

Colorectal Cancer Screening and Quality Improvement in Perioperative Care

Howard S. Kaufman, MD, MBA, FACS

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Huntington Cancer Center, an Affiliate of Cedars-Sinai Cancer
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State Chair, Zone S, American College of Surgeons Commission on Cancer

September 21, 2024 L.A. Care Quality Improvement Conference
Hilton Woodland Hills, CA

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- Medtronic – Consultant, Advisory Board; ROMTech – Investor; and Pacira Biosciences – Consultant. Dr. Howard Kaufman is a consultant and on the Advisory Board of Medtronic; an investor of ROMTech and a consultant for Pacira Biosciences.

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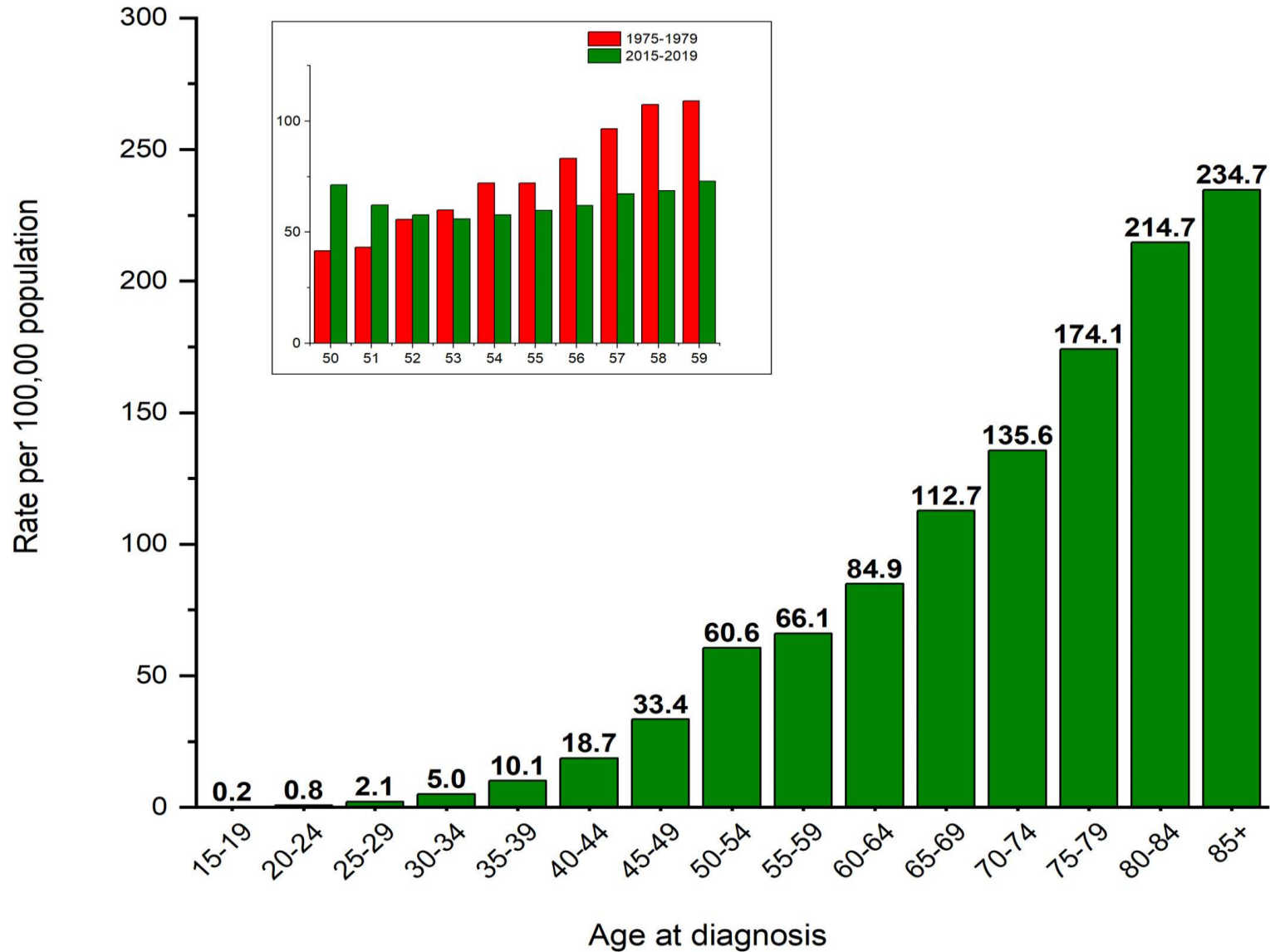
1. Summarize options for colorectal cancer screening.
2. Identify differences and disparities in colorectal cancer incidence and mortality by sex, race, and ethnicity.
3. Apply at least two (2) strategies to eliminate disparities in colorectal cancer screening outcomes.
4. Identify components of Enhanced Recovery Pathways that are associated with reduced length of stay in patients undergoing minimally invasive colorectal surgery.

- Colorectal cancer burden
- Age, ethnic, and racial disparities
- Screening – recommendations and options
- Reducing disparities in colorectal cancer screening and outcomes
- Prevention

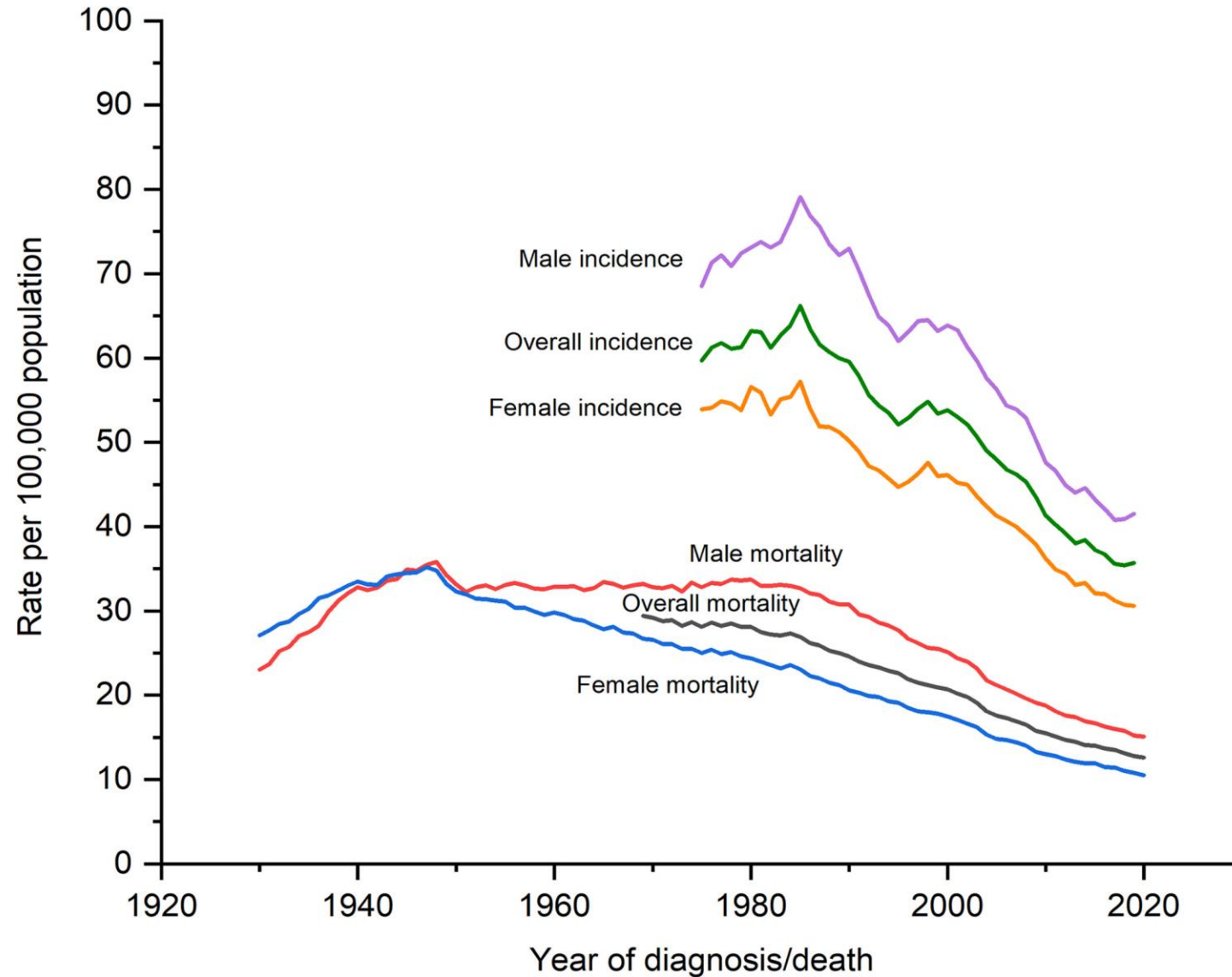
- QI in colorectal surgery
 - Minimally invasive surgery (MIS)
 - Enhanced Recovery Pathways (ERP)
 - Enhanced Recovery after Surgery (ERAS)

- 3rd most common cancer in US
- 2nd leading cause of cancer death
- Incidence - 2024
 - Colon cancer – 106,590
 - Rectal cancer – 46,220
- Deaths – 53,010
- Adults born ~1990 have 2X risk of colon cancer and 4X risk of rectal cancer than those born in 1950

