High-value intervention: Providing colorectal cancer screening



Jaesu Han, MD

Associate Professor Department of Psychiatry and Behavioral Medicine University of California, Davis Sacramento, California

Cerrone Cohen, MD

Medical Instructor Department of Community and Family Medicine Department of Psychiatry Duke University School of Medicine Durham, North Carolina

Mortality from colorectal cancer is elevated among the mentally ill; watchfulness helps level risk

ancer screening is an important example of secondary prevention—the aim being to detect disease at an early stage, when treatment can prevent symptomatic disease. Over the years, screening tests for breast cancer, colorectal cancer (CRC), cervical cancer, and, most recently, lung cancer have been developed and recommended by the U.S. Preventive Services Task Force (USPSTF). Among breast cancer, cervical cancer, and CRC, the screening rate for CRC remains lowest, at 58.6%.¹

The importance of screening for CRC is highlighted by the facts that:

- CRC is the third most commonly diagnosed form of cancer in the United States among both men and women
- CRC is the second leading cause of cancer-related death.²

The overall decrease in the incidence of CRC in the United States has been credited to improvements in screening and removal of potentially precancerous lesions.³

Harmful disparity puts the mentally ill at exceptional risk

Screening patterns for CRC among patients with mental illness are poorly characterized, but it is known that the overall cancer screening rate among patients with severe psychiatric illness lags significantly behind the rate in the general population.^{4,5} In addition, studies have shown that mortality among patients with CRC who have a mental disorder is elevated, compared with CRC patients who do not have a psychiatric diagnosis.⁶

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7 Tools for colorectal cancer screening: Benefits and limitations

Test	Benefits	Limitations	
Highly sensitive	Bowel preparation is not	Poorly detects polyps	
fecal occult	required	Multiple stool specimens are required	
blood test	Test is completed at home	Higher rate of false positives than other tests	
	Inexpensive	Pre-test dietary limitations	
	Noninvasive	Colonoscopy is necessary if abnormalities are	
	Substantial research supports a	detected	
	reduction in cancer mortality		
Fecal	Bowel preparation is not	Poorly detects polyps	
test	Tequired	More expensive than traditional fecal occult	
	Noninvesive	Colonoscony is possessory if obnormalities are	
	Fewer dietany restrictions than	detected	
	fecal occult blood test		
	More specific for human blood		
	than guaiac-based tests		
Stool DNA test	Bowel preparation is not	Poorly detects polyps	
	required	More costly than other stool tests	
	Test is completed at home	Still being researched; uncertainty about	
	Noninvasive	adequate screening intervals	
	Only 1 stool specimen usually is	Colonoscopy is necessary if abnormalities are	
FIEXIDIE	Minimal bowel preparation is	Examines only the distal colon	
Significaceopy	Sedation is not required	Bowei preparation is required	
	Substantial research supports a		
	reduction in cancer mortality	Cannot remove large polyps	
	Lower risk of complications than		
	colonoscopy	abnormalities are detected	
	Can be performed by less-		
	specialized providers		
Colonoscopy	Examines entire colon	Full bowel cleansing is required	
	Polyps can be removed and biopsied	More expensive than stool testing	
	Can diagnose other colon	Requires sedation and someone to accompany patient on the day of the procedure	
	pathology	Patients might miss a day of work	
	Longest interval between screenings	Highest risk of complications when compared with other methods	
	Can remove potentially	No randomized trials documenting benefit on	
	precancerous polyps	mortality	
Double-contrast	Can view entire colon	Largely discarded in favor of newer methods	
barium enema	Sedation is not required	Full bowel preparation is required	
		Cannot remove polyps	
		Often misses small polyps	
		Exposes patients to radiation	
		Colonoscopy is necessary if abnormalities are detected	
CT colonography	Examines entire colon	Full bowel preparation is required	
0, , ,	Performance is similar to that of optical colonoscopy for large	Cannot remove polyps	
		Exposes patients to radiation	
	polyps and invasive cancers	Expensive	
	Few complications	May not be readily available in smaller centers	
	Sedation is not required	or rural areas	
	Noninvasive	Colonoscopy is necessary if abnormalities are	
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Common modifiable risk factors for colorectal cancer include obesity, smoking, and alcohol consumption

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Why this disparity? It might be that CRC is more likely to be diagnosed at an advanced stage among these patients, or that they are less likely to receive cancer treatment after diagnosis, or are more likely to have a longer delay between diagnosis and initial treatment than patients who do not have a psychiatric diagnosis.⁷

Regardless, psychiatric practitioners can make a significant impact on reducing this health disparity by leveraging their unique therapeutic relationship to educate patients about screening options and dispel myths about cancer screening. In this article, we outline practical strategies for CRC screening and weigh the advantages and disadvantages for the use of several tools and guidelines in psychiatric patients.

What is the pathogenesis of colorectal cancer?

Most cases of CRC evolve from *polyps*, abnormal growths on the lining of the colon or rectum. Constituting an estimated 96% of all polyps, *adenomas* are by far the most common form in the colon and rectum.

Adenomas also are most likely to transform over time to dysplasia, and then to progress to cancer.⁸ Although all adenomas have malignant potential, <10% evolve to adenocarcinoma. This proposed adenoma→carcinoma sequence is not well understood; however, it is known that CRC usually develops slowly—over 10 to 15 years.⁹ Detection and removal of adenomas and treatable, localized carcinomas form the basis of screening for CRC.

Risk factors for colorectal cancer

A number of risk factors for CRC have been identified.

Specific heritable conditions, such as Lynch syndrome and familial adenomatous polyposis, pose the greatest risk of CRC, particularly at younger ages and compared with people without such a history.¹⁰

Family history. One of the strongest risk factors for CRC remains a family history of

the disease. People who have a first-degree relative with a diagnosis of CRC are at 2 to 3 times the risk of CRC, compared with people without a family history of the disease. This risk increases further if multiple family members are affected or if the diagnosis was made in a relative at a young age.^{11,12}

Other non-modifiable risk factors include a personal history of inflammatory bowel disease, type 2 diabetes mellitus, male sex, African American heritage, and increasing age.¹³⁻¹⁵

Common modifiable risk factors include obesity, smoking, and alcohol consumption.¹⁶⁻¹⁸

What is the role of screening?

CRC screening is only appropriate for patients who are asymptomatic. CRC generally is asymptomatic in early stages. Prognosis also is most favorable when CRC is detected in the asymptomatic stage.

As lesions of CRC grow, the presentation might include hematochezia, melena, abdominal pain, weight loss, occult anemia, constipation or diarrhea, and changes in stool caliber.¹⁹ These signs and symptoms are not highly specific for CRC, however, and might be indicative of other gastrointestinal pathology, including inflammatory bowel disease, diverticulitis, irritable bowel syndrome, infectious colitis, hemorrhoids, and mesenteric ischemia.

Symptomatic patients should be referred directly for diagnostic evaluation. Colonoscopy with biopsy is the standard for diagnosing CRC. Once a diagnosis of CRC is made, patients should be referred to a specialist to discuss treatment; options largely depend on the stage of the cancer at diagnosis.

What screening tests are available?

Unlike screening for other cancers, there are a number of reasonable options for CRC screening; *Table 1*¹⁵ (*page 35*) compares their relative pros and cons. Each test has its benefits and drawbacks, allowing the



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Each screening test has benefits and drawbacks, allowing the screening strategy to be customized by patient preference and characteristics



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Refer patients who are symptomatic for colorectal cancer directly for diagnostic evaluation with colonoscopy/biopsy

- Box Key points: When to begin screening for colorectal cancer

- Average-risk patients: Begin colorectal cancer (CRC) screening at age 50
- Higher-risk patients (history of a hereditary syndrome associated with CRC, radiation exposure, history of CRC or inflammatory bowel disease, multiple first-degree relatives with CRC, family history of CRC at a young age): Refer before age 50 to establish an individualized timeline for screening
- Patients (any age) with signs or symptoms suspicious for CRC: Refer directly for a diagnostic workup; these patients are not a candidate for CRC screening

screening strategy to be customized based on patient preference and characteristics, but this variability also can lead to confusion by patient and provider about those options.

Stool-based tests detect trace amounts of blood from early-stage treatable cancers. Highly sensitive fecal occult blood testing (FOBT) has been shown specifically to decrease mortality from CRC.²⁰ Stoolbased tests are inexpensive and noninvasive, but require:

- more frequent testing
- that the patient collect the stool specimen
- follow-up colonoscopy when test results are positive.

Endoscopic and imaging tests detect polyps and early-stage treatable cancers; all require some degree of bowel preparation, and some require sedation. Testing intervals vary but, as a group, are longer than the interval between stool-based tests because polyps grow slowly. Because colonoscopy with biopsy is the preferred screening method for diagnosing CRC, it is the only screening option that also is a diagnostic procedure.

Where can screening guidelines be found?

Several professional organizations have developed guidelines for CRC screening.

The 2 major U.S. guidelines come from USPSTF and a joint guideline from The American Cancer Society, Multi-Society Task Force, and American College of Radiology (ACS-MSTF-ACR).

An update to both guidelines was released in 2008. *Table* $2^{21,22}$ (*page* 46) summarizes their recommendations.

Both guidelines recommend that screening begin at age 50 (*Box*). The primary differences between the 2 guidelines lie in the scope of recommended options for screening and the time frame for discontinuing screening:

• **USPSTF** requires a higher level of evidence for screening options and limits recommended options to FOBT, sigmoidoscopy combined with FOBT, and colonoscopy.

• ACS-MSTF-ACR emphasizes options that detect premalignant polyps, and generally is more inclusive of testing options; it also delineates tests as useful for either (1) early detection of cancer (stool-based studies) or (2) cancer prevention (endoscopic and imaging tests).

On the question of when to *stop* screening, ACS-MSTF-ACR bases its recommendations on life expectancy; USPSTF sets a specific age for ending screening.^{21,22}

Recommendations of a third entity, the American College of Gastroenterology (ACG), are similar to those of ACS-MSTF-ACR; however, ACG (1) recommends beginning screening African American patients at age 45 because of their increased risk of CRC and (2) gives preference to colonoscopy as the preferred screening modality.²³

Guidelines vary for high-risk patients (those with a history of familial adenomatous polyposis or another inherited syndrome associated with CRC; those with a family history of CRC in the young; those with a history of radiation exposure, history of CRC, or inflammatory bowel disease; and those with several first-degree relatives with CRC). Patients who fall into any of these categories should be referred for specialty care to establish the time of initial screening and the interval of subsequent screening.



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For a patient with moderate or severe psychiatric symptoms, optimize treatment of the underlying disorder before starting screening

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– Table 2 A summary of 2 colorectal screening guidelines

Organization	When to begin screening	Method	Screening interval
USPSTF ^a	Age 50 (average-risk adults)	Highly sensitive guaiac FOBT Combined flexible sigmoidoscopy and highly sensitive guaiac FOBT Colonoscopy	Annually Sigmoidoscopy every 5 years, with FOBT every 3 years Every 10 years
ACS-MTF-ACR ^b	Age 50 (average-risk adults)	For early detection Highly sensitive guaiac FOBT FIT Stool DNA testing To prevent colorectal cancer Flexible sigmoidoscopy Colonoscopy CT colonography Double-contrast barium enema	Annually Annually Unknown Every 5 years Every 10 years Every 5 years Every 5 years

^aThis guideline does not apply to patients with inflammatory bowel disease or specific hereditary syndromes, such as familial adenomatous polyposis and Lynch syndrome

^bThis guidelines is designed for patients deemed to be at average risk of colorectal cancer

ACS-MTF-ACR: Joint guidelines from the American Cancer Society, the United States Multi-society Task Force for Colorectal Cancer, and the American College of Radiology; FIT: fecal immunochemical test; FOBT: fecal occult blood test; USPSTF: United States Preventive Task Force

Source: Adapted from references 21 and 22

CRC screening in the presence of psychiatric illness

Psychiatrists have an opportunity to support their patients when considering potentially confusing CRC screening recommendations. This opportunity might occur during a discussion about general preventive care, or a patient might come to an appointment after visiting a primary care provider, and ask for advice about screening options.

The potential benefits of CRC screening are negated if a patient is unable or unwilling to complete the test or undergo timely follow-up of positive results. It is important, therefore, to individualize screening recommendations—keeping in mind the degree of impairment from mental illness and the patient's preferences and reliability to engage in follow-up. To date, there are no agreed-on screening guidelines specifically for patients with comorbid mental illness.

Adapting USPSTF guidelines for CRC screening of average-risk patients with mental illness, we offer the following recommendations:

Recommend screening. Begin routine screening at age 50. Patients with well-controlled or mild symptoms should be screened with a stool study with or without flexible sigmoidoscopy. Stool studies are safe, noninvasive, and require no bowel preparation; when used alone, however, they need to be performed yearly.

Screening accuracy is increased when a stool-based test is combined with flexible sigmoidoscopy; screening then can be performed less often. Unlike colonoscopy, flexible sigmoidoscopy does not involve sedation; a high-functioning patient might find this appealing and tolerate the greater frequency of screening. On the other hand, some patients might not accept the inconvenience of collecting the stool sample with the kit provided and returning it to the lab for processing.

Manage psychiatric illness optimally. For a patient with moderate or severe psychiatric symptoms, first attempt to optimize treatment of the underlying

When to discontinuing screening

Between age 76 to 85, although this decision should be individualized, based on personal risk and health status

Do not screen after age 85

When life expectancy is <10 years

psychiatric condition before establishing a CRC screening program. If control of symptoms is likely to improve over the next 1 or 2 visits, it might be reasonable to defer screening until symptoms are better controlled and then reassess the patient before making specific screening recommendations. Screening should not be delayed, however, if significant improvement in symptoms is not expected in the near future. Lengthy delay might lead to failure in initiating screening at all.

We recommend that patients with persistent moderate or severe symptoms be screened with traditional colonoscopy. The sedation associated with colonoscopy (1) may be preferable to some patients with more severe illness and (2) allows for screening and diagnostic biopsy if needed during the same procedure. Screening with colonoscopy also:

• avoids the yearly adherence to a screening program that is needed with stool cards alone

• does not rely on patients collecting and returning stool kits for processing.

A potential challenge for patients with limited social support is the requirement to have someone accompany the patient on the day of colonoscopy.

Take steps to improve the screening rate. In addition to specific recommendations based on symptom severity, there are systems-level interventions that should be considered to improve the screening rate. These include:

• addressing transportation issues that are a barrier to screening

• considering the use of health navigators or peer advocates to help guide patients through the sometimes complex systems of care.

A more comprehensive systems-level intervention for mental health clinics that work primarily with persistent and severe mentally ill populations might include employing a care coordinator to organize referrals to primary care or even exploring *reverse integration*. In reverse integration, primary care providers co-locate within the mental health clinic, (1) allowing for "one-stop shopping" of mental health and primary care needs and (2) facilitating collaboration and shared treatment planning between primary care and mental health for complex patients.

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Potential benefits of screening are negated if a patient is unable or unwilling to complete the test or undergo timely follow-up of results



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A potential challenge for patients with limited social support is the need to have someone accompany them on the day of colonoscopy

Related Resources

- American Cancer Society. www.cancer.org.
- National Cancer Institute. Colorectal cancer–patient version. www.cancer.gov/types/colorectal.
- United States Preventive Services Task Force. www. uspreventiveservicestaskforce.org.
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Bottom Line

Screening tests for colorectal cancer (except colonoscopy) are useful for the asymptomatic stage only. Stool-based studies, flexible sigmoidoscopy, and colonoscopy are the best studied screening options, and vary in recommended screening intervals. The specific screening modality to recommend should (1) be individualized and (2) take into account patient preference, likelihood of adherence to the schedule, and how well psychiatric symptoms are controlled.