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Patient and provider factors associated with colorectal cancer screening among average risk health plan enrollees in the US, 2015–2018



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Abstract

Background To assess patient and primary care provider (PCP) factors associated with adherence to American Cancer Society (ACS) and United States Preventive Services Task Force (USPSTF) guidelines for average risk colorectal cancer (CRC) screening.

Methods Retrospective case-control study of medical and pharmacy claims from the Optum Research Database from 01/01/2014 - 12/31/2018. Enrollee sample was adults aged 50 - 75 years with ≥ 24 months continuous health plan enrollment. Provider sample was PCPs listed on the claims of average-risk patients in the enrollee sample. Enrollee-level screening opportunities were based on their exposure to the healthcare system during the baseline year. Screening adherence, calculated at the PCP level, was the percent of average-risk patients up to date with screening recommendations each year. Logistic regression modelling was used to examine the association between receipt of screening and enrollee and PCP characteristics. An ordinary least squares model was used to determine the association between screening adherence among the PCP's panel of patients and patient characteristics.

Results Among patients with a PCP, adherence to ACS and USPSTF screening guidelines ranged from 69 to 80% depending on PCP specialty and type. The greatest enrollee-level predictors for CRC screening were having a primary/ preventive care visit (OR = 4.47, p < 0.001) and a main PCP (OR = 2.69, p < 0.001).

Conclusions Increased access to preventive/primary care visits could improve CRC screening rates; however, interventions not dependent on healthcare system contact, such as home-based screening, may circumvent the dependence on primary care visits to complete CRC screening.

Keywords Screening, Colorectal cancer, Patient characteristics, Provider characteristics

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Introduction

Approximately 4% of the US population will develop colorectal cancer (CRC) over their lifetime [1]. Early-stage CRC is highly treatable; however, only 37% of people are diagnosed with localized CRC [2]. When diagnosed with localized disease, the a 5-year survival rate is approximately 91%; however, this drops to 15% when distant metastases are present at diagnosis [3].

The American Cancer Society (ACS) and the United States Preventive Services Task Force (USPSTF) recommend CRC screening for average-risk individuals beginning at 45 years of age using stool, imaging, or endoscopy tests. [4, 5] Screening can detect neoplasia at an earlier, more treatable stage and prevent CRC through identification and removal of pre-invasive lesions (e.g., adenomatous polyps); however, 30.3% of eligible US adults are not up to date with CRC screening [6]. In previous studies, fear, lack of provider recommendation, and lack of patient knowledge about screening were cited as the top patient-reported barriers to CRC screening. [7, 8] CRC screening adherence increased when patients were given a choice of screening modality [9].

Several studies have investigated patient- and providerlevel characteristics associated with CRC screening adherence. In previous studies, patient characteristics associated with higher CRC screening participation rates included male sex, white race, being married, non-obese, higher education levels, and higher socioeconomic status. [10–14] Provider factors associated with higher adherence to CRC screening guidelines include a larger panel of patients eligible for screening, working in a facility that is a shorter distance to a colonoscopy center, and providing time and reminders for CRC screening discussions with patients [15, 16].

The purpose of this study was to assess and update patient and provider factors associated with adherence to ACS and USPSTF guidelines for average risk CRC screening in the context of currently available screening modalities, including the newer Cologuard test (FDA approved in 2014). Large-scale survey studies, including data on CRC screening rates collected by the Behavior Risk Factor Surveillance System (BRFSS) [17] and National Health Interview Survey (NHIS) [18], rely on patient self-report on whether they've received screening and the timing of screening. Previous studies have suggested self-report of CRC screening has a moderate level of validity [19-21] and reliability [22] with a tendency towards over-reporting. The data used in the current study provides a more accurate assessment of the timing of screening tests with confirmed dates of service.

Methods

Study design and data source

In this retrospective case control study, medical and pharmacy claims and enrollment information from January 01, 2014 to December 31, 2018 (study period) were obtained from US commercial and Medicare Advantage health plan members in the Optum Research Database (ORD). The ORD is a de-identified, diverse, and nationally representative administrative claims database containing approximately 9% of the US commercially insured population as of 2017. Institutional review board approval or waiver of approval was not required for this study because the study data were secondary and deidentified in accordance with the United States Department of Health and Human Services (HHS Privacy Rule's requirements for de-identification codified at 45 C.F.R. § 164.514(b)). Two study samples were identified: a population of eligible health plan enrollees (enrollees), and a population of primary care providers. Medical claims for screening tests were identified through International Classification of Diseases, 9th and 10th Revisions, Clinical Modification (ICD-9-CM/ICD-10-CM) procedure codes (ICD-0-PCS/ICD-10-PCS) and Current Procedural Terminology (CPT) codes.

Enrollee sample selection

The enrollee study sample consisted of adults aged 50 to 75 years during the calendar year of the analysis with continuous enrollment in the health plan with medical and pharmacy benefits for ≥ 12 months (up to 10 years) prior to the year of the analysis (baseline period) through the entire year of the analysis (minimum of 24 months of enrollment [follow-up period]). The most recent, available, complete year of enrollment was used as the year of analysis for each enrollee. Enrollees were excluded if they had evidence of conditions indicating a higher risk for CRC (i.e., ≥ 1 medical claim during the baseline period with a diagnosis code in any position on the claim for adenoma, sessile serrated polyp, prior diagnoses of CRC) or a personal history of or diagnosis of inflammatory bowel disease at any time during the study period; or a family history of CRC/gastrointestinal cancer.

