asymptomatic early in the disease course due to the slow growth of precancerous polyps to invasive cancer. Screening allows for the opportunity for early detection, removal of precancerous polyps and prevention of CRC.^{3 4 18 19} The faecal immunochemical test (FIT) is a less expensive, non-invasive alternative to colonoscopy,^{20 21} which uses antibodies specific for human haemoglobin to reveal haemoglobin in faecal occult blood.^{22–25} Currently, FIT is the most commonly used method for CRC screening in average-risk patients^{26 27}; greater adherence to this test is due to fewer stool samplings and lack of dietary or medication restrictions.^{22 24 28 29} The aim of this QI was to increase CRC screening in the IMC from the baseline rate of 50%–70% in patients between the ages of 50 years and 75 years over 12 months with the introduction of FIT.

METHODS

Setting

We conducted a QI project in an academic IMC, located within a tertiary care safety-net hospital, ECMC. The IMC patient population consists of mostly urban, underprivileged and African-Americans (68.42%). Patients use the



© Author(s) (or their employer(s)) 2018. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

¹Department of Internal Medicine, University at Buffalo, The State University of New York, Buffalo, New York, USA

²Department of Medicine, University at Buffalo, The State University of New York, Buffalo, New York, USA

Correspondence to

Dr Smita Bakhai;
sybakhai@buffalo.edu

IMC as a longitudinal primary care clinic; the IMC has an average of 700 monthly visits. The IMC is composed of a multidisciplinary care team including 35 residents from the University at Buffalo's Internal Medicine Residency Program and four attending physicians.

Design

We designed this QI based on the Plan–Do–Study–Act (PDSA) model of healthcare improvement.^{30–31} The QI team included a physician champion, nursing and ancillary staff, residents, attending physicians, a social worker, gastroenterology (GI) nurse practitioner (NP), a patient navigator, patients, administrative and IT staff and a project liaison from the ACS. The QI team performed a root cause analysis using a fishbone diagram and identified the materials/methods, physician and patient-related barriers to the acceptance of FIT and colonoscopy (figure 1). The QI team identified primary and secondary drivers, brainstormed about potential change ideas and created a driver diagram in order to accomplish our aim^{32–33} (figure 2). We identified strengths and prioritised change ideas to overcome the challenges to improve screening rates (table 1). We developed a new process flow map to optimise opportunity to improve CRC screening (figure 3).

Inclusion criteria consisted of asymptomatic male and female patients, between the ages of 50 years and 75 years for CRC screening with (A) no prior colonoscopy, (B) had a colonoscopy greater than 10 years ago and due for a rescreen and (C) had a colonoscopy less than 10 years ago with a diagnosis of precancerous polyp; therefore, patient was due for a rescreen. Patients were excluded if they had a colonoscopy with normal findings within 10 years or had a negative FIT within 1 year. In December 2016, the ECMC laboratory implemented a hema-screen FIT. The sensitivity cut-off is 50 µg Hb/g of faeces.³⁴ The sensitivity, specificity, positive likelihood ratio and

negative likelihood ratio of a single FIT for cancer has been shown to 0.79, 0.94, 13.10 and 0.23 µg Hb/g of faeces, respectively.³⁵ Physicians discussed the FIT-positive results with the patients and referred the patients for subsequent diagnostic colonoscopy. The physician scheduled a clinic visit to evaluate the barriers to acceptance of colonoscopy after patients missed scheduled colonoscopy appointment. Our CRC screening rates for FIT or colonoscopy were based on the review of completed test reports by the physician.

Measurements

We created an electronic patient database in collaboration with the information technology (IT) department. The baseline CRC screening rate was 50% in December 2016, obtained by retrospective review of the database of active, eligible patients seen at least once within the past 18 months in IMC. The outcome measures included: (1) increase in CRC screening rates to 70% with either FIT or colonoscopy from the baseline rate and (2) diagnostic colonoscopy completion rate in FIT-positive patients. The process measures included FIT order and completion rates. The balance measures were an increase in patient wait time during the IMC visit and poor access for timely screening colonoscopy. The majority of average risk patients refused colonoscopy and preferred the FIT. Therefore, we selected FIT order and completion rates as process measures. CRC screening by FIT is only effective in identifying early CRC or preventing CRC by timely diagnostic colonoscopy in FIT-positive patients.^{17–19} Therefore, we included subsequent colonoscopy in positive FIT patients for our outcome measures. The QI team identified multiple challenges for colonoscopy including lack of patient's preference and a long wait time due to limited capacity from increased demand for CRC screening for high-risk patients.

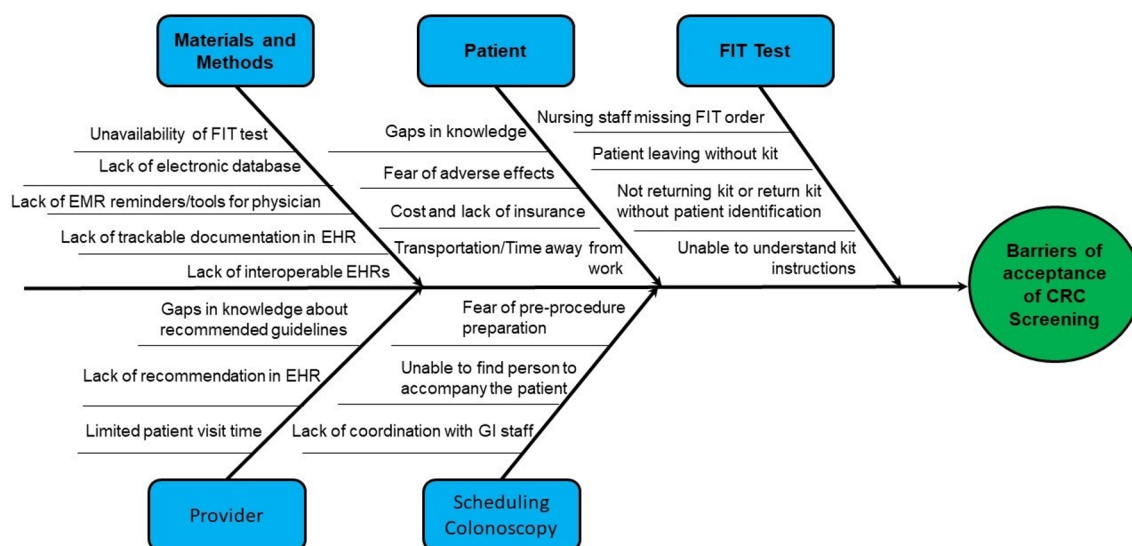


Figure 1 Fishbone diaphragm: root cause analysis identifying barriers to acceptance of CRC screening. CRC, colorectal cancer; FIT, faecal immunochemical test; EHR, electronic health record; EMR, electronic medical record.

