Effects of Academic Detailing, Panel Management and Mailed Multi-Target Stool-DNA Testing on Colorectal Cancer Screening

Arturas Klugas, MD, Sara Elsayed, MD, Michael Rodriguez, MD, Shourya Verma, BBA, Andre' Bateman, PhD, and Matthew Stack, DHA

Introduction: Academic detailing, patient-panel management, and mailed, stool-based testing have each been utilized to increase colorectal cancer (CRC) screening in rural clinics. The effectiveness of combining these interventions to increase CRC screening during COVID-19 restrictions was unclear.

Methods: We explored the effects of a multi-component intervention including academic detailing, active patient panel management, and mailed MT-sDNA testing on colorectal cancer screening in our rural family medicine clinic. Baseline interventions included EMR-based provider alerts and mailed patient reminders. Our intervention (March–May 2020) and follow-up periods (June–August 2020) coincided with the initial COVID-19 surge, giving us the opportunity to observe the effects of our intervention during COVID-19 restrictions.

Results: A total of 407 patients were eligible and overdue for colorectal cancer screening. Our clinic's CRC screening rate increased significantly after intervention (69.7%) as compared with before (64.3%) (P = <0.01; 95%CI = 5.39-5.4). Our clinic's CRC screening rates increased significantly during the initial 3 months of the COVID-19 surge (67.8%) compared with the same period the prior year. (62.3%) (P = .003; 95%CI = 3.4-7.6). Our CRC screening rates increased after intervention (69.7%) compared with our regional health system (67%) (P = <0.01; 95%CI = 2.6-2.77). Our weekly stool-based CRC screening increased (94% increase) compared with other health systems nationally (61 to 83% decrease).

Discussion: A multi-component intervention, including academic detailing, panel management, and mailed MT-sDNA testing, can lead to significant increases in CRC screening in a rural family medicine clinic, empowering providers to maintain an effective CRC screening outreach during COVID-19 related restrictions. (J Am Board Fam Med 2023;36:000–000.)

Keywords: Academic Detailing, Cancer Screening, Colorectal Cancer, COVID-19, DNA, Family Medicine, Rural Health

Introduction

Colorectal cancer (CRC) has the third highest cancer incidence among both men and women in the United States with 80,690 new cases in men and 70,340 new cases in women based on 2022 estimates.¹ Colorectal cancer is the third most common cause of cancer mortality for males (28,400 deaths) and females (24,180 deaths) in the US based on 2022 estimates.¹

Screening for colorectal cancer is recommended for asymptomatic men and women ages 50–75 (ages 45–49 grade B recommendation) at average risk via the following: colonoscopy every 10 year; flexible sigmoidoscopy every 10 years + Fecal Immunochemical Test (FIT) every year; flexible sigmoidoscopy every 5 years; CT colonography every 5 years; multitarget stool DNA test (sDNA-FIT) every 1–3 years or highsensitivity guaiac fecal occult blood test (gFOBT) or Fecal Immunochemical Test (FIT) test annually.²

Despite multiple effective screening methods, CRC screening is often underutilized. Only 63.4%

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From the MSU/MyMichigan Medical Center Alma Family Medicine Residency Program, Alma, MI (AK, MS); Family Medicine Residency-McLaren Flint (SE); Western Michigan University Homer Stryker M.D. School of Medicine–Sports Medicine Fellowship, Kalamazoo, MI (MR); Michigan State University College of Human Medicine (SV); University of the West Indies at Mona, Mona, Saint Andrews, Jamaica (AB).