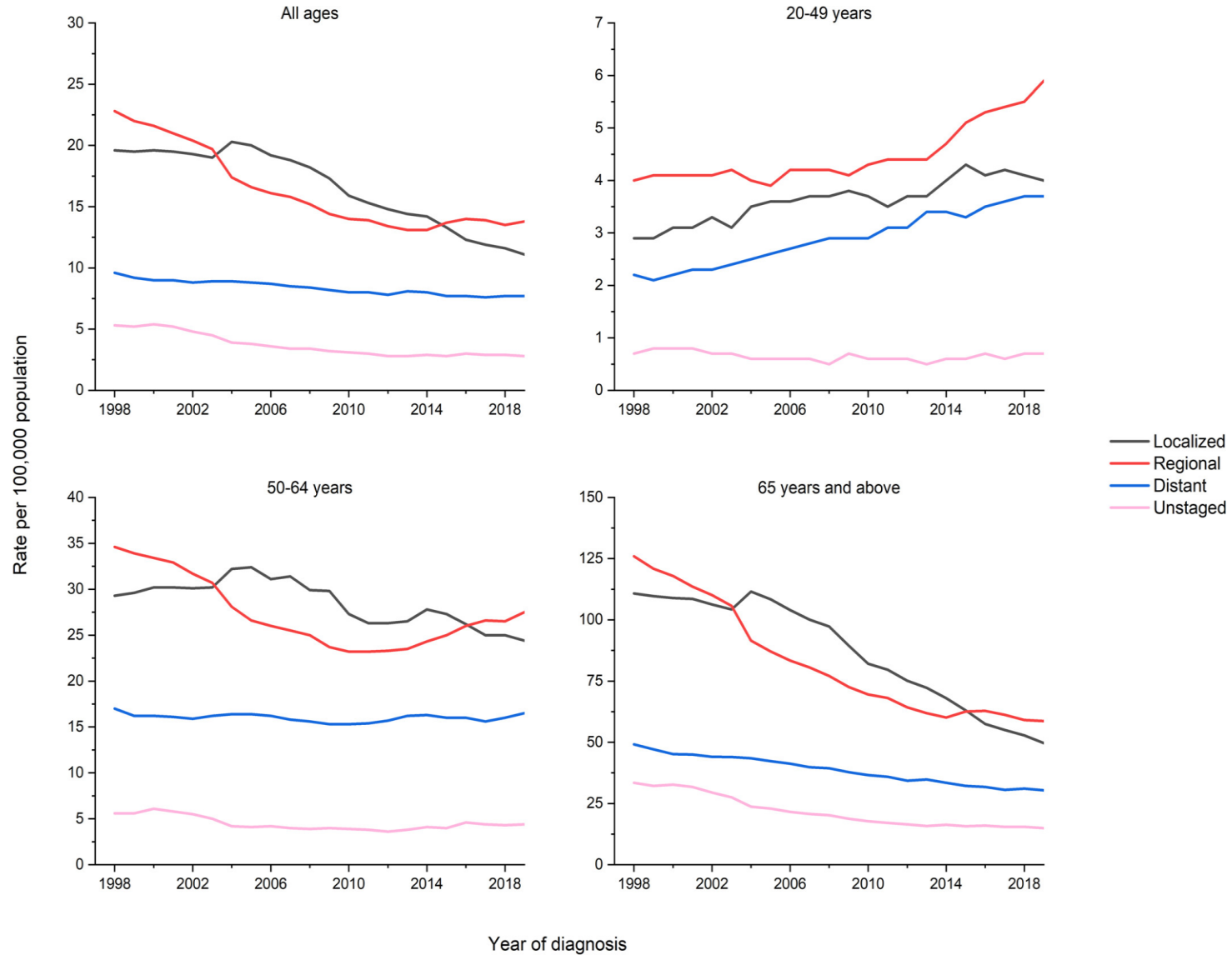
Colorectal Cancer Statistics - 2023



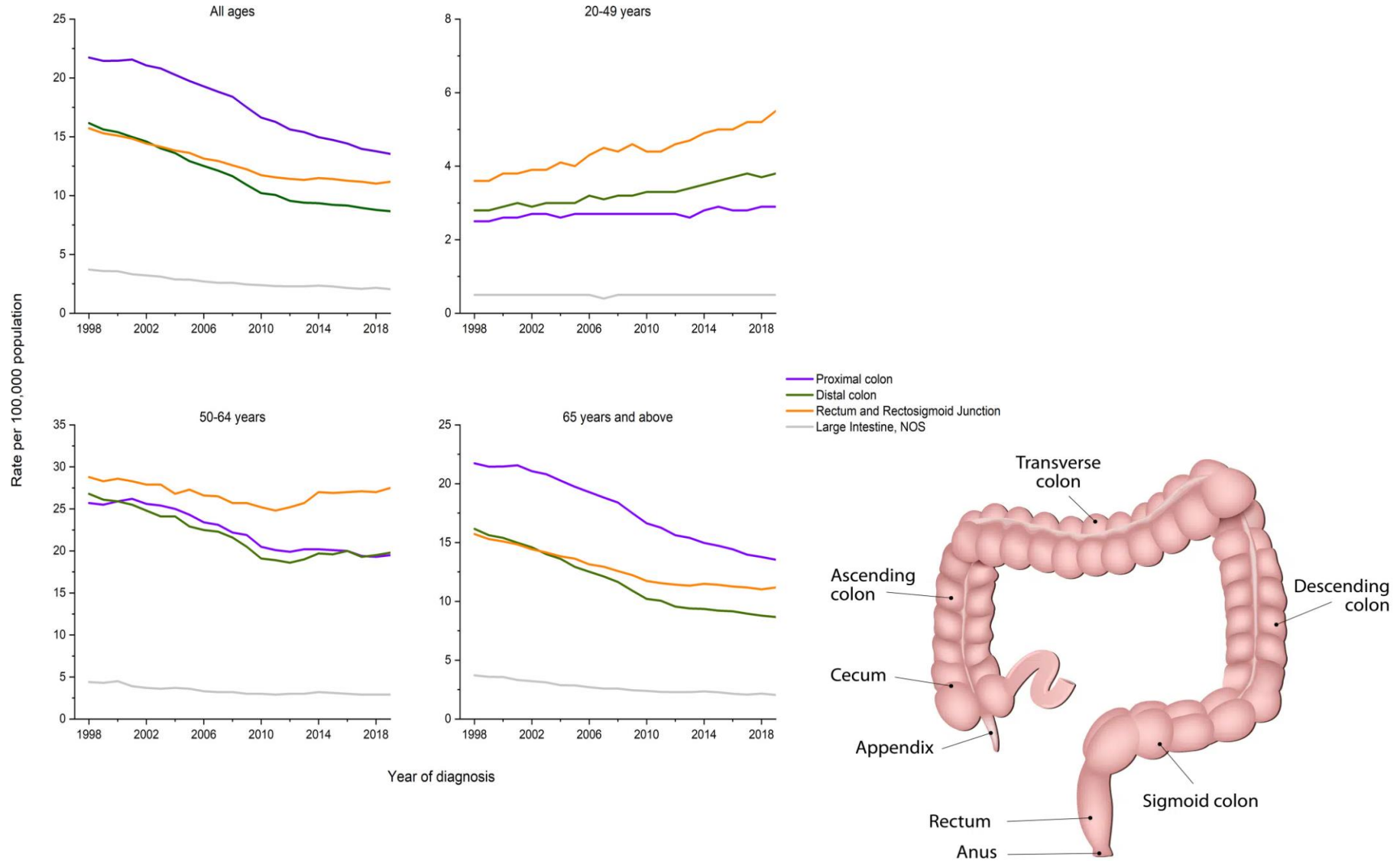
Trends in Incidence and Mortality



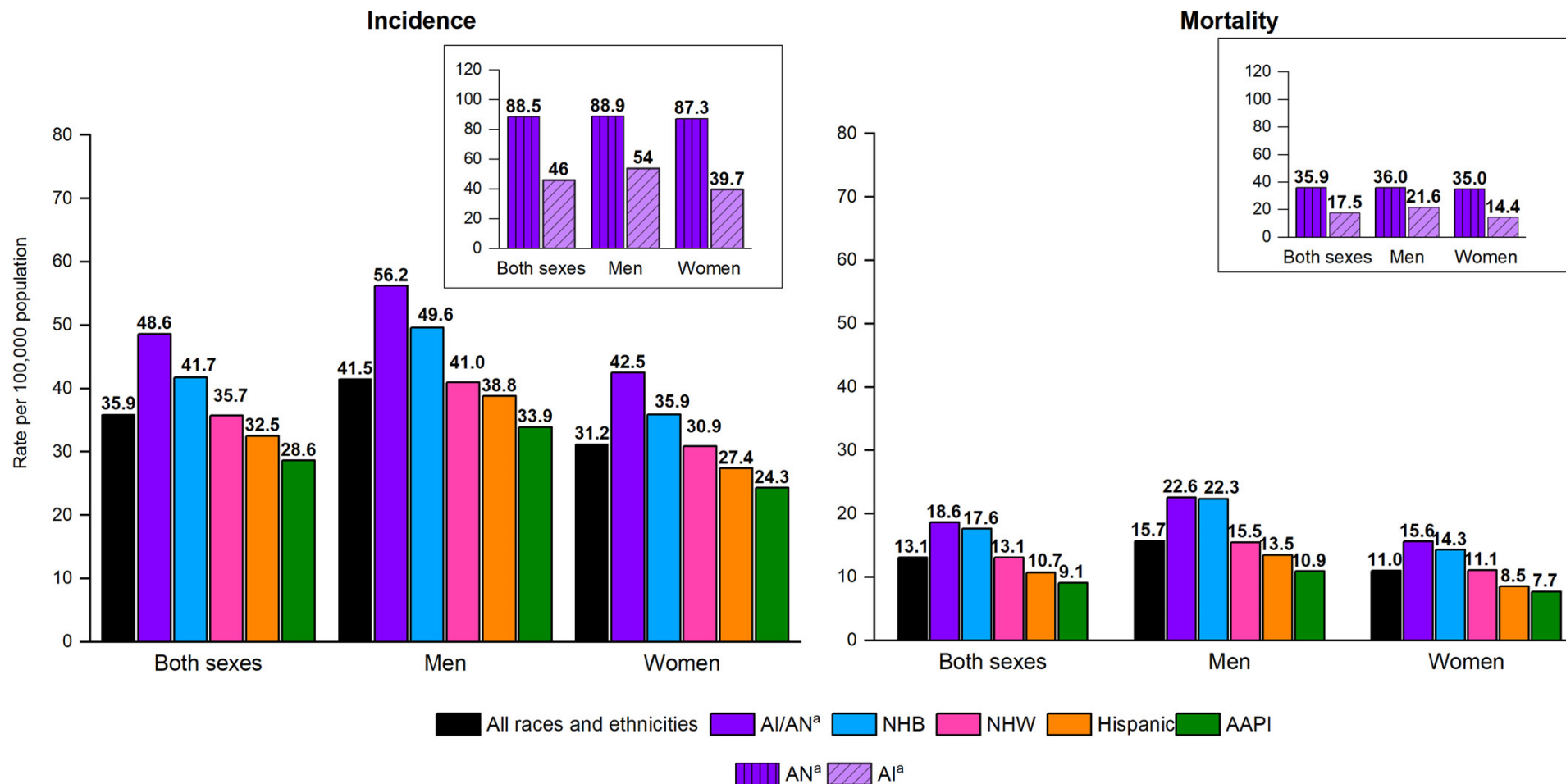
Trends in CRC by Age and Stage



Trends in CRC by Age and Subsite

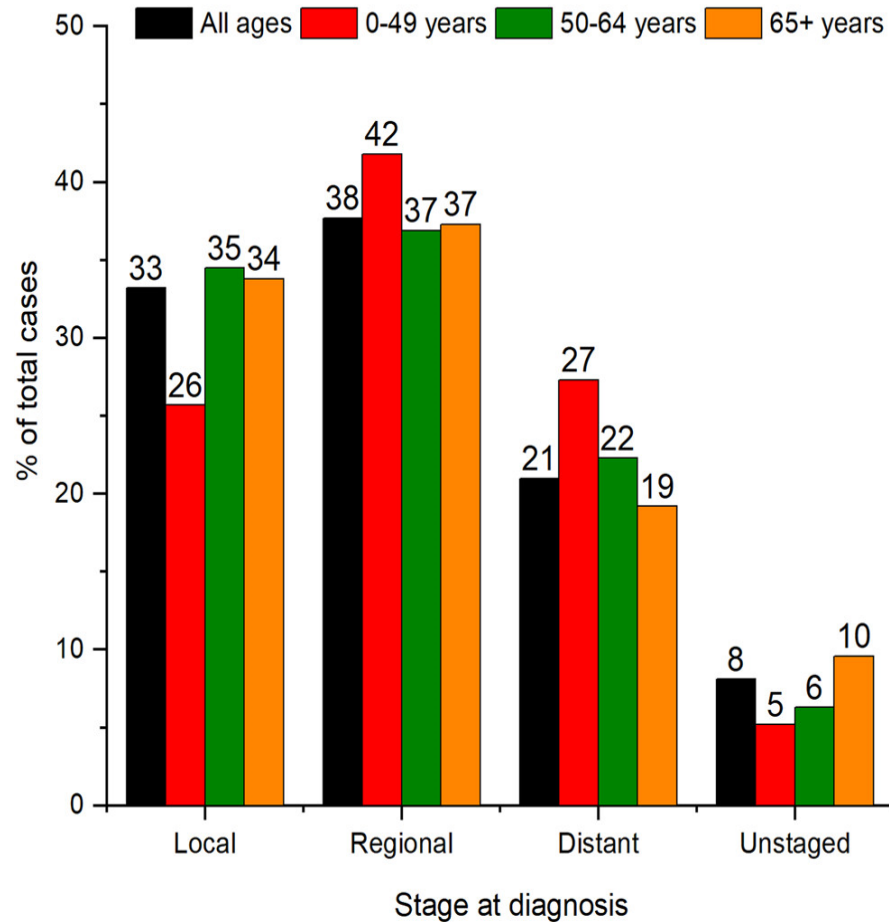


CRC Incidence and Mortality by Sex, Race, Ethnicity

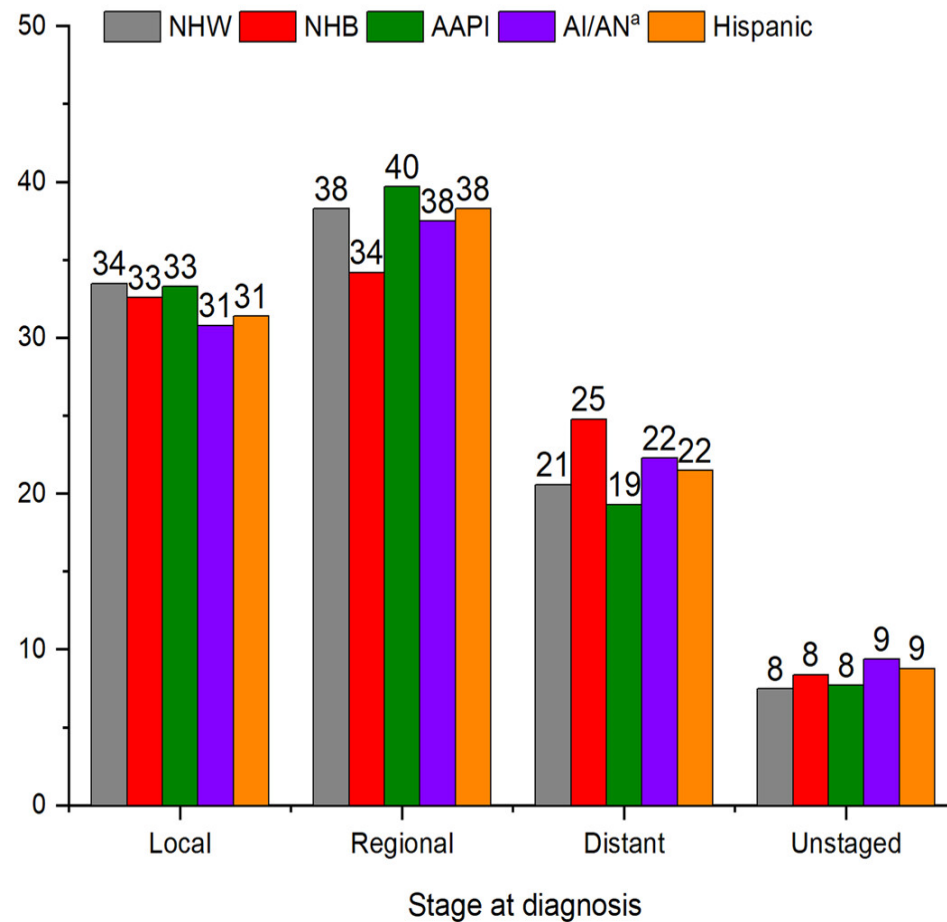


CRC Stage Distribution by Age, Race, and Ethnicity (2015-2019)

A. By age



B. By race/ethnicity



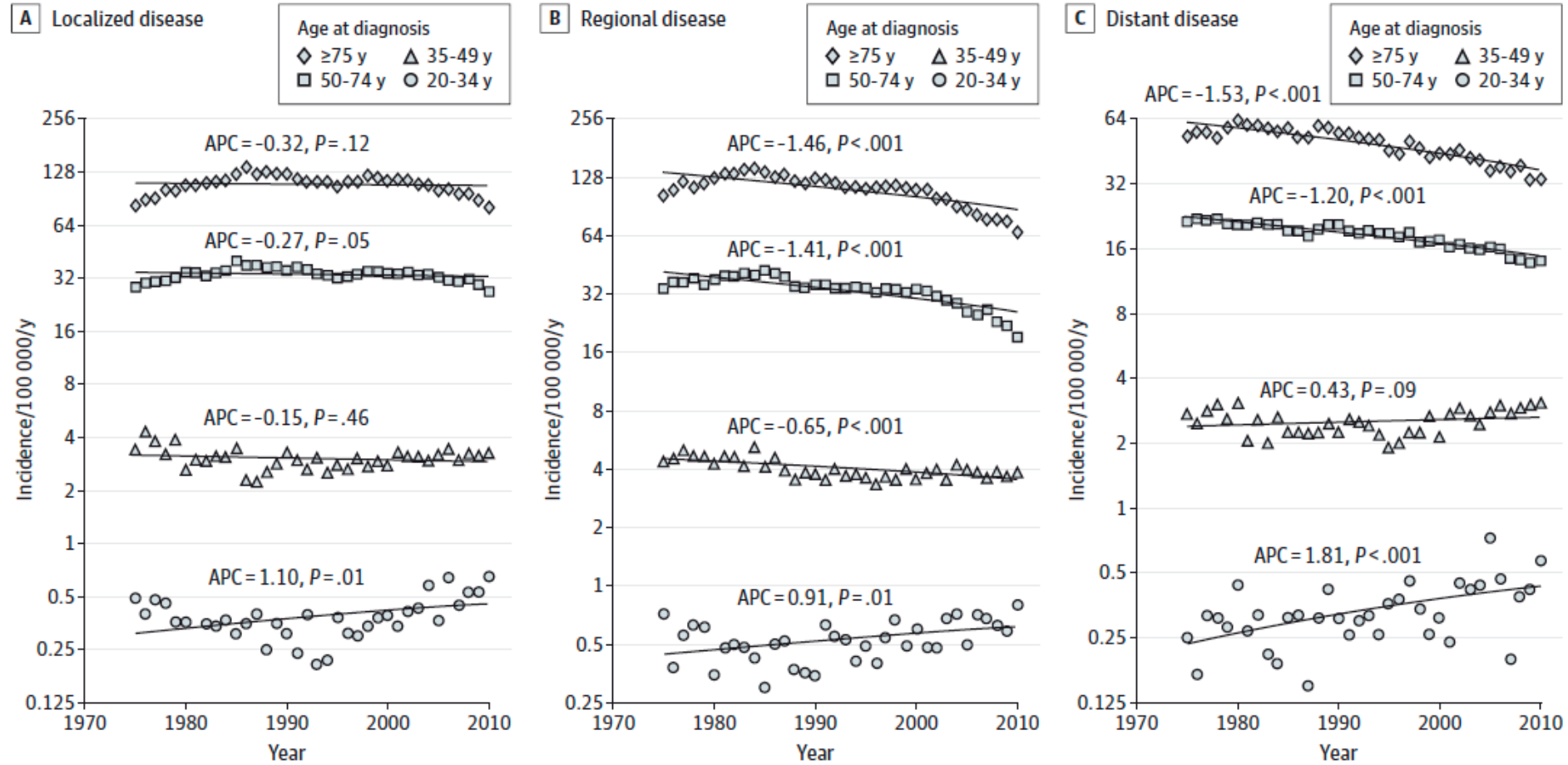
Original Investigation

Increasing Disparities in the Age-Related Incidences of Colon and Rectal Cancers in the United States, 1975-2010

Christina E. Bailey, MD, MSCI; Chung-Yuan Hu, MPH, PhD; Y. Nancy You, MD, MHSc; Brian K. Bednarski, MD; Miguel A. Rodriguez-Bigas, MD; John M. Skibber, MD; Scott B. Cantor, PhD; George J. Chang, MD, MS

Increasing Age Disparities in CRC

Figure 1. Annual Incidence Rates of Colon Cancer From 1975 to 2010



Rates are per 100 000 and age adjusted to the 2000 US standard population for localized disease (A), regional disease (B), and distant disease (C). The trend lines are logarithmic. APC indicates annual percentage change.

Figure 2. Annual Percentage Change–Based Predicted Incidence Rates of Colon Cancer by Age Compared With Incidence Rate in 2010

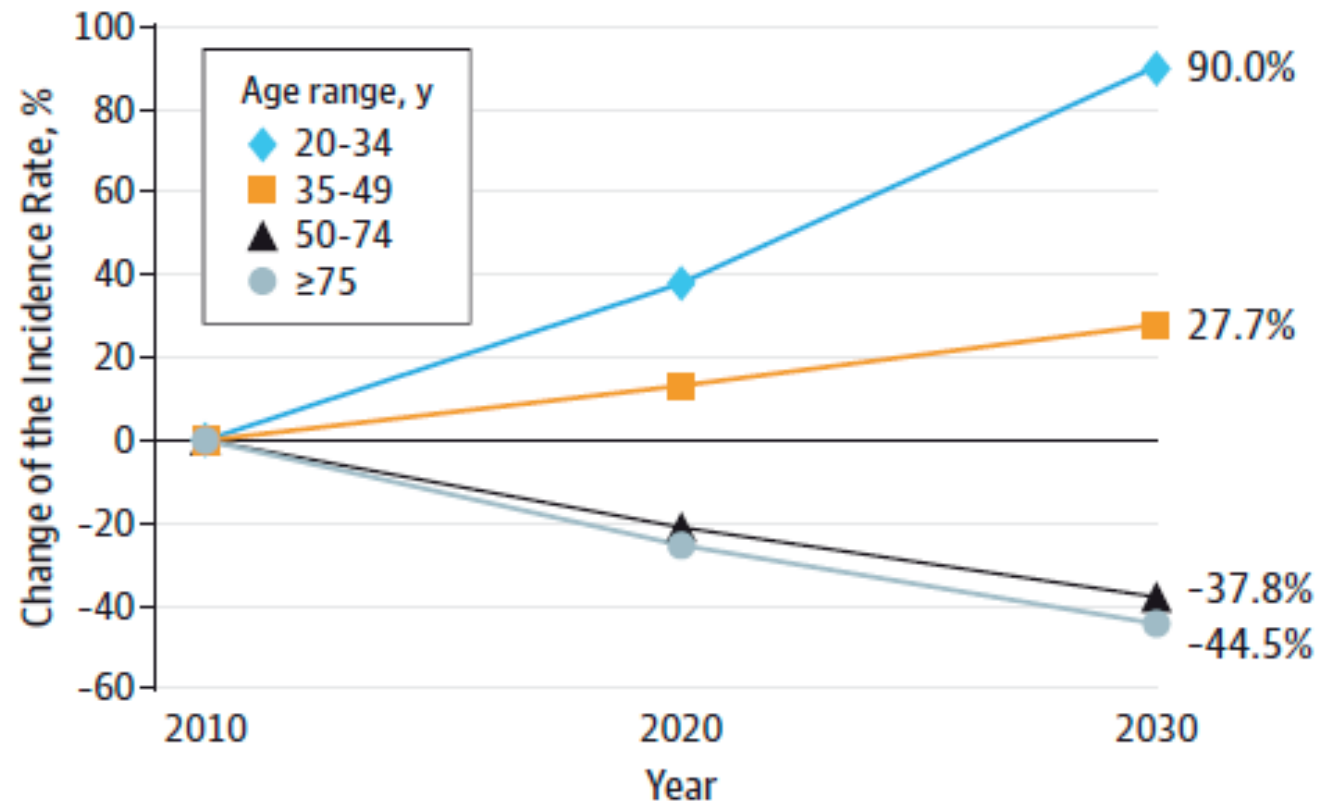
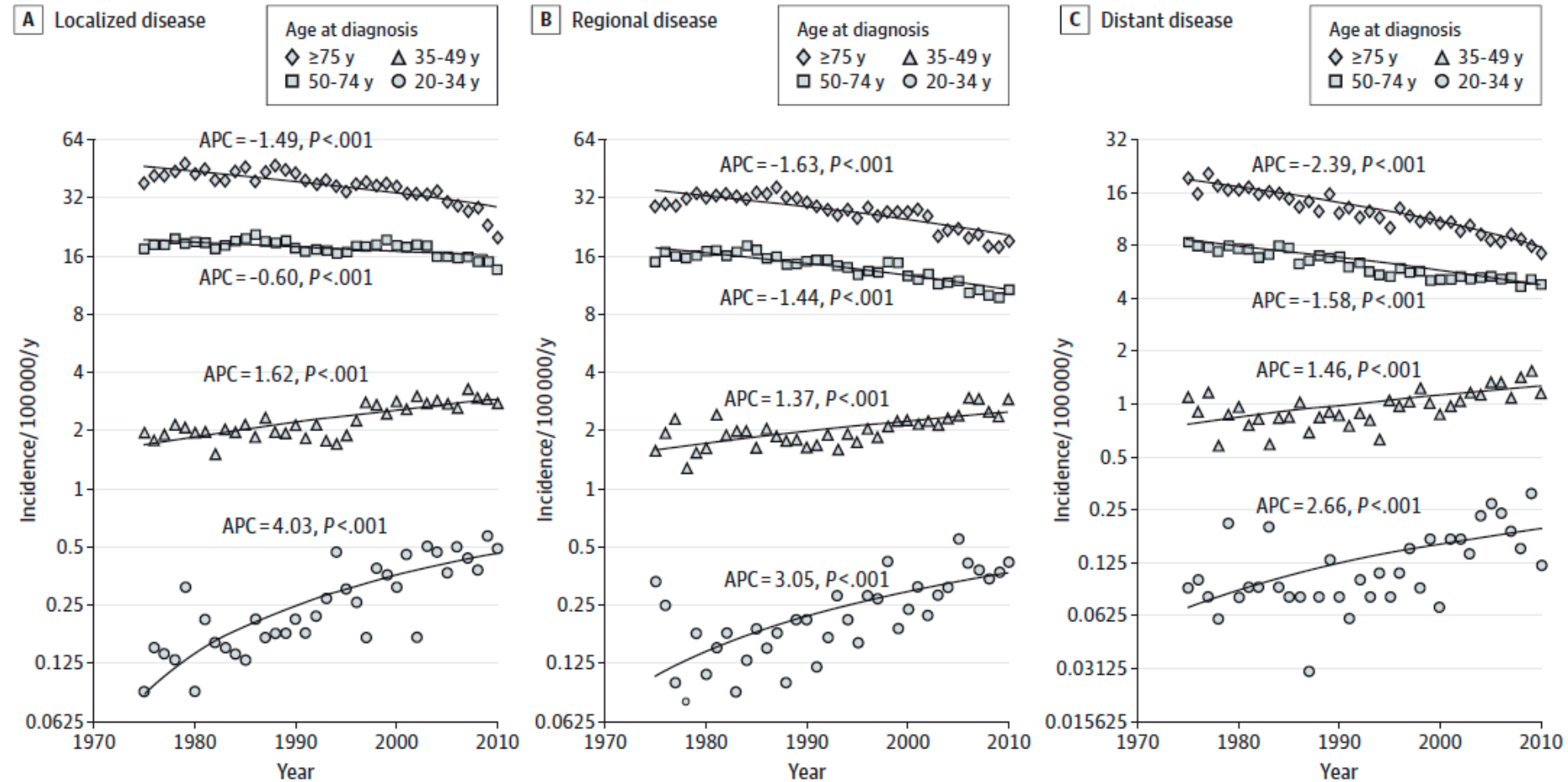


Figure 3. Annual Incidence Rates of Rectosigmoid and Rectal Cancers From 1975 to 2010



Rates are per 100 000 and age adjusted to the 2000 US standard population for localized disease (A), regional disease (B), and distant disease (C). The trend lines are logarithmic. APC indicates annual percentage change.

Figure 4. Annual Percentage Change–Based Predicted Incidence Rates of Rectosigmoid and Rectal Cancers by Age Compared With Incidence Rate in 2010

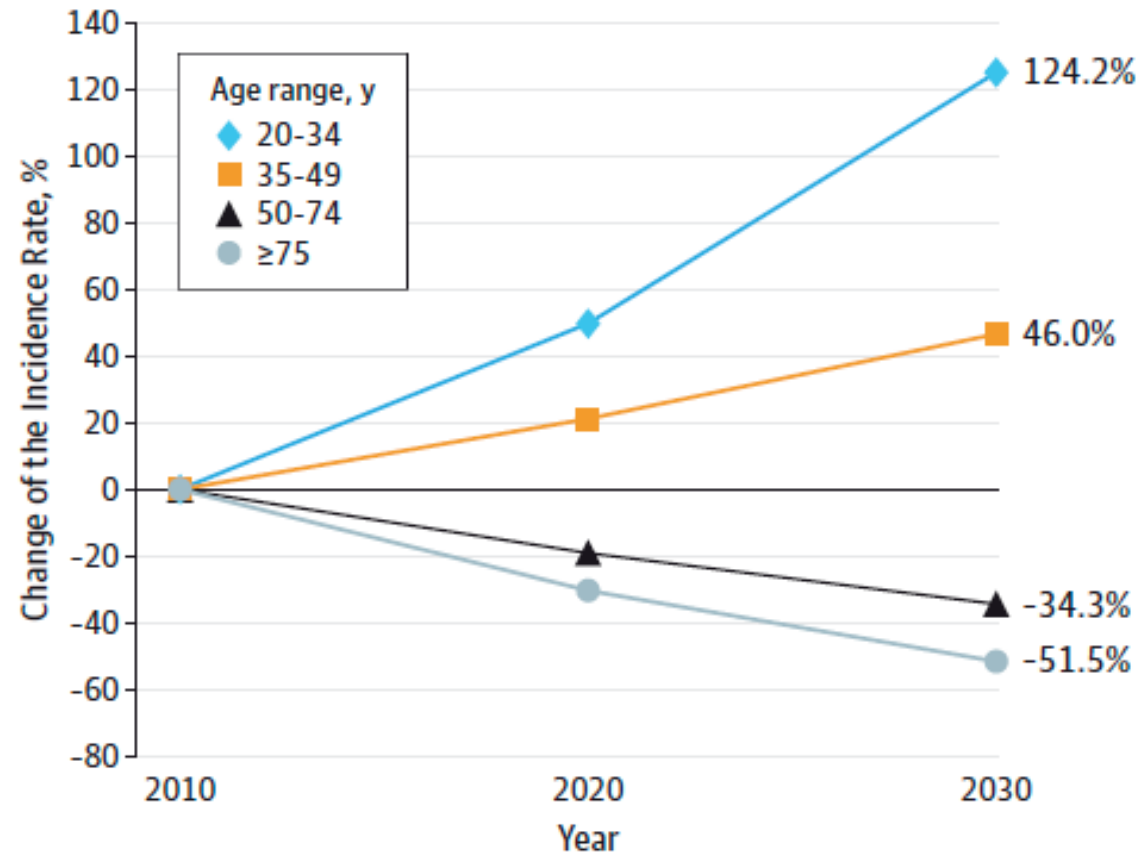


Figure 2. Annual Percentage Change–Based Predicted Incidence Rates of Colon Cancer by Age Compared With Incidence Rate in 2010