Enrollee screening status cohorts

The most recent full calendar year of available data was used to assign screening status to the enrollee population. Screening was based on the most recent ACS and USPSTF guidelines using a stool-based test (i.e., fecal immunochemical test [FIT], guaiac-based fecal occult blood test [gFOBT], multi-targeted stool DNA test [mt-sDNA]) or visual exams of the colon and rectum (i.e., colonoscopy, CT colonography, flexible sigmoidoscopy) [4]. Enrollees in the screened cohort had \geq 1 CPT/ICD-9-CM/ICD-10-CM procedural code during the analysis

year for colonoscopy, mt-sDNA, FIT, gFOBT, CT colonography or flexible sigmoidoscopy, or barium enema. The index year was the last available full calendar year with evidence of screening for the enrollee. Enrollees in the unscreened cohort were a control population of randomly selected enrollees who did not have high-risk conditions for CRC and did not have a claim for a CRC screening test during the analysis year or any previous study year. The index date for enrollees in the unscreened cohort was selected to mirror the month/year distribution of index dates among enrollees in the screened cohort. Enrollees who did not have CRC screening in the year of analysis but had screening in previous years were not included in the analysis.

Provider sample

The provider sample included primary care providers (i.e., family/general practice, internal medicine, obstetrics/gynecology [OB/GYN], geriatrician, nurse practitioner, physician assistant) who were listed on the claims of average-risk patients (i.e., no evidence of high-risk CRC conditions) in the enrollee sample between January 01, 2014 through December 31, 2018. Nurse practitioners and physician assistants were included if they were listed on ≥ 1 claim for a preventive health care service. Each provider was required to have seen ≥ 10 patients from the enrollee sample to ensure the sample was representative of the provider's broader patient caseload of insured patients and to provide a sufficient patient sample for calculating adherence rates unique to each provider.

Study measures

Enrollee-level measures

Enrollee demographic and clinical characteristics were measured during the baseline period and included age, gender, insurance type, geographic region, urbanicity, race/ethnicity, education level, income, net worth, health plan type, indication of consumer-driven healthcare (i.e., health reimbursement arrangement, health savings account), Charlson comorbidity score, and Agency for Healthcare Research and Quality comorbidities.

Screening opportunities were measured based on the enrollee's exposure to the health care system during the baseline year and included having at least one primary or preventive care visit, the presence of a main primary care provider (visits to only one provider or, if multiple providers, visited one provider more than others), and the presence of claims for preventive care visits.

Provider-level measures

Provider characteristics measured during the baseline period included provider specialty, gender, ethnicity, total number of patients in the ORD, and mean number of visit days per patient. Patient characteristics by provider were also measured and included the count of patients with any claim for CRC and the percent of patients by ethnicity category, income and net worth level, rural versus urban living, gender/age category/mean age, geographic region, and influenza vaccination status during the baseline calendar year. The count of patients with a CRC diagnosis was a measure of provider exposure to CRC to determine whether providers who were exposed to patients with CRC would be more adherent to CRC screening guidelines. Influenza vaccination was used as an indicator of general preventive care.

Screening adherence was calculated at the provider level as the percent of average-risk patients who were up to date with screening recommendations each calendar year. A patient was considered up to date if they had evidence of a colonoscopy within 10 years, CT colonography or flexible sigmoidoscopy within 5 years, mt-sDNA within 3 years, or FIT/gFOBT within 1 year.

Analysis

Enrollee-level analysis

Results were stratified by screening status. Bivariate comparisons of baseline characteristics and outcomes measures were calculated using the appropriate test based on the distribution of the measure (e.g., odds ratio [OR], t-test, f-test, Wilcoxon rank-sum, chi-square test).

A logistic regression model was developed to determine the association between receipt of screening and enrollee demographic and clinical characteristics, enrollee socioeconomic status (SES), and provider characteristics. An additional model was also estimated including only women to assess the impact of provider type, specifically OB/GYN, on adherence rates.

Provider-level analysis

In the provider analyses, the cumulative incidence was calculated for each provider to assess the proportion of eligible patients who received CRC screening for each provider, and the characteristics of the provider and their patient panel (as observed in the claims data) were assessed. Patients who saw multiple providers of different types in the selected provider population counted towards each providers' numbers.

An ordinary least squares (OLS) model was estimated to determine the association between screening adherence among the physician's panel of patients and physician demographics, patient demographic and clinical characteristics, and patient SES.

Results

Enrollee-level results

Baseline demographic and clinical characteristics

A total of 664,234 screened and 548,758 unscreened enrollees were included in the enrollee analysis (Fig. 1).



Fig. 1 Health plan enrollee and provider sample selection

Baseline enrollee demographics and clinical characteristics are shown in Fig. 2.