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Corresponding author: Matthew Stack, DHA, MidMichigan Medical Center-Gratiot, 300 E. Warwick Drive, Alma, MI 48801 (E-mail: Matthew.Stack@mymichigan.org).

of females and 61.9% of males are up to date on CRC screening.³ In addition, state to state and rural/urban colorectal cancer screening disparities are recognized. At the state level, screening rates ranged from a high of 76.3% (Massachusetts) to a low of 58.8% (New Mexico). Nationally, screening rates for urban populations were 68.2% compared with a rural screening rate of 65.5%. States with the highest screening rates had the smallest urbanrural disparities (74.6 vs 73.0%) whereas states with the lowest screening rates had the highest urbanrural disparities (61.3% vs 56.9%).⁴

Interventions to increase CRC screening can be categorized as: interventions to increase provider delivery of screening services (provider assessment and feedback; provider reminders); Interventions to increase community demand (patient reminders; small media; group education) and interventions to increase community access (reducing structural barriers; reducing patient costs).⁵ The largest screening increases were seen among multicomponent interventions that combined approaches from each of the 3 strategies.⁵

One approach to increasing provider delivery is academic detailing (AD). Academic detailing is defined as an "interactive educational outreach to physicians to provide unbiased, noncommercial, evidence-based information about medications and other therapeutic decisions with the goal of improving patient care."⁶ This approach parallels the "marketing" strategies used by pharmaceutical sales representative ("detailers") to increase use of their company's products. However, academic detailing is a peer to peer educational outreach in which medical professionals go to colleagues offices and provide brief, objective information to optimize implementation of evidence-based guidelines.⁶ Academic detailing involved a 6 step process: introductions; needs assessment; key message and benefits; handling objections; summary; closing the visit.⁶ Academic detailing can influence breast cancer screening,⁷ pediatric developmental screening,8 HIV testing9 and family physician prescribing behavior.¹⁰ Academic detailing can lead to increased rates of screening colonoscopies and fecal occult blood testing in rural settings.¹¹ Academic detailing to increase CRC screening has proven both feasible and acceptable in a metropolitan¹² and rural¹³ setting. However, although AD may be clinically effective in improving CRC screening, some have questioned its cost effectiveness.14

Patient panel management is one approach to increasing community demand and access to screening

services. Panel management is defined as "a set of tools and processes for population care that are applied systematically at the level of a primary care panel, with PCP's directing proactive care for their empaneled patients."¹⁵ Panel management involves "identifying and reaching out to patients in the panel of a primary care practice who have unmet preventative and chronic condition care needs."¹⁶ Active panel management is regarded as one of the key building blocks to high -performing primary care practices.¹⁷ Active panel management may be a central component of a continuous quality improvement approach to increasing CRC screening.¹⁸

One important intervention shown to increase community access to CRC screening is the mailed FIT test. Interventions using mailed FIT significantly increase CRC screening completion rates compared with usual care.¹⁹ Provision of CRC screening kits by direct mail, use of preaddressed stamped return envelopes and patient reminders seem to be effective in increasing CRC screening in rural and low income populations.²⁰ Multi-target Stool DNA (MT-sDNA) tests combine the features of direct mail, preaddressed postage paid return packaging and patient reminders with increased sensitivity but slightly reduced specificity compare with FIT testing.²¹

In March 2020, the World Health Organization (WHO) designated the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) outbreak a pandemic.²² In addition in March 2020, the American College of Surgeons issued recommendations that hospitals and health systems minimize, postpone or cancel elective operations and endoscopies in an effort to minimize unnecessary SARS-CoV-2 transmissions and direct health care resources toward caring for the surge of SARS-CoV-2 patients.²³ Subsequent to these and similar guidelines, issued in other nations, colorectal cancer screening declined significantly both in the United States and internationally.²⁴

We began our intervention to increase our clinics colorectal cancer screening in March of 2020, coincidental to and concurrent with the initial SARS-CoV-2 surge. Therefore, we faced the added challenge of trying to increase CRC screening in our rural, family medicine clinic during a time when CRC screening was declining nationally and internationally.²⁴ The purpose of this study was to examine the effects of a multicomponent intervention, centered on academic detailing, patient-panel management and mailed MT-sDNA tests, on CRC screening in a rural family medicine clinic during SARs-CoV-2 related restrictions on endoscopic procedures.