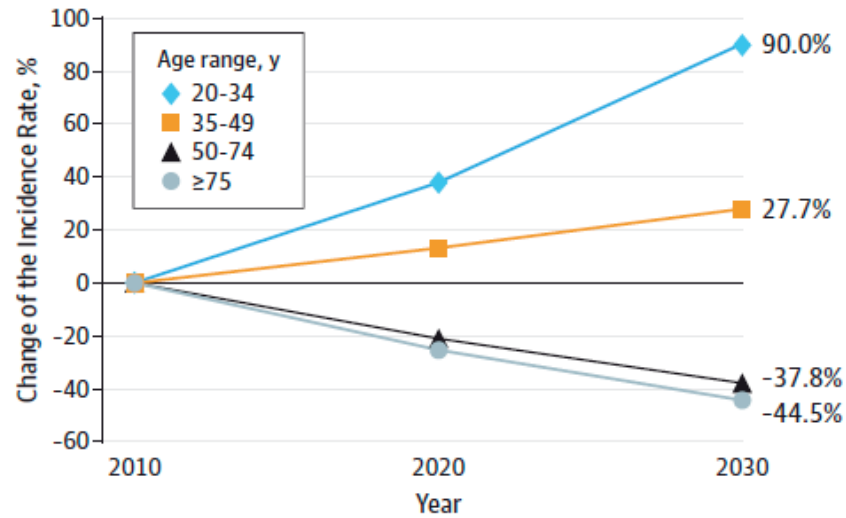
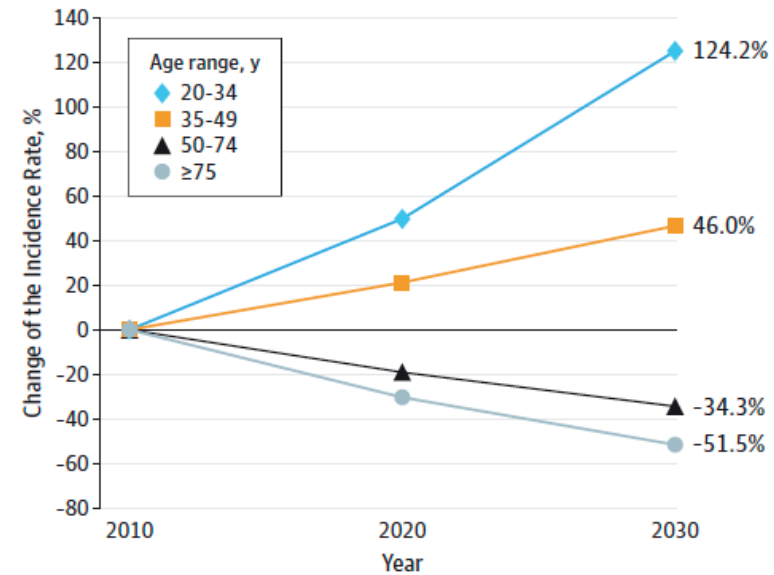
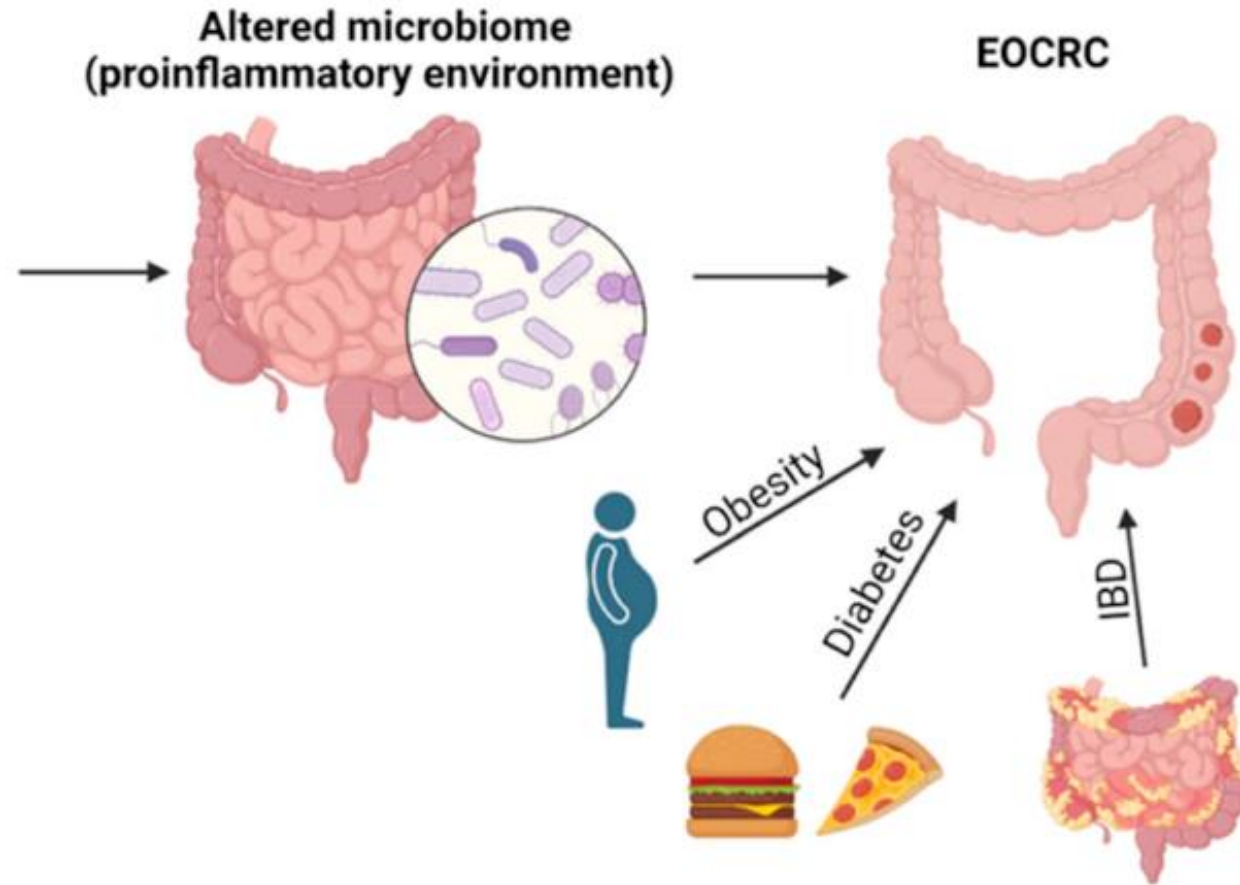


Figure 4. Annual Percentage Change–Based Predicted Incidence Rates of Rectosigmoid and Rectal Cancers by Age Compared With Incidence Rate in 2010



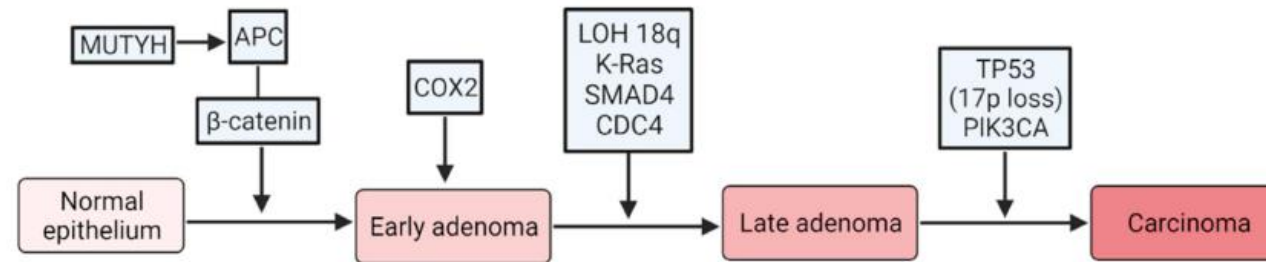
Risk Factors for EOCRC

- Western diet
- Red and processed meats
- Synthetic dyes
- High-fructose corn syrup
- Smoking & alcohol
- Physical inactivity
- Antibiotic exposure
- Genetic (hereditary cancer syndromes)

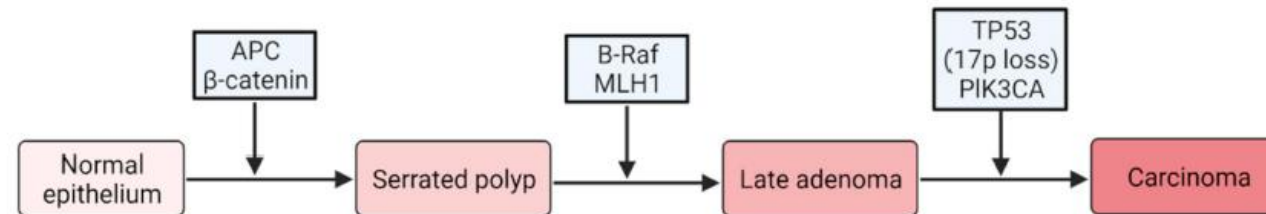


Molecular Subtypes in CRC

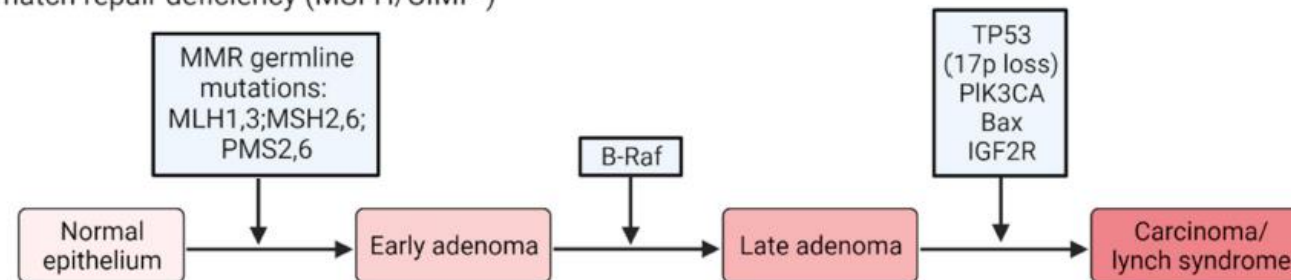
- a Adenoma-carcinoma sequence: 70-90% CRC
(chromosomal instability pathway (CIN)/CIMP-)



- b Serrated polyp pathway: 10-20% CRC
(CpG-island methylator (CIMP)/MSS/MSI-H)



- c Microsatellite instability pathway: 2-7% CRC
(Mismatch repair deficiency (MSI-H/CIMP-)



Colorectal Cancer Screening for Average-Risk Adults: 2018 Guideline Update From the American Cancer Society

Andrew M. D. Wolf, MD¹; Elizabeth T. H. Fontham, MPH, DrPH²; Timothy R. Church, PhD³; Christopher R. Flowers, MD, MS⁴; Carmen E. Guerra, MD⁵; Samuel J. LaMonte, MD⁶; Ruth Etzioni, PhD⁷; Matthew T. McKenna, MD⁸; Kevin C. Oeffinger, MD⁹; Ya-Chen Tina Shih, PhD¹⁰; Louise C. Walter, MD¹¹; Kimberly S. Andrews, BA¹²; Otis W. Brawley, MD¹³; Durado Brooks, MD, MPH¹⁴; Stacey A. Fedewa, PhD, MPH¹⁵; Deana Manassaram-Baptiste, PhD, MPH¹⁶; Rebecca L. Siegel, MPH¹⁷; Richard C. Wender, MD¹⁸; Robert A. Smith, PhD¹⁹

People at average risk

- Men and women should start regular screening at **age 45**
- People who are in good health and with a life expectancy of more than 10 years should continue regular colorectal cancer screening through **age 75**
- For people **ages 76 through 85**, the decision to be screened should be based on their preferences, life expectancy, overall health, and prior screening history
- People over **age 85** should no longer get colorectal cancer screening

Test Options for Colorectal Cancer Screening

Visual exams:

- Colonoscopy every 10 years, OR
- CT colonography (virtual colonoscopy)* every 5 years, OR
- Flexible sigmoidoscopy* every 5 years

* If a person chooses to be screened with a test other than colonoscopy, any abnormal test result should be followed up with colonoscopy.

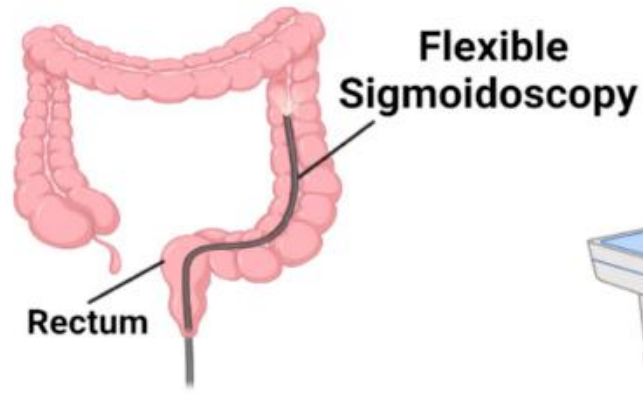
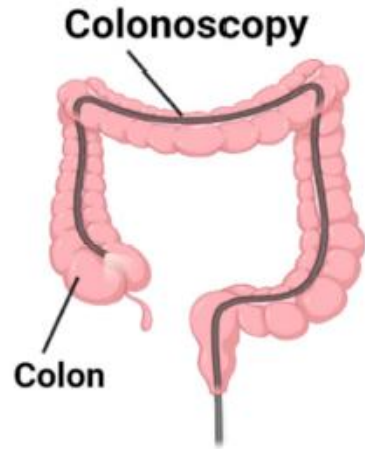
Test Options for Colorectal Cancer Screening

Stool-based tests:

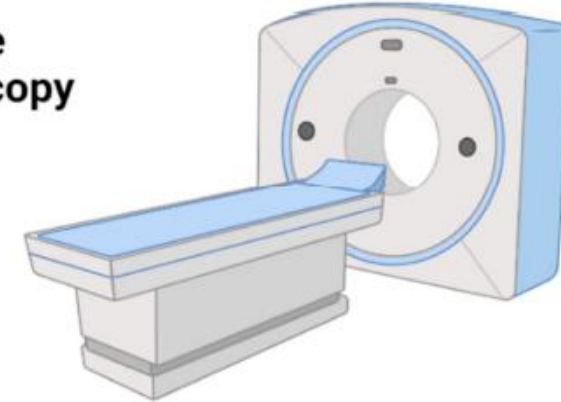
- Highly sensitive fecal immunochemical test (FIT)* every year, OR
- Highly sensitive guaiac-based fecal occult blood test (gFOBT)* every year, OR
- Multi-targeted stool DNA test (MT-sDNA)* every 3 years

* If a person chooses to be screened with a test other than colonoscopy, any abnormal test result should be followed up with colonoscopy.

Screening Options



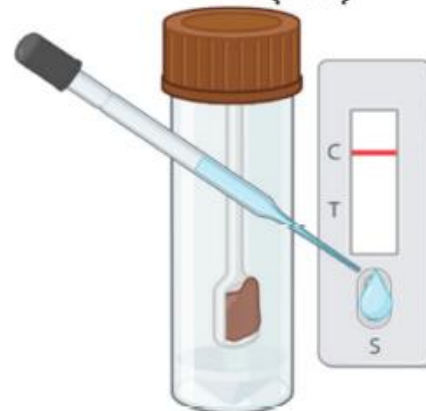
CT colonography



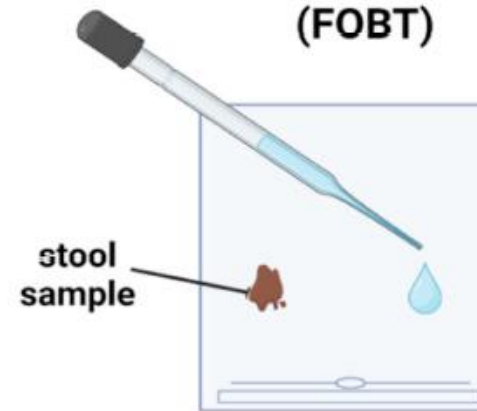
Stool DNA test



Fecal Immunochemical test (FIT)



Fecal occult blood test (FOBT)





Colorectal Cancer: Screening

May 18, 2021

Recommendation Summary

Population	Recommendation	Grade
Adults aged 50 to 75 years	<p>The USPSTF recommends screening for colorectal cancer in all adults aged 50 to 75 years.</p> <p>See the "Practice Considerations" section and Table 1 for details about screening strategies.</p>	A
Adults aged 45 to 49 years	<p>The USPSTF recommends screening for colorectal cancer in adults aged 45 to 49 years.</p> <p>See the "Practice Considerations" section and Table 1 for details about screening strategies.</p>	B
Adults aged 76 to 85 years	<p>The USPSTF recommends that clinicians selectively offer screening for colorectal cancer in adults aged 76 to 85 years. Evidence indicates that the net benefit of screening all persons in this age group is small. In determining whether this service is appropriate in individual cases, patients and clinicians should consider the patient's overall health, prior screening history, and preferences.</p>	C

Screening Test Characteristics

Test	Sensitivity for CRC	Sensitivity for adv. Adenoma	Specificity for CRC	Evidence	Risk	Deaths averted per 1000 screened
High sensitivity guaiac FOBT	62-79%	7%	87%-96%	Strong	Low	26
FIT	76-95%	27%-47%	89%-96%	Weak	Low	26
FIT-DNA (Cologuard)	93%	43%	85%	Early	Low	28 (yearly) 25 (Q 3 years)
CT Colonography	96%	67%-94% (>10mm) 73%-98% (6mm)	86%-98% (>10mm) 80%-93% (>6mm)	Weak	Low	26
Flexible Sigmoidoscopy	58-75%	72%-86%	92%	Strong	Intermediate	24 (28 with FIT)
Colonoscopy	95%	89%-98% (>10mm) 75%-93% (>6mm)	90%	Intermediate	High	28

Patient Considerations

	HSgFOBT	FIT	FIT-DNA	CT Colonography	FS (+ FIT)	Colonoscopy
Invasiveness	+	+	+	++	++	+++
Home test	Yes	Yes	Yes	No	No	No
Dietary restrictions	Yes	No	No	Yes	Yes	Yes
Interval	1 year	1 year	1-3 years	5 years	5 (10 years)	10 years (if normal)
Complications	Negligible	Negligible	Negligible	Few	Few	Low (0.1%)
Patient Participation	Moderate	Moderate	Moderate	Moderate	Moderate	Lowest
Cost	\$	\$	\$\$	\$\$	\$	\$\$

Robertson et al. AJG; 2017: 112; 37-53.

Inadomi. NEJM; 2017; 376:1598-1600.

May F. Huntington Health grand rounds Sept 6, 2024

Blood tests for colorectal cancer?



Epi proColon[®] is an FDA-approved blood test for colorectal cancer screening. This test is intended for persons age 50 and older who are unwilling or unable to be screened by recommended methods.

Rx Only

ABOUT GALLERI

A new way to screen for more cancers

Go further with Galleri[®]. Adding Galleri, a multi-cancer early detection test, to your cancer screenings allows you to go beyond what's currently possible. Now you can screen for a signal shared by more than 50 types of cancer with Galleri.¹

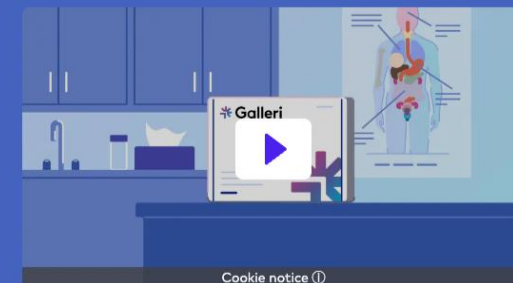
Request Galleri

The Galleri test does not detect a signal for all cancers and not all cancers can be detected in the blood. Galleri should be used in addition to healthcare provider recommended screening tests.



Galleri screens for a signal associated with active cancer

Galleri checks more than 100,000 DNA regions and over a million specific DNA sites to screen for a signal shared by cancers that could be hiding.² The Galleri test looks for cell-free DNA and identifies whether it comes from healthy or cancer cells.³ DNA from cancer cells has specific methylation patterns that identify it as a cancer signal. Methylation patterns also contain information about the tissue type or organ associated with the cancer signal to guide next steps.³



A comprehensive clinical studies program

Our clinical research program drives discovery, development, and validation of our early cancer detection tests.

[The PREEMPT CRC Study](#)

[The PROACT LUNG Study](#)

[The Vallania Study](#)

Our colorectal cancer screening study: The PREEMPT CRC® Study

The PREEMPT CRC Study is the largest clinical study validating a blood-based colorectal screening test. The PREEMPT CRC Study included more than 200 study sites across urban and rural communities, enrolling more than 40,000 participants across a range of racial, ethnic, and socioeconomic backgrounds.

We are thankful for the time and commitment of our participants and investigators to help Freenome develop tests to detect cancer early.

 Recruitment complete

.....

Get more information on our PREEMPT CRC Study.

 [Learn more at **clinicaltrials.gov**](#)

Blood tests for colorectal cancer?