CRC screening and odds of healthcare utilization

Enrollees who were screened had greater odds of having a primary or preventive care visit and having a main primary care provider (Fig. 2). The odds of having ≥ 1 primary care visit or preventive care visit was 4.47 (95% CI=4.42-4.51, p<0.001) and 2.97 (95% CI=2.95-3.00, p<0.001) times greater, respectively, among screened versus unscreened enrollees. Additionally, screened enrollees had 2.69 times greater odds of having a main primary care provider than unscreened enrollees (95% CI=2.67-2.71, p<0.001). Visits to an internal medicine provider or an OB/GYN were associated with 1.22 (95% CI=1.21-1.23, p<0.001) and 1.94 (95% CI=1.91-1.96, p < 0.001) greater odds of being screened, respectively. Conversely, visits to a family practice or advanced practice provider were associated with lower odds of being screened (p < 0.001 for both comparisons).

Multivariate model results were consistent, showing the same general effects found in the descriptive analyses (Fig. 3). Of note, male enrollees were slightly more likely than female enrollees to be screened after adjusting for other characteristics. Having visited a primary care provider or having a preventive care visit in the baseline year were positively associated with screening. The effect of the primary care provider varied across provider type, with higher odds of CRC screening among enrollees who were patients of OB/GYNs and internal medicine providers (Fig. 3 and Supplementary Table 1).

Provider-level results

Provider demographic characteristics

Most providers were family practice (39.1%) or internal medicine (38.0%) and had a mean age of 54.4 years (Table 1). Gender varied by specialty with more female OB/GYNs (57.9%) and advanced practice providers (77.1%) than males. Most providers were in an individual practice, which ranged from 98.3% of advanced practice providers to 90.6% of geriatricians.

Patient demographic characteristics by provider type

Demographic and clinical characteristics of the PCP's patient panels are shown in Table 2. Primary care providers saw a mean of 56.7 average-risk patients and 23.4 high-risk patients during the calendar year (data not shown). Family medicine providers had greater mean average-risk patient counts (62.5) than other providers (p<0.001 for all comparisons) and internal medicine providers had greater high-risk patient counts (26.0) compared to other providers.

The mean cumulative incidence of CRC diagnoses was 0.52% among all primary care providers, with rates ranging from a low of 0.28% among OB/GYNs and a high of 0.77% among geriatricians (Table 3). As a measure of preventive care utilization, the incidence rate of patient



Fig. 2 Unadjusted health plan enrollee demographic and clinical characteristics

receipt of influenza vaccination was 51.1% among all primary care providers (Table 3).

Screening adherence

Among patients with a PCP, mean overall adherence with ACS and USPSTF screening guidelines varied by specialty, ranging from a low of 63.9% among geriatricians to a high of 79.5% among OB/GYNs (Table 3). Compared to family medicine providers, internal medicine, OB/GYN, and advanced practice providers all had significantly higher adherence rates.

In the multivariate model, provider factors associated with screening adherence in the enrollee patient sample included provider age, geographic region, and specialty (Supplementary Table 2). Specific characteristics of the Discussion

This retrospective, claims-based study examined patient and provider factors associated with CRC screening adherence, as defined by ACS and USPSTF guidelines, for average-risk individuals. Enrollees who had a primary care visit, a preventive care visit, or a main primary care provider had the greatest odds of being screened. Additionally, screened enrollees were more likely to be commercially insured, living in the South, residing in an urban setting, and having at least a high school diploma; however, given the large sample sizes and smaller effect

enrollee patient panel were also associated with adher-

ence levels, including an increasing adherence rate with

higher proportions of high-income patients.



Footnotes

Footnotes 1 Additional covariates not shown: index year; other/missing region; other race/ethnicity, missing/unknown race/ethnicity, or no SES information for race/ethnicity, missing/unknown/SES information for race/ethnicity; missing/unknown/SES information for race/ethnicity; missing/unknown/SES information for race/ethnicity; missing/unknown/SES information for net worth; missing urbanicity; multiple/unknown/missing health plan type; missing indicator for or no indication of consumer driven health care; baseline AHRQ measures: thyroid disorders, bening neoplasms, nutritional deficiencies, disorders of lipid metabolism, other nutritional; endocrine; and metabolic disorders, anemia, eye disorders, ear conditions, diseases of versions, other uper respiratory indectiones, other uper respiratory indectiones, uper again and upper lipid metabolism other untritional; endocrine; and metabolic disorders, anemia, eye disorders, ear condition of skin, osteoporosis, other connective tissue disease, other bone disease and musculoskeletal deformities. Abbreviations: OR, dots ratio; Cl. confidence interval; CDHC, consumer driven health care; EPO, exclusive provider organization; HMO, health plan organization; IND, indemnity health plan; POS, point of service health plan; PPO, preferred provider organization

Fig. 3 Logistic regression model of the association between health plan enrollee characteristics and CRC screening status

sizes, differences in screening rates by these characteristics may not be clinically meaningful. Female patients appeared more likely to have been screened versus males; however, after adjustment for other demographic and clinical factors, males were more likely to be screened. In Weiss et al., patient-level predictors of screening included increasing age, White race, being married, primarily English-speaking, having commercial insurance, not having congestive heart failure or diabetes, and utilizing more healthcare resources [15]. Consistent with the results from our study, enrollees who had more contact with the healthcare system were more likely to receive preventive care and CRC screening.