Methods

Our study was a within- groups (prepost) and between groups (intervention – control) design and was not randomized. We "controlled" our study by comparing our clinic results against: 1) our clinic's CRC screening data from the year before the SARs-CoV-2 pandemic (March–June 2019); 2) our overall health system CRC screening rates; 3) CRC screening data from other health systems for the same time period as reported in the literature.^{25–27} All researchers were trained in Good Clinical Practice for the protection of human subjects and our study was approved by our Institutional Review Board (IRB).

Our population was age eligible adults in our rural, Midwest family medicine clinic. Inclusion criteria were asymptomatic male and female patients, ages 50-75, at average risk of CRC who were due/ overdue for CRC screening (our protocol was established before the USPSTF 2021 grade B recommendation² that CRC screening begin at age 45). Frequency of recommended screening depended on the method used. Exclusion criteria included: previous gastrointestinal cancer; Lynch syndrome; family adenomatous polyposis; inflammatory bowel disease; patients symptomatic on presentation (rectal bleeding) requiring diagnostic colonoscopies; patients with abnormal results on previous CRC screening necessitating more frequent colonoscopies; and patients on hospice.²

Our health system already had 2 interventions in place to increase CRC screening before our study: EMR based provider alerts of care gaps and reminders mailed to patients overdue for screening. Our baseline (preintervention) CRC screening rates are reflective of these 2 interventions already being in place.

We conducted a phone and e-mail survey among medical providers in our health system who had the highest CRC screening rates (exemplar²⁸ providers) to identify their best practices. Most of them were skilled in active patient panel management, using the EMR to identify patients on their panel overdue for health screenings and to facilitate these screenings. Many of these exemplar providers also utilized mailed stool-based CRC screening that included addressed, postagepaid return packaging.

We conducted 2, brief online academic detailing sessions with our clinic providers in early March 2020 which included a review of: colorectal cancer screening guidelines; how to access their panel in the EMR and identify patients overdue for screening; how to order the MT-sDNA test and track patient completion and results. Providers were encouraged to dedicate 1 hour each week to panel management, during which they: identified patients overdue for screening, called and educated them about CRC screening and, if appropriate, recommend home based MT-sDNA testing. The intervention period was March through end of May 2020 and outcome data were tracked through August 2020.

Our multi-component intervention involved several strategies in accordance with the Community Preventive Service Task Force Recommendations⁵ and is summarized in Table 1.

Table 1. Multi-Component CRC Screening Intervention in Rural Family Medicine Clinic March-May 2020

Interventions to increase provider delivery of screening services

- Surveyed "exemplar" providers in our health system to identify their best practices
- Academic detailing of all clinic providers
- Providers dedicating time each week to panel management
- Each provider audited monthly and received a statement about their individual CRC screening rates
- Overall clinic CRC screening rates were posted and updated monthly on a white board inside the staff clinic entrance Interventions to increase community demand
 - Follow up phone call and letter to each patient at 1- and 2-months post MT-sDNA order
 - Staff and clinicians wore CRC awareness ribbons
 - CRC informational posters and brochures (small media) were originally planned for clinic waiting areas but removed due to COVID-19 infectious disease control measures

Interventions to increase community access

- Prioritization of use of mailed MT-sDNA tests rather than colonoscopy/ sigmoidoscopy
- Phone call from provider to overdue patients recommending CRC screening and offering MT-sDNA screening
- One staff designated for patient follow up and navigation on all MT-sDNA orders

Abbreviation: CRC, colorectal cancer.

Our independent (predictor) variable was our multi-component CRC screening intervention. Our dependent (outcome) variables included: Stool-Based CRC screenings ordered per month; Stool-Based CRC screenings completed per month; Stoolbased CRC screenings ordered or completed per week; and overall colorectal cancer screening rates. The colorectal cancer screening rate statistic was based on the number of eligible patients who completed screening/number of patients eligible for screening during the time period. Stool based CRC screenings ordered and completed per month were compared before (Feb/March) and after (April/May) intervention. Stool based CRC screenings ordered/completed per week were compared within-groups (before and after) for our clinic and between groups (our clinic results compared with results from other clinics cited in the literature from the same time period $^{25-27}$). Change in overall CRC screening rates are compared with-in groups (pre and post intervention), between our clinic (intervention) and our regional health system (control) and between our clinic during the initial

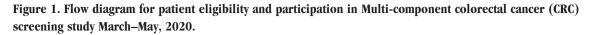
4 months (March–June, 2020) of the SARS-CoV-2 surge and our clinics data during the same time period the year prior.