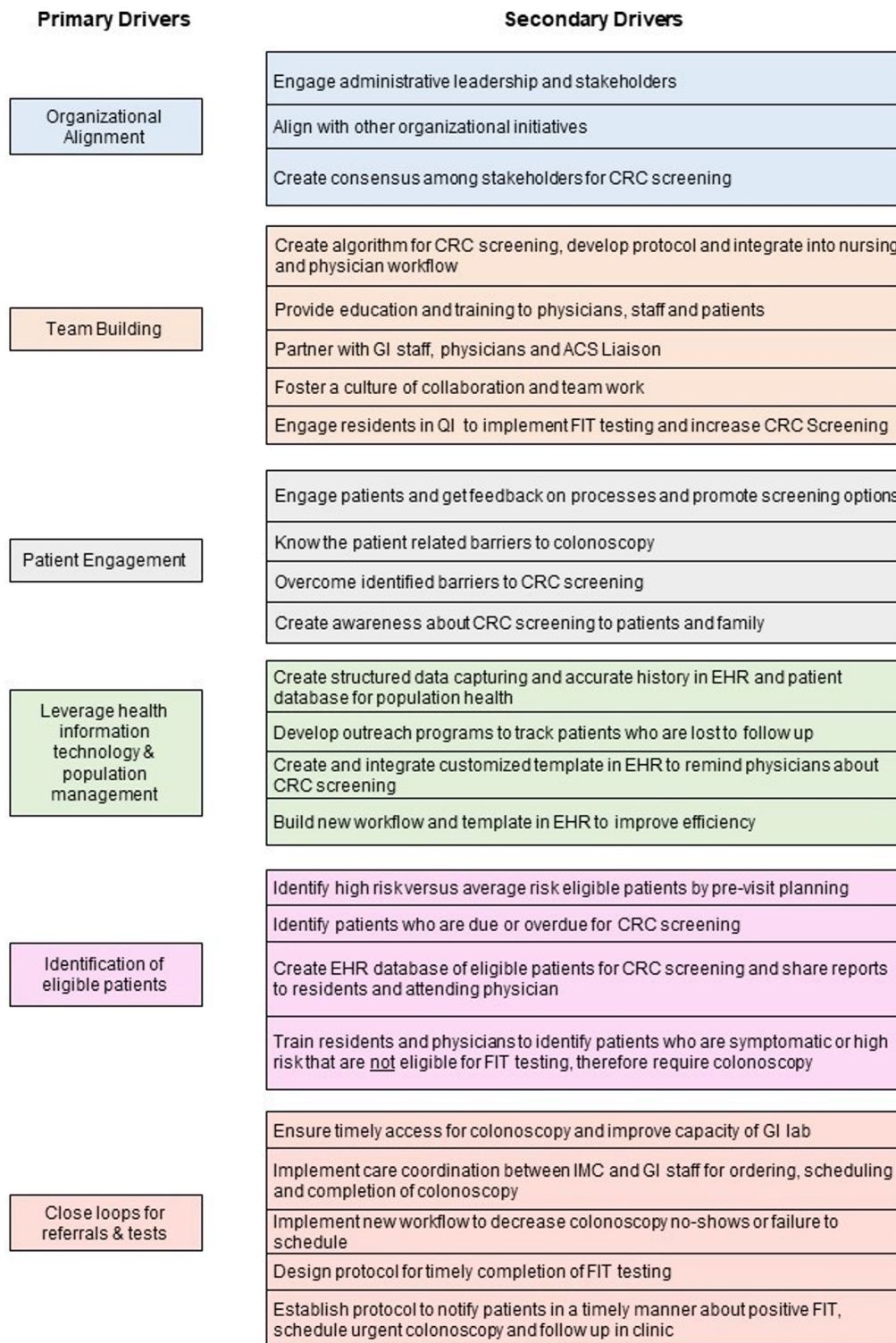


Figure 2 CRC screening driver diagram. ACS, American Cancer Society; CRC, colorectal cancer; FIT, faecal immunochemical test; IMC, internal medicine clinic; QI, quality improvement; GI, gastroenterology; EHR, electronic health record.

Strategy

PDSA cycle 1 (December 2016–January 2017): customised template in the EHR and physician training

We created a customised EHR template that included a FIT check list for the physician. This list included: (1) FIT or colonoscopy option; (2) discussion with patient about the need for colonoscopy after a positive

FIT; and (3) discussion with patient about annual FIT after a negative FIT. We identified gap in resident's knowledge of CRC screening. We educated physicians and clinic staff through PowerPoint presentations in a small group discussion. This curriculum included current CRC guidelines and EHR workflow integration for FIT.