Among PCPs, adherence of the patient panel to ACS and USPSTF screening guidelines varied by specialty and ranged from 69.3% among geriatricians to 79.5% among OB/GYNs. In a previous survey study of primary care providers, 77.5% reported using national CRC screening guidelines, but only 51.7% cited recommendations consistent with those guidelines [23]. Additionally, in Weiss et al., CRC screening rates among primary care clinics ranged from 51 to 80%, where, after controlling for multiple patient and clinical factors, an increasing panel size of eligible patients was the only significant predictor of CRC screening [15]. Screening rates reported in our study were also similar to those obtained through patient

Demographics	Overall Primary (n = 226,534)	Family (n=88,600)	Internal Medicine (n=86,164)	OB/Gyn (n = 23,097)	Advanced Prac- tice (n = 28,035)	Geri- atrician (n=638)
Age, mean (SD)	54.4 (11.2)	55.2 (11.3)	54.4 (11.1)	56.1 (10.9)	49.1 (10.4)	54.1 (9.7)
Male gender, n (%)	111,360 (49.2)	45,524 (51.4)	50,342 (58.4)	9,230 (40.0)	5,945 (21.2)	319 (50.0)
Region, n (%)						
Northeast ¹	40,021 (17.7)	11,671 (13.2)	20,802 (24.1)	4,437 (19.2)	2,968 (10.6)	143 (22.4)
Midwest ²	61,636 (27.2)	26,121 (29.5)	20,195 (23.4)	5,429 (23.5)	9,738 (34.7)	153 (24.0)
South ³	87,052 (38.4)	34,526 (39.0)	31,570 (36.6)	9,786 (42.4)	10,910 (38.9)	260 (40.8)
West ⁴	37,825 (16.7)	16,282 (18.4)	13,597 (15.8)	3,445 (14.9)	4,419 (15.8)	82 (12.9)
Race/ethnicity, n (%)						
White	21,304 (9.4)	8,906 (10.1)	7,181 (8.3)	3,320 (14.4)	1,867 (6.7)	30 (4.7)
African American	494 (0.2)	160 (0.2)	234 (0.3)	73 (0.3)	26 (0.1)	1 (0.2)
Asian	3,590 (1.6)	933 (1.1)	2,252 (2.6)	332 (1.4)	51 (0.2)	22 (3.5)
Hispanic	1,728 (0.8)	651 (0.7)	727 (0.8)	254 (1.1)	88 (0.3)	8 (1.3)
Other	1,033 (0.5)	296 (0.3)	617 (0.7)	91 (0.4)	26 (0.1)	3 (0.5)
Missing	198,385 (87.6)	77,654 (87.7)	75,153 (87.2)	19,027 (82.4)	25,977 (92.7)	574 (90.0)
Practice type, n (%)						
Individual	217,700 (96.1)	83,639 (94.4)	83,355 (96.7)	22,568 (97.7)	27,560 (98.3)	578 (90.6)
Hospital	81 (0.04)	38 (0.04)	38 (0.04)	3 (0.01)	2 (0.01)	0 (0.0)
Group practice	7,730 (3.4)	4,266 (4.8)	2,499 (2.9)	488 (2.1)	419 (1.5)	58 (9.1)
Other facility	786 (0.4)	563 (0.6)	173 (0.2)	17 (0.1)	33 (0.1)	0 (0.0)
Unknown/missing	237 (0.1)	94 (0.1)	99 (0.1)	21 (0.1)	21 (0.1)	2 (0.3)

Table 1 Provider Demographics

¹Connecticut, Massachusetts, Maine, New Hampshire, Rhode Island, Vermont, New Jersey, New York, Pennsylvania

²Illinois, Indiana, Michigan, Ohio, Wisconsin, Iowa, Kansas, Minnesota, Missouri, North Dakota, Nebraska, South Dakota

³Washington DC, Delaware, Florida, Georgia, Maryland, North Carolina, South Carolina, Virginia, West Virginia, Alabama, Kentucky, Mississippi, Tennessee, Arkansas, Louisiana, Oklahoma, Texas

⁴Arizona, Colorado, Idaho, Montana, New Mexico, Nevada, Utah, Wyoming, Alaska, California, Hawaii, Oregon, Washington

surveys of US adults aged 50–75 years using both BRFSS data (71.6%) [17] and NHIS data (67.1%) [18].

The lower screening adherence rate among patients of geriatricians in our study was likely due to the medical complexity of those seeing this provider type. Providers must balance the risk of mortality from CRC against that for other comorbidities. A patient with life expectancy estimated at less than 10 years would not likely experience a mortality benefit from CRC screening. [24, 25] Additional factors such as frailty, cognitive function, and patient priorities also play a role in the decision to screen elderly patients [26]. Several studies have documented higher rates of adverse events during and post colonoscopy that increase with age and the presence of comorbid conditions. [27-29] Patients of geriatricians are typically \geq 65 years of age with complex health care needs and numerous comorbidities that require individualized care beyond that given by a standard family or internal medicine provider.