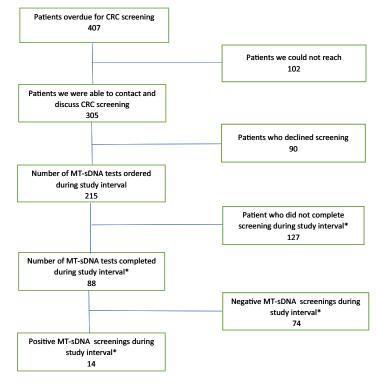
Within-group (before and after) analysis was performed using paired sample t test. Between group analysis was performed by comparing our study outcomes with regional and national test values cited in the literature^{25–27} using 1 -sample t test. Statistical analysis was conducted with the Statistical Package for Social Sciences (SPSS) version 25.

Results

Our study population consisted of 407 patients in our rural Midwest family medicine clinic who were eligible and overdue for colorectal cancer screening. Patient eligibility, participation and disposition is illustrated in Figure 1.

Patient demographics were obtained via a selfreport survey. Overall, our study population tended to be older, white, middle class, female, and more likely to report higher education and having a





*An additional 37 MT-sDNA tests results were received in the 3 months post study interval

primary care provider compared with county wide, U.S Census Bureau statistics. The older age of our sample is likely a result of age criteria to be eligible for CRC screening. The higher rate of having a PCP may be due to the fact our sample population was drawn from our electronic medical record. The gender, educational and socio-economic differences in of our sample may be reflective of self- sampling bias and/or differences between our definition of terms and that of the U.S Census Bureau.²⁹ Patient demographic of our sample and for the county wide population are shown in Table 2.

We observed a significant increase in the number of stool- based CRC screenings *ordered* per month in our clinic after intervention (n = 64) as compared with before (n = 1) (t = 4.49; P = <0.01; 95% CI = 1.96-5.46) We observed a significant increase

Table 2. Demographic Characteristics of Multi-Component Study Sample and County-Wide PopulationMarch 2020

Demographic Category	Study Sample Percentages	County-Wide Percentages*	
Average age	63.3	40	
Gender			
Man	45.2	54.1	
Woman	54.1	45.9	
Location of Residence			
Urban	0.7	0	
Small town/Suburb	44.4	49.1	
Rural	54.8	50.9	
Ethnicity			
White	97.8	84.8	
Black/African American	0	6.4	
Hispanic	1.5	6.8	
Other	0.7	2.0	
Education			
Elementary/Junior high only	1.5	10.5	
High school	42.6	41.3	
College/Vocational training	42.6	42.8	
Graduate/Professional degree	13.2	5.4	
Socioeconomic status			
Lower	28.7	46	
Middle	64.3	51	
Upper	7	3	
Smoker			
Yes	20.9	22.1	
No	79.1	77.9	
Primary care provider			
Yes	97.8	83.6	
No	2.2	16.4	

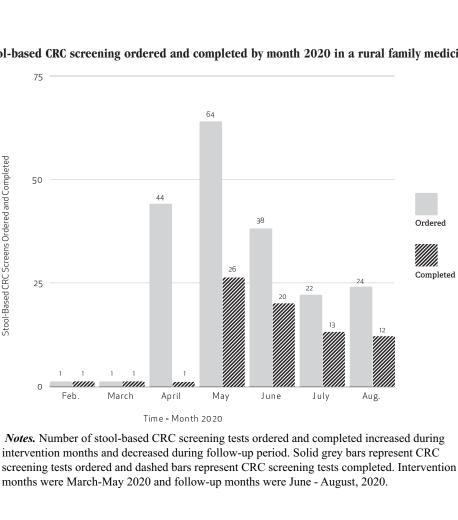
Notes. *Based on U.S. Census county population data.

in the number of stool based CRC screening *completed* per month in our clinic after intervention (n = 26) as compared with before (n = 1) (t = 2.71; P = .02; 95% CI = 0.32-2.62). (See Figure 2).