Table 1 Change ideas tested by IMC

Drivers	Change ideas
Organisational alignment	<ul style="list-style-type: none"> ▶ ECMC leadership pledged '80% by 2018 CRC screening' initiative by NCCRT.¹⁰ ▶ Schedule regular meetings with key members to share successes, discuss opportunities for improvement and request allocation of resources to overcome barriers. ▶ Implement FIT by ECMC Director of Laboratory Services. ▶ Offer FIT as an option to colonoscopy in average risk patients. ▶ Increase access for colonoscopy.
Team building	<ul style="list-style-type: none"> ▶ Train nursing staff to dispense FITs and provide patient education on CRC screening options and FIT instructions. ▶ Appoint patient navigator to track FIT orders, FIT completion and urgent colonoscopy referrals. ▶ Nursing staff to identify eligible patients for CRC screening during previsit planning. ▶ Engage information technology staff to confirm accuracy of EHR database. ▶ Prepare monthly run charts to inform team including administrative leadership. ▶ Place stamper on discharge sheet to remind nursing staff about FIT order by physician. ▶ Develop workflow whereby GI coordinator notifies physicians of colonoscopy no-shows. ▶ Improve process to optimise adherence to scheduled colonoscopy appointment. ▶ Develop curriculum to enhance motivational interview techniques among physicians and staff and implement interactive workshops.
Patient engagement	<ul style="list-style-type: none"> ▶ Physicians review current processes for CRC screening with patients and offer new option of FIT or colonoscopy for average risk patients. ▶ Outline patient-related barriers to colonoscopy and FIT and develop plans to overcome. ▶ Provide education to patients about CRC screening. ▶ Engage patients in designing simplified instructions for FIT. ▶ Assign social worker to improve patient related barriers of transportation for colonoscopy. ▶ Patient navigator assesses barriers for FIT completion by calling patients. ▶ Discuss CRC screening at every visit or at least annually for patients who refused in the past. ▶ Offer precolonoscopy appointment by GI staff to discuss procedure and bowel preparation to improve understanding and adherence. ▶ Create workflow to ensure that a physician reviews abnormal FIT results and discuss with patient in a timely manner. ▶ Offer incentive to patients for FIT completion.
Leverage health information technology	<ul style="list-style-type: none"> ▶ Design new EHR patient database to identify and track patients for CRC screening. ▶ Design new nursing workflow for CRC screening to improve physicians' efficiency during clinic visit. ▶ Design new FIT workflow in EHR to document discussion with patients about FIT and colonoscopy options. ▶ Call and send letters to those patients who are lost to follow-up for population health. ▶ Track completed FIT and colonoscopy and ensure physicians' acknowledgement and follow-up. ▶ Generate registry to track patients who refused CRC screening and allocate resources to overcome barriers. ▶ Send automated letter to notify patient about negative FIT result. ▶ Enter colonoscopy results as a structured data field in EHR to improve accuracy of database. ▶ Enter refusal for FIT or colonoscopy as a structured data in EHR. ▶ Enter EHR reminder for high risk patients due or overdue for colonoscopy.
Identification of eligible patients	<ul style="list-style-type: none"> ▶ Physician and nurse documents family history of CRC and tracks pathology report of polyps from previous colonoscopy in EHR. ▶ Medical office assistant retrieves and scans previous colonoscopy report and enters it as a structured data and also scans pathology report in EHR.
Close loops for referrals and tests	<ul style="list-style-type: none"> ▶ Ensure clear indication for FIT or colonoscopy. ▶ Patient navigator tracks FIT orders for completion. ▶ Schedule urgent colonoscopy and track completion in FIT-positive patients. ▶ Track follow-up clinic appointments and improve adherence. ▶ Track no shows for colonoscopy and notification to PCP. ▶ Develop tracking system for positive FIT and abnormal colonoscopy report. ▶ Establish a protocol to notify patients in a timely manner about positive FIT; schedule urgent colonoscopy and follow-up in clinic. ▶ Improve communication with patients when unable to reach by phone; send a letter to remind of scheduled appointment to review positive FIT report.

CRC, colorectal cancer; ECMC, Erie County Medical Center; FIT, faecal immunochemical test; IMC, internal medicine clinic; NCCRT, National Colorectal Cancer Roundtable; PCP, primary care provider

PDSA cycle 2 (February 2017): physician and staff reminder, FIT instruction and patient navigator *Physician and staff reminder*

There was a lack of automated EMR chart alerts for the physician to identify patients that needed CRC screening. We implemented a new workflow that consisted of completion of the CRC-2 questionnaire by the patients with facilitation of the nursing staff prior to the physician evaluation. The CRC-2 questionnaires included the following questions: (1) Have you ever had a colonoscopy?, (2) Are you willing to discuss CRC screening with your physician

today? This workflow was designed to identify patients and to remind the physicians to discuss the CRC screening. Initially, patients completed CRC-2 questionnaire during registration and the nursing staff entered this information into the EHR. After a few weeks, the nursing staff administered and entered the CRC-2 questionnaire into the EHR and also entered a chart alert for the physicians. The QI team developed a 'FIT' stamper, and physicians were instructed to stamp the billing discharge paper to remind the nursing staff to dispense the FIT KIT prior to the patient being discharged.

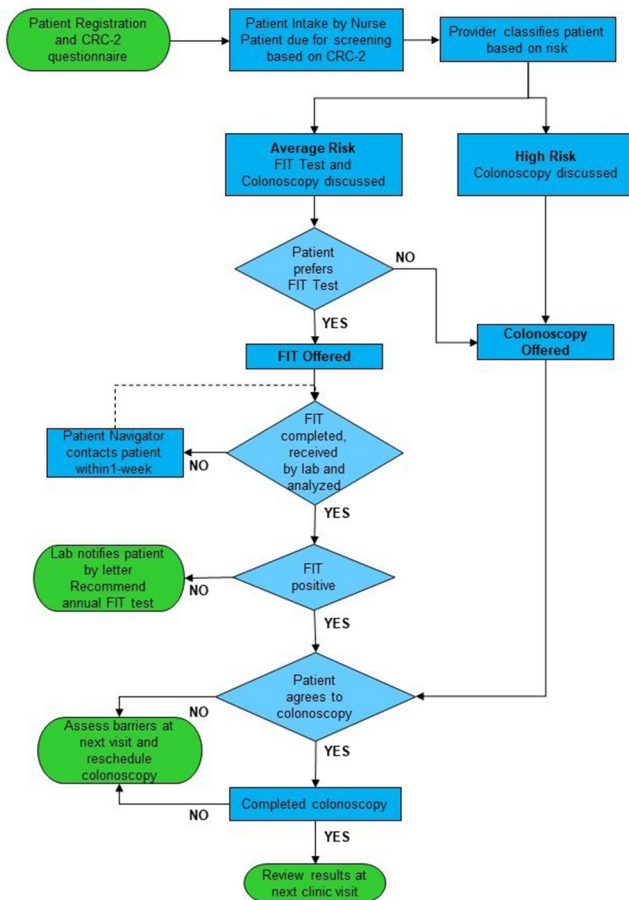


Figure 3 Process flow map. FIT, faecal immunochemical test; CRC-2, colorectal cancer screen 2

FIT instruction

Most of the patients left without a FIT kit due to inability to watch the 5 min video regarding CRC screening and FIT instruction. We replaced the video with a simplified one-page paper instruction that described the three steps of the FIT and also included a reminder to return FIT kit within 3 days.

Patient reminders by navigator

A patient navigator tracked FIT orders in the EHR database and contacted patients weekly with a reminder phone call if they did not return FIT kits.

PDSA cycle 3 (March 2017): interactive workshop

The physician champion conducted an interactive workshop^{36–38} using the NCCRT handbook,³⁹ engaging the IMC multidisciplinary team. A pretest and post-test was administered to identify gaps in the physician's knowledge and to evaluate improvement. The pretest consisted of five questions about CRC statistics, barriers, target audiences and top-rated clinical messages to screening. Professional role play scenarios were created by the IMC staff; and residents were assigned to motivate patient to make a shared patient-centred decision for CRC screening.

PDSA cycle 4 (April–May 2017): EHR outage

On 9 April 2017, ECMC faced an EHR outage for 2 months. FIT was not available for 1 month. The QI team continued to share progress of this QI with IMC staff.

