

# Impact of the COVID-19 pandemic on follow-up colonoscopy rates after a positive stool-based screening test for colorectal cancer among U.S. health care organizations

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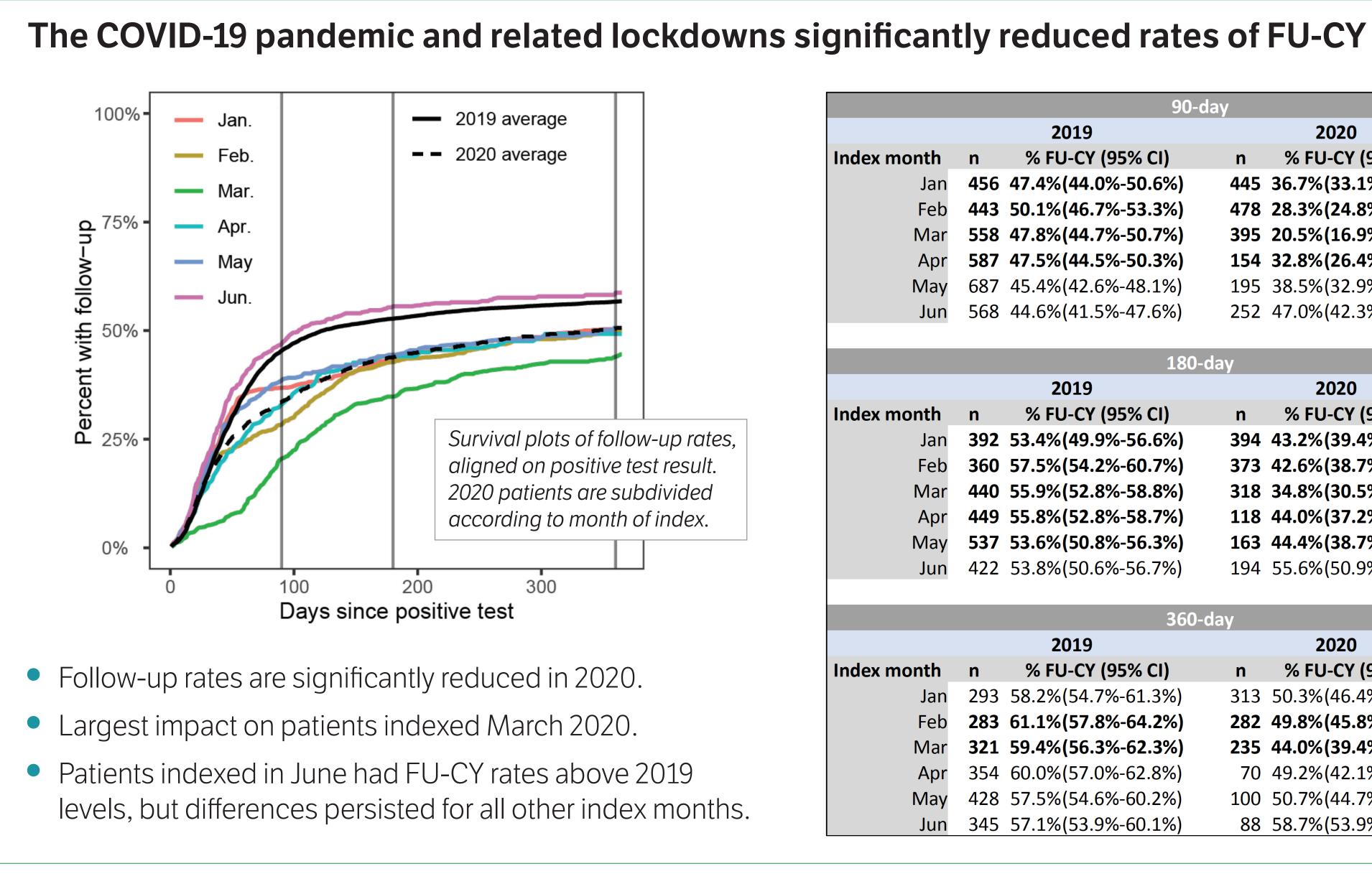
## **Background and Objective**

- The COVID-19 pandemic has disrupted cancer detection, diagnosis, and treatment including colorectal cancer (CRC) screening.<sup>1</sup>
- There was a sharp decline in screening colonoscopies during the COVID-19 pandemic, while utilization of stool-based screening tests (SBTs) such as fecal immunochemical test (FIT) or multitarget stool DNA test (mt-sDNA) increased.<sup>2-3</sup>
- A positive SBT result requires a follow-up colonoscopy (FU-CY) to complete the screening paradigm.
- Simultaneous increased use of SBTs for screening and decrease in colonoscopy accessibility creates a potential care gap if patients fail to follow up after a positive SBT.
- This study evaluated the impact of the COVID-19 pandemic on FU-CY rates within 90, 180, and 360 days of a positive SBT (FIT or mt-sDNA).