There are several limitations to this study, including those inherent to claims database analyses. The presence of a diagnosis code on a medical claim does not indicate a positive presence of disease as the diagnosis code may be incorrectly coded or included as rule-out criteria. Information not readily available in the claims data could have affected study outcomes, such as certain clinical and disease parameters. As the claims database included only enrollees with commercial and Medicare Advantage health insurance, the results of this study may not be applicable to the uninsured population or those covered through Medicaid. Uninsured patients face an additional cost barrier and thus, may be less likely to receive screening. The counter-intuitive association between lower education levels and higher rates of CRC screening within this insured population may have been due to confounding with the type of insurance plan available to people in that education category, such as manufacturing jobs with union or contractually ensured levels of health benefits [30]. Information about the plan characteristics were not available for this study. We did not find significant differences in influenza vaccination (a measure of utilization of preventive care measures) by provider type and rates of vaccination were lower than that of CRC screening; however, this may have been an unreliable measure given that influenza vaccinations are frequently received outside of a health plan. Lastly, although advanced practice providers were included as potential providers of primary care, our ability to capture the effect for patients treated by nurse practitioners or physician assistants was limited.

Table 2 Demographic characteristics of patient panels by primary care provider type¹

Demographics	Overall Primary	Family $(n = 88.600)$	Internal Medicine	OB/Gyn (n = 23.097)	Advanced Practice	Geri- atrician
	(n=226,534)	(11 - 00,000)	(n=86,164)	(11-25)0577	(n = 28,035)	(n=638)
Number of patients in each provider's panel ²	252,545	96,784	95,970	25,921	33,159	711
Age	63.0 (4.1)	62.4 (3.9)	64.2 (3.8)	59.9 (3.7)	63.6 (4.1)	66.5 (3.9)
Male gender	39.2 (18.5)	42.7 (13.7)	45.0 (13.8)	2.1 (8.5)	40.7 (15.8)	41.1 (12.0)
Region						
Northeast ³	17.1 (36.6)	12.7 (32.2)	23.5 (41.1)	18.7 (38.1)	10.3 (29.7)	21.3 (39.9)
Midwest ⁴	27.5 (43.4)	29.8 (44.5)	23.8 (41.1)	23.6 (41.5)	35.2 (46.5)	25.3 (42.1)
South⁵	39.0 (47.1)	39.5 (47.2)	37.5 (46.4)	42.8 (48.0)	39.3 (47.3)	40.7 (47.2)
West ⁶	16.3 (35.8)	18.0 (37.4)	15.2 (34.7)	14.9 (34.8)	15.2 (34.8)	12.6 (32.2)
Race/ethnicity						
White	69.6 (22.4)	71.1 (22.7)	66.6 (22.4)	69.7 (21.8)	73.9 (20.3)	62.0 (22.5)
African American	11.2 (16.6)	10.1 (16.4)	13.1 (17.5)	10.4 (15.2)	9.9 (15.5)	15.7 (19.5)
Asian	2.8 (7.4)	2.4 (6.6)	3.4 (8.7)	3.3 (8.0)	1.4 (3.8)	3.8 (9.1)
Hispanic	8.6 (14.5)	8.7 (15.5)	8.9 (13.9)	9.0 (14.5)	6.7 (12.7)	10.2 (14.8)
Other	0.7 (2.1)	0.7 (2.2)	0.8 (2.1)	0.7 (1.9)	0.6 (1.9)	0.8 (2.1)
Missing	7.1 (7.8)	7.0 (7.9)	7.3 (7.5)	6.9 (8.1)	7.4 (7.9)	7.6 (7.6)
Education						
<12 grade	0.4 (2.3)	0.4 (2.6)	0.4 (2.1)	0.3 (1.9)	0.3 (2.1)	0.4 (1.6)
High school diploma	29.5 (23.7)	31.2 (25.5)	28.9 (22.0)	23.1 (21.0)	31.2 (23.8)	30.9 (20.3)
Some college/Associate degree	50.4 (21.4)	51.4 (23.0)	49.1 (19.9)	50.4 (20.0)	51.8 (21.5)	48.8 (18.3)
Bachelor/graduate degree	15.3 (19.2)	12.8 (17.9)	17.1 (20.0)	22.3 (22.3)	12.0 (15.6)	15.2 (17.6)
Education missing	4.4 (7.2)	4.3 (7.4)	4.5 (7.0)	4.0 (7.6)	4.8 (7.3)	4.7 (6.8)
Net worth						
<\$25,000	24.6 (17.2)	25.3 (17.3)	24.9 (17.2)	18.7 (15.4)	26.0 (17.2)	27.3 (17.0)
\$25,000-\$149,999	20.4 (11.1)	21.9 (11.4)	18.7 (10.3)	19.2 (11.4)	21.7 (11.1)	18.5 (9.5)
\$150,000-\$249,999	10.2 (6.8)	10.6 (7.0)	9.6 (6.5)	10.3 (7.0)	10.6 (7.2)	9.6 (6.7)
\$250,000-\$499,999	14.6 (9.3)	14.5 (9.4)	14.3 (9.0)	16.2 (9.2)	14.5 (9.6)	13.8 (9.2)
\$500,000-\$999,999	13.0 (11.3)	12.1 (11.4)	13.4 (11.1)	16.8 (11.9)	11.6 (10.5)	11.7 (10.4)
≥\$1,000,000	7.9 (12.0)	6.3 (10.6)	9.3 (13.2)	11.6 (14.3)	5.8 (8.9)	6.7 (10.1)
Net worth missing	9.4 (9.4)	9.2 (9.4)	9.9 (9.3)	7.3 (8.7)	9.9 (9.8)	12.5 (11.6)
Insurance type						
Commercial	54.2 (33.1)	58.6 (32.5)	46.8 (32.3)	72.5 (27.6)	48.8 (33.4)	31.5 (29.4)
Medicare	45.8 (33.1)	41.4 (32.5)	53.2 (32.3)	27.5 (27.6)	51.2 (33.4)	68.5 (29.4)
Urbanicity						
Rural	6.4 (16.9)	8.5 (21.1)	4.5 (12.2)	4.0 (11.1)	8.1 (17.6)	3.0 (9.2)
Urban	93.5 (16.9)	91.5 (21.1)	95.4 (12.2)	95.9 (11.1)	91.8 (17.6)	96.9 (9.3)
Urbanicity missing	0.1 (1.0)	0.1 (0.9)	0.1 (1.0)	0.1 (1.1)	0.1 (1.3)	0.1 (0.6)

