LabCorp

COLORECTAL CANCER PREVENTION, DETECTION, AND MANAGEMENT

Helping you provide better patient care

Colorectal cancer is the **third most commonly diagnosed cancer among women and men** in the United States, with 90% of cases occurring after age 50,¹ with a 4.5% (women) to 4.8% (men) chance of individuals, over their lifetime, developing colorectal cancer.²

Current guidelines emphasize routine screening beginning at age 45.³ According to the Centers for Disease Control and Prevention, screening is an important tool in earlier detection of colorectal and other cancer, "when cancers might respond better to treatment, thereby reducing deaths."⁴

The National Cancer Institute estimates more than 140,000 new cases of colorectal cancer will be diagnosed in 2018, while more than 50,000 people will die from the disease.⁵ In contrast, nearly twice the number of new breast cancer cases are estimated to be diagnosed, and 10,000 fewer deaths are anticipated compared to colorectal cancer.⁵ (See Table 1)

Colonoscopies are the screening standard for individuals at risk for colorectal cancer, performed once every 10 years.⁶

However, the invasiveness of a colonoscopy can result in patients refusing the procedure. One study found that reasons for individuals rejecting a colonoscopy included being uncomfortable with preparing for the test, as well as fearing that a colonoscopy would be painful. That same study found that 97% of patients refusing a colonscopy accepted a non-invasive colorectal cancer screening option.⁷

Non-invasive testing also provides another benefit: cost savings. A recent cost analysis by the Centers for Disease Control and Prevention found that, on average, FOBT/FIT testing was almost four times less costly than colonoscopies, which could "significantly impact the timeliness of the initial screen offered as a much larger number of individuals can be screened quicker."⁸

	Estimated New Cases⁵	Estimated Deaths⁵	Annual Screening Rates ⁸	5-year Survival Rate⁵
Breast	266,120	40,920	72%	89.7%
Colorectal	140,250	50,630	59%	64.5%

Table 1: Breast v. Colorectal Cancer: By The Numbers



Colorectal Cancer Prevention, Detection, and Management

LabCorp offers physicians colorectal cancer screening tools that are cost effective, easy to interpret, and provide high sensitivity and specificity.

An Improved Fecal Occult Blood Test (iFOBT)	An FDA-approved blood-based test for colorectal cancer screening
Test name: Occult Blood, Fecal, Immunoassay Test number: 182949	Test name: Epi ProColon [®] , Septin 9 Gene Methylation Detection Test number: 481160
 Annual screening with iFOBT has been shown in peer- reviewed literature to detect a majority of prevalent CRC in an asymptomatic population at the time of testing, and is an acceptable option for CRC screening in average-risk adults who are 45 years and older. No dietary or medicinal restrictions⁹ Lower GI specific¹⁰ Easy-to-use Greater sensitivity and specificity than GT¹⁰ LabCorp iFOBT: Sensitivity: 98.8% / Specificity 99.6%¹¹ Improves patient participation in screening¹⁰ Specific to human globin¹⁰ Annual screening test³ DRE sample indication⁹ 	 Epi proColon is a qualitative in vitro diagnostic method for the detection of methylated Septin 9 DNA in EDTA plasma derived from patient whole blood specimens. Methylation of the target DNA sequence in the promoter region of the SEPT9_v2 transcript has been associated with the occurrence of CRC. The test is indicated to screen adults of either sex, 50 years or older, defined as average risk for CRC, who have been offered and have a history of not completing CRC screening, and results should be used in combination with physician's assessment and individual risk factors in guiding patient management. Reported with sensitivity of 68.2%, 73.3% and 72.2%, and specificity of 78.8%, 81.5% and 80.8% when compared to fecal immunochemical test (FIT).¹² Option for a blood-based alternative to get a non-compliant patient tested. Tests that are available and recommended in the USPSTF 2008 CRC screening guidelines should be offered and declined prior to offering the Epi proColon® test. Patients with a positive Epi proColon® test result should be referred for diagnostic colonoscopy.

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