# **Study Design**

- A retrospective analysis of de-identified administrative claims and electronic health record data between June 1, 2015, and June 30, 2021, obtained from the Optum Labs Data Warehouse.
- The study population included 14,623 average-risk patients aged 50-75 years old with positive SBT (FIT or mt-sDNA) results in the years 2019 or 2020.
- The index date was the date of the first positive SBT result in 2019 or 2020.
- Patients were included if they had a primary care visit within 15 months prior to the index date and had documented activity at least 90 days after the index date.
- Patients were excluded if they were at higher-than-average CRC risk, had a prior CRC diagnosis, or had recent CRC screening tests prior to the index date.
- Patients were clustered by month of the index date, allowing a comparison of the evolving pandemic impact over time.
- The Kaplan-Meier method was used to compare FU-CY rates at 90, 180, and 360 days of the index date. Patients were censored on death or on the date of last known activity (e.g., visit, prescription, or procedure).
- A difference of differences analysis was used to compare the impact across patient characteristics, including race, Charlson Comorbidity Index (CCI), and insurance type.

# **Principal Findings**



#### FU-CY rates disproportionately impacted certain patient subpopulations Patient Race Charlson Comorbidity Index (CCI) African American Caucasiar σ 20 % 100 200 300 200 Time from index (days) Time from index (days)

- Rates of follow-up over time are compared for patients indexed in 2020 vs 2019 (absolute change).
- No significant difference across African American and Caucasian patients (other patient races had insufficient data in 2020).
- Patients with 1-4 CCI (likely those with 1-2 chronic conditions) had the largest drop-off and the least recovery in FU-CY rates as compared to patients in other CCI categories.
- Medicare patients were overall less impacted than commercially insured patients, but this difference was small.
- No subpopulation recovered to 2019 rates within one year.

90-day										
		2019		2020						
ndex month	n	% FU-CY (95% CI)	n	% FU-CY (95% CI)	p-value					
Jan	456	47.4%(44.0%-50.6%)	445	36.7%(33.1%-40.2%)	0.001					
Feb	443	50.1%(46.7%-53.3%)	478	28.3%(24.8%-31.7%)	<0.001					
Mar	558	47.8%(44.7%-50.7%)	395	20.5%(16.9%-24.0%)	<0.001					
Apr	587	47.5%(44.5%-50.3%)	154	32.8%(26.4%-38.6%)	0.001					
May	687	45.4%(42.6%-48.1%)	195	38.5%(32.9%-43.6%)	0.087					
Jun	568	44.6%(41.5%-47.6%)	252	47.0%(42.3%-51.3%)	0.534					

180-day									
		2019		2020					
ndex month	n	% FU-CY (95% CI)	n	% FU-CY (95% CI)	p-value				
Jan	392	53.4%(49.9%-56.6%)	394	43.2%(39.4%-46.7%)	0.004				
Feb	360	57.5%(54.2%-60.7%)	373	42.6%(38.7%-46.3%)	<0.001				
Mar	440	55.9%(52.8%-58.8%)	318	34.8%(30.5%-38.9%)	<0.001				
Apr	449	55.8%(52.8%-58.7%)	118	44.0%(37.2%-50.2%)	0.022				
May	537	53.6%(50.8%-56.3%)	163	44.4%(38.7%-49.6%)	0.039				
Jun	422	53.8%(50.6%-56.7%)	194	55.6%(50.9%-59.9%)	0.671				
360-day									
		2019		2020					
ndex month	n	% FU-CY (95% CI)	n	% FU-CY (95% CI)	p-value				
Jan	293	58.2%(54.7%-61.3%)	313	50.3%(46.4%-53.9%)	0.053				
Feb	283	61.1%(57.8%-64.2%)	282	49.8%(45.8%-53.5%)	0.007				
Mar	321	59.4%(56.3%-62.3%)	235	44.0%(39.4%-48.3%)	<0.001				

