Identifying the right patient for hereditary cancer testing

Do any of the following apply to your patient?

Patient history

- Cancer 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

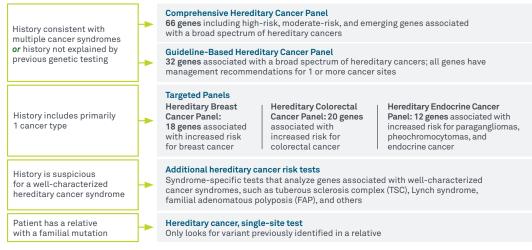
- ≥2 relatives with related cancer,° with 1 diagnosed under age 50
- ≥3 relatives with related cancer^c on the same side of the family
- Relative with a known familial mutation
- Ashkenazi Jewish ancestry^b

^a Male breast cancer, metastatic prostate cancer, ovarian cancer, pancreatic 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* and *BRCA2* mutations

 $^\circ\mbox{For a complete list of related cancer types, see the Hereditary Cancer Reference Guide$





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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