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## **INTRODUCTION**

Colorectal cancer (CRC) is the third most common cancer and the second most common cause of cancer death in the U.S., with over 135,000 new cases and over 50,000 deaths expected in 2017 Identifying and removing pre-cancerous polyps car greatly reduce invasive disease, emphasizing the need for effective CRC screening. Yet, in 2015 only 63% of adults aged 50+ were up-to-date on CRC screening. This rate falls short of targets set by the National Colorectal Cancer Roundtable (80% by 2018) and Healthy People 2020 (70.5%). Although improving, adult CRC screening rates in federally gualified health centers (FQHCs) remain well below those of non-FQHC populations, at 38.3% in 2015. Low screening utilization delays CRC detection and leads to higher CRC-related morbidity and mortality.

Fecal immunochemical testing (FIT) may be a lowcost alternative, especially well-suited to FQHC patients, and an effective population-based CRC screening approach when combined with colonoscopy following positive tests. Strategies and Opportunities to STOP Colon Cancer In Priority Populations (STOP CRC) is a clusterrandomized pragmatic study of an intervention designed to increase CRC screening rates within FQHCs by promoting use of low-cost FIT. We present development, implementation, and maintenance costs of the STOP CRC intervention as adopted by participating FQHCs.



KAISER PERMANENTE **Center for Health Research** 

# **Costs of Establishing a Colorectal Cancer Screening Program in Safety Net Clinics**

## INTERVENTION

Eight FQHC organizations, seven in Oregon and one in California, and comprising 26 individual clinics, participated in STOP CRC (Table 1). Eligible patients were 50-74 years old with at least one office visit in the last year, and were not current for CRC screening (colonoscopy within nine years, no flexible sigmoidoscopy within four years, no FIT or fecal occult blood test (FOBT) within 11 months, no referral for colonoscopy, flexible sigmoidoscopy, or gastroenterology within one year or FIT order within six months), and were not living in a nursing home or assisted living facility within 12 months. Permanent exclusions were prior CRC diagnosis, prior colectomy, ulcerative or inflammatory colitis, history of end stage renal disease or evidence of hospice care.

STOP CRC was an automated electronic health record (EHR) data-driven program for mailing FIT kits to patients due for CRC screening. Eligible patients received an introductory letter with a number to call to decline participation, address clinical concerns, etc. Remaining eligible patients were mailed FIT kits, including illustrated instructions and return postage. Non-completers received reminder letters. EHR mailing status data were updated nightly. Four to six months after staff training, a plan-do-study-act (PDSA) improvement cycle was facilitated with participating clinics to identify strategies to improve reach or effectiveness.

|                                  |            |               | Table 3. Intervention Activities  |  |           |  |  |
|----------------------------------|------------|---------------|---|--|-----------|--|--|
|                                  |            |               | DEVELOPMENT   | IMPLEMENTATION                             | MAI       |  |  |
|                                  |            |               | Data management   | Data management                            | Data      |  |  |
|                                  |            |               | Updating claims data  | <ul> <li>Lab orders tracking</li> </ul>    | • E⊦      |  |  |
|                                  |            |               | Initial EHR training  | <ul> <li>Results pool tracking</li> </ul>  | • Re      |  |  |
|                                  |            |               | Testing EHR tools   |  | • La      |  |  |
|                                  |            |               |   |  | • Re      |  |  |
|                                  |            |               | Staff training  | Staff training                             |           |  |  |
| Table 1.                         | STOP CRO   | C Federally   | • Staff (e.g., medical  | <ul> <li>On-going training, new</li> </ul> |           |  |  |
| Qualified Health Centers (FQHCs) |            |               | assistant) training   | staff on-boarding                          |           |  |  |
|                                  |            |               |   | Dissemination process                      | Diss      |  |  |
| FQHC                             | Number     | Patients Ages |   | <ul> <li>Adapting mailings</li> </ul>      | • Ma      |  |  |
|                                  | of Clinics | 50-74         |   | <ul> <li>Mailing intro letters</li> </ul>  | • Ma      |  |  |
| Α                                | 2          | 3.747         |   | Mailing FIT kits                           | • Ma      |  |  |
|                                  |            | 1.015         |   | Mailing reminders                          |           |  |  |
| В                                | 2          | 1,215         |   | • IN-CIINIC FIT KIT                        |           |  |  |
| С                                | 6          | 7,334         | Program management  | Program management                         | Prog      |  |  |
| D                                | 2          | 2,498         | Execution of lab  | Billing adjustments                        | • Bil     |  |  |
|                                  |            | //75          | interface agreements  | Conducting PDSAs                           | • Cc      |  |  |
| E                                | 4          | 0,0/5         | Research team   | <ul> <li>Provider engagement</li> </ul>    | • Wo      |  |  |
| F                                | 3          | 3,267         | meetings  | meetings                                   | on        |  |  |
| G                                | 3          | 3.810         |   | č  | • Sta     |  |  |
|                                  |            | 2,010         | *FHR: electronic medical reco   | rd: FIT: fecal immunochemical test: PC     | )SA· plan |  |  |
| н                                | 3          | 2,085         | Link, electronic medical record, Fir, lecal minuhochemical lest; PDSA; plan |  |           |  |  |