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July 29, 2024

- *Shield is first blood test approved by FDA as a primary screening option for colorectal cancer and that meets performance requirements for Medicare coverage*
- *Blood test offers easy, convenient and pleasant screening option with potential for improving colorectal cancer screening rates*
- *Guardant Health to host investor conference call and webcast Monday, July 29, at 8:30 a.m. ET / 5:30 a.m. PT*

PALO ALTO, Calif.--(BUSINESS WIRE)-- Guardant Health, Inc. (Nasdaq: GH), a leading precision oncology company, today announced the U.S. Food and Drug Administration (FDA) has approved the company's Shield™ blood test for colorectal cancer (CRC) screening in adults age 45 and older who are at average risk for the disease. It is the first blood test to be approved by the FDA as a primary screening option for CRC, meaning healthcare providers can offer Shield in a manner similar to all other non-invasive methods recommended in screening guidelines. Shield is also the first blood test for CRC screening that meets the requirements for Medicare coverage. ¹

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March 14, 2024

A Cell-free DNA Blood-Based Test for Colorectal Cancer Screening

Daniel C. Chung, M.D., Darrell M. Gray II, M.D., M.P.H., Harminder Singh, M.D., Rachel B. Issaka, M.D., M.A.S., Victoria M. Raymond, M.S., Craig Eagle, M.D., Sylvia Hu, Ph.D., Darya I. Chudova, Ph.D., AmirAli Talasaz, Ph.D., Joel K. Greenson, M.D., Frank A. Sinicrope, M.D., Samir Gupta, M.D., M.S.C.S., and William M. Grady, M.D.

Blood tests for colorectal cancer?

Table 2. Sensitivity and Specificity of the Cell-free DNA (cfDNA) Blood-Based Test for the Most Advanced Findings on Colonoscopy.*

Variable	Most Advanced Finding on Colonoscopy <i>no.</i>	cfDNA Blood-Based Test	
		Positive Test <i>no.</i>	Sensitivity (95% CI) %
Colorectal cancer			
Any	65	54	83.1 (72.2–90.3)
Stage I, II, or III*	48	42	87.5 (75.3–94.1)
Advanced precancerous lesions†	1116	147	13.2 (11.3–15.3)
			Specificity (95% CI)
Nonadvanced adenomas, nonneoplastic findings, and negative colonoscopy	6680	698	89.6 (88.8–90.3)
Nonneoplastic findings and negative colonoscopy	4514	457	89.9 (89.0–90.7)

* Excluded were 10 stage IV and 7 pathologically confirmed, incompletely staged colorectal cancers.

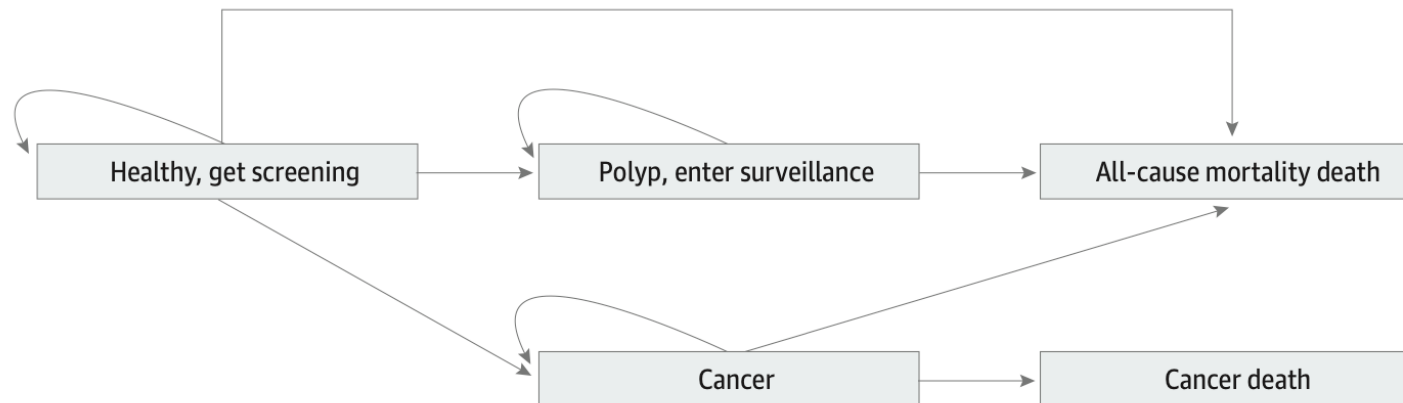
† Advanced precancerous lesions include advanced adenomas and sessile serrated lesions at least 10 mm in the largest dimension.

Original Investigation | Gastroenterology and Hepatology

Cost-Effectiveness of Liquid Biopsy for Colorectal Cancer Screening in Patients Who Are Unscreened

Zainab Aziz, BS; Sophie Wagner, BS; Alice Agyekum, BS; Yoanna S. Pumpalova, MD; Matthew Prest, MS; Francesca Lim, MS; Sheila Rustgi, MD; Fay Kastrinos, MD, MPH; William M. Grady, MD; Chin Hur, MD, MPH

Figure 1. Markov Model Schematic



Discussion

In this study, we used a Markov simulation to analyze the cost-effectiveness of LB, used both as a novel first or second-line screening modality. We present the first analysis that integrates novel LB into paradigms for CRC screening and systematically explores scenarios to determine the cost-effectiveness of LB.

The most cost-effective screening strategy in our base-case model was colonoscopy, with an ICER of \$28 071 per LYG. While C-LB had the highest number of LYG and prevented the most cancers, the cost of LB would have to reduce by 66% (from \$949 to \$324) for the C-LB strategy to become cost-effective in our model. Compared with NH, the cost of LB would have to be reduced by 94% for its ICER to drop below the WTP threshold of \$100 000 per LYG. When compared with stool-based tests, the cost of LB would have to decrease by 43% to 80% to be cost-effective. LB and C-LB had more LYG when polyp detection was introduced, but they did not achieve cost-effectiveness at LB's current price even with perfect performance.

- Most people with early cancer have no symptoms
- Change in stool – shape, color, blood
- Change in bowel habits (persists)
- Abdominal pain, cramping
- Urges to have bowel movements
- Unintentional weight loss

Prevention (AICR)



Analysing research on cancer prevention and survival



Diet, nutrition, physical activity
and **colorectal cancer**

2017

Revised 2018

Prevention (AICR)

2017	DIET, NUTRITION, PHYSICAL ACTIVITY AND COLORECTAL CANCER		
		DECREASES RISK	INCREASES RISK
STRONG EVIDENCE	Convincing	Physical activity ^{1,2}	Processed meat ³ Alcoholic drinks ⁴ Body fatness ⁵ Adult attained height ⁶
	Probable	Wholegrains Foods containing dietary fibre ⁷ Dairy products ⁸ Calcium supplements ⁹	Red meat ¹⁰
LIMITED EVIDENCE	Limited – suggestive	Foods containing vitamin C ¹¹ Fish Vitamin D ¹² Multivitamin supplements ¹³	Low intakes of non-starchy vegetables ¹⁴ Low intakes of fruits ¹⁴ Foods containing haem iron ¹⁵
	Limited – no conclusion	Cereals (grains) and their products; potatoes; animal fat; poultry; shellfish and other seafood; fatty acid composition; cholesterol; dietary n-3 fatty acid from fish; legumes; garlic; non-dairy sources of calcium; foods containing added sugars; sugar (sucrose); coffee; tea; caffeine; carbohydrate; total fat; starch; glycaemic load; glycaemic index; folate; vitamin A; vitamin B6; vitamin E; selenium; low fat; methionine; beta-carotene; alpha-carotene; lycopene; retinol; energy intake; meal frequency; dietary pattern	
STRONG EVIDENCE	Substantial effect on risk unlikely		

- Age
- Family history of colon cancer
- Personal history
- Alcohol
- Cigarette smoking
- Obesity

- Physical activity
- Aspirin
- Combination hormone replacement therapy
- Polyp removal

- Unclear affects on risk:
 - NSAIDs other than aspirin
 - Calcium
 - Diet
- No affects on risk
 - HRT with estrogen only
 - Statins

- Get screened regularly
- Maintain a healthy weight
- Adopt a physically active lifestyle
- Consume a healthy diet
 - ≥ 5 daily servings of fruits/vegetables
 - Limit red meat/processed meat
 - Choose whole grains instead of processed
- Limit alcohol consumption



HHS Public Access

Author manuscript

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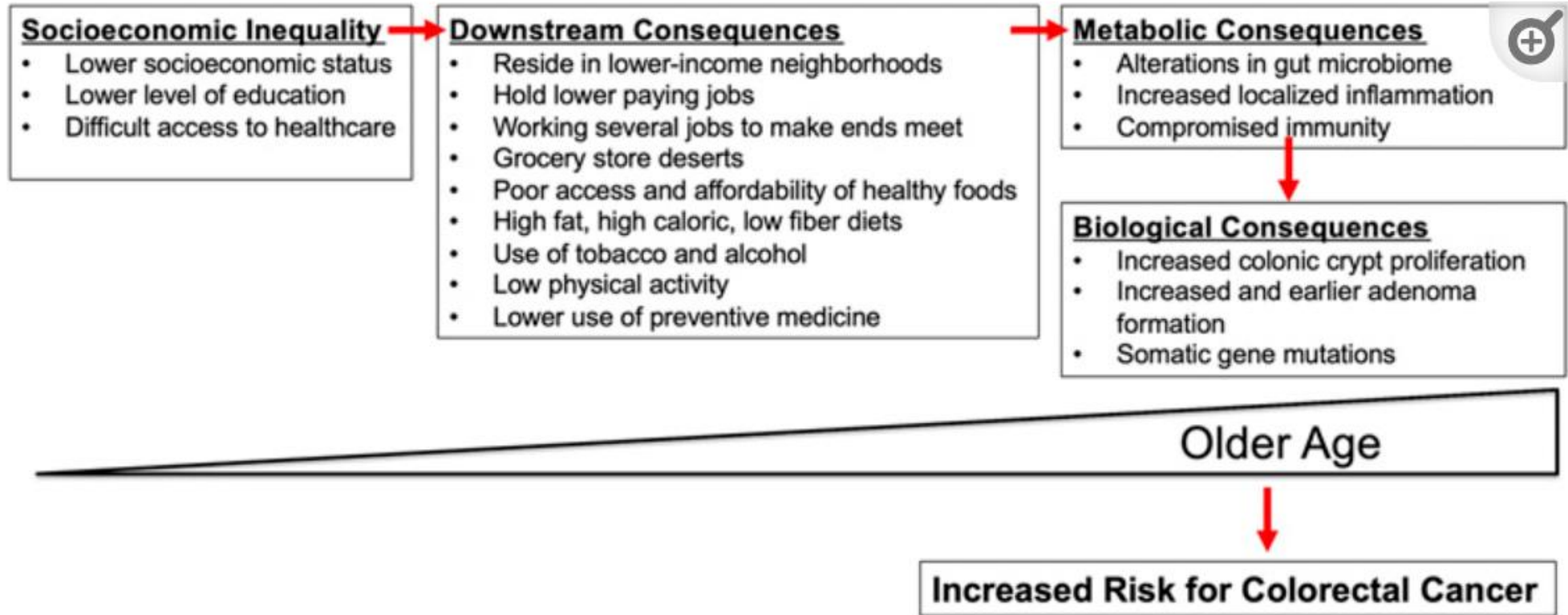
Adv Cancer Res. 2021 ; 151: 197–229. doi:10.1016/bs.acr.2021.02.007.

Racial and Ethnic Disparities in Colorectal Cancer Incidence and Mortality

John M. Carethers

Division of Gastroenterology and Hepatology, Department of Internal Medicine, and Department of Human Genetics and Rogel Cancer Center, University of Michigan, Ann Arbor, Michigan

Socioeconomic Disparities for CRC Risk Connections and Consequences





HHS Public Access

Author manuscript

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Framework and Strategies to Eliminate Disparities in Colorectal Cancer Screening Outcomes

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BOX 1:

MAJOR STRATIFICATIONS OF DISPARITIES IN COLORECTAL CANCER SCREENING OUTCOMES

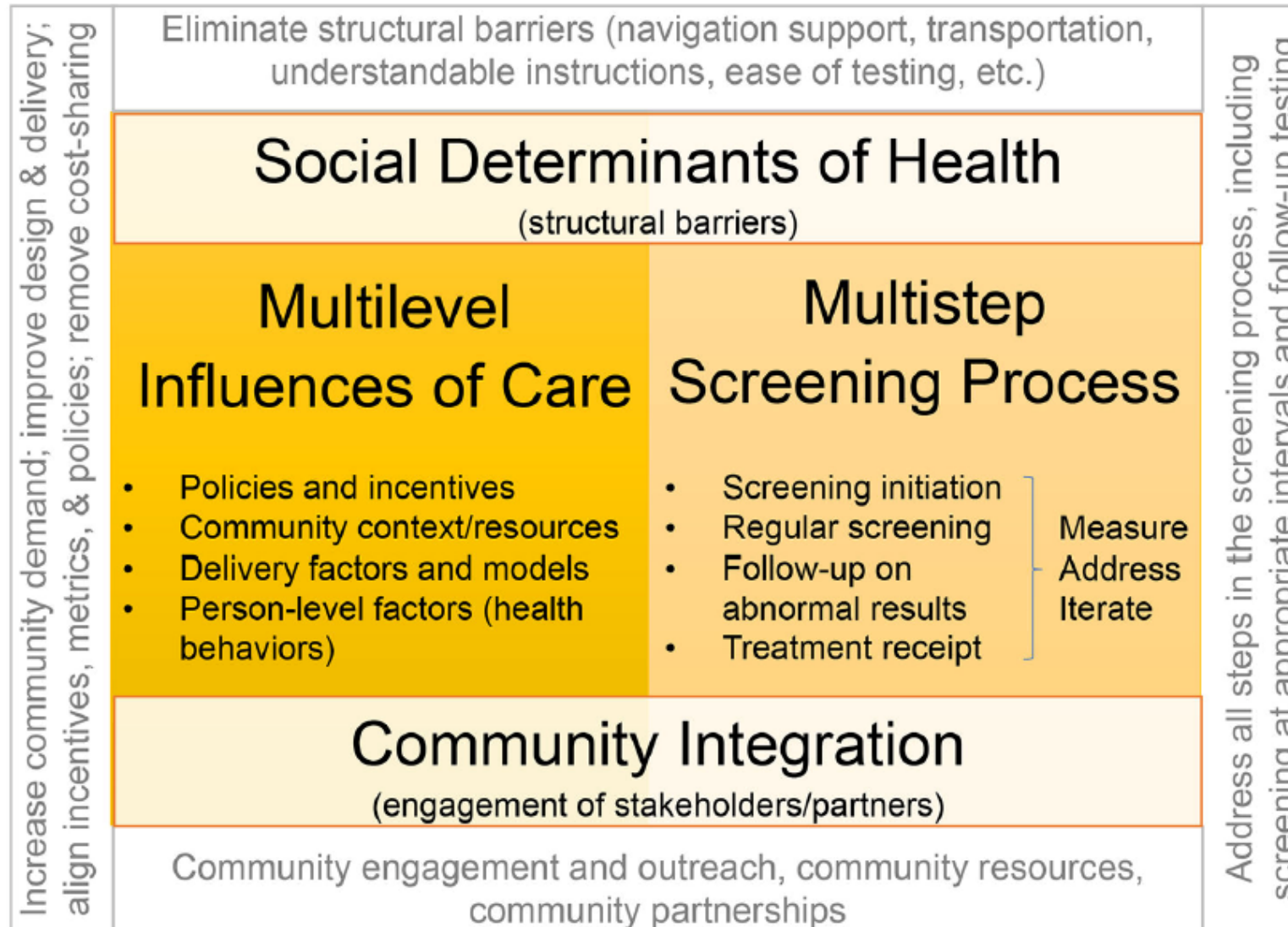
1. Race/ethnicity
2. English proficiency/Language
3. Immigrant status
4. Educational level
5. Income
6. Insurance coverage
7. Occupation
8. Age
9. Sex/Gender
10. Geography (neighborhoods, county, state, rural vs. urban, etc.)
11. Behavioral risk factors (e.g., obesity)

BOX 2:

PRINCIPLES OF COMMUNITY ENGAGEMENT

1. Have shared goals with the community
2. Understand the community and its history of engagement
3. Build trust and seek commitment from stakeholders
4. Respect diverse perspectives within the community
5. Identify and mobilize community assets
6. Partner with the community
7. Assure community ownership and control of actions
8. Long-term commitment

Framework/Strategies to Eliminate Disparities in CRC Screening Outcomes



Journal of Clinical Oncology®

An American Society of Clinical Oncology Journal

[J Clin Oncol.](#) 2013 Jun 1; 31(16): 1928–1930.

PMCID: PMC3661932

Published online 2013 Apr 15. doi: [10.1200/JCO.2012.47.8412](https://doi.org/10.1200/JCO.2012.47.8412)

PMID: [23589553](https://pubmed.ncbi.nlm.nih.gov/23589553/)

Eliminating Racial Disparities in Colorectal Cancer in the Real World: It Took a Village

[Stephen S. Grubbs](#), [Blase N. Polite](#), [John Carney, Jr](#), [William Bowser](#), [Jill Rogers](#), [Nora Katurakes](#), [Paula Hess](#), and [Electra D. Paskett](#)

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The Delaware Model...

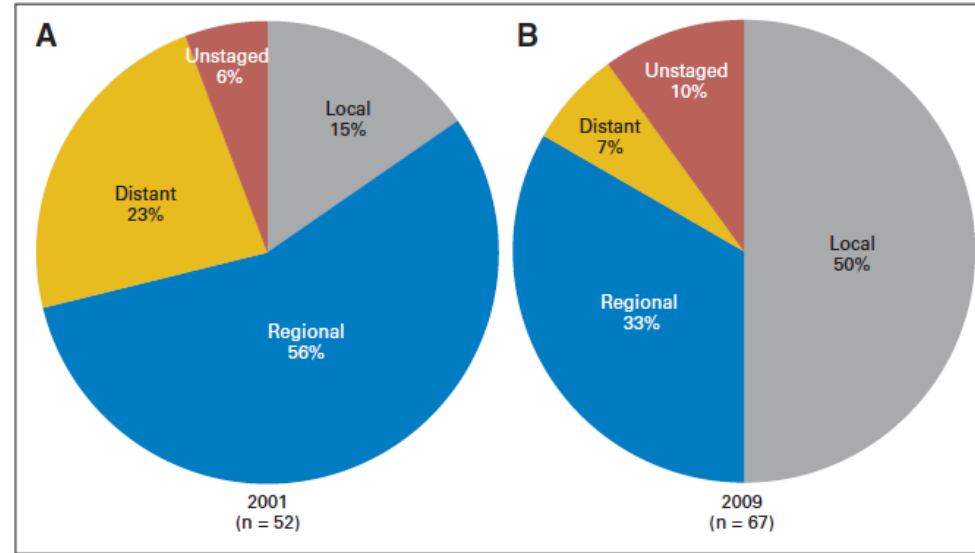


Fig 1. Colorectal cancer by stage of diagnosis among African Americans in Delaware (A) 2001 and (B) 2009.

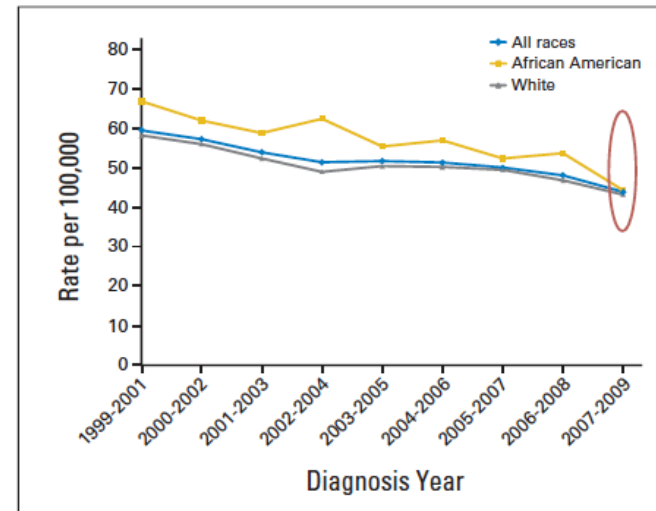


Fig 2. Age-adjusted colorectal cancer incidence rates (rolling 3-year averages) by race in Delaware from 1999 to 2009.

The Delaware Model...

Trends in CRC Screening, Incidence, and Mortality Rates by Race in Delaware: 2001 and 2009

Trend	2001*		2009		>Change From 2001 to 2009 (%)	
	Black	White	Black	White	Black	White
Ever had screening colonoscopy, %	47.8	58.0	73.5	74.7	54	29
CRC incidence rate per 100,000 [‡]	66.9	58.2	44.3	43.2	-34	-26
Total No. of cases [‡]	205	1,206	235	1,149		
CRC mortality rate per 100,000 [‡]	31.2	19.5	18.0	16.9	-42	-13
Total No. of cases [§]	88	398	75	420		



HHS Public Access

Author manuscript

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Published in final edited form as:

Adv Cancer Res. 2021 ; 151: 197–229. doi:10.1016/bs.acr.2021.02.007.

