

GENETICS & WOMEN'S HEALTH

BRCAssure® BRCA1 and BRCA2 analysis

Because knowledge is a powerful tool



Hereditary breast and ovarian cancer

Patients with BRCA mutations are at increased risk for breast, ovarian and other cancers.

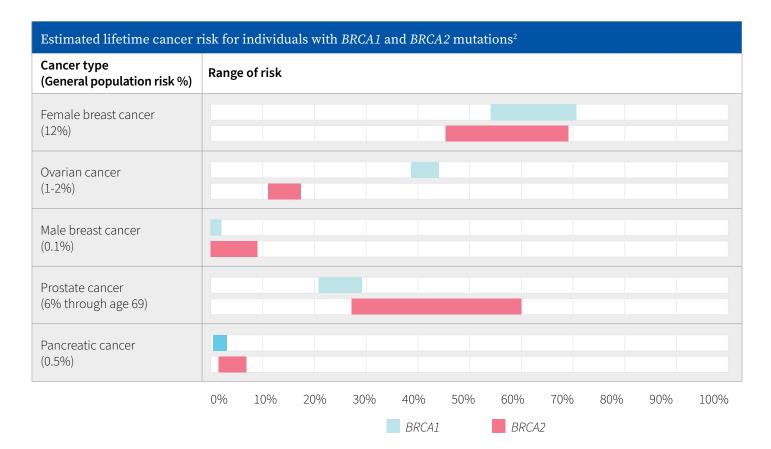
- Mutations in BRCA1 and BRCA2 account for the vast majority of families with hereditary breast and ovarian cancer syndrome (HBOC).¹
- The estimated lifetime risk of breast cancer in women with *BRCA1* mutations is 55%-72% by age 70; for *BRCA2*, the risk is 45%-69%.²
- The risk of ovarian cancer is 39%-44% for a woman with *BRCA1* and 11%-17% with *BRCA2*.²
- Both men and women with mutations in BRCA1/2 may also be at increased risk of other cancers, including pancreatic, prostate and melanoma.²

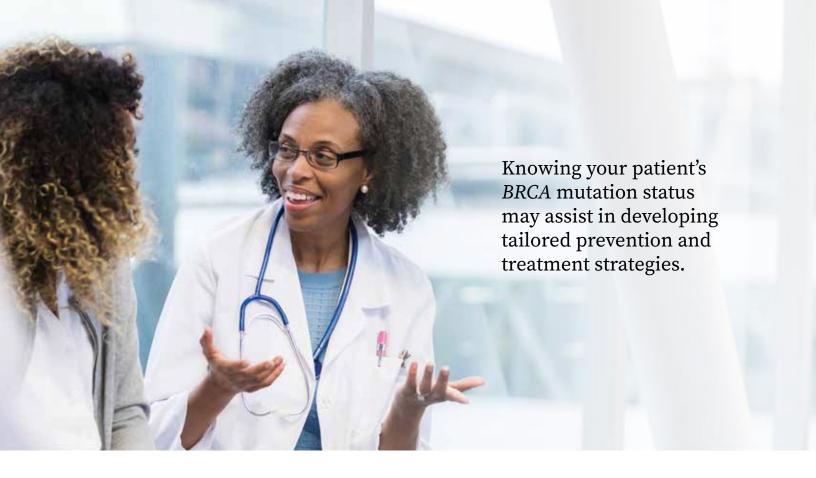


Approximately 1 in 40 Ashkenazi Jewish individuals carries one of three founder mutations.²



The overall prevalence of BRCA1 and BRCA2 mutations in the general population is estimated at 1 in 400^2 and varies with ethnicity.





Indications for BRCAssure testing^{1,3}

Consider BRCAssure testing for your patients if their personal or family history has one of the following:

- Breast cancer diagnosed at age 50 or younger
- Ovarian cancer
- Both breast and ovarian cancer
- Multiple primary breast cancers
- Male breast cancer

- Triple-negative breast cancer
- Pancreatic cancer
- Metastatic prostate cancer
- A previously identified BRCA1 or BRCA2 mutation in the family
- Ashkenazi Jewish ancestry
- Three or more relatives with breast cancer at any age

Notes: "Breast cancer" includes both invasive and ductal carcinoma in situ (DCIS). "Ovarian cancer" includes epithelial ovarian cancer, fallopian tube and primary peritoneal cancer.³ Complete Guidelines may be found at www.nccn.org; www.ACOG.org.

NCCN® is a trademark owned by the National Comprehensive Cancer Network, Inc.

Genetic counseling is highly recommended when BRCA testing is offered and after results are provided¹

Labcorp provides nationwide access to genetic counseling expertise through our telegenetic counseling program and provides convenience to patients with online scheduling. Your patients can quickly schedule an appointment at womenshealth.labcorp.com/genetic-counseling.

Call **855-GC-CALLS** (**855-422-2557**) to learn more about our exceptional services or visit womenshealth.labcorp.com/genetic-counseling.

Test/Panel Name	Test No.
BRCAssure: Comprehensive BRCA1/2 Analysis	485030
BRCAssure: Ashkenazi Jewish Panel	485097
BRCAssure: BRCA1 Targeted Analysis	485066
BRCAssure: BRCA2 Targeted Analysis	485081



Specimen requirements:

7 mL whole blood lavender-top (EDTA) tube

2 mL saliva Oragene®•Dx saliva collection kit

A continuity of care, pioneering science, professional service

We provide the scientific expertise you need and the customer experience patients want.

Rapid results

Samples have a typical turnaround time of 3 weeks after a test arrives at our lab.

Extensive managed care contracts

Help patients maximize their benefits. Where applicable, Labcorp can initiate prior authorization for hereditary cancer testing with or without a specimen.

Convenient blood draws

We have a nationwide network of patient service centers, allowing for convenient access to sample collection. Visit labcorp.com to find your nearest location.

Confirmatory testing for direct-to-consumer results

Our hereditary cancer portfolio provides the testing options needed to confirm patient results from direct-to-consumer genetics laboratories as appropriate.

Women's health and wellness

Labcorp provides access to a complete women's health offering featuring a full range of testing and services that support the continuity of care for your patients through a single laboratory.

References

- 1. American College of Obstetricians and Gynecologists. Hereditary Breast and Ovarian Cancer Syndrome. ACOG Practice Bulletin. No. 182. Obstet Gynecol. 2017; Sep;130(3):e-110-e126.
- 2. Petrucelli, N et al. BRCA1 and BRCA2 Hereditary Breast and Ovarian Cancer. Gene Reviews. Available at: http://www. ncbi.nlm.nih.gov/books/NBK1247. Accessed November 9, 2022.
- $3.\,Genetic/Familial\,High-Risk\,Assessment:\,Breast,\,Ovarian,\,and\,Pancreatic.\,NCCN\,Guidelines\,Version\,1.2023.\,Available$ at: http://www.nccn.org/professionals/physician_gls/pdf/genetics_screening.pdf. Accessed October 4, 2022.

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