

Disclaimer: *This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.*

TABLE OF CONTENTS

Average Risk	Page 2
Increased Risk	Page 3
High Risk	Page 4
Suggested Readings	Page 5
Development Credits	Page 6

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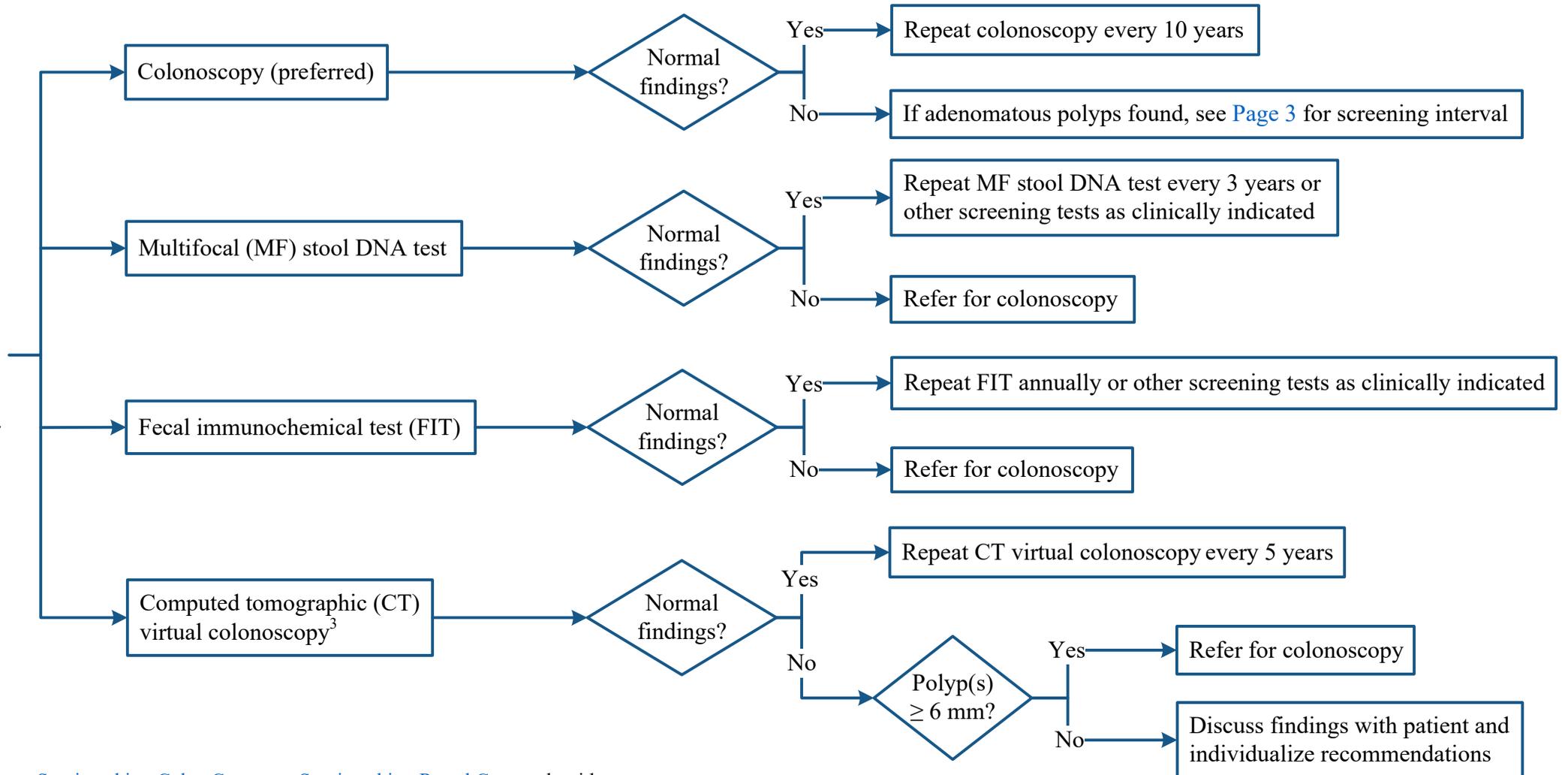
Note: This algorithm is not intended for individuals with a personal history of colorectal cancer¹.

Screening for adults ages 76-85 years old should be evaluated on an individual basis to assess the risks and benefits of screening. Colorectal cancer screening is not recommended over age 85 years.