¹Data presented as the mean (SD) of the proportion in each category for each provider's patient panel; age is presented as the mean (SD) of the mean age of the patient panel

²Only includes patients in the Optum Research Database

³Connecticut, Massachusetts, Maine, New Hampshire, Rhode Island, Vermont, New Jersey, New York, Pennsylvania

⁴Illinois, Indiana, Michigan, Ohio, Wisconsin, Iowa, Kansas, Minnesota, Missouri, North Dakota, Nebraska, South Dakota

⁵Washington DC, Delaware, Florida, Georgia, Maryland, North Carolina, South Carolina, Virginia, West Virginia, Alabama, Kentucky, Mississippi, Tennessee, Arkansas, Louisiana, Oklahoma, Texas

⁶Arizona, Colorado, Idaho, Montana, New Mexico, Nevada, Utah, Wyoming, Alaska, California, Hawaii, Oregon, Washington

While United States-focused data sources may not translate directly to other countries' health care systems and populations, similar themes may emerge as additional non-invasive CRC screening methods become available. In a survey-study of European Union countries, screening rates were highest among countries with organized screening programs (from 29.7% in Croatia to 66.7% in the United Kingdom) and those offering both fecal tests and colonoscopy (from 22.7% in Greece to 70.9% in Germany) [31]. A younger age (50–54 years), a longer time since the last physician visit, and a life-style score indicating higher CRC risk were associated with lower utilization of screening tests. Patients who reported not having a physician visit within the previous

	Overall Primary (n = 226,534)	Family (n=88,600)	Internal Medicine (n=86,164)	OB/Gyn (n = 23,097)	Advanced Prac- tice (n = 28,035)	Geri- atrician (n=638)
Overall adherence ²	70.0 (16.7)	68.4 (16.6)	69.2 (16.4)	79.5 (14.5)	69.8 (17.5)	63.9 (18.0)
CRC diagnoses ³	0.5 (1.6)	0.4 (1.2)	0.7 (1.9)	0.3 (1.3)	0.6 (1.9)	0.8 (1.9)
Influenza vaccinations ³	39.2 (18.5)	42.7 (13.7)	45.0 (13.8)	2.1 (8.5)	40.7 (15.8)	41.1 (12.0)

 Table 3
 Cumulative incidence rates (mean [standard deviation]) for provider adherence, CRC diagnoses, and influenza vaccination¹

¹p < 0.001 for all overall primary provider rates and family versus other provider rates

²Assumes 10-year fixed rate

³Values based on %

12 months were 40-60% less likely to have undergone a CRC screening test than patients who had an office visit.

Conclusions

Among PCPs, adherence of the patient panel to ACS and USPSTF screening guidelines ranged from 69-80% depending on the specialty and provider type. The greatest enrollee-level predictors for CRC screening were having a primary or preventive care visit and having a main primary care provider. These results suggest that increased access to preventive/primary care visits, such as via telemedicine, could improve CRC screening rates. More dependence on home-based screening methods that are non-invasive and non-procedural may reduce some of the burden on patients and providers to complete CRC screening.

List of abbreviations

ACS	American Cancer Society
CI	Confidence interval
CPT	Current Procedural Terminology
CRC	Colorectal cancer
FIT	Fecal immunochemical test
gFOBT	Guaiac-based fecal occult blood test
HHS	Health and Human Services
ICD-9-CM/ICD-10-CM	International Classification of Diseases 9th and 10th
	revisions Clinical Modification
mt-sDNA	Multi-targeted stool DNA test
OB/Gyn	Obstetrics/gynecology
OLS	Ordinary least squares
OR	Odds ratio
ORD	Optum Research Database
PCP	Primary care provider
SES	Socioeconomic status
USPSTF	United States Preventive Services Task Force

Supplementary Information

The online version contains supplementary material available at https://doi. org/10.1186/s12913-023-09474-9.