Racial and Ethnic Disparities in Colorectal Cancer Incidence and Mortality

John M. Carethers

Division of Gastroenterology and Hepatology, Department of Internal Medicine, and Department of Human Genetics and Rogel Cancer Center, University of Michigan, Ann Arbor, Michigan

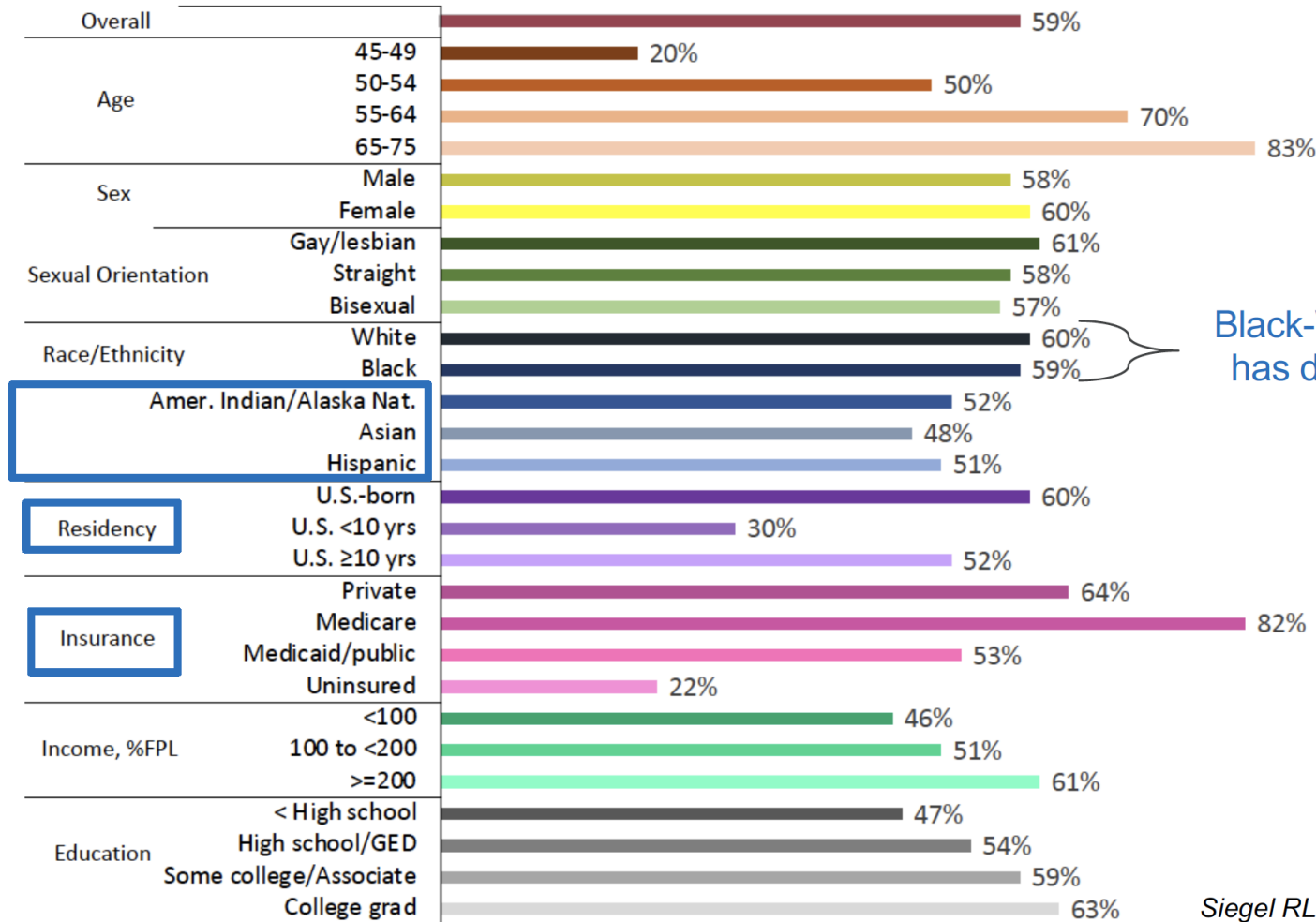
Adv Cancer Res. 2021 ; 151: 197–229. doi:10.1016/bs.acr.2021.02.007.

Socioeconomic Disparities for CRC Risk Mitigation

Colonoscopy	Non-invasive Screening
Navigation personnel's racial/ethnic background similar to patient's background; use native language	Telemedicine (phone or video) instructions and advice from virtual assistants
Multifaceted points of communication and execution: (a) provides general education about the procedure, including its importance in reducing cancer risk, (b) ensure prep is picked up and/or delivered, (c) instructions and coaching on prep utilization and completion, (d) arrange transportation to and from colonoscopy site, (e) arrange observer post-procedure with follow-up contact within hours post procedure	Post-navigation follow-up after test evaluation for transmission of results and next steps
Mitigates screening costs through insurance and other means for underinsured patients	Move to colonoscopy navigation if non-invasive test is positive
With healthcare provider, communicates results of colonoscopy and any pathology, and next steps	Persistent community education presence on importance of colorectal cancer screening for racial/ethnic groups

Disparities in Screening

Variation in Screening Participation



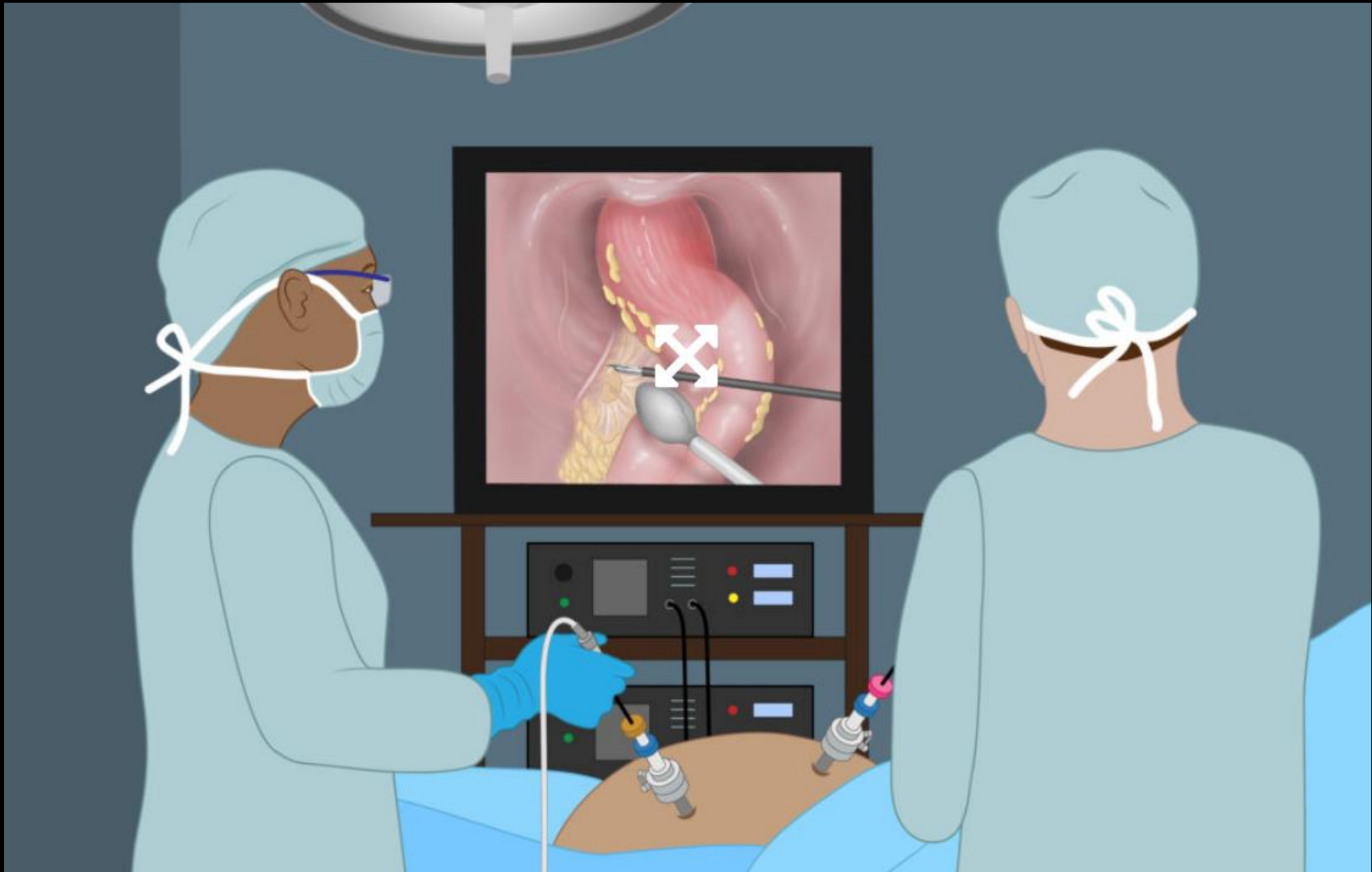
Black-White screening gap has decreased over time

Up-to-date with CRC screening (%)

Siegel RL CA Cancer J Clin 2023; National Health Interview Survey 2021; Health Resources Services Administration Uniform Data Set; Ladabaum U Gastro 2023.

- What's new(er) in treatment
 - Minimally invasive surgery (MIS)
 - Enhanced recovery pathways
 - Opiate reducing strategies
 - Reduced complications
 - Shorter hospital stay
 - Reduced costs





Randomized Controlled Trial

> [Ann Surg.](#) 2007 Oct;246(4):655-62; discussion 662-4.

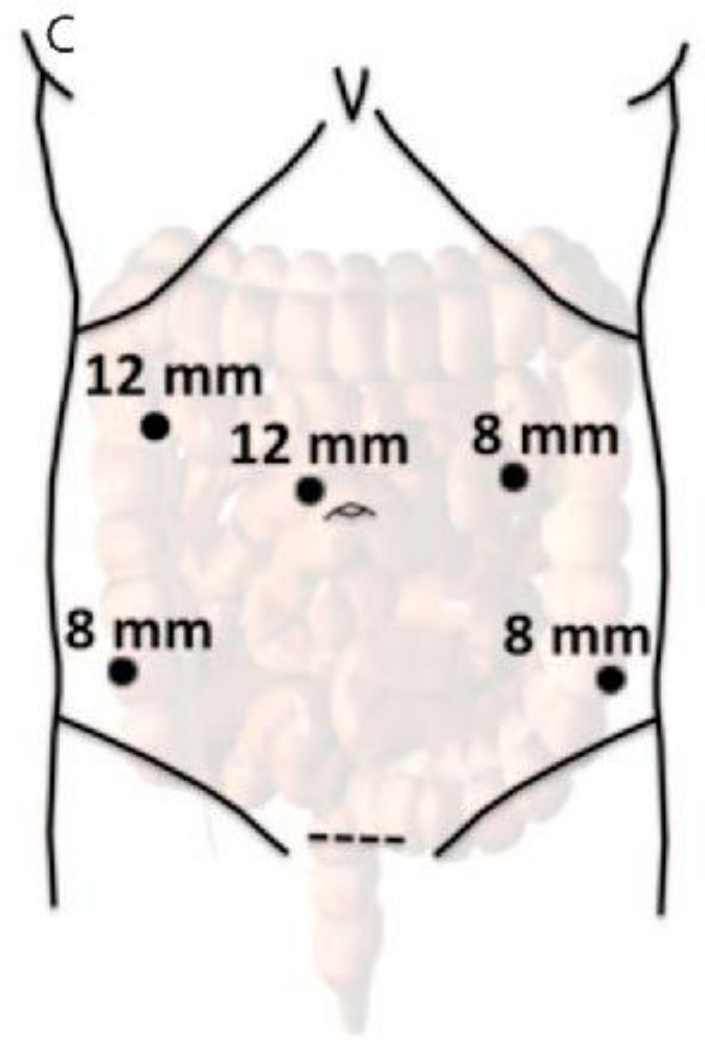
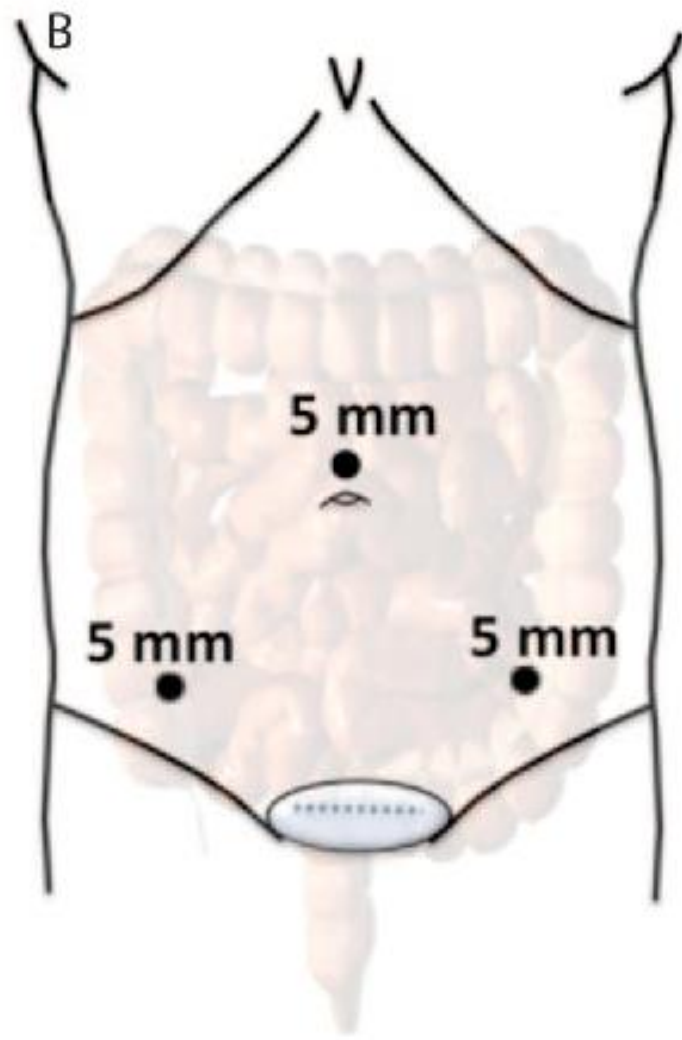
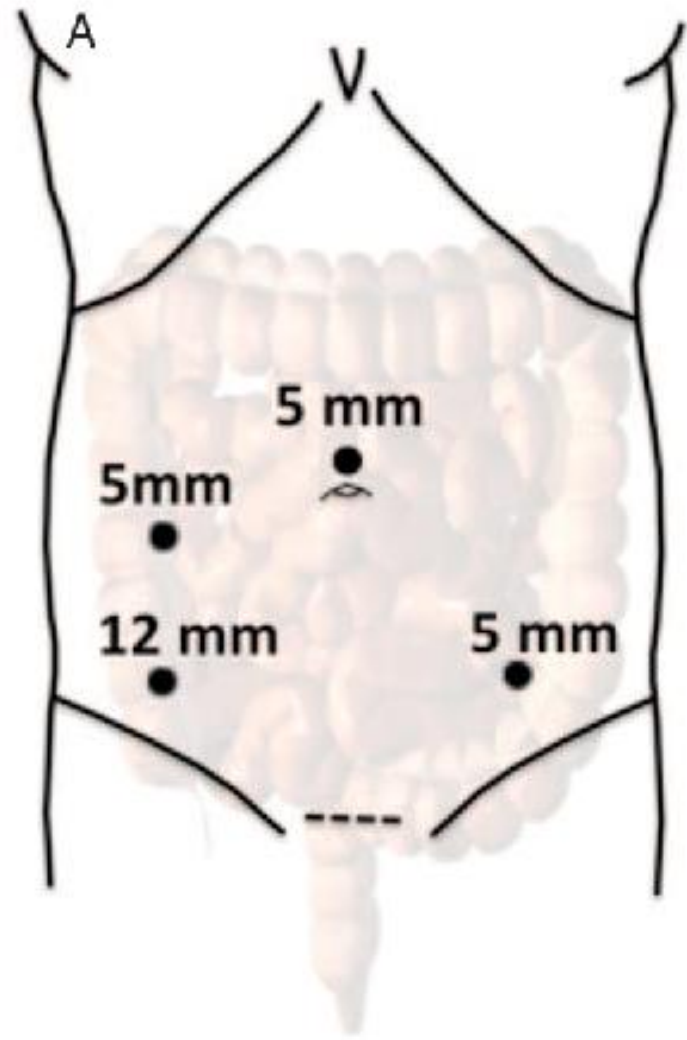
doi: [10.1097/SLA.0b013e318155a762](#).

Laparoscopic colectomy for cancer is not inferior to open surgery based on 5-year data from the COST Study Group trial

[James Fleshman](#) ¹, [Daniel J Sargent](#), [Erin Green](#), [Mehran Anvari](#), [Steven J Stryker](#),
[Robert W Beart Jr](#), [Michael Hellinger](#), [Richard Flanagan Jr](#), [Walter Peters](#), [Heidi Nelson](#),
[Clinical Outcomes of Surgical Therapy Study Group](#)

Affiliations + expand

PMID: [17893502](#) DOI: [10.1097/SLA.0b013e318155a762](#)