PRESENTATION

RECOMMENDED SCREENING INTERVAL²

- Average Risk:
- Age ≥ 45 years
 - No personal history of adenoma, inflammatory bowel disease, or family history of colorectal cancer



¹ See the [Colon Cancer](#), [Rectal Cancer](#), [Survivorship - Colon Cancer](#), or [Survivorship - Rectal Cancer](#) algorithms

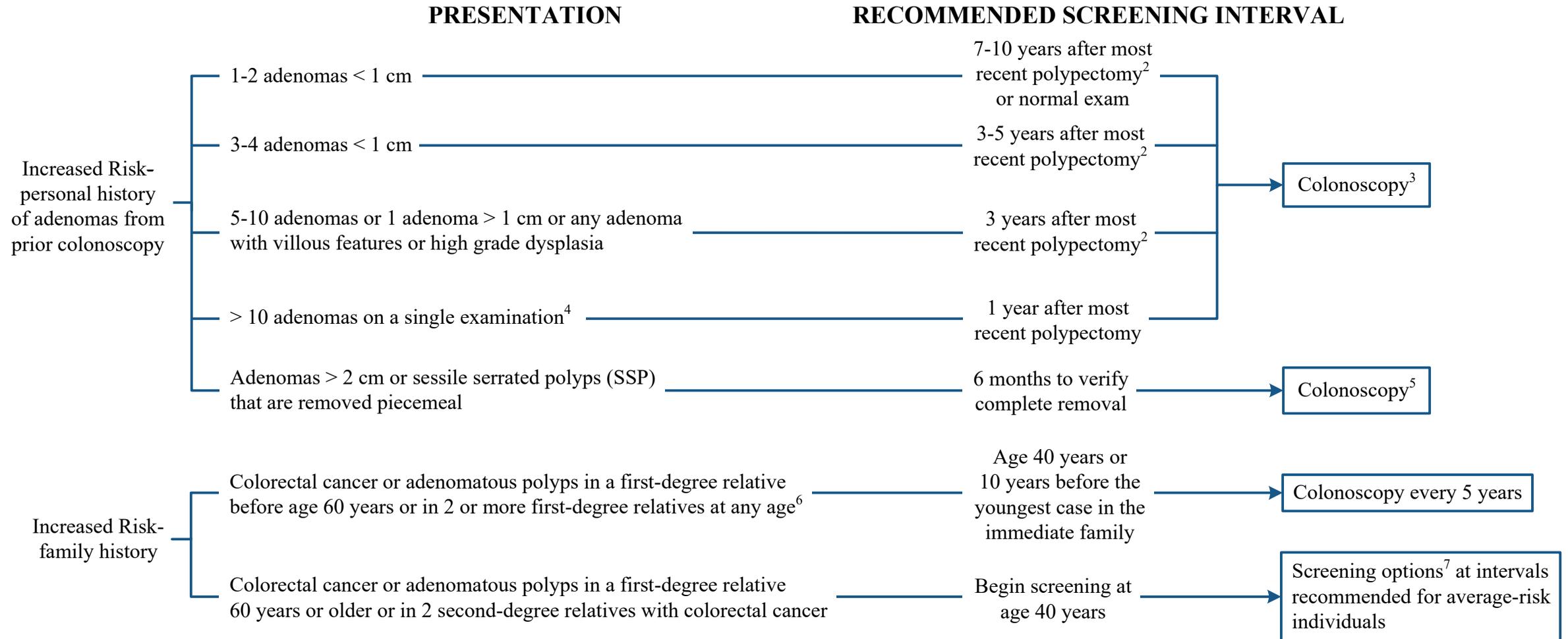
² Because there are multiple options for screening, the choice of a particular screening modality should include a conversation with the patient concerning their preference and availability. Colonoscopy and CT virtual colonoscopy are utilized in identification and removal of precancerous polyps. FIT and MF stool DNA are utilized in early detection of colon cancer.

³ Preauthorization with patient’s insurance carrier is advised

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¹ See the [Colon Cancer, Rectal Cancer, Survivorship - Colon Cancer](#), or [Survivorship - Rectal Cancer](#) algorithms

² Precise timing based on clinical factors, patient and physician preference

³ Subsequent follow-up is based on the number and size of polyps at the time of colonoscopy as well as the degree of dysplasia. If negative for adenomatous polyps, follow-up in 5 years is recommended.

⁴ Genetic testing for a polyposis syndrome is recommended for patients with > 10 adenomas