Additional File 1 Supplementary Table 1. Logistic regression model of the association between female health plan enrollee characteristics and CRC screening status

Additional File 2 Supplementary Table 2. Ordinary least squares regression model of the association between the characteristics of the provider and their patient panel and adherence with CRC screening guidelines¹

Acknowledgements

Authors would like to acknowledge Felix Cao, Lynn Wacha, Priyanka S. Koka, Yiyu Fang, Jaycee L. Karl, and Damon L. Van Voorhis for their assistance with programming and construction of the analytic datasets; Timothy Bancroft and Lee Brekke for consultation on the analyses; and Deja Scott-Shemon for medical writing assistance.

Authors' contributions

NEN participated in funding acquisition, conceptualization of the study, methodology, investigation, project administration and supervision, and reviewing and editing the manuscript. LAMW, PL, and DF were involved in conceptualization of the study, interpretation of the study data, and reviewing and editing the manuscript. LL participated in formal analysis and validation, programming, and reviewing and editing the manuscript.

Funding

This work was funded by Exact Sciences Corporation.

Data Availability

The data contained in the Optum Research Database contains proprietary elements owned by Optum and, therefore, cannot be broadly disclosed or made publicly available at this time. The disclosure of this data to third party clients assumes certain data security and privacy protocols are in place and that the third-party client has executed our standard license agreement which includes restrictive covenants governing the use of the data.

Declarations

Competing interests

NMEN and LL are employees of Optum and shareholders in UnitedHealth Group. LAMW is an employee and shareholder of Exact Sciences at the time the study was conducted. PL served as Chief Medical Officer for Screening at Exact Sciences through a contracted services agreement with Mayo Clinic at the time the study was conducted; PL and Mayo Clinic had contractual rights to receive royalties through that agreement. PL is currently an employee of Exact Sciences. DAF has been a consultant with Exact Sciences and Guardant Health. DAF was employed at Duke University at the time of the study and is currently employed by Eli Lilly and Company.

Ethics approval

Institutional review board approval or waiver of approval was not required for this study because the study data were secondary and de-identified in accordance with the United States Department of Health and Human Services (HHS Privacy Rule's requirements for de-identification codified at 45 C.F.R. § 164.514(b). Throughout the process, patient privacy was preserved, and researchers complied strictly with all applicable Health Insurance Portability and Accountability Act data management rules and the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.

Consent for publication

Not applicable

Received: 10 August 2022 / Accepted: 28 April 2023 Published online: 26 May 2023

- References
- 1. American Cancer Society. Key Statistics for Colorectal Cancer. https://www. cancer.org/cancer/colon-rectal-cancer/about/key-statistics.html. Accessed 2/11/2022.
- American Society for Clinical Oncology. Colorectal Cancer: Statistics. https:// www.cancer.net/cancer-types/colorectal-cancer/statistics. Accessed 2/11/2022.
- Siegel RL, Miller KD, Fuchs HE, Jemal A. Cancer statistics, 2022. CA Cancer J Clin. 2022;72(1):7–33. https://doi.org/10.3322/caac.21708.
- 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(4):250–81. https://doi.org/10.3322/caac.21457.
- Force USPST, Davidson KW, Barry MJ, et al. Screening for Colorectal Cancer: US Preventive Services Task Force Recommendation Statement. JAMA. 2021;325(19):1965–77. https://doi.org/10.1001/jama.2021.6238.
- Centers for Disease Control and Prevention. Use of Colorectal Cancer Screening Tests: 2020 Behavioral Risk Factor Surveillance System. https://www.cdc. gov/cancer/colorectal/statistics/use-screening-tests-BRFSS.htm. Accessed 5/19/2022.
- Muthukrishnan M, Arnold LD, James AS. Patients' self-reported barriers to colon cancer screening in federally qualified health center settings. Prev Med Rep. 2019;15:100896. https://doi.org/10.1016/j.pmedr.2019.100896.
- Zhu X, Parks PD, Weiser E, Jacobson DJ, Limburg PJ, Finney Rutten LJ. Barriers to utilization of three colorectal cancer screening options - data from a national survey. Prev Med Rep. 2021;24:101508. https://doi.org/10.1016/j. pmedr.2021.101508.
- Inadomi JM, Vijan S, Janz NK, et al. Adherence to colorectal cancer screening: a randomized clinical trial of competing strategies. Arch Intern Med. 2012;172(7):575–82. https://doi.org/10.1001/archinternmed.2012.332.
- Power E, Miles A, von Wagner C, Robb K, Wardle J. Uptake of colorectal cancer screening: system, provider and individual factors and strategies to improve participation. Future Oncol. 2009;5(9):1371–88. https://doi.org/10.2217/ fon.09.134.
- Steele CBRS, Joseph DA, King JB, Seeff LC. Colorectal cancer incidence and screening - United States, 2008 and 2010. MMWR Suppl. 2013;62:53–60.
- Peterson NB, Murff HJ, Ness RM, Dittus RS. Colorectal cancer screening among men and women in the United States. J Womens Health (Larchmt). 2007;16(1):57–65. https://doi.org/10.1089/jwh.2006.0131.
- Anderson JC, Fortinsky RH, Kleppinger A, Merz-Beyus AB, Huntington CG 3rd, Lagarde S. Predictors of compliance with free endoscopic colorectal cancer screening in uninsured adults. J Gen Intern Med. 2011;26(8):875–80. https:// doi.org/10.1007/s11606-011-1716-7.
- Neilson AR, Whynes DK. Determinants of persistent compliance with screening for colorectal cancer. Soc Sci Med. 1995;41(3):365–74. https://doi. org/10.1016/0277-9536(94)00329-r.
- Weiss JM, Smith MA, Pickhardt PJ, et al. Predictors of colorectal cancer screening variation among primary-care providers and clinics. Am J Gastroenterol. 2013;108(7):1159–67. https://doi.org/10.1038/ajg.2013.127.
- Gimeno Garcia AZ. Factors influencing colorectal cancer screening participation. Gastroenterol Res Pract. 2012;2012:483417. https://doi. org/10.1155/2012/483417.
- 17. Centers for Disease Control and Prevention. Use of colorectal cancer screening tests: 2020 behavioral risk factor surveillance system. Available at https://