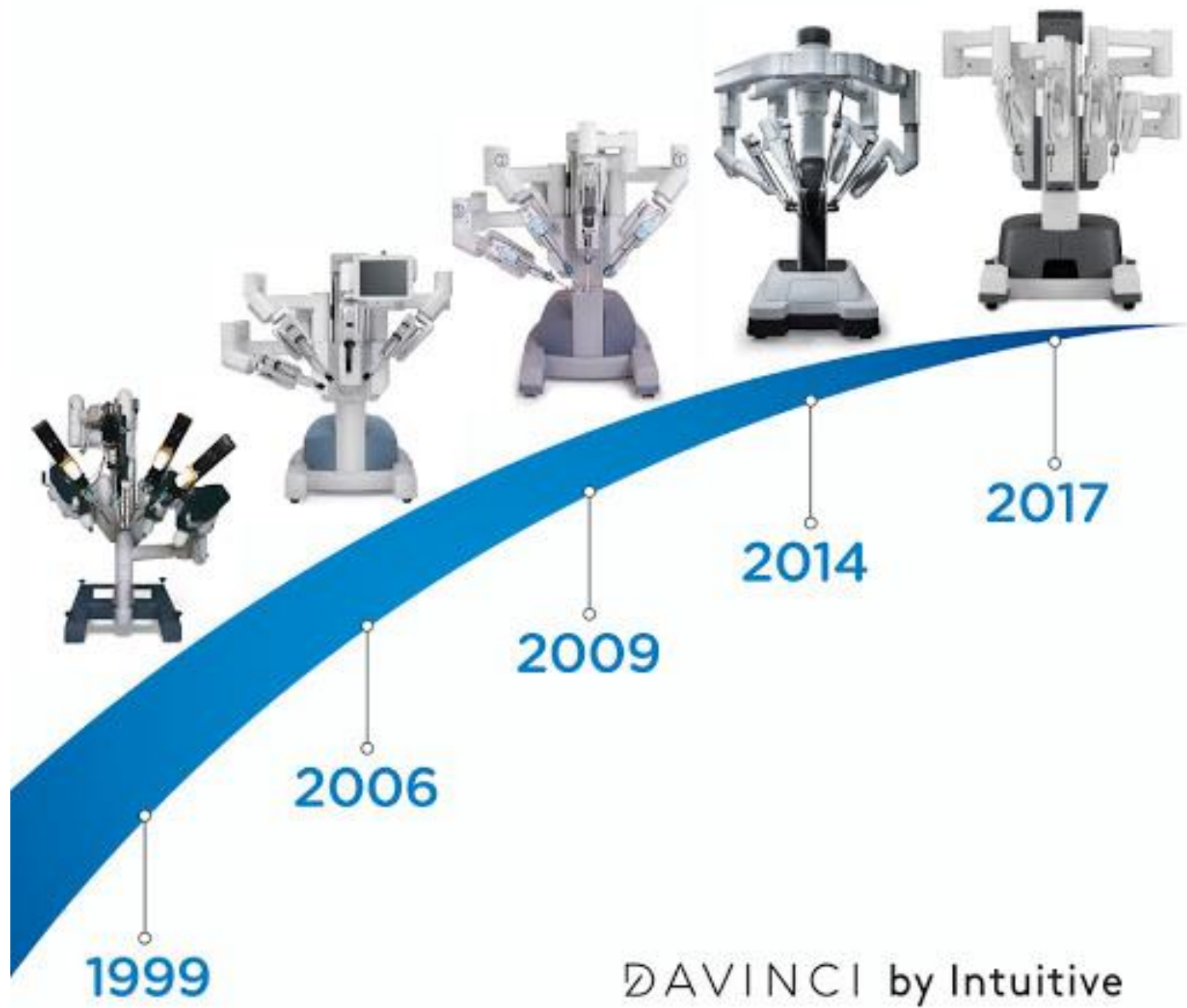




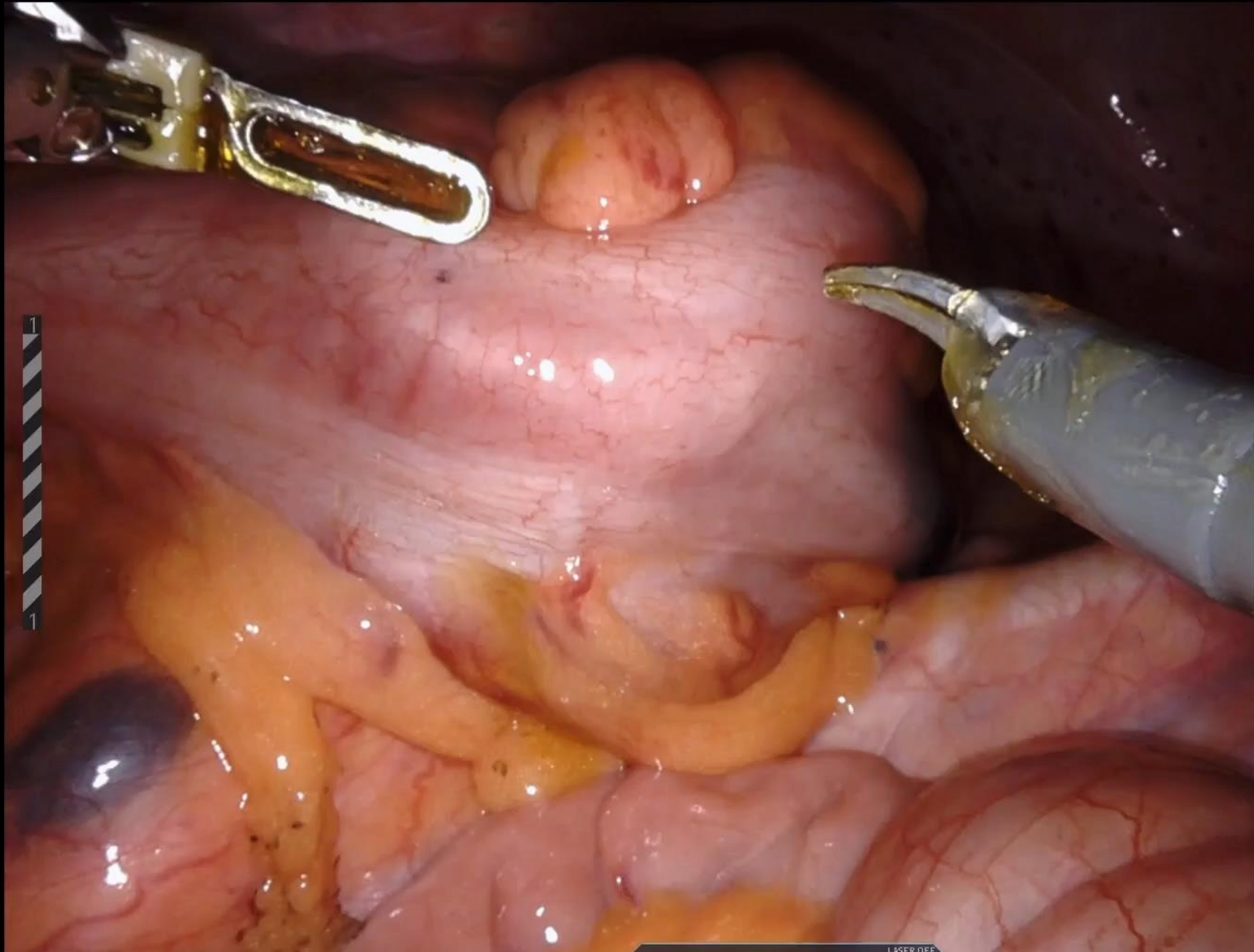












1 SMALL GRASPING RETRACTOR

2 FENESTRATED BIPOLAR FORCEPS

L BIPOLAR

3 27° 10° R 1x 30°

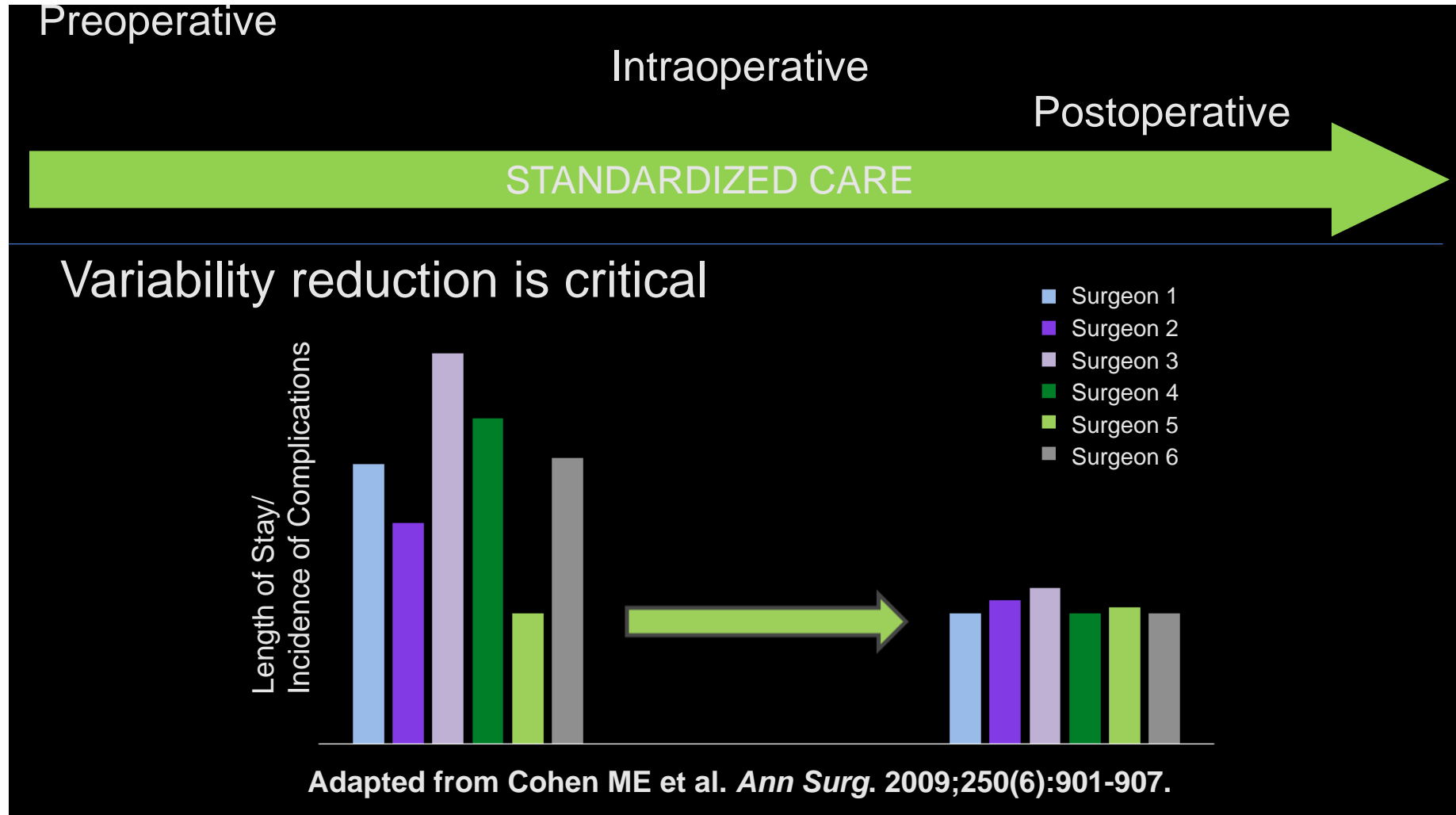
LASER OFF

4 MONOPOLAR CURVED SCISSORS

R CUT

R COAG

Current Surgical Challenges



What Is Enhanced Recovery After Surgery?^{1,2}

ERAS protocols are:

- Multidisciplinary care pathways
 - From decision of need for procedure to return to baseline level of function

ERAS key characteristics:

- Evidence-based, patient-centered care
- Designed to reduce patients' stress response to surgery
- Includes prehabilitation
- Patient involved in his/her own preparation/recovery

Integrated continuum from:

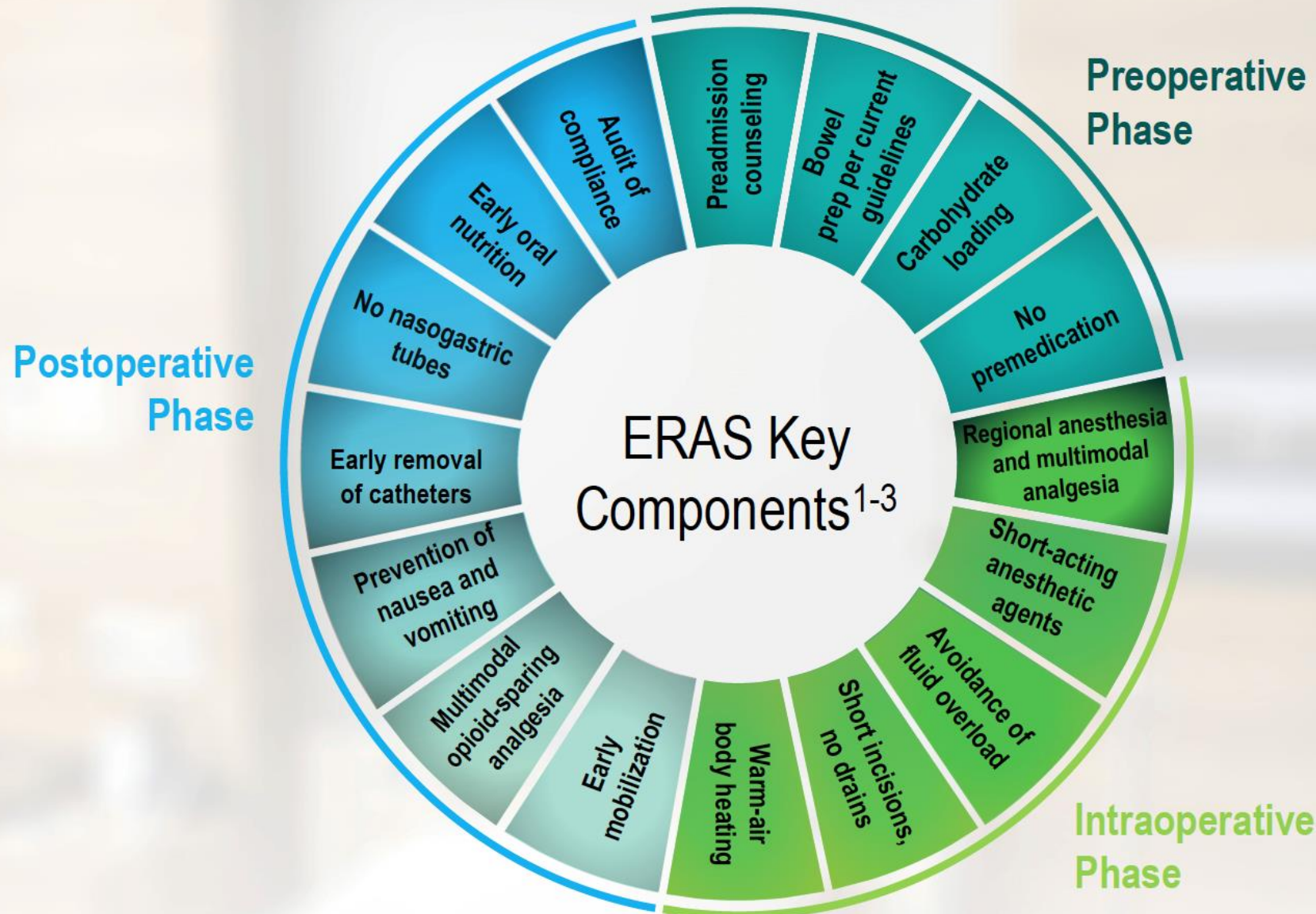
Home

Preoperative

Intraoperative

Postoperative

Home



1. Modified from Melnyk M, et al. *Can Urol Assoc J.* 2011;5(5):342-348. 2. Ljungqvist O, et al. *JAMA Surg.* 2017;E1-E7. Published online January 11, 2017. Accessed July 8, 2018. 3. AANA. <https://www.aana.com/practice/clinical-practice-resources/enhanced-recovery-after-surgery>. Accessed July 8, 2018.

ANNALS OF SURGERY

ORIGINAL ARTICLES

A Clinical Pathway to Accelerate Recovery After Colonic Resection

Basse, Linda MD; Hjort Jakobsen, Dorthe RN; Billesbølle, Per MD;
Werner, Mads MD, PhD; Kehlet, Henrik MD, PhD

[Author Information](#) 

Annals of Surgery 232(1):p 51-57, July 2000.

Early Studies – Open Surgery

- Prospectively study (n=60), mean age 74
- Defined anesthesia and analgesia
- Enforced early mobilization
- Early enteral nutrition
- Planned 48-hour postop stay

Postoperative Regimen

- No NG tube
- Mobilization
- Cisapride, magnesium oxide
- Epidural discontinued at 44 hours
- Patient discharged at 48 hours

Results

- 53 patients - no risk factors precluding 2d stay
- Right hemicolectomy (23), transverse (2), left (34), subtotal (1)
- Median stay - 2 days
- 57 patients defecated within 2 days
- 2 deaths (3.3%), 5 serious complications (8.3%)
- 73% of patients were satisfied with short stay

World J Surg (2019) 43:659–695
<https://doi.org/10.1007/s00268-018-4844-y>



CrossMark

SCIENTIFIC REVIEW

Guidelines for Perioperative Care in Elective Colorectal Surgery: Enhanced Recovery After Surgery (ERAS[®]) Society Recommendations: 2018

U. O. Gustafsson¹ · M. J. Scott^{2,3} · M. Hubner⁴ · J. Nygren⁵ · N. Demartines⁴ · N. Francis^{6,7} ·
T. A. Rockall⁸ · T. M. Young-Fadok⁹ · A. G. Hill¹⁰ · M. Soop¹¹ · H. D. de Boer¹² · R. D. Urman¹³ ·
G. J. Chang¹⁴ · A. Fichera¹⁵ · H. Kessler¹⁶ · F. Grass⁴ · E. E. Whang¹⁷ · W. J. Fawcett¹⁸ ·
F. Carli¹⁹ · D. N. Lobo²⁰ · K. E. Rollins²⁰ · A. Balfour²¹ · G. Baldini¹⁹ · B. Riedel²² · O. Ljungqvist²³

Published online: 13 November 2018

© The Author(s) 2018

Fig. 1 Preadmission items

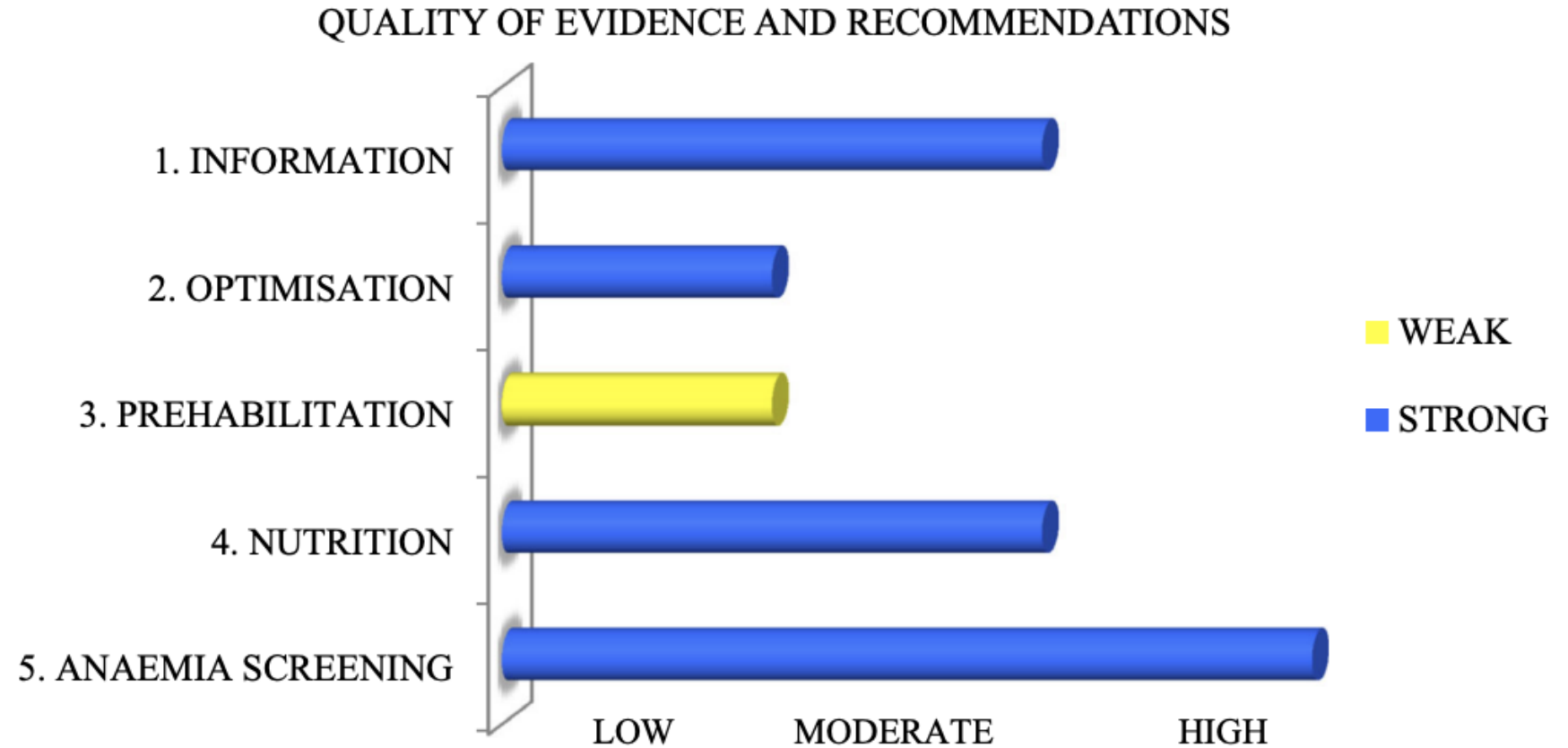


Fig. 2 Preoperative items

QUALITY OF EVIDENCE AND RECOMMENDATIONS

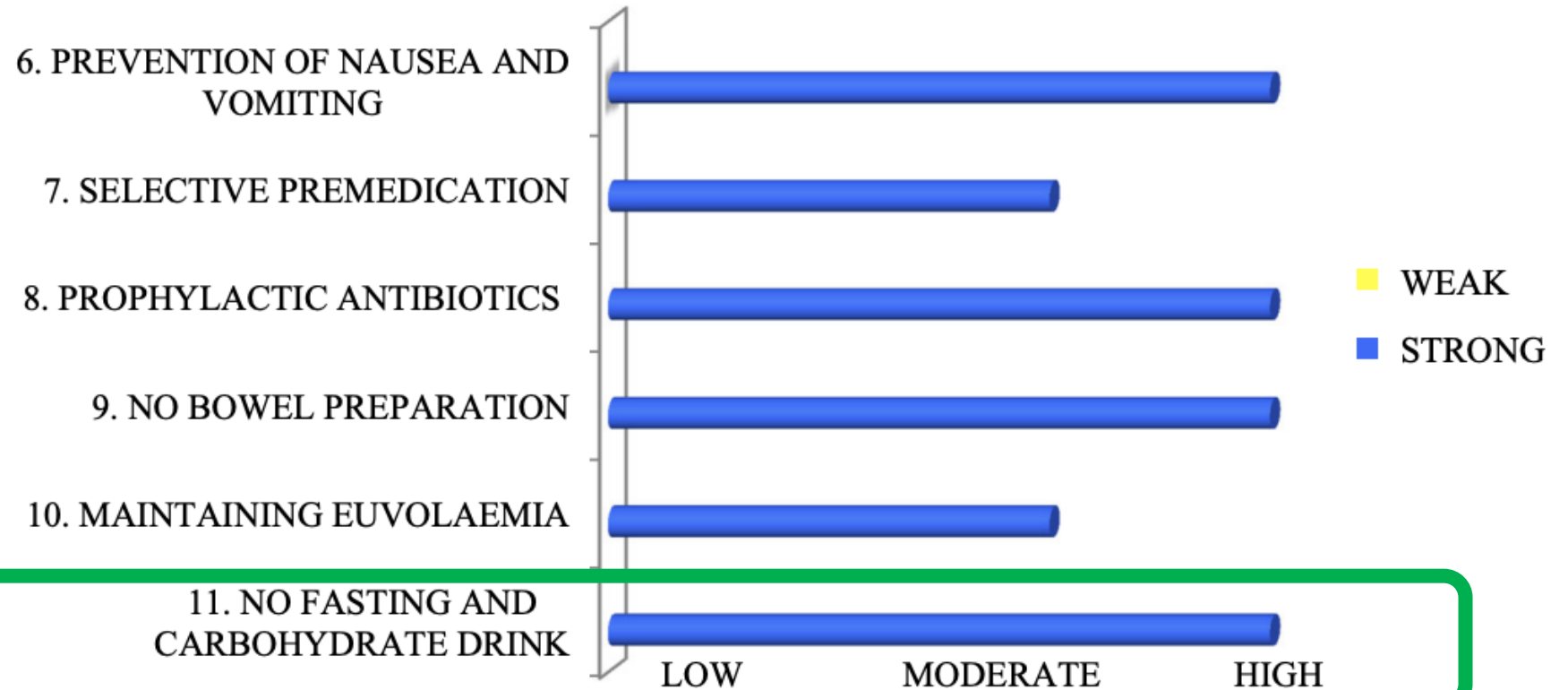


Fig. 3 Intraoperative items

QUALITY OF EVIDENCE AND RECOMMENDATIONS

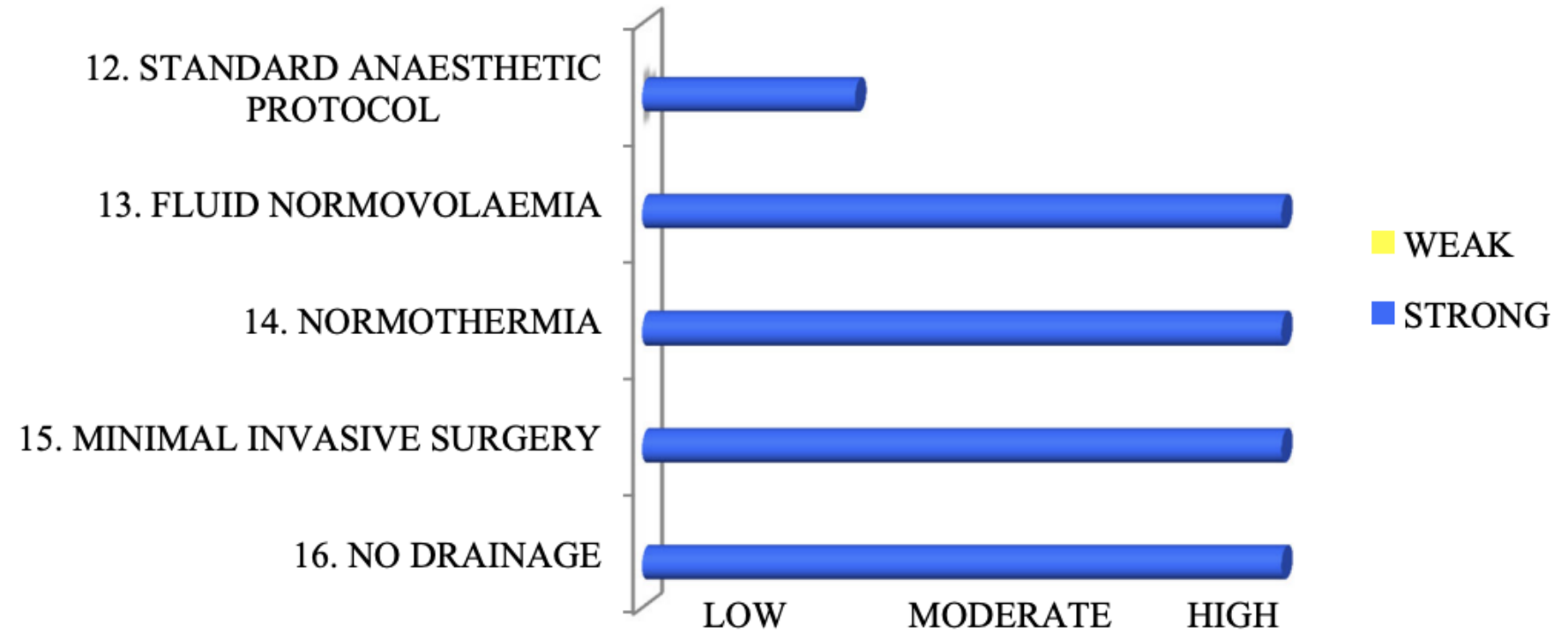
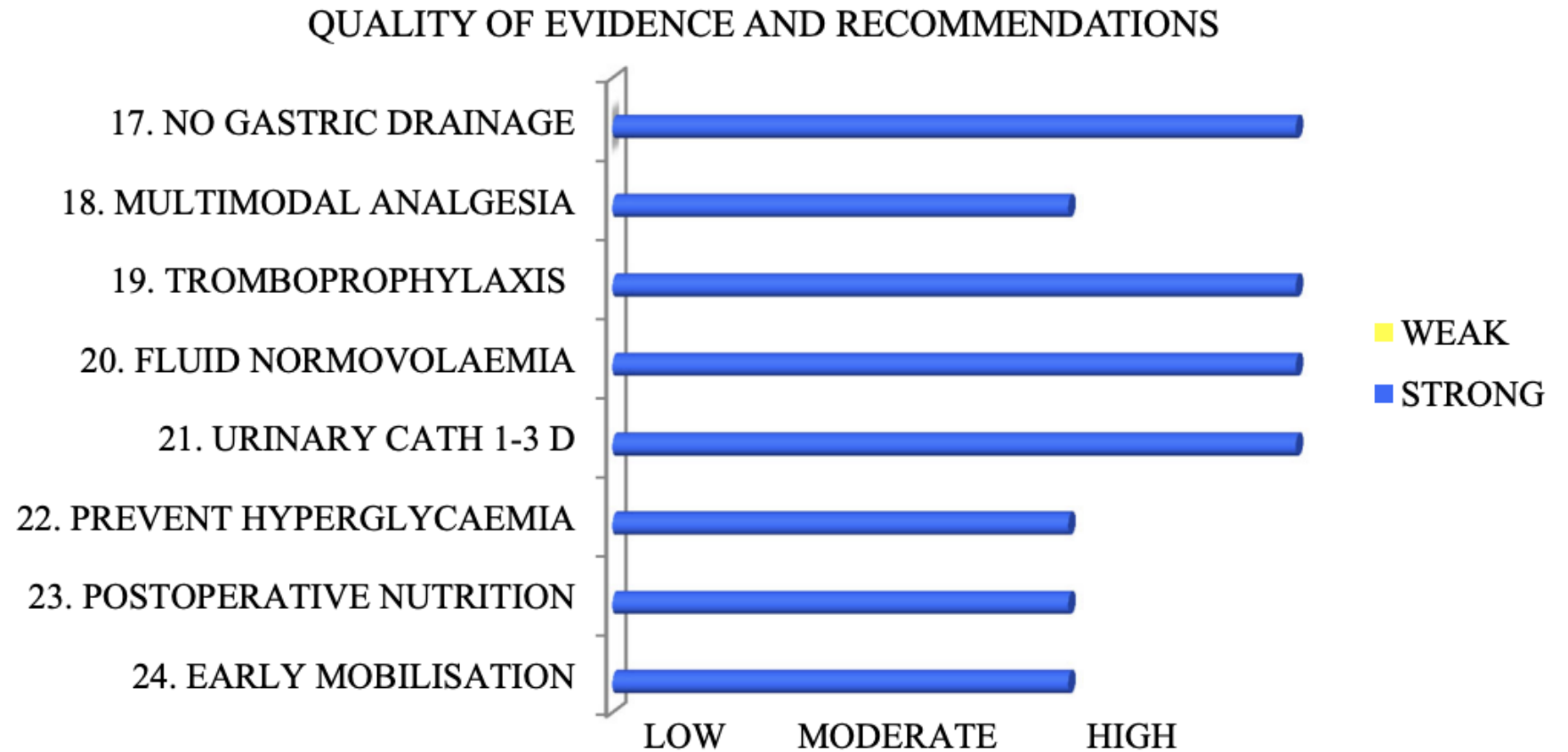



Fig. 4 Postoperative items



Impact of structured multicentre enhanced recovery after surgery (ERAS) protocol implementation on length of stay after colorectal surgery

Zubair Bayat^{1,2,3}, Anand Govindarajan^{1,2,3,4}, J. Charles Victor^{2,4} and Erin D. Kennedy^{1,2,3,*} 

¹Division of General Surgery, Department of Surgery, University of Toronto, Toronto, Ontario, Canada

²Institute of Health Policy Management and Evaluation, University of Toronto, Toronto, Ontario, Canada

³Department of Surgery, Mount Sinai Hospital, Sinai Health System, Toronto, Ontario, Canada

⁴Institute for Clinical Evaluative Sciences, Toronto, Ontario, Canada

*Correspondence to: Erin D. Kennedy, Department of Surgery, Mount Sinai Hospital, Suite 449, 600 University Avenue, Toronto, Ontario, M5G 1X5, Canada (e-mail: erin.kennedy@sinaihealth.ca)

Fig. 1 Results of the interrupted time series analysis for the impact of the introduction of ERAS protocols on LOS

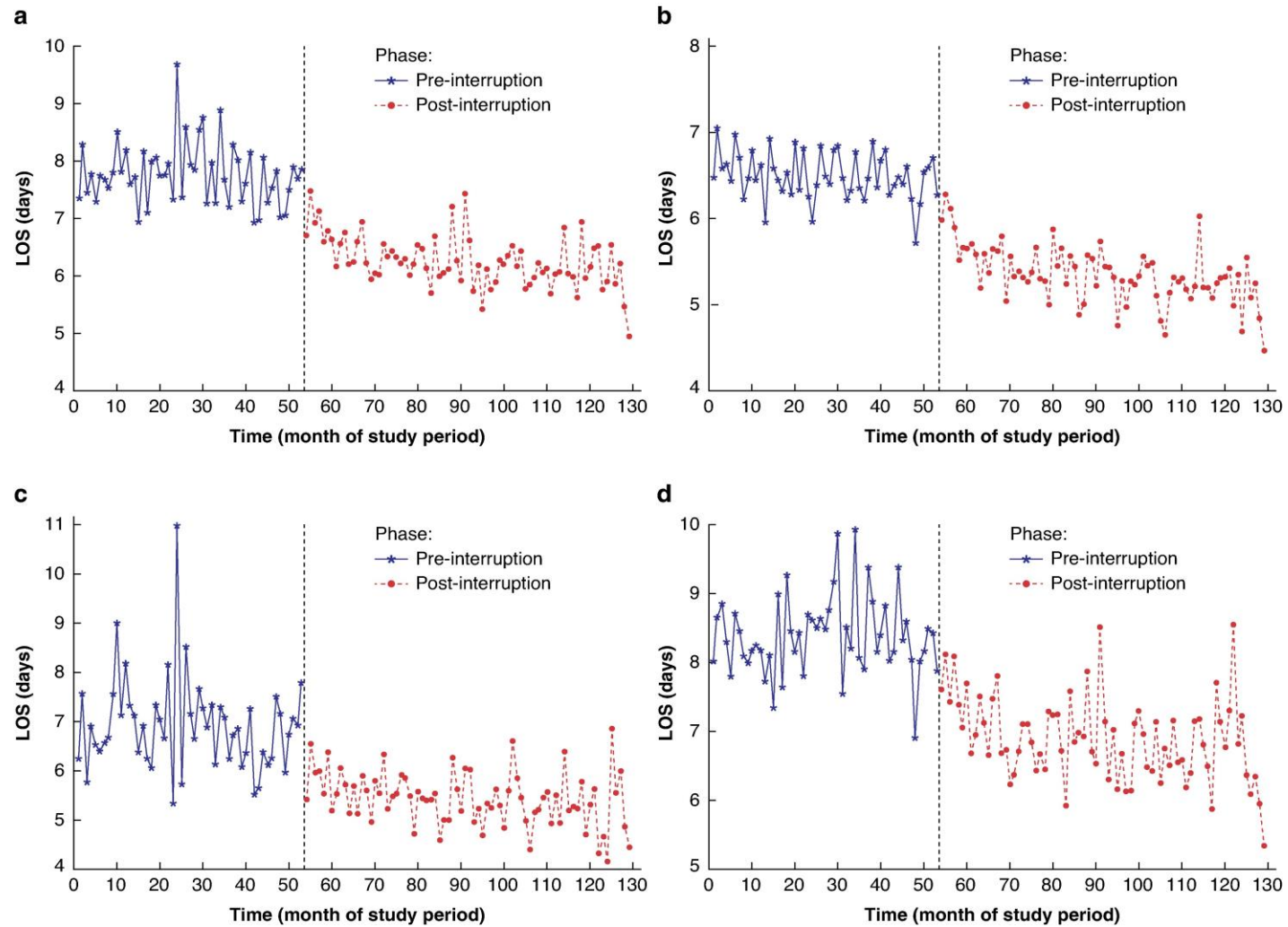


Fig. 2 Results of the interrupted time series analysis for the impact of the introduction of ERAS protocols on adjusted LOS

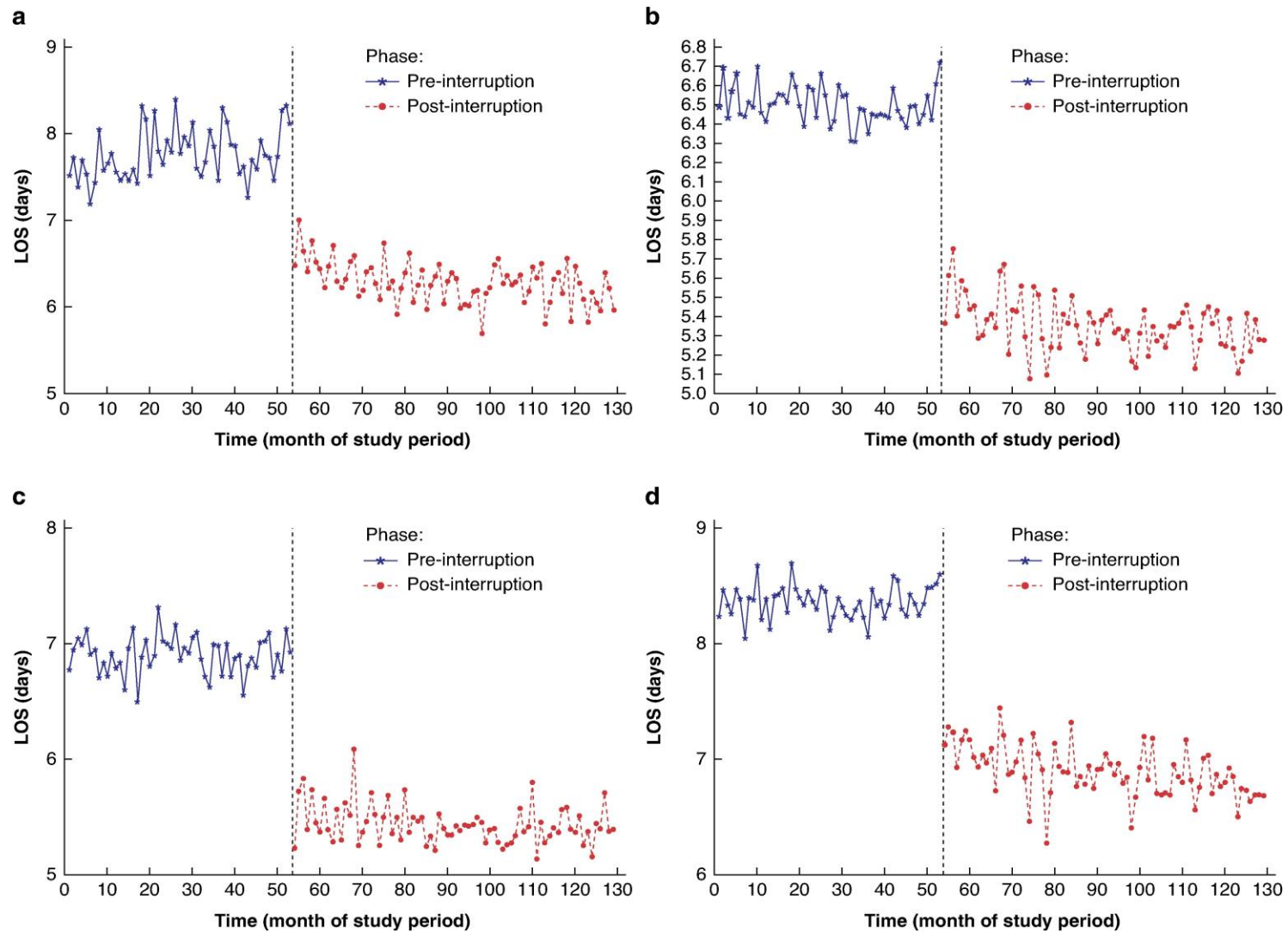
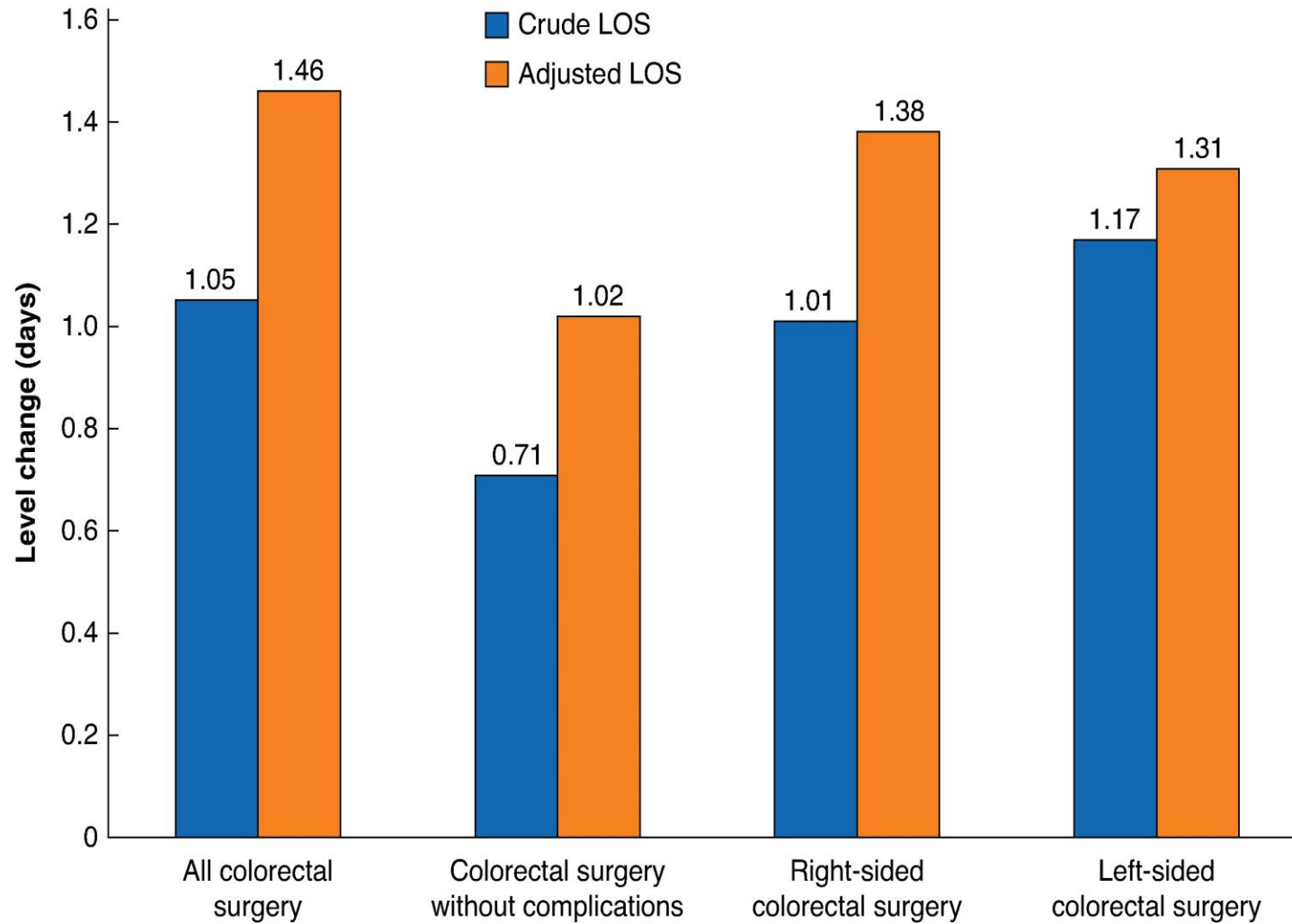