70 49.2%(42.1%-55.4%) 0.094

100 50.7% (44.7%-56.0%) 0.215

88 58.7%(53.9%-63.0%) 0.783

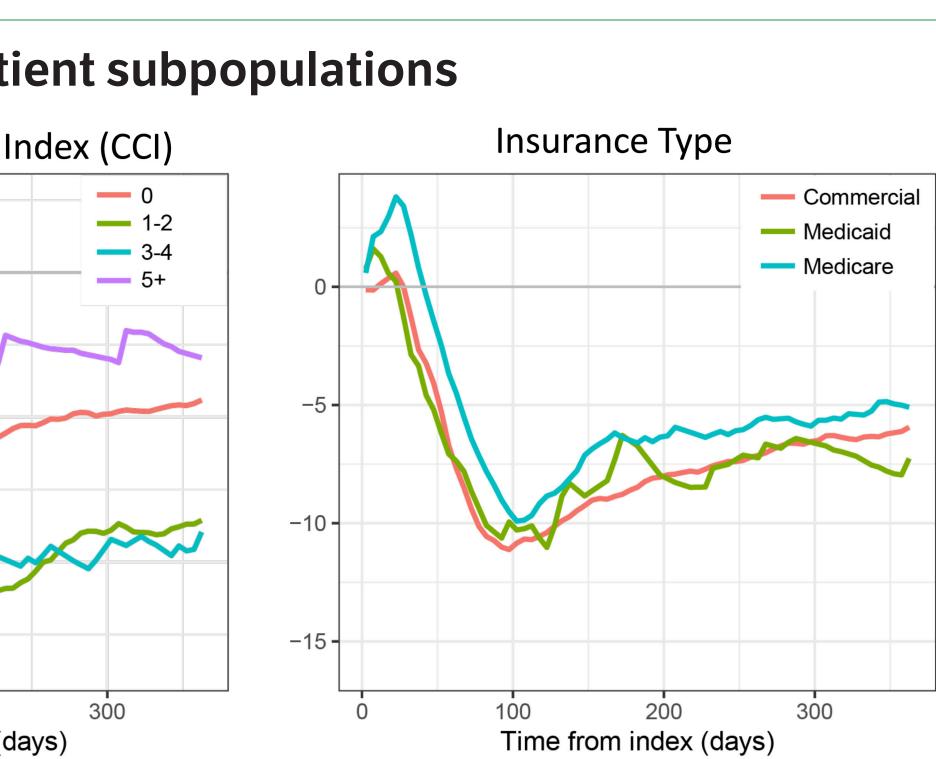
354 60.0% (57.0%-62.8%)

428 57.5% (54.6%-60.2%)

Jun 345 57.1%(53.9%-60.1%)

Apr

May



### 50% of patients with a positive SBT CRC screening test result in 2020 failed to complete a follow-up colonoscopy within 1 year.

- years.

### Patients at higher risk of mortality had lower FU-CY rates during the pandemic

### References

# **Implications for Policy and Practice**

 Our study demonstrates the long-lasting impact of the COVID-19 pandemic on follow-up colonoscopy rates after a positive SBT result.

 Patients with positive SBT had much lower rates of follow-up in 2020 relative to 2019, which created a potential backlog of patients at high risk of CRC that must be addressed.

 Patients and health systems utilize SBTs for initial CRC screening as a convenient way to increase population-level screening rates; however, a lack of follow-up after a positive SBT during and after the COVID-19 pandemic needs to be addressed.

# Conclusions

This failure rate was approximately 10% higher than the prior

 Patients with a positive screening test in March were disproportionately affected and never recovered to the same (lower) level as other months, potentially due to the disruption at the initial scheduling stage.

• By June 2020, FU-CY rates recovered to 2019 levels.

 A higher risk of mortality (as measured by CCI) was associated with lower FU-CY rates, perhaps due to difficulty in providing care to more complex patients.

• Overall, the differential impact of the pandemic across patient groups was modest, though this issue is separate from overall disparities in screening rates.

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Stool-based Colorectal Cancer Screening in the COVID-19 Era. Available at https://blogs. cdc.gov/genomics/2021/04/12/stool-based-colorectal/. Accessed May 3, 2022. . Mazidimoradi A, T. A. (2021). Impact of the COVID-19 Pandemic on Colorectal Cancer Screening: a Systematic Review. J Gastrointest Cancer, 18:1–15.

3. Myint A, R. L. (2021). Noninvasive Colorectal Cancer Screening Tests Help Close Screening Gaps During Coronavirus Disease 2019 Pandemic. Gastroenterology, Aug;161(2):712-714.e1.