We observed a significant increase in weekly stool-based CRC screenings in our clinic after intervention (average 16 screens per week) as compared with before (average 1 screen per week) (t =10.94; $P = \langle 0.01; 95\%$ CI = 0.71-1.05). We observed a significant increase in our clinics weekly stool - based CRC screening (average of 1 screen per week to 16 per week) when compared with our regional health system²⁵ (average of 210 per week to 51 per week) (t = 2719; P = <0.01; 95% CI = 159.8 to160.1). We observed a significant increase in our clinic's average weekly stool-based CRC screening when compared with health systems in the San Francisco²⁶ (370 per week to 60 per week) (t = 5286; P = <0.01; 95% CI = 310.8 to 311) and Los Angeles areas²⁷ (154 per week to 60 per week) (t = 1614; P = <0.01; 95% CI = 94.8 to 9). Although the total weekly stool-base CRC screening of these larger health systems was still higher than our individual clinic, our stool-based CRC screens were trending up during COVID-19 restrictions (94% increase) whereas our regional and the national health system rates were trending down (61 to 83% decrease) during the same time period. See Table 3.

We observed a significant increase in the overall CRC screening rate in our clinic after intervention (69.7%) as compared with before intervention. (64.3%) (t = 1231.9; P = <0.01; 95% CI = 5.39-5.4;). We observed a significant increase in the overall CRC screening rates in our clinic during the initial 4 months (March-June 2020) of the SARS-CoV-2 surge (67.8%) compared with our clinic CRC screening rate from the same period the year prior (62.3%) (t = 8.5; P = .003; 95% CI = 3.4 to 7.6). Our health system experience a significant decrease in overall CRC screening rates during the initial 4 months of the SARS-CoV-2 surge (68.3%) compared with the same 4 month time period the year prior (74.5%) (t = 7.3; P = .005; 95% CI = 3.5 - 8.7) We observed a significant increase in our overall clinic CRC screening rates (69.7%) as compared with our regional health system rates (67%) (t = 74.2; P = <0.01; 95% CI = 2.6-2.77). See Figure 3.

During our 3-month postintervention follow up (June–August 2020) we observed a decrease in some





64

May

April

Time - Month 2020

44

months were March-May 2020 and follow-up months were June - August, 2020. clinic outcome measures including: stool-based CRC screens orders (38, 22, and 24 orders placed) and stool-based CRC screens completed (20, 13, and 12 completed). After our intervention period, our clinicians focused their academic detailing and panel management efforts onto other preventive services

March

Discussion

Our multi-component intervention, centered on academic detailing, patient-panel management and mailed MT-sDNA, led to a significant increase in CRC screening in our rural family medicine clinic. Of additional importance, we were able to achieve this significant increase during a time when CRC screening was declining regionally,²⁵ nationally,^{26,27} and internationally²⁴ due to SARS-CoV-2 restrictions. Our study parallels and furthers the evidence that

Table 3. Clinic Stool-Based CRC Screening Orders per Week Compared with Other Health Systems Pre and Post Initial COVID-19 Surge, 2020

Study	Stool-Based Screen/Week Pre	Stool-Based Screen/Week Post	Paired-Sample t Test	One-Sample <i>t</i> Test	% Change in CRC Screening
Klugas et al. 2023	1	16	$t = 10.95^{***}$		94% more
Gorin ^a et al. 2021	210	51		t = 2719***	75% less
Patel et al. 2020	370	60		$t = 5286^{***}$	83% less
Myint et al. 2021	154	60		<i>t</i> = 1614***	61% less

Notes: ^aClinic's regional health system. ***P = <0.01.