PDSA cycle 5 (June 2017): patient education and tracking of FIT orders

Patient education materials were placed in the examination rooms to create awareness about CRC screening. The QI team outlined the IMC protocol for tracking of FIT orders by the patient navigator. The patient navigator contacted the patient weekly for three attempts, followed by a letter to remind the patient to return the FIT kit. The patient navigator cancelled the FIT order after no response and notified the physician about FIT order status. The patient navigator documented this process in the patient's EHR.

PDSA cycle 6 (July–August 2017): physician education and patient reminder

Initial and refresher training for the residents was conducted for CRC screening. The patient navigator continued to track FIT orders and remind patients to return FIT kits.

PDSA cycle 7 (September–October 2017): FIT incentive and tracking FIT orders

We introduced a FIT incentive programme (\$5 gift card) for timely completion of the FIT kits.

PDSA cycle 8 (November–December 2017): interactive workshop, patient education and physician reminder

The interactive workshop was conducted for the residents. We created a pocket card for physicians that included a CRC screening algorithm and the process flow map for CRC screening. Patient education pamphlets were placed in the examination room to create CRC awareness.

Data analysis

Data were analysed using statistical process control charts for process measures and run chart for outcome measure of CRC Screening. A paired samples t-test was used to compare the mean percentage of correct answers between pretest and post-test.

RESULTS

Demographics

A total of 407 patients received FIT kits. This population had a mean age of 61.3, was 49.4% female, 64.9% were African-Americans, 30.7% were white while 4.4% were classified as other race. We performed a demographic comparison of patients that completed the FIT (group 1) versus those that did not (group 2); 252 completed the test, while 155 did not complete the test. Those in group 1 were 51.2% female, with a mean age of 61.5, 65.5% were African-American, 29.4% were white and 5.2% were classified as other race. Furthermore, in group 1, 17.1% of patients had a history of serious mental illness (schizophrenia or bipolar

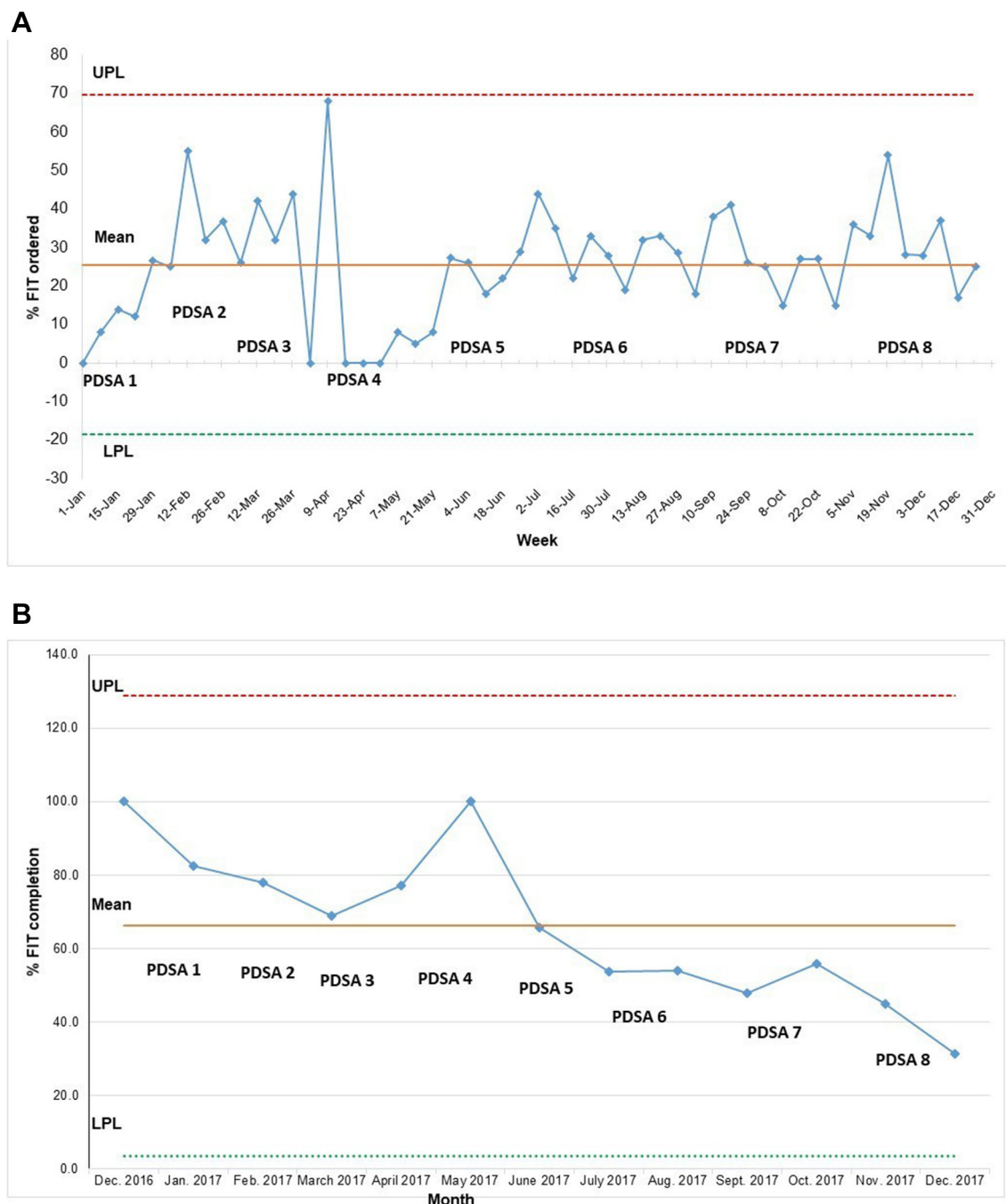


Figure 4 (A) Weekly statistical process control (SPC) chart showing percentage of FIT order rate. (B) Monthly SPC chart showing percentage of FIT completion rate. CL, control limit; FIT, faecal immunochemical test; LCL, lower control limit; PDSA, Plan-Do-Study-Act; UCL, upper control limit.

disorder). For group 2, patients were 46.4% female, with a mean age of 60.9, 63.9% were African-American, 32.9% were white and 32% were classified as other race. Additionally, in group 2, 11.0% of patients had a history of serious mental illness. We did not observe any significant difference in age, gender, race and history of serious mental illness in patients who completed FITs.

Process measures

FIT order rate

The mean FIT order rate was 25.6% (figure 4A). We observed significant weekly variations in FIT order rates

during various PDSA cycles and a sustainable increase in specific weeks within 12 months.