⁵ Surveillance individualized based on Endoscopist’s judgment

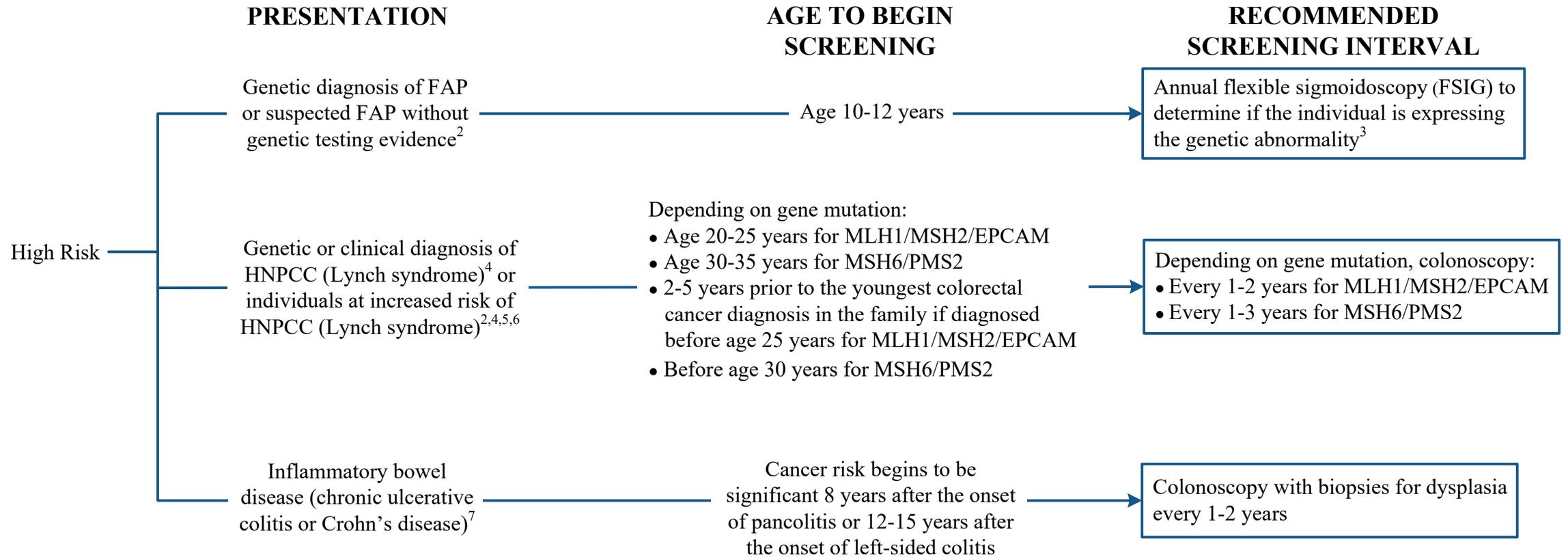
⁶ Counseling to consider genetic testing. See the [Genetic Counseling algorithm](#) for additional information.

⁷ Individuals may be screened with any recommended form of testing

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FAP = familial adenomatous polyposis
 HNPCC = hereditary nonpolyposis colorectal cancer

¹ See the [Colon Cancer, Rectal Cancer, Survivorship - Colon Cancer](#), or [Survivorship - Rectal Cancer](#) algorithms

² Counseling to consider genetic testing. See the [Genetic Counseling algorithm](#) for additional information.

³ If the genetic test is positive, colectomy should be considered

⁴ Lynch syndrome represents a heterogeneous group depending on the specific genetic alteration. Screening and surveillance should be individualized based on expert consultation, including review by a Genetic Counselor.

⁵ First degree relative of known mutation carriers, obligate carriers of a family history concerning for HNPCC (Lynch syndrome)

⁶ Genetic testing for HNPCC (Lynch syndrome) should be offered to first-degree relatives of persons with a known inherited MMR gene mutation. It should also be offered when the family mutation is not known, but 1 of the first 3 of the [modified Bethesda Criteria](#) is present.

⁷ These patients are best referred to a center with experience in the surveillance and management of inflammatory bowel disease

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SUGGESTED READINGS

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DEVELOPMENT CREDITS

This screening algorithm is based on majority expert opinion of the Colorectal Cancer Screening workgroup at the University of Texas MD Anderson Cancer Center. It was developed using a multidisciplinary approach that included input from the following:

Core Development Team Leads

Therese Bevers, MD (Cancer Prevention)
Robert Bresalier, MD (Gastroenterology, Hepatology & Nutrition)
Gottumukkala Raju, MD (Gastroenterology, Hepatology & Nutrition)
Mehnaz Shafi, MD (Gastroenterology, Hepatology & Nutrition)
David Vining, MD (Abdominal Imaging)

Workgroup Members

Heather Alexander Dahl, PgDip, BA (Community Alliances)
Joyce Dains, DrPH, JD, DNur, FNP-BCNAP (Nursing)
Wendy Garcia, BS♦
Ernest Hawk, MD, MPH (Cancer Prevention)
Thoa Kazantsev, MSN, RN, OCN♦
Scott Kopetz, MD, PhD (Gastrointestinal Medical Oncology)
Brittnee Macintyre, MSN, APRN, FNP-C♦
Maureen Mork, MS (Cancer Genetics)
Ana Nelson, DNP, RN, FNP (Cancer Prevention)
Eduardo Vilar Sanchez, MD, PhD (Cancer Prevention)
Brian Weston, MD (Gastroenterology, Hepatology & Nutrition)
Tonya Whitlow, MSPA (Gastroenterology, Hepatology & Nutrition)

♦ Clinical Effectiveness Development Team