75

50

25

Feb

(breast cancer screening). It is possible that our

decline in CRC screening during the follow up phase

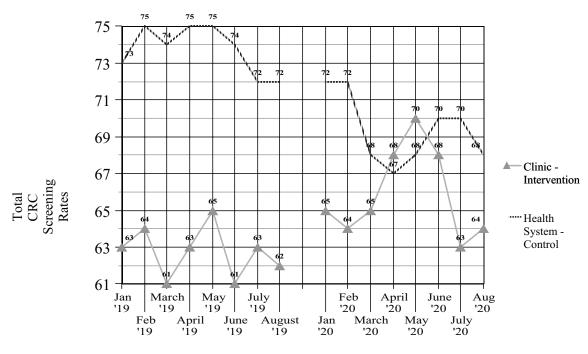
was due to this abrupt redirection of our preventive

Stool-Based CRC Screens Ordered and Completed

Abbreviation: CRC, colorectal cancer.

screening efforts.





Months 2019 and 2020

Notes. Clinic CRC screening rates increased during intervention while health system CRC screening rates decreased during the same period. CRC screening rates for both clinic and health system trended towards baseline after intervention. Changes in screening rates during initial COVID-19 surge are compared with the same time period the year prior (Jan.- August, 2019). Clinic rates represented by solid line and health system rates represented by dashed line.

increased use of mailed stool -based CRC screening may counterbalance and offset SARS-CoV-2 induced decreases in screening endoscopies.³⁰

Our findings confirm and extend the evidence a that multi-component intervention, focused on increasing provider delivery, community demand and community access, can significantly increase CRC screening.⁵ The individual and separate effects of academic detailing¹¹ and mailed FIT^{19,20} on CRC screening have been studied previously. However,

Table 4. Strengths and Limitations of Our Multi-Component Intervention to Increase CRC Screening Rates in aRural Family Medicine Clinic During Initial COVID-19 Surge, 2020

Strengths

Our survey of exemplar²⁸ providers' strategies for achieving higher CRC screening rates lends qualitative support to our intervention strategy

We used a multi-component intervention strategy in line with Community Preventive Service Task Force recommendations We attempted to "control" our study by comparing our intervention clinic data with regional and national outcomes as well as our own clinic data from year before the pandemic

The fact that our CRC screening rates increased during a time when rates were decreasing regionally,²⁵ nationally^{26,27} and internationally²⁴ adds evidence that our intervention had an impact

Limitations

Inability to differentiate the individual contribution of each component of our intervention to the overall treatment effect Our study may be more subject to effects of confounding variables due to the lack of randomized, controlled design

Based on our self-reported demographics, persons of color and persons without a primary care provider may be under represented in our results

Our results based on one, rural Midwest clinic may limit external validity and generalizability

Abbreviation: CRC, colorectal cancer.

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the literature on the effects of patient-panel management on CRC screening was limited.¹⁸ Our study demonstrates that active panel management can be an important component, when combined with other strategies, to increased colorectal screening. To our knowledge, ours is the first study to demonstrate the effectiveness of combining academic detailing, panel management and mailed MT-sDNA testing during SARS-Cov-2 restrictions.

Our interventions are applicable to busy primary care providers and under-resourced patients in medically underserved settings. Academic detailers come to the clinic, eliminating the time and expense of CME travel. Medical assistants can be trained to facilitate panel management. Mailed MT-sDNA tests reduce the financial, scheduling, and transportation barriers that colonoscopies pose for patients. Other strengths and limitations of our study are detailed in Table 4.

Future studies could recruit multiple clinics from a broader geographic area and incorporate stratified sampling, so the sample was more representative of the US population distribution. A design which randomized clinics into intervention and "treatment as usual" (control) clinics would reduce the effects of confounding variables. Future studies could be designed so that the unique contributions of academic detailing, panel management and mailed MT-sDNA could be individually analyzed via multiple regression.

Conclusion

Our study demonstrates that a multi-component intervention, centered on academic detailing, active patient-panel management and mailed MT-sDNA can lead to significant increases in CRC screening in a rural family medicine clinic even during SARS-CoV-2 induced restrictions on endoscopic screening.

To see this article online, please go to: http://jabfm.org/content/ 00/00/000.full.

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