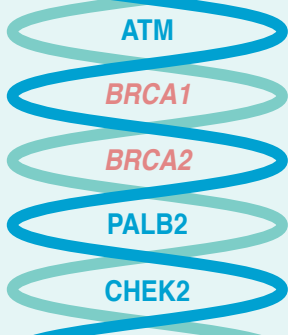


Understanding Hereditary Breast Cancer

Some breast cancers are caused by genes with mutations, or changes, passed down from either parent to their daughters or sons.

Genetic testing at any stage, even when the disease has spread beyond the breast to other parts of the body, known as metastatic disease, can determine if breast cancer is hereditary. Genetic test results and other disease variables, including breast cancer subtype, are important factors in helping inform treatment decisions.

Some genes linked to hereditary breast cancer include:



As of 2017, *BRCA1* and *BRCA2* mutations are the most common cause of hereditary breast cancer, responsible for approximately:



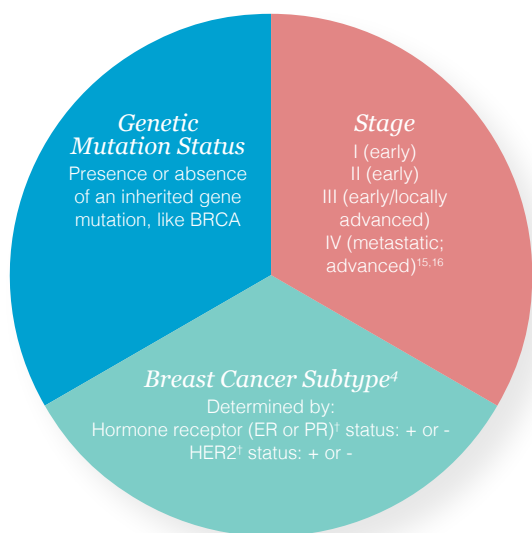
3-6% of all breast cancers¹⁻⁴

25-30% of hereditary breast cancer cases⁵

People with inherited mutations in *BRCA1/2* genes are often younger than the overall breast cancer population, and can be diagnosed in their 30s-40s.⁶

Know the Status

Understanding the *hereditary status, subtype, and stage* of breast cancer can help patients and their physicians make informed decisions:



Any breast cancer at any stage or subtype can have an inherited gene mutation

Know the Risk Factors

According to epidemiologic studies and national guidelines, people with any of these *criteria* should speak with a cancer genetics professional* about getting a test to see if their breast cancer carries a hereditary mutation⁷⁻⁹:

Age

Diagnosed with breast cancer at age 50 or earlier

Multiple Breast Cancers

Cancer in both breasts or a second cancer in the same breast

Triple-Negative Breast Cancer

Diagnosed with triple-negative (ER-/PR-/HER2-)[†] breast cancer at age 60 or earlier

Recurrent Or Metastatic Breast Cancer

People with an inherited *BRCA1* or *BRCA2* mutation may be eligible for a certain type of targeted therapy

Heritage

Ashkenazi (Eastern European) Jewish, African American, or Hispanic heritage, among others⁹

Family History

Relatives with male breast cancer, ovarian, pancreatic, or metastatic prostate cancer, or breast cancer diagnosed at an early age

*These are just some of the criteria that should be considered. Individuals should consult their healthcare provider or a cancer genetics professional for more information.

[†]ER: estrogen receptor; PR: progesterone receptor; HER2: human epidermal growth factor receptor 2

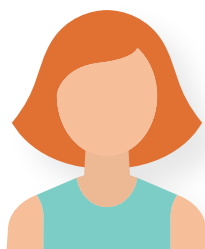
Know the Emotional Impact

It is important to know if breast cancer is hereditary to help people living with breast cancer *think through emotional considerations*, including^{10,11}:

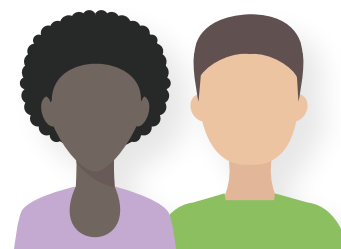
If, when, and how to share genetic test results with loved ones



Whether or not to encourage family members to be tested



Anxiety about the potential health and emotional impact on family



There is a 50% chance... that first-degree relatives—*children, siblings, or a parent of a person with hereditary breast cancer*—have the same mutated gene, increasing their risk of getting breast or other types of cancer.¹² Other, more distant relatives also are at risk of carrying the mutation.¹³



Speaking with a *genetic counselor, medical geneticist, trained physician, or advanced practice professional* can help patients at risk for hereditary breast cancer*, make decisions about genetic testing.¹⁴

These individuals can also help people who test positive for a hereditary mutation feel empowered, cope with the results and determine a treatment plan.

*Please refer to the "Know the Risk Factors" section.

To learn more about metastatic breast cancer, visit www.StoryHalfTold.com

References

1. Fasching PA, Hu C, Hart SN, et al. Cancer predisposition genes in metastatic breast cancer - association with metastatic pattern, prognosis, patient and tumor characteristics. Paper presented at: San Antonio Breast Cancer Symposium; December 5-9, 2017; San Antonio, TX. **2.** Tung N, Lin NU, Kidd J, et al. Frequency of germline mutations in 25 cancer susceptibility genes in a sequential series of patients with breast cancer. *J Clin Oncol.* 2016;34(13):1460-1468. **3.** Nelson HD, Fu R, Goddard K, et al. Risk assessment, genetic counseling, and genetic testing for BRCA-related cancer: systematic review to update the U.S. preventive services task force recommendation. Report No. 12-05164-EF-1. Rockville, MD: Agency for Healthcare Research and Quality; 2013. <https://www.ncbi.nlm.nih.gov/pubmed/24432435>. Accessed September 2020. **4.** Meynard G, Villanueva C, Thiery-Vuillemin A, et al. 284P Real-life study of BRCA genetic screening in metastatic breast cancer. *Ann Oncol.* 2017;28(5). doi:10.1093/annonc/mdx365.047. **5.** Kleibl Z, Kristensen VN. Women at high risk of breast cancer: molecular characteristics, clinical presentation and management. *The Breast.* 2016;28:136-144. **6.** Kuchenbaecker KB, Hopper JL, Barnes DR, et al. Risks of breast, ovarian, and contralateral breast cancer for BRCA1 and BRCA2 mutation carriers. *JAMA.* