Colorectal Cancer Screening and Surveillance

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Colorectal cancer is the third most common cancer in men and women. The incidence and mortality rate of the disease have been declining over the past two decades because of early detection and treatment. Screening in persons at average risk should begin at 50 years of age; the U.S. Preventive Services Task Force recommends against routine screening after 75 years of age. Options for screening include high-sensitivity fecal occult blood testing annually, flexible sigmoidoscopy every five years with high-sensitivity fecal occult blood testing every three years, or colonoscopy every 10 years. In 2012, the U.S. Multi-Society Task Force on Colorectal Cancer updated its surveillance guidelines to promote the appropriate use of colonoscopy resources and reduce harms from delayed or unnecessary procedures; these guidelines provide recommendations for when to repeat colonoscopy based on findings. Adenomatous and serrated polyps have malignant potential and warrant early surveillance colonoscopy. Patients with one or two tubular adenomas that are smaller than 10 mm should have a repeat colonoscopy in five to 10 years. Repeat colonoscopy at five years is recommended for patients with nondysplastic serrated polyps that are smaller than 10 mm. Patients with three to 10 adenomas found during a single colonoscopy, an adenoma or serrated polyp that is 10 mm or larger, an adenoma with villous features or high-grade dysplasia, a sessile serrated polyp with cytologic dysplasia, or a traditional serrated adenoma are at increased risk of developing advanced neoplasia during surveillance and should have a repeat colonoscopy in three years. More than 10 synchronous adenomas warrant surveillance colonoscopy in less than three years. Colonoscopy may be repeated in 10 years if distal, small (less than 10 mm) hyperplastic polyps are the only finding. (Am Fam Physician. 2015;91(2):93-100. Copyright © 2015 American Academy of Family Physicians.)

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► Patient information: http://www.aafp.org/ afp/2015/0115/p93-s1.html.

olorectal cancer is the third most common cancer in men and women. Despite a reduction in incidence and mortality over the past two decades from early detection and treatment, an estimated 137,000 new diagnoses and 50,000 deaths were expected in 2014.1-3 Screening and surveillance detect cancer in its early stages when symptoms are not typically present.1 Approximately 30% of patients have nonmodifiable risk factors that increase their risk of colorectal cancer,4 including a family or personal history of colorectal cancer or advanced adenomas, personal history of inflammatory bowel disease, and personal history of hereditary polyposis syndromes.1 Modifiable risk factors include obesity, inactivity, smoking, and heavy alcohol use.5

This article reviews the 2012 U.S. Multi-Society Task Force on Colorectal Cancer consensus guidelines for surveillance of persons with colonoscopy findings, which aim to promote the appropriate use of colonoscopy resources and reduce harms from delayed or unnecessary procedures.⁶

Colorectal Cancer Screening

The U.S. Preventive Services Task Force and other organizations recommend that colorectal cancer screening begin at 50 years of age in persons at average risk.⁷⁻¹⁰ The American College of Gastroenterology recommends that black persons receive initial screening at 45 years of age because of an increased incidence of colorectal cancer in this population, although no patient-oriented outcome studies currently support this recommendation.^{8,11}

The U.S. Preventive Services Task Force recommends against routine screening after 75 years of age because the potential benefits of screening may be outweighed by harms and competing causes of mortality;

Table 1. Colorectal Cancer Screening Recommendations in Asymptomatic Adults at Average Risk

Organization	Screening test and interval	Patient age
U.S. Preventive Services Task Force*7	The following options are equally acceptable High-sensitivity FOBT annually Flexible sigmoidoscopy every 5 years with high-sensitivity FOBT every 3 years	Start at 50 years; individualize after 75 years
American College of Gastroenterology† ⁸	Colonoscopy every 10 years Preferred Colonoscopy every 10 years Fecal immunochemical test annually (if colonoscopy is declined) Alternative, prevention Flexible sigmoidoscopy every 5 to 10 years Computed tomography colonography every 5 years Alternative, cancer detection High-sensitivity FOBT annually	Start at 50 years, or 45 years in blacks
American Cancer Society, U.S. Multi-Society Task Force, American College of Radiology‡ ⁹	Stool DNA test every 3 years Tests that detect adenomas and cancer Flexible sigmoidoscopy every 5 years Colonoscopy every 10 years Double-contrast barium enema every 5 years Computed tomography colonography every 5 years Tests that primarily detect cancer High-sensitivity FOBT annually Fecal immunochemical test annually Stool DNA test, interval uncertain	Start at 50 years

NOTE: Table 2 summarizes screening recommendations based on risk factors.

FOBT = fecal occult blood testing.

†—A family history of small tubular adenomas in first-degree relatives is not considered to increase the risk of colorectal cancer. Patients with a family history of colorectal cancer in a single first-degree relative at 60 years or older can be screened on the same schedule as average-risk persons. ‡—Guidelines are not for those with a personal or family history of colorectal cancer or adenomas, inflammatory bowel disease, or high-risk genetic syndromes.

Information from references 7 through 9.

physicians and patients should individualize screening decisions after this age.7 Similarly, the U.S. Multi-Society Task Force believes the decision to continue surveillance should be based on a patient's estimated risk, benefit, and comorbidities.6 Table 1 summarizes screening recommendations for asymptomatic adults at average risk.7-9 Patients with a positive test result should undergo full visualization of the colon with colonoscopy.

Screening recommendations for patients with a family history of colorectal cancer vary based on the relative's relationship to the patient, findings, and age when the cancer was diagnosed (Table 2).8 Patients with a firstdegree relative who has had colorectal cancer or an advanced adenoma (i.e., an adenoma that is 10 mm or larger, has villous elements, or has high-grade dysplasia) diagnosed before 60 years of age, or two first-degree relatives with colorectal cancer or an advanced adenoma diagnosed at any age, should receive a screening colonoscopy every five years starting at 40 years of age or 10 years younger than the relative's age when diagnosed, whichever is earlier. Patients who have one first-degree relative who has had colorectal cancer or an advanced adenoma diagnosed at 60 years of age or older can be screened on the same schedule as average-risk individuals.

Guidelines for follow-up surveillance colonoscopy are summarized in Table 3.6,8,12

Colon Polyps

A colon polyp is a growth protruding into the lumen from the colonic mucosa. The

^{*—}Recommendations also apply to those with first-degree relatives with colorectal cancer or adenomas, but the guideline notes that earlier screening is reasonable in those with a first-degree relative who developed colorectal cancer at a young age. Recommendations exclude those with inherited syndromes or inflammatory bowel disease.

Table 2. Colonoscopy Screening Recommendations Based on Risk Factors

Risk factor	Age to initiate screening	Interval if normal (years)
Single first-degree relative with colorectal cancer or an advanced adenoma diagnosed at ≥ 60 years of age	50 years (may start at 45 years in blacks)	10
Single first-degree relative with colorectal cancer or an advanced adenoma diagnosed at < 60 years of age	40 years or 10 years younger than affected relative's age when diagnosed, whichever is earlier	5
Two first-degree relatives with colorectal cancer or an advanced adenoma diagnosed at any age	40 years or 10 years younger than the youngest affected relative's age when diagnosed, whichever is earlier	5

NOTE: An advanced adenoma is defined as an adenoma that is 10 mm or larger, has villous elements, or has highgrade dysplasia.

Information from reference 8.

two main categories of polyps are neoplastic and nonneoplastic. Hyperplastic polyp is the most common nonneoplastic polyp. Common neoplastic polyps include adenomas and serrated polyps.

HYPERPLASTIC POLYPS

Hyperplastic polyps (Figure 1), which account for up to 50% of sigmoid and rectal polyps, are typically small, sessile polyps that measure 1 to 5 mm in size.¹³ Because small (less than 10 mm) hyperplastic polyps in the rectum and sigmoid rarely exhibit dysplasia or develop into colon cancer, colonoscopy may be repeated in 10 years if they are the only finding. No increased risk has been shown three years after baseline colonoscopy in patients with hyperplastic polyps and coexisting adenomas.14 Therefore, small, distal hyperplastic polyps do not alter colonoscopy surveillance guidelines.

ADENOMAS

Adenomas are polyps that have malignant potential and require early surveillance colonoscopy.15 They are classified by their glandular histology and level of dysplasia, which determine their malignant potential and interval for repeat colonoscopy. Tubular adenomas (Figures 2 and 3) account for 80% of adenomas and have a malignant transformation rate at diagnosis of 4.8%. Tubulovillous and villous adenomas (Figure 4) are less common but have more malignant potential (19.0% for tubulovillous, 38.4% for villous).16 Although all adenomas are dysplastic, the degree of dysplasia is classified

as low or high grade. High-grade dysplasia (Figure 5) suggests that the polyp is evolving toward malignancy. An adenoma that is 10 mm or larger, has villous elements, or has high-grade dysplasia, or the presence of three or more adenomas during a single examination, has a strong association with advanced neoplasia on future colonoscopies. 6,17-19

SERRATED POLYPS

Sessile serrated polyps are thought to be the principal precursor of hypermethylated gene cancers; 20% to 30% of colorectal cancers can arise from this pathway.6 These polyps are often difficult to detect during colonoscopy because they can be flat and indiscrete, and have adherent mucus^{6,20} (Figures 6 and 7). All sessile serrated polyps require early follow-up colonoscopy; those that have cytologic dysplasia, are 10 mm or larger, or are located proximal to the sigmoid colon may be associated with a higher risk of developing cancer.6

Colonoscopy Surveillance After Polypectomy

Patients who have adenomas are more likely to have additional adenomas or colorectal cancer on subsequent examinations.6 Endoscopic follow-up of patients with adenomas is referred to as a surveillance colonoscopy. The presence of low- or high-risk adenomas determines the surveillance interval.6

LOW-RISK POLYPS

Low-risk polyps include one or two small (less than 10 mm) tubular adenomas or serrated

Table 3. Guidelines for Follow-Up Surveillance Colonoscopy

Initial colonoscopy findings	Follow-up interval
Normal ⁸	
No polyps or normal biopsy results	10 years
Hyperplastic polyps ⁶	
Small (< 10 mm) hyperplastic polyps in rectum or sigmoid	10 years
Low-risk polyps ⁶	
1 or 2 small (< 10 mm) tubular adenomas	5 to 10 years
Small sessile serrated polyp (< 10 mm) without dysplasia	5 years
High-risk polyps ⁶	
3 to 10 tubular adenomas	3 years
Tubular adenoma or serrated polyp that is \geq 10 mm	
Adenoma with villous features or high-grade dysplasia	
Sessile serrated polyp with cytologic dysplasia	
Traditional serrated adenoma	
Other circumstances ⁶	
More than 10 adenomas	< 3 years
Serrated polyposis syndrome*	1 year
Following piecemeal removal of a large (> 15 mm) sessile adenoma or serrated polyp	Consider repeat in < 1 year if question of residual polyp
Following curative resection of colorectal cancer ¹²	1 year after resection, then 3 and 5 years if normal

^{*—}Criteria for serrated polyposis syndrome: at least 5 serrated polyps proximal to the sigmoid with 2 or more that are > 10 mm, any serrated polyp proximal to sigmoid with a family history of serrated polyposis syndrome, or > 20 serrated polyps of any size throughout the colon.

Information from references 6, 8, and 12.

polyps without cytologic dysplasia. Patients with low-risk tubular adenomas should have a repeat colonoscopy in five to 10 years. Evidence suggests that a surveillance interval of more than five years for low-risk tubular adenomas is reasonable in most patients; however, a five-year interval is prudent if the bowel preparation is inadequate or the cecum is not reached during colonoscopy. ^{17,18,21,22} Repeat colonoscopy at five years is recommended for patients with nondysplastic serrated polyps that are smaller than 10 mm. ²⁰

HIGH-RISK POLYPS

High-risk polyps include three to 10 tubular adenomas found during a single colonoscopy, at least one tubular adenoma or serrated polyp that is 10 mm or larger, at least one adenoma with villous features or high-grade



Figure 1. Hyperplastic polyp.



Figure 2. Tubular adenoma (pedunculated).



Figure 3. Tubular adenoma (sessile).



Figure 4. Tubulovillous adenoma.



Figure 5. Tubular adenoma with high-grade dysplasia.

dysplasia, a sessile serrated polyp with cytologic dysplasia, or a traditional serrated adenoma. Surveillance colonoscopy is recommended at three years for high-risk polyps.^{6,17,18}

The U.S. Multi-Society Task Force recommends surveillance colonoscopy earlier than three years in some situations. Patients with more than 10 synchronous adenomas warrant surveillance colonoscopy in less than three years. MUTYH mutation analysis or genetic consultation should also be considered to evaluate for MUTYHassociated polyposis, an autosomal recessive disease associated with increased risk of colorectal cancer.23 The U.S. Multi-Society Task Force has also released guidelines on the genetic evaluation and management of Lynch syndrome, an autosomal dominant condition and the most common cause of inherited colorectal cancer.24 Patients diagnosed with serrated polyposis syndrome should have surveillance colonoscopy in one year because of a significant risk of colorectal cancer.6,25 A repeat colonoscopy should be considered in less than one year for a large (more than 15 mm) sessile adenoma or serrated polyp removed by piecemeal resection if there is concern about incomplete removal. A repeat colonoscopy in two to six months is no longer required for a polyp removed by piecemeal resection if the endoscopist is certain that the resection was complete.6

The second surveillance interval relies on findings from the baseline and first surveillance colonoscopy (*Figure 8*).⁶ Patients with a history of low-risk adenomas and negative findings on their first surveillance colonoscopy are very low risk.⁶ Patients with high-risk adenomas on baseline colonoscopy



Figure 6. Sessile serrated polyp.

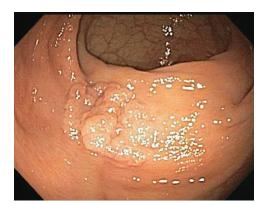


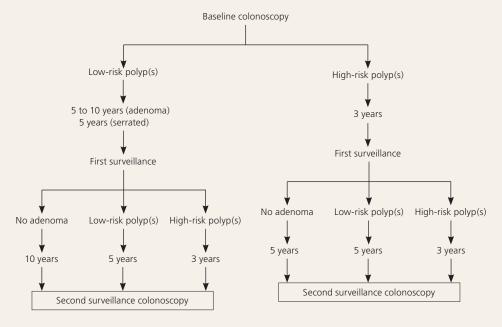
Figure 7. Traditional serrated adenoma.

should continue to have shorter follow-up intervals. ^{26,27}

Quality of Colonoscopy

The recommended colorectal cancer screening and surveillance intervals assume that high-quality colonoscopy has been performed. There is a direct relationship between published quality indicators for colonoscopy and interval cancer risk.^{28,29} Interval cancers are less likely when colonoscopy is performed by endoscopists with high cecal intubation and high adenoma detection rates.30,31 Highquality endoscopists are expected to reach the cecum in 95% of screening procedures performed, and detect adenomas in 15% of women and 25% of men undergoing screening colonoscopy after 50 years of age.²⁸ A repeat colonoscopy should be considered in one year for patients with a poor bowel preparation because this is associated with missed polyps.6 The use of fecal occult blood testing between surveillance colonoscopies is currently not recommended, but it requires further study. 32,33 If a patient has a positive finding on fecal occult blood testing, the decision to perform an early colonoscopy

Surveillance Colonoscopy



NOTE: Low-risk = 1 or 2 tubular adenomas (< 10 mm) or sessile serrated polyp (< 10 mm) without dysplasia; high-risk = 3 to 10 adenomas, tubular adenoma or serrated polyp \geq 10 mm, adenoma with villous features or high-grade dysplasia, a sessile serrated polyp with cytologic dysplasia, or a traditional serrated adenoma.

Figure 8. Algorithm for surveillance colonoscopy.

Information from reference 6.

should be based on the quality of the previous colonoscopy.

Surveillance After Colorectal Cancer

If colorectal cancer is found during colonoscopy, the remainder of the colon should be viewed proximal to the cancer before surgery. If there is an obstructing cancer, computed tomography colonography with

intravenous contrast media or a double-contrast barium enema should be used to view the proximal colon followed by a colonoscopy three to six months after resection. After curative resection of the cancer, patients should undergo colonoscopy one, three, and five years after the initial colonoscopy if findings on these surveillance colonoscopies remain normal.¹² Intervals may be shortened after the one-year examination based on adenomatous findings or hereditary causes of colon cancer.

BEST PRACTICES IN PREVENTIVE MEDICINE: RECOMMENDATIONS FROM THE CHOOSING WISELY CAMPAIGN

Recommendation Sponsoring organization Do not recommend screening for breast or American Geriatrics colorectal cancer, nor prostate cancer (with Society the prostate-specific antigen test) without considering life expectancy and the risks of testing, overdiagnosis, and overtreatment. Do not repeat colorectal cancer screening (by American Gastroenterological any method) for 10 years after a high-quality colonoscopy is negative in average-risk Association individuals.

Source: For more information on the Choosing Wisely Campaign, see http://www.choosingwisely.org. For supporting citations and to search Choosing Wisely recommendations relevant to primary care, see http://www.aafp.org/afp/recommendations/search.htm.

Adherence to Guidelines

Adherence to surveillance guidelines is variable with reports of overutilization in low-risk groups and underutilization in high-risk groups.³⁴ Following evidence-based guidelines can help reduce the cost and risk of unnecessary procedures, and prevent cancer in high-risk individuals. The Choosing Wisely campaign aims to promote patient-physician conversations by helping patients choose care that is supported by evidence, not duplicative of other tests or procedures already received, free from harm, and truly necessary.³⁵ Through

Clinical recommendation	Evidence rating	References
Colorectal cancer screening should begin at 50 years of age in average-risk individuals.	А	7-9
The American College of Gastroenterology recommends that colorectal cancer screening begin at 45 years of age in black patients.	С	8
Average-risk patients with normal findings on colonoscopy should have repeat colonoscopy in 10 years.	C	8
Patients with small, distal hyperplastic polyps are considered to have a normal colonoscopy result and should have repeat colonoscopy in 10 years.	С	6, 14
Patients with 1 or 2 small (< 10 mm) tubular adenomas should have repeat colonoscopy in 5 to 10 years.	C	6, 17, 18, 21, 2
Patients with small (< 10 mm) serrated polyps without dysplasia should have repeat colonoscopy in 5 years.	C	6, 20
Patients with 3 to 10 tubular adenomas, a tubular adenoma or serrated polyp ≥ 10 mm, an adenoma with villous features or high-grade dysplasia, a sessile serrated polyp with cytologic dysplasia, or a traditional serrated adenoma should have repeat colonoscopy in 3 years.	С	6, 17, 18

A = consistent, good-quality patient-oriented evidence; B = inconsistent or limited-quality patient-oriented evidence; C = consensus, diseaseoriented evidence, usual practice, expert opinion, or case series. For information about the SORT evidence rating system, go to http://www.aafp. org/afpsort.

Choosing Wisely, the American Gastroenterological Association recommends against repeating colorectal cancer screening (by any method) for 10 years after a negative colonoscopy result, and screening no earlier than five years after removal of a low-risk adenoma in asymptomatic patients.³⁶

Data Sources: A PubMed search was completed in Clinical Queries using the key terms colonic carcinoma, colonic neoplasms, colonoscopy, and colorectal neoplasms. The search included meta-analyses, randomized controlled trials, controlled trials, and reviews. Searches were also performed using Essential Evidence Plus, the Cochrane database, the National Guideline Clearinghouse database, the Trip Database, DynaMed, and the Agency for Healthcare Research and Quality evidence reports. Search dates: November 2, 2013, and October 4, 2014.

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