FIT completion rate

The mean FIT monthly completion rate was 66.2% (figure 4B). The mean time from FIT dispense to results was 27.1±37.0 days and median time from FIT dispense to results was 14.0±23.0 days (IQR). Sixty-two per cent (n=252/407) of patients completed FIT. Out of these patients, about 68% (n=172) were screened for initial CRC and 32% (n=80) were screened for repeat CRC.

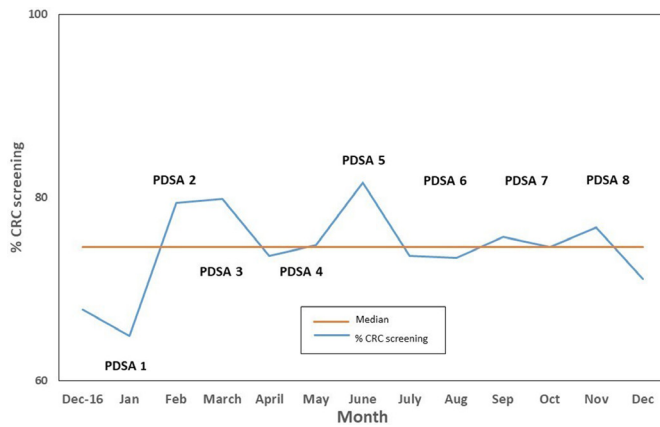


Figure 5 Monthly CRC combined Screening Run chart 2017. CRC, colorectal cancer; PDSA, Plan–Do–Study–Act.

Financial incentives did not significantly improve the FIT completion rate.

Improvement in physician knowledge after interactive workshop

We conducted a pretest and post-test to assess objective evidence of improvement in residents' knowledge. Results showed a statistically significant improvement in the mean percentage of correct responses between pretest and post-test answers ($n=32$). The mean percentage of correct pretest answers was $48.1\% \pm 25.8$, which improved to $96.3\% \pm 7.9$ ($p<0.001$) post-test.

Outcome measures

CRC combined (FIT and colonoscopy) screening rate

The median CRC combined rate (with either FIT or colonoscopy) in the run chart was 74.6% from December 2016 to December 2017 (figure 5). We achieved a sustainable increase in the CRC screening rate to 74.94% (3428/4574 visits), with either FIT or colonoscopy within 12 months (January–December 2017). Positive FIT patients who did not get subsequent diagnostic colonoscopy were not counted for CRC screening rates, since their screening status remained incomplete. We achieved 70.6% ($n=1238/1754$) CRC screening by FIT or colonoscopy in the active (seen within past 18 months) IMC patient population by December 2017. We are able to sustain a CRC screening rate of 75% during the 6-month post-project period (January–July 2018).

Colonoscopy completion rate

There was an increase in referrals for screening colonoscopy for high-risk patients during 12 months. The colonoscopy completion rate was 64% ($n=228/358$) from January 2017 to December 2017. The colonoscopy completion rate also included diagnostic colonoscopy. However, the majority of patients for diagnostic colonoscopy did not have any prior CRC screening.

FIT-positive rate

The FIT-positive rate was 8.7% ($n=22/252$) and 14 of those (63.6%) underwent subsequent diagnostic colonoscopy. All the patients completed colonoscopy within 3

months, except two patients completed within 8 months. One patient was not appropriate for a repeat colonoscopy due to recent colonoscopy in 2015, and one patient has pending colonoscopy. Colonoscopy appointments were rescheduled in six patients after the IMC physicians re-evaluated and discussed the need for diagnostic colonoscopy with patients. Patients had agreed for the colonoscopy; however, they did not keep scheduled colonoscopy appointments. The majority of the patients (4/6) had serious mental illness (schizophrenia or bipolar disorder), and they declined colonoscopy due to the fear of the procedure. Two patients had agreed for an alternate radiology test (double contrast barium CT scan), and one completed the test, but it was a poor study due to large amount of stool present and the other patient did not keep the scheduled appointment. In the FIT-positive IMC population, we found no active cancer, 50% average risk patients had precancerous polyps. Diagnostic colonoscopy results showed tubular adenoma in six patients, tubulovillous adenoma in one patient, diverticulosis in three patients, normal findings in two patients and normal colon with haemorrhoids in one patient. One patient needed a repeat colonoscopy due to poor preparation. One hundred and twenty-five FITs were completed, and 8.0% (10/125) were positive FIT during the 6 month postproject period. Fifty per cent (5/10) of these FIT-positive patients had subsequent colonoscopy, and the other patients are scheduled for colonoscopy.

Balance measures

Increase in wait time for patient

The extra time used on the screening tool did not result in any increase in patient wait time in the clinic. This was determined by anecdotal reports of lack of patient's complaints for a long wait time and lack of overtime for clinic staff during this project. The average time spent in the clinic during a follow-up visit was about 1 hour prior to this QI. The time did not change during project. We did not measure actual time wait time during this QI project.

Increase in wait time for screening colonoscopy

Due to an increase in demand for screening colonoscopy and lack of sufficient GI physicians, there was an increase in the wait time. Before this QI, the average wait time for a screening colonoscopy at ECMC was about 3 months, and it doubled initially after the implementation of this QI due to increase in colonoscopy referrals.

DISCUSSION

We identified a gap in CRC screening at safety-net IMC at ECMC. This QI project was implemented in response to the NCCRT '80% by 2018' CRC screening initiative.¹⁰ The aim of this QI was to increase CRC screening from the baseline rate of 50%–70% in the IMC population over the period of 12 months with the introduction of the FIT. We were able to demonstrate meaningful impact from the different PDSA cycles during this QI. We achieved a sustainable increase in CRC screening rate with either FIT

or colonoscopy to 75% and surpassed our goal of 70% within a short period of 12 months (January–December 2017). In the IMC, 8.7% of the patients screened for CRC had positive FITs. Of those FIT-positive patients, 63.6% completed colonoscopy within 3–6 months. Studies have shown that the time to colonoscopy can result in a negative impact on the patient's health.^{40–41} Delays of up to 12 months after a FIT-positive result can produce up to nearly 10% in losses in overall screening benefits.⁴⁰

The success of this QI project was attributed to patient education, interactive workshops and patient outreach by a navigator. There was objective evidence of increased physician knowledge related to CRC screening after implementation of interactive workshops. Several other initiatives have had relatively positive success rates in improving CRC screening using FIT,^{42–46} patient^{5–47–48} and physician education.^{49–51} Patient outreach by a navigator had a positive impact on FIT completion rates similar to previous studies.^{46–52–54} A decline in the FIT completion rate in the last few months in 2017 may be attributed to the lack of sufficient time allocated to the patient navigator due to shortage of staffing in the IMC.