## Table 2. Implementation Activities by Organization

|                            | ORGANIZATION                           |  |                                     |                                  |  |                                  |                                      |  |
|----------------------------|--|--|-------------------------------------|----------------------------------|--|----------------------------------|--------------------------------------|--|
| Clinic Activity            | A                                      | В  | С                                   | D                                | E                                      | F                                | G                                    |  |
| Chart Scrubbing            | YES                                    | YES  | YES                                 | YES                              | YES                                    | YES                              | YES                                  |  |
| Mailed Intro<br>Letters    | Single batch                           | YES at start/<br>with kits later               | YES at start/<br>with kits later    | YES at start/<br>with kits later | With kits from start                   | YES at start/<br>with kits later | YES                                  |  |
| Mailed FIT Kits            | Single batch                           | Monthly  | Monthly                             | Monthly                          | Monthly on<br>birthdays                | Randomly when time allowed       | Single lar<br>batch                  |  |
| FIT Return                 | Direct to lab<br>at start              | Return to clinic<br>for in-house<br>processing | Return to<br>clinic prior<br>to lab | Direct to lab<br>at start        | Patient hand<br>delivered to<br>clinic | Direct to lab                    | Return to<br>for in-hou<br>processin |  |
|                            | Patient hand delivered to clinic later |  |                                     | Return to clinic first later     |  |                                  |                                      |  |
| Mailed Reminder<br>Letters | NO                                     | YES  | YES                                 | NO                               | YES                                    | NO                               | NO                                   |  |
| Phone Calls                | NO                                     | YES  | NO                                  | YES at control clinic            | NO                                     | NO                               | NO                                   |  |

Organizations implemented STOP CRC as appropriate for their individual systems (Table **2)**. The greatest variety across organizations involved handling of the FITs themselves, which were variously mailed to participants, sometimes monthly, in single large batches, or as staff time permitted. Most organizations mailed the FIT and introductory letter together. Processing of completed FITs was perhaps even more varied. FITs were returned to the clinic for lab processing either in-house (four organizations) or at an outside lab (one); at three organizations, FITs were sent directly to the outside lab. Usual care was existing clinic activities that supported CRC screening.

## **METHODS**

in 2016–2017.

Clinics at each organization were split between intervention and usual care for 18 months (February 2014–August 2015). From August 2015 to February 2016, intervention clinic staff were trained in implementation. Three organizations did not participate in the maintenance phase, so maintenance costs are limited to the five remaining organizations. Organization staff completed spreadsheets using an activity-based costing format (Table 3), assigning labor hours by intervention activity and wage rates by job position. Non-labor costs were obtained from study data. Data were from 2014–2015, and analyses were performed

## RESULTS

Estimated average development costs were nearly \$6K (Figure 1) with 54% allocated to data management and 37% to program management. Overall implementation costs averaged \$14K, allocated mostly to mailing tests (54%) and staff training (28%). Development costs per clinic averaged nearly \$3.8K (Figure 2). Implementation costs per clinic averaged almost \$9.7K; one organization (C)'s costs were \$24.1K, primarily due to extensive training and processing of returned FITs. Per-clinic implementation costs for the seven other organizations averaged \$7.6K.

## NTENANCE

management HR scrubbing port generation b orders tracking sults pool tracking

#### mination process ailing intro letters ailing FIT kits ailing reminders

ram managemen<sup>:</sup>

ing adjustments onducting PDSAs orking with sponsor issues taff check-ins -do-study-act



# Figure 1. Development and Implementation Costs by Activity Category



# Figure 2. Development and Implementation Costs Per Clinic, by Organization\*



Figure 3 illustrates per-clinic activity categories across implementation and maintenance phases The largest reported cost category for each organization was preparing mailings, including printing letters, affixing labels, and placing lab orders.

Six-month maintenance costs, averaging nearly \$20K across five reporting organizations, were much more heterogeneous than implementation costs. One organization reported program management as its most resource-intensive category, while two others reported data





management and test distribution, respectively, as their most resource-intensive. No organization reported staff training during the maintenance phase. Figure 4 presents crude evidence of economies of scale, i.e., cost per person screened appears to decline with the number of individuals screened.

# CONCLUSIONS

Our results highlight the cost implications of implementing a standard CRC screening intervention within a pragmatic trial setting involving multiple FQHCs with varied patient populations, clinical structures, and resource availability. We plan to integrate these findings into a more comprehensive economic evaluation that contributes to our knowledge of how to introduce such programs to underscreened populations most effectively and efficiently.

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