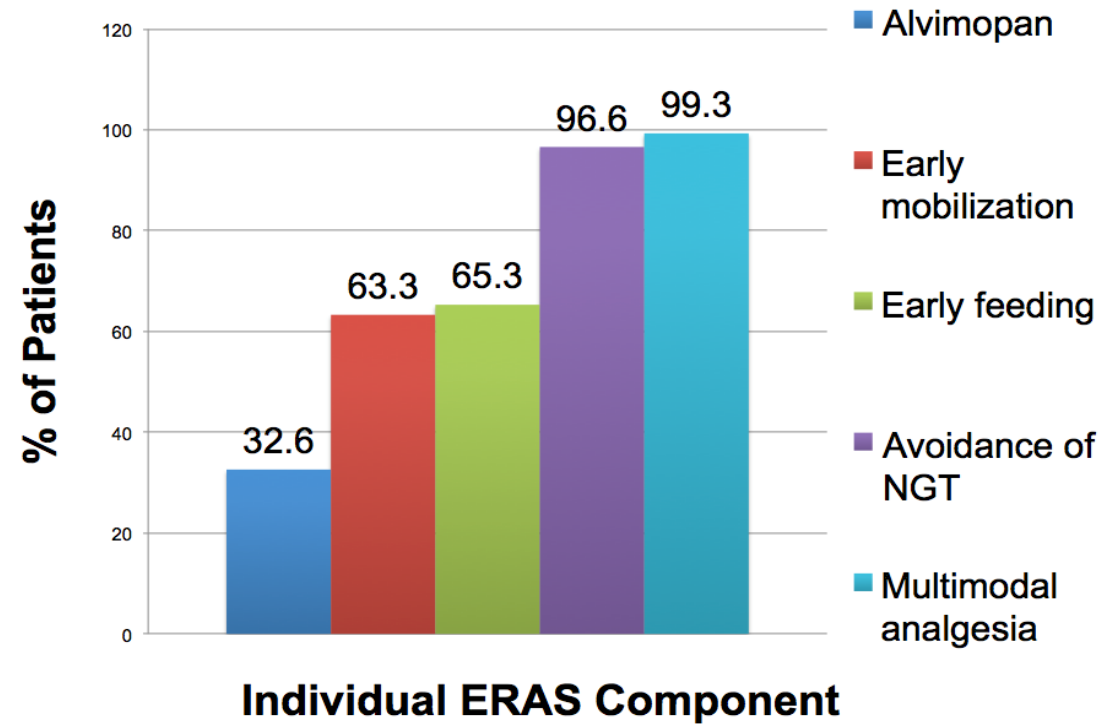
Fig. 3 Graph depicting the level change (decrease in LOS, measured in days) after ERAS protocol implementation for patients in each cohort



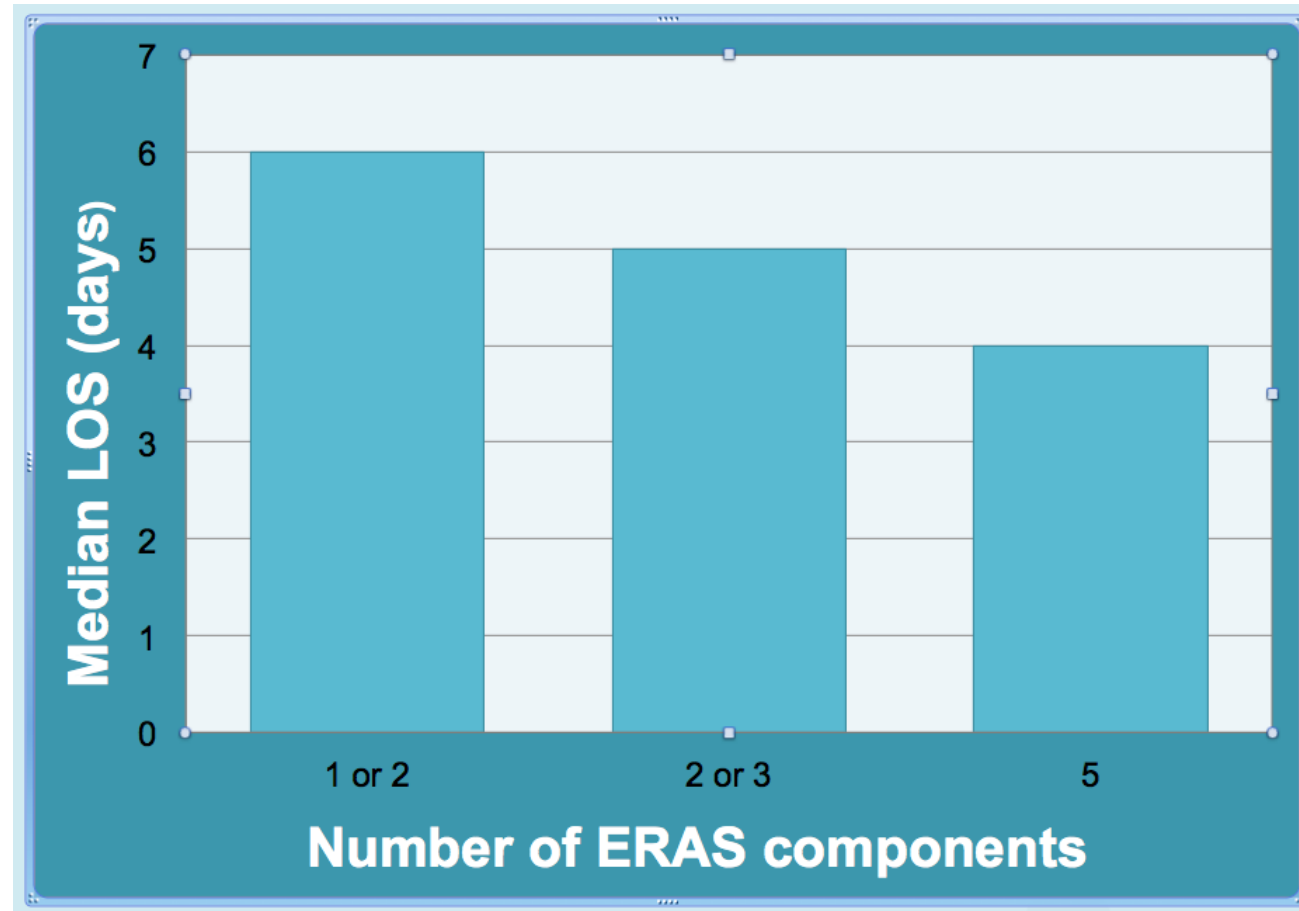
Audit of ERP adoption

Post EHR (Cerner)
126 Consecutive patients
MIS colorectal surgery 3/14-12/15

Use of ERAS Components



Audit of ERP adoption



Ambulatory Colectomy?



Identification of Eligibility Criteria for Ambulatory Colectomy

Seija Maniskas, MD, MS; Dena Nasir, MD; Allison McCurdy, MD;
Juliane Y. Golan, MD; Gabriel Akopian, MD; Howard S. Kaufman, MD

QI - Ambulatory Colectomy?



Patients identified using ICD 9/10 and CPT codes

Data collected (2017-21):

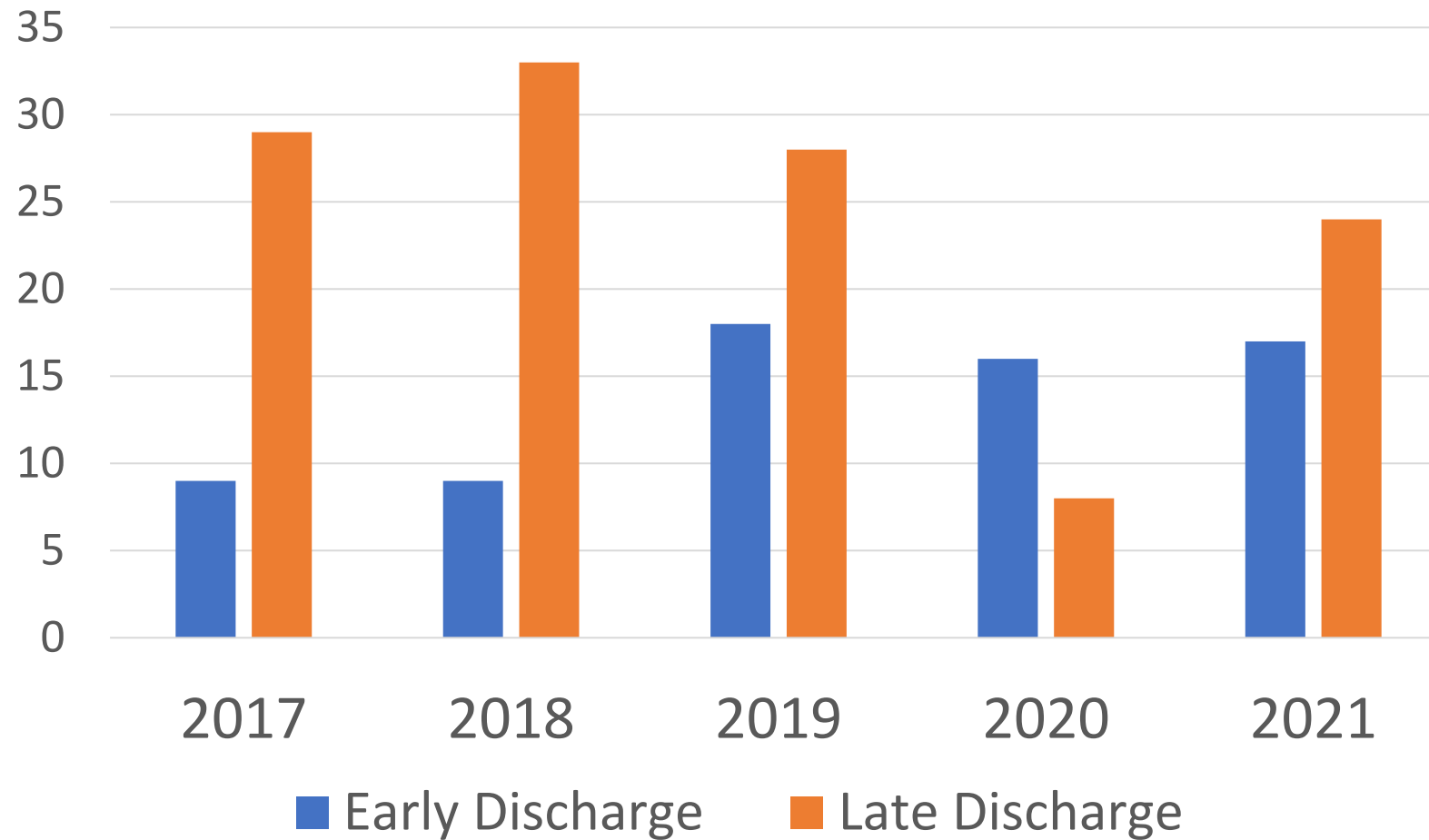
- Demographics
- Disease data
- Procedural data
- Perioperative/Post operative data

Patients were divided into groups based on LOS:

Early Discharge ($\leq 2d$, n=70) vs. Late Discharge ($>2d$, n=125)

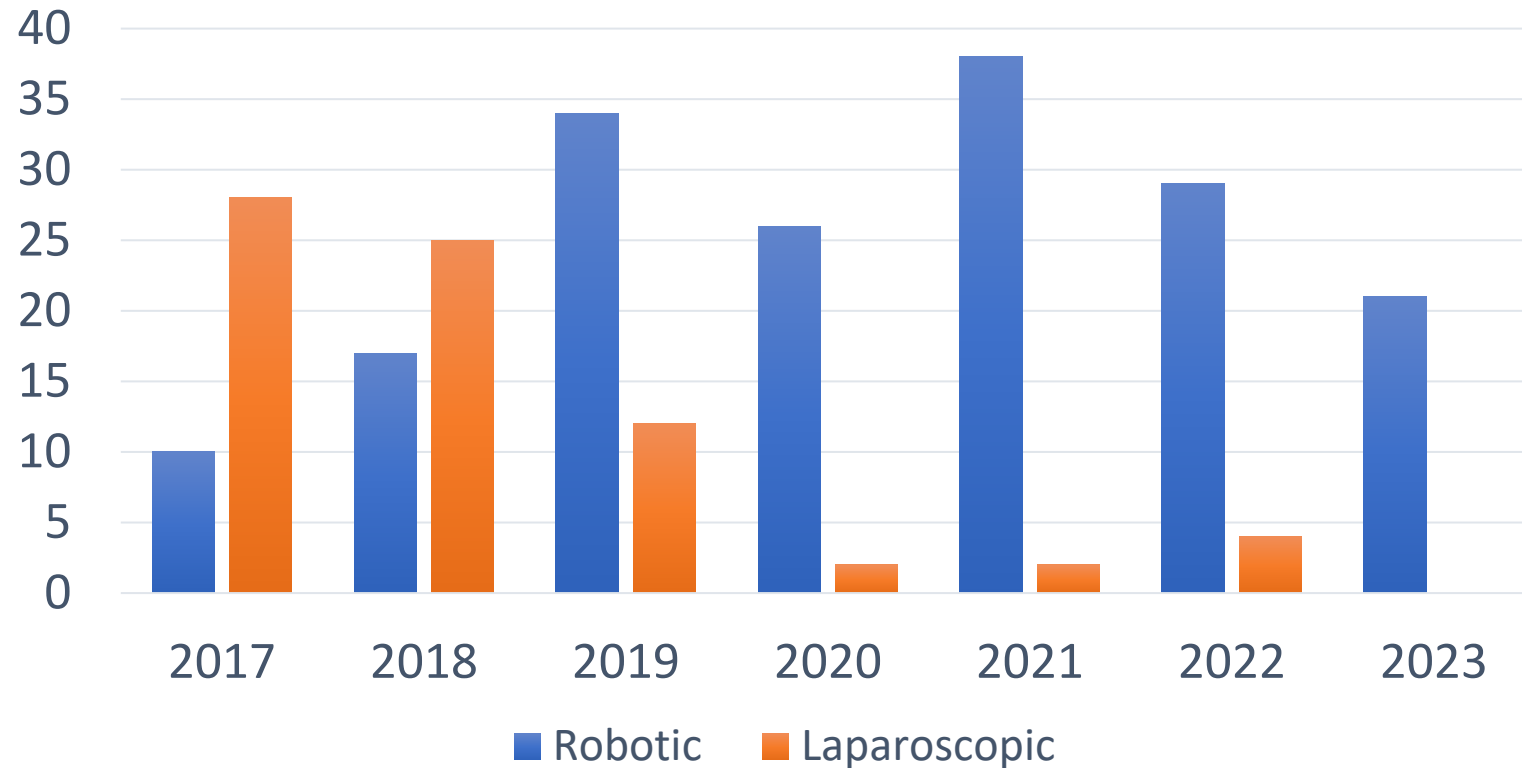
Analyses performed on SPSS

QI - Ambulatory Colectomy?



Robotics

Robotic Trend



Patient Factors

	Early Discharge (70)	Late Discharge (125)	
	<u>Median</u>	<u>Median</u>	<i>p</i>
Age	63	68	0.006
BMI	27	26	0.76
	<u>n (%)</u>	<u>n (%)</u>	
DM	10 (14)	25 (20)	.32
Prior Abdominal/Pelvic Surgery	27 (39)	63 (50)	.11
Steroids	2 (3)	1 (1)	.29
Anticoagulation	4 (6)	11 (9)	.58

Procedural Variables

	Early Discharge	Late Discharge	<i>p</i>
	n (%)	n (%)	
Robotic	55 (79)	71 (57)	0.002
Extraction Incision			0.004
Pfannenstiel	55 (79)	69 (55)	
Minilap (midline)	12 (17)	50 (40)	
Other	3 (4)	6 (5)	
Anastomosis			0.006
Intracorporeal	55 (79)	74 (59)	
Extracorporeal	15 (21)	51 (41)	
LOA	11 (15)	24 (19)	0.54

Procedural Variables

	Early Discharge	Late Discharge	
	n (%)	n (%)	<i>p</i>
TAP Block	59 (84)	97 (78)	0.26
Alvimopan	59 (84)	69 (55)	<.001
Physical Therapy	14 (20)	43 (34)	0.03
Complications	2 (3)	13 (10)	0.06
30 Day Readmission	3 (4)	4 (3)	0.68
	Median	Median	<i>p</i>
MME POD 0	16	20	0.19
MME POD 1	18	33	0.02
MME POD 0 + 1	39	55	0.02

Procedural Variables

	Early Discharge	Late Discharge	
	n (%)	n (%)	<i>p</i>
Side of Resection	18	18	0.33
Left	30 (43)	41 (33)	
Right	39 (56)	83 (66)	
Subtotal	1 (1)	1 (1)	
Surgeon Specialty			0.001
Colorectal	43 (61)	40 (32)	
Surgical Oncology	22 (31)	68 (54)	
Minimally Invasive	1 (1)	7 (6)	
General	4 (6)	10 (8)	
Intraoperative Complication	1 (1)	1 (1)	0.68

- ChatGPT (2019-2023)
 - Opioid prescription reduction ↓ 30-50%
 - Effective nonopioid analgesia
 - Fewer opioid-related side effects
 - Complication rates ↓ 40%
 - Pneumonia/DVT ↓ 5-10%
 - Average cost/patient ↓ 15-25%
 - Patient satisfaction ↑ 20% on average

- Common, 2nd leading US cause of cancer death
- Screening saves lives, multiple options exist
- Disparities in age, race, ethnicity
- System- and community-wide efforts are needed to address disparities
- Improved outcomes including opioid reduction through MIS and ERPs
- More to come...
 - Outpatient colorectal surgery?

1. What are the 2018 American Cancer Society recommendations for colorectal cancer screening for average-risk adults?

- a. Begin screening at age 50.
- b. Continue to screen all individuals after age 85.
- c. Screen all average-risk individuals age 45 to 75; selectively offer screening up to age 85.

2. Which is a true statement regarding colorectal cancer in younger individuals?

- a. Individuals born in the 1990s have an approximately fourfold greater risk of developing rectal cancer than those born in the 1950s
- b. A personal history of inflammatory bowel disease is not a risk factor for developing colorectal cancer
- c. Rectal bleeding in a young individual can always be attributed to hemorrhoids and does not require further evaluation.

3. Which outcomes should be tracked to identify disparities in colorectal cancer screening?

- a. Rates of screening participation, income, and geography
- b. Follow up for abnormal results
- c. Incidence of colorectal cancer by age, race, and ethnicity
- d. All of the above

4. Which of the following statements are true regarding enhanced recovery for colorectal surgery?

- a. Enhanced recovery pathways are limited to postoperative care
- b. Diets should be started only after the return of bowel sounds
- c. Multimodality pain management is associated with decreased postoperative opioid needs

References

- Bailey CE, Hu C, You YN, et al. Increasing Disparities in the Age-Related Incidences of Colon and Rectal Cancers in the United States, 1975-2010. *JAMA Surg.* 2015;150(1):17–22. doi:10.1001/jamasurg.2014.1756
- Ullah F, Pillai AB, Omar N, Dima D, Harichand S. Early-Onset Colorectal Cancer: Current Insights. *Cancers.* 2023; 15(12):3202. <https://doi.org/10.3390/cancers15123202>
- Wolf AMD, Fontham ETH, Church TR, Flowers CR, Guerra CE, LaMonte SJ, Etzioni R, McKenna MT, Oeffinger KC, Shih YT, Walter LC, Andrews KS, Brawley OW, Brooks D, Fedewa SA, Manassaram-Baptiste D, Siegel RL, Wender RC, Smith RA. Colorectal cancer screening for average-risk adults: 2018 guideline update from the American Cancer Society. *CA Cancer J Clin.* 2018 Jul;68(4):250-281. doi: 10.3322/caac.21457. Epub 2018 May 30. PMID: 29846947.
- Doubeni CA, Selby K, Gupta S. Framework and Strategies to Eliminate Disparities in Colorectal Cancer Screening Outcomes. *Annu Rev Med.* 2021 Jan 27;72:383-398. doi: 10.1146/annurev-med-051619-035840. Epub 2020 Nov 18. PMID: 33208026; PMCID: PMC7846969.
- Siegel RL, Wagle NS, Cercek A, Smith RA, Jemal A. Colorectal cancer statistics, 2023. *CA Cancer J Clin.* 2023; 73(3): 233-254. doi:[10.3322/caac.21772](https://doi.org/10.3322/caac.21772)
- Aziz Z, Wagner S, Agyekum A, et al. Cost-Effectiveness of Liquid Biopsy for Colorectal Cancer Screening in Patients Who Are Unscreened. *JAMA Netw Open.* 2023;6(11):e2343392. doi:10.1001/jamanetworkopen.2023.43392
- Grubbs SS, Polite BN, Carney J Jr, Bowser W, Rogers J, Katurakes N, Hess P, Paskett ED. Eliminating racial disparities in colorectal cancer in the real world: it took a village. *J Clin Oncol.* 2013 Jun 1;31(16):1928-30. doi: 10.1200/JCO.2012.47.8412. Epub 2013 Apr 15. PMID: 23589553; PMCID: PMC3661932.

Thank you!

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