

Do any of the following apply to your patient?

Patient history

- Colorectal, uterine, pancreatic, ovarian, stomach, or triple-negative breast cancer at any age
- Other cancers diagnosed under age 50
- Cancer diagnosed at any age in the context of significant family history (see right)
- Bilateral or multiple primary cancers
- Rare cancer^a
- Ashkenazi Jewish ancestry^b

Family history

- **≥2 relatives** with related cancers,^c with 1 diagnosed under age 50
- **≥3 relatives** with related cancers^c on the same side of the family
- Relative with a known familial variant
- Ashkenazi Jewish ancestry^b

^a Eg. Male With: breast cancer, metastatic prostate cancer, colorectal or uterine cancer with abnormal microsatellite instability/immunohistochemistry (MSI/IHC), pheochromocytoma, paraganglioma, 10 or more gastrointestinal polyps

^b Increased risk for specific *BRCA1*, *BRCA2*, *APC*, *CHEK2*, *GREM1*, *MSH2*, and *MSH6* mutations

^c For a complete list of related cancer types, see the Hereditary Cancer Reference Guide

Choosing the right test: A quick guide to help with collaborative, informed decision-making

History consistent with multiple cancer syndromes **or** history not explained by previous genetic testing

Comprehensive Hereditary Cancer Panel

66 genes, including high-risk, moderate-risk, and emerging genes associated with a broad spectrum of hereditary cancers

Guideline-Based Hereditary Cancer Panel

32 genes associated with a broad spectrum of hereditary cancers; all genes have management recommendations for 1 or more cancer sites

History includes primarily 1 cancer type

Targeted Panels

Hereditary Breast Cancer Panel:

18 genes associated with increased risk for breast cancer

Hereditary Colorectal Cancer Panel: 20 genes

associated with increased risk for colorectal cancer

Hereditary Endocrine Cancer Panel: 12 genes

associated with increased risk for paragangliomas, pheochromocytomas, and endocrine cancer

History is suspicious for a well-characterized hereditary cancer syndrome

Additional hereditary cancer risk tests

Syndrome-specific tests that analyze genes associated with well-characterized cancer syndromes, such as tuberous sclerosis complex (TSC), Lynch syndrome, familial adenomatous polyposis (FAP), and others

Patient has a relative with a familial mutation

Hereditary cancer, single-site test

Only looks for variant previously identified in a relative

For assistance, contact Genomic Client Services at **1.866.GENE.INFO (1.866.436.3463)** or **QuestHereditaryCancer.com**

The information presented is not intended to be a complete source for cancer risk in any patient. The patient's physician must determine the patient's risk based upon the patient's clinical assessment, history, and the physician's training and experience.

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