www.cdc.gov/cancer/colorectal/statistics/use-screening-tests-BRFSS.htm. Accessed 3/17/2023.

- Cancer Trends Progress Report. National Cancer Institute, NIH, DHHS, Bethesda MD. July 2021. Available at: https://progressreport.cancer.gov. Accessed3/17/2023.
- Baier M, Calonge N, Cutter G, et al. Validity of self-reported colorectal cancer screening behavior. Cancer Epidemiol Biomarkers Prev. 2000;9(2):229–32. https://www.ncbi.nlm.nih.gov/pubmed/10698488.
- Hiatt RA, Perez-Stable EJ, Quesenberry C Jr, Sabogal F, Otero-Sabogal R, McPhee SJ. Agreement between self-reported early cancer detection practices and medical audits among hispanic and non-hispanic white health plan members in northern California. Prev Med. 1995;24(3):278–85. https://doi. org/10.1006/pmed.1995.1045.
- Lipkus IM, Rimer BK, Lyna PR, Pradhan AA, Conaway M, Woods-Powell CT. Colorectal screening patterns and perceptions of risk among africanamerican users of a community health center. J Community Health. 1996;21(6):409–27. https://doi.org/10.1007/BF01702602.
- Bradbury BD, Brooks DR, Brawarsky P, Mucci LA. Test-retest reliability of colorectal testing questions on the Massachusetts behavioral risk factor Surveillance System (BRFSS). Prev Med. 2005;41(1):303–11. https://doi. org/10.1016/j.ypmed.2004.11.015.
- Nodora JN, Martz WD, Ashbeck EL, Jacobs ET, Thompson PA, Martinez ME. Primary care physician compliance with colorectal cancer screening guidelines. Cancer Causes Control. 2011;22(9):1277–87. https://doi.org/10.1007/ s10552-011-9801-0.
- Lin OS, Kozarek RA, Schembre DB, et al. Screening colonoscopy in very elderly patients: prevalence of neoplasia and estimated impact on life expectancy. JAMA. 2006;295(20):2357–65. https://doi.org/10.1001/jama.295.20.2357.
- Inadomi JM, Sonnenberg A. The impact of colorectal cancer screening on life expectancy. Gastrointest Endosc. 2000;51(5):517–23. https://doi.org/10.1016/ s0016-5107(00)70282-3.
- Blaum CS, Rosen J, Naik AD, et al. Feasibility of implementing patient priorities care for older adults with multiple chronic conditions. J Am Geriatr Soc. 2018;66(10):2009–16. https://doi.org/10.1111/jgs.15465.
- 27. Causada-Calo NBK, Albashir S, Al Mazroui A, Armstrong D. Association between age and complications after outpatient colonoscopy. JAMA Netw Open. 2020;3:e208958.
- Day LW, Kwon A, Inadomi JM, Walter LC, Somsouk M. Adverse events in older patients undergoing colonoscopy: a systematic review and metaanalysis. Gastrointest Endosc. 2011;74(4):885–96. https://doi.org/10.1016/j. gie.2011.06.023.
- Warren JL, Klabunde CN, Mariotto AB et al. Adverse events after outpatient colonoscopy in the Medicare population. Ann Intern Med 2009;150(12):849 – 57, W152. DOI: https://doi.org/10.7326/0003-4819-150-12-200906160-00008.
- Congressional Research Service. Job creation in the manufacturing revival. Available at: https://crsreports.congress.gov/product/pdf/R/R41898/22. Accessed 5/24/2022.
- Cardoso R, Guo F, Heisser T, Hoffmeister M, Brenner H. Utilisation of Colorectal Cancer Screening tests in european countries by type of screening offer: results from the european health interview survey. Cancers (Basel). 2020;12(6). https://doi.org/10.3390/cancers12061409.

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