Patients were actively engaged in this QI project in identification of barriers to CRC screening, education, shared decision making for CRC screening options and feedback on various change ideas. Physicians understood patients' related challenges to CRC screening by embracing open communication with the patients. A social worker interacted with the patient during the clinic visit and offered solutions to overcome barriers. Additionally, a patient navigator assessed the patient-related barriers for FIT completion by calling patients. We identified two major barriers to the completion of FIT: lack of interest due to knowledge gaps and forgetting to do the test. The nursing staff and physicians educated patients on CRC and screening options and facilitated a shared decision to determine the method of CRC screening. After receiving feedback from the patients, the QI team simplified FIT instructions and offered a patient incentive for FIT completion. We were able to update family history of CRC and retrieve reports of prior colonoscopy after physician's interaction with patients during clinic visit. The team implemented a new workflow of precolonoscopy consultation with a GI NP to overcome the barrier of fear that helped to improve patient's understanding and adherence to colonoscopy.

Lessons learnt

Critical factors for success

We attributed the success of this QI to the three T's: (1) technology, (2) tactics and (3) team. Engagement of a multidisciplinary team in performing the root cause analysis and creating the driver diagram was crucial to the design of this QI.

Learning from failures

We offered a \$5 financial incentive, which did not increase our FIT completion rates. Financial incentives lower than \$50 has been shown not to increase FIT

completion rates.^{55–57} Established barriers to colonoscopy such as transportation and financial considerations were not major barriers in this QI.^{19–58} Viewing of videos on FIT instructions by patients was not successful in helping the patient to understand the procedure; this led to the creation of simplified paper instructions, designed with patient's feedback. Initially, the nurses were missing FIT orders in the EHR; therefore, the team created a visual reminder using a red colour 'FIT' stamper on the discharge paper for the nursing staff. We were unable to reach patients for notification of positive FIT results due to inaccurate phone numbers, so we reminded staff to update the patients' contact information during clinic visit. Diagnostic colonoscopy for FIT-positive patients were not scheduled in a timely manner by placing urgent referral in EHR, therefore in addition to a referral, the team designed a new workflow where the physician sent a task directly to the GI NP to facilitate urgent colonoscopy. Initially, we observed multiple no-shows for scheduled colonoscopy, so the team implemented a precolonoscopy consultation appointment with the GI NP to review bowel preparation and procedure with patients. The team improved communication between the GI staff and IMC physicians by adding a new process of notification of no-shows of colonoscopy to the physicians by sending task in the EHR. Due to an increase in demand for colonoscopy, the ECMC leadership added two additional physicians to assist in colonoscopy to improve timely access. During this QI, we discovered inaccuracies in the database for CRC screening, through collaboration with a highly engaged IT staff we were able to ensure accuracy of the database.

Challenges

A few team members proposed using an iPad in the examination room for patient education to show videos and PowerPoint slides for messages for CRC screening. However, due to the cost and safety of iPad in the clinic, the team decided to purchase digital photo frames. Digital photo frames worked well; patients were engaged in watching CRC educational material in the examination room while waiting for the physician. Open and honest communication among multidisciplinary team members facilitated challenging discussions, and the team agreed to try new changes as a pilot test even after there was disagreement among members. The team learnt about various new ideas and embraced successes and failures. The team meetings with IMC residents, physicians and interdisciplinary team, held every 5 weeks, enhanced communication and learning that led to redesigning different interventions.

Previsit planning by the nursing staff to identify patients due for CRC screening was not successful due to time constraints and limited resources. The team agreed to change this workflow and instructed physicians to review and retrieve medical records of prior colonoscopy. Furthermore, we implemented a medical office assistant (MOA) workflow whereby the MOA

entered colonoscopy and FIT reports, refusal for both tests and a reminder for a repeat colonoscopy as a structured data field in the EHR to improve accuracy of EHR database. However, due to limited resources, this workflow was not sustainable. Physicians were trained to enter this information in the EHR, and due to physician's time constraints, it was not entered in the majority of patients. The team planned to reach out to patients who were lost to follow-up for population health, but due to lack of resources, we were unable to accomplish patient outreach. The lack of clinical decision support tool for the physician to identify patients and order appropriate CRC screening test was found to be the greatest IT challenge. Multiple attempts were made by physicians to review a need for subsequent colonoscopy in FIT-positive patients; however, physicians identified major barrier of serious mental illness (schizophrenia and bipolar disorder) to the acceptance to colonoscopy.

This QI has evidence of internal validity. The average time spent by the patient navigator was about 4 hours per week. There was minimal cost involved in conducting this QI. This QI project has several limitations. We reported the data for this QI at 12 months, and it was difficult to have significant improvements in CRC screening rates using colonoscopy due to a long waiting period for the test. This QI was performed in a safety-net primary care clinic in patients with multiple comorbidities, so the barriers and interventions identified in this QI may not be generalisable to other settings.

Sustainability and spread

Processes and workflows designed during this QI project for improving CRC screening in the IMC have become the standard of care and routine part of the clinic visit. IMC physicians continue to offer FIT or colonoscopy options to average risk patients and colonoscopy to high-risk patients. ECMC leadership developed a culture supportive of QI and innovation and assisted in improving access for screening colonoscopy. Continuous training, education and feedback to IMC physicians, residents and staff have fostered team work. We were able to sustain a CRC screening rate of 75% during the 6-month postproject period (January–July 2018). This QI was expanded to new providers (NPs, physician assistants and physicians) in the IMC hired in 2017–2018. The option for FIT for CRC screening is available to patients in the family medicine clinic at ECMC; however, due to physician time constraints, it is not optimal. The team plans to share QI tools, strategies, successes and challenges learnt in this QI project with the family medicine clinic at ECMC and at academic primary care clinic sites at other hospitals. The physician champion presented seminars on this QI project at the University at Buffalo and shared team experience with the residents and faculty from various hospitals. Strategies that were implemented in this QI can be replicated in other settings to improve CRC screening.

Annual FIT screening is feasible for CRC screening in a safety-net clinic. For average-risk patients, FIT was the preferred method of CRC screening in the IMC. We exceeded our goal, achieving 75% CRC screening with either FIT or colonoscopy within a short period of 12 months. NCCRT's goal of 80% by 2018 is attainable in the IMC patient population. Successful strategies included engaging leadership, the front-line staff, a highly effective multidisciplinary team and leveraging health information technology. We were able to sustain CRC screening rate of 75% during the 6-month postproject period. Longitudinal adherence with annual FIT is critical to reduce CRC incidence and for early diagnosis of CRC. Future directions will include strategies to promote repeat annual FIT and outreach of patients that were lost to follow-up in the IMC for population health CRC screening.

Acknowledgements We would like to thank Gregory D. Gudleski, PhD, and staff from the Internal Medicine Clinic, Information Technology and the Gastroenterology clinic.

Contributors SYB: study oversight, study concept and design, acquisition of data, analysis and interpretation of data, drafting of manuscript, critical revision of the manuscript for important intellectual content and finalisation of manuscript. GA and NN: acquisition of data, analysis and interpretation of data. AM: acquisition of data. JR: interpretation of data, drafting of manuscript, critical revision of the manuscript for important intellectual content and finalisation of manuscript.

Funding ACS for the CHANGE grant to increase CRC screening (SB). Research reported in this publication was supported by the National Center for Advancing Translational Sciences of the National Institutes of Health under award Number UL1TR001412.

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.

Ethics approval This study was approved by the Human Subjects Institutional Review Board (HSIRB) of the University at Buffalo and was exempt from patient consent. The work was deemed a quality improvement project and not a study on human subjects.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement An electronic database was setup; contact authors for access.

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

1. National Cancer Institute S. Epidemiology, and end results program. Cancer stat facts: colon and rectum cancer. <https://seer.cancer.gov/statfacts/html/colorect.html>
2. Lin JS, Piper MA, Perdue LA, *et al*. Screening for colorectal cancer: updated evidence report and systematic review for the US Preventive services task force. *JAMA* 2016;315:2576–94.
3. Inadomi JM. Screening for Colorectal Neoplasia. *N Engl J Med* 2017;376:149–56.
4. Brenner H, Stock C, Hoffmeister M. Colorectal cancer screening: the time to act is now. *BMC Med* 2015;13:262.
5. Christy SM, Perkins SM, Tong Y, *et al*. Promoting colorectal cancer screening discussion: a randomized controlled trial. *Am J Prev Med* 2013;44:325–9.
6. Gupta S, Sussman DA, Doubeni CA, *et al*. Challenges and possible solutions to colorectal cancer screening for the underserved. *J Natl Cancer Inst* 2014;106:dju032.

7. Shahidi N, Cheung WY. Colorectal cancer screening: Opportunities to improve uptake, outcomes, and disparities. *World J Gastrointest Endosc* 2016;8:733–40.
8. Wu CA, Mulder AL, Zai AH, et al. A population management system for improving colorectal cancer screening in a primary care setting. *J Eval Clin Pract* 2016;22:319–28.
9. Meester RG, Doubeni CA, Zauber AG, et al. Public health impact of achieving 80% colorectal cancer screening rates in the United States by 2018. *Cancer* 2015;121:2281–5.
10. Roundtable NCC, 2018. Working toward the shared goal of 80% screened for colorectal cancer. <http://nccrt.org/what-we-do/80-percent-by-2018/>
11. Toes-Zoutendijk E, van Leerdam ME, Dekker E, et al. Real-Time monitoring of results during first year of dutch colorectal cancer screening program and optimization by altering fecal immunochemical test cut-off levels. *Gastroenterology* 2017;152:767–75.
12. van der Vlugt M, Grobbee EJ, Bossuyt PMM, et al. Interval colorectal cancer incidence among subjects undergoing multiple rounds of fecal immunochemical testing. *Gastroenterology* 2017;153:439–47.
13. Bibbins-Domingo K, Grossman DC, Curry SJ, et al. Screening for Colorectal Cancer: US Preventive services task force recommendation statement. *JAMA* 2016;315:2564–75.
14. Lin JS, Piper MA, Perdue LA, et al. U.S. Preventive services task force evidence syntheses, formerly systematic evidence reviews. *Screening for colorectal cancer: a systematic review for the US Preventive Services Task Force*. Rockville (MD: Agency for Healthcare Research and Quality (US), 2016.
15. (USPSTF) USPSTF, 2016. Colorectal cancer: screening. <https://www.uspreventiveservicestaskforce.org/Page/Document/UpdateSummaryFinal/colorectal-cancer-screening2>
16. Davis SN, Christy SM, Chavarria EA, et al. A randomized controlled trial of a multicomponent, targeted, low-literacy educational intervention compared with a nontargeted intervention to boost colorectal cancer screening with fecal immunochemical testing in community clinics. *Cancer* 2017;123:1390–400.
17. Wolf AMD, Fontham ETH, Church TR, et al. Colorectal cancer screening for average-risk adults: 2018 guideline update from the American Cancer Society. *CA Cancer J Clin* 2018;68:250–81.
18. Hawley ST, Lafata JE. Colon cancer screening: tackling a multifaceted challenge. *J Natl Cancer Inst* 2014;106:dju383.
19. ACS) ACS. Colorectal cancer facts & figures 2017–2019. <https://www.cancer.org/content/dam/cancer-org/research/cancer-facts-and-statistics/colorectal-cancer-facts-and-figures/colorectal-cancer-facts-and-figures-2017-2019.pdf> (accessed 8 Dec 2017).
20. Goede SL, Rabeneck L, van Ballegooijen M, et al. Harms, benefits and costs of fecal immunochemical testing versus guaiac fecal occult blood testing for colorectal cancer screening. *PLoS One* 2017;12:e0172864.
21. Greuter MJE, de Klerk CM, Meijer GA, et al. Screening for colorectal cancer with fecal immunochemical testing with and without postpolypectomy surveillance colonoscopy: a cost-effectiveness analysis. *Ann Intern Med* 2017;167:544–54.
22. Pham R, Cross S, Fernandez B, et al. Finding the right FIT: rural patient preferences for fecal immunochemical test (FIT) characteristics. *J Am Board Fam Med* 2017;30:632–44.
23. Rex DK, Boland CR, Dominitz JA, et al. Colorectal cancer screening: recommendations for physicians and patients from the U.S. Multi-Society task force on colorectal cancer. *Am J Gastroenterol* 2017;112:1016–30.
24. Robertson DJ, Lee JK, Boland CR, et al. Recommendations on fecal immunochemical testing to screen for colorectal neoplasia: a consensus statement by the US Multi-Society task force on colorectal cancer. *Am J Gastroenterol* 2017;112:37–53.
25. Shapiro JA, Bobo JK, Church TR, et al. A Comparison of fecal immunochemical and high-sensitivity guaiac tests for colorectal cancer screening. *Am J Gastroenterol* 2017;112:1728–35.
26. Akram A, Juang D, Bustamante R, et al. Replacing the guaiac fecal occult blood test with the fecal immunochemical test increases proportion of individuals screened in a large healthcare setting. *Clin Gastroenterol Hepatol* 2017;15:1265–70.
27. Katsoula A, Paschos P, Haidich AB, et al. Diagnostic accuracy of fecal immunochemical test in patients at increased risk for colorectal cancer: a meta-analysis. *JAMA Intern Med* 2017;177:1110–8.
28. Kim NH, Park JH, Park DI, et al. The fecal immunochemical test has high accuracy for detecting advanced colorectal neoplasia before age 50. *Dig Liver Dis* 2017;49:557–61.
29. Soraya GV, Nguyen TC, Abeyrathne CD, et al. A Label-free, quantitative fecal hemoglobin detection platform for colorectal cancer screening. *Biosensors* 2017;7:19.
30. Taylor MJ, McNicholas C, Nicolay C, et al. Systematic review of the application of the plan-do-study-act method to improve quality in healthcare. *BMJ Qual Saf* 2014;23:290–8.
31. Coury J, Schneider JL, Rivelli JS, et al. Applying the Plan-Do-Study-Act (PDSA) approach to a large pragmatic study involving safety net clinics. *BMC Health Serv Res* 2017;17:411.
32. Reed JE, McNicholas C, Woodcock T, et al. Designing quality improvement initiatives: the action effect method, a structured approach to identifying and articulating programme theory. *BMJ Qual Saf* 2014;23:1040–8.
33. Provost LBB. What's your theory? Driver diagram serves as tool for building and testing theories for improvement. *Quality Progress* 2015:36–43.
34. Immunostics Inc, 2018. Fecal Occult Blood 2012. http://www.immunostics.com/fecal_occult_blood_test (accessed 1 Sep 2018).
35. Lee JK, Liles EG, Bent S, et al. Accuracy of fecal immunochemical tests for colorectal cancer: systematic review and meta-analysis. *Annals of internal medicine* 2014;160:171.
36. Vujovic P. Improving teaching skills: from interactive classroom to applicable knowledge. *Adv Physiol Educ* 2016;40:1–4.
37. Ivarsson J, Rystedt H, Asplund S, et al. The application of improved, structured and interactive group learning methods in diagnostic radiology. *Radiat Prot Dosimetry* 2016;169:416–21.
38. Javadi M, Kargar A, Gholami K, et al. Didactic Lecture Versus Interactive Workshop for Continuing Pharmacy Education on Reproductive Health: A Randomized Controlled Trial. *Eval Health Prof* 2015;38:404–18.
39. NCCRT. 80% by 2018, 2016 Communications Guidebook. 2016. <http://nccrt.org/resource-center/>
40. Meester RG, Zauber AG, Doubeni CA, et al. Consequences of Increasing Time to Colonoscopy Examination After Positive Result From Fecal Colorectal Cancer Screening Test. *Clin Gastroenterol Hepatol* 2016;14:1445–51.
41. Corley DA, Jensen CD, Quinn VP, et al. Association between time to colonoscopy after a positive fecal test result and risk of colorectal cancer and cancer stage at diagnosis. *JAMA* 2017;317:1631–41.
42. van der Steen A, Knudsen AB, van Hees F, et al. Optimal colorectal cancer screening in states' low-income, uninsured populations—the case of South Carolina. *Health Serv Res* 2015;50:768–89.
43. Crosby RA, Stradtman L, Collins T, et al. Community-Based colorectal cancer screening in a rural population: who returns fecal immunochemical Test (FIT) Kits? *J Rural Health* 2017;33:371–4.
44. Green BB, Fuller S, Anderson ML, et al. A Quality Improvement Initiative to Increase Colorectal Cancer (CRC) Screening: Collaboration between a Primary Care Clinic and Research Team. *J Fam Med* 2017;4.
45. Myers RE, Sifri R, Daskalakis K, et al. Increasing colon cancer screening in primary care among African Americans. *J Natl Cancer Inst* 2014;106:dju344.
46. Honeycutt S, Green R, Ballard D, et al. Evaluation of a patient navigation program to promote colorectal cancer screening in rural Georgia, USA. *Cancer* 2013;119:3059–66.
47. Christy SM, Davis SN, Williams KR, et al. A community-based trial of educational interventions with fecal immunochemical tests for colorectal cancer screening uptake among blacks in community settings. *Cancer* 2016;122:3288–96.
48. Christy SM, Sutton SK, Gwede CK, et al. Examining the durability of colorectal cancer screening awareness and health beliefs among medically underserved patients: baseline to 12 months post-intervention. *Journal of Cancer Education* 2017;16.
49. Hountz D, Coddington J, Folli KJ, et al. Increasing colorectal cancer screening using a quality improvement approach in a nurse-managed primary care clinic. *Journal for Healthcare Quality* 2017;39:379–90.
50. Arsenault PR, John LS, O'Brien LM. The use of the whole primary-care team, including community health workers, to achieve success in increasing colon cancer screening rate. *Journal for Healthcare Quality* 2016;38:76–83.
51. Schiff GD, Bearden T, Hunt LS, et al. Primary care collaboration to improve diagnosis and screening for colorectal cancer. *Jt Comm J Qual Patient Saf* 2017;43:338–50.
52. Martin RL, Tully M, Kos A, et al. Increasing colorectal cancer screening at an Urban FQHC Using iFOBT and patient navigation. *Health Promot Pract* 2017;18:741–50.
53. Thamarasseril S, Bhuket T, Chan C, et al. The need for an integrated patient navigation pathway to improve access to colonoscopy after positive fecal immunochemical testing: a safety-net hospital experience. *J Community Health* 2017;42:551–7.
54. Sly JR, Jandorf L, Dhulkifl R, et al. Challenges to replicating evidence-based research in real-world settings: training African-American peers as patient navigators for colon cancer screening. *J Cancer Educ* 2012;27:680–6.

55. Gupta S, Miller S, Koch M, *et al.* Financial incentives for promoting colorectal cancer screening: a randomized, comparative effectiveness trial. *Am J Gastroenterol* 2016;111:1630–6.
56. Kullgren JT, Dicks TN, Fu X, *et al.* Financial incentives for completion of fecal occult blood tests among veterans: a 2-stage, pragmatic, cluster, randomized, controlled trial. *Ann Intern Med* 2014;161:S35–43.
57. Mehta SJ, Feingold J, Vandertuyn M, *et al.* Active choice and financial incentives to increase rates of screening colonoscopy—a randomized controlled trial. *Gastroenterology* 2017;153:1227–9.
58. Filippi MK, Perdue DG, Hester C, *et al.* Colorectal cancer screening practices among three American Indian Communities in minnesota. *J Cult Divers* 2016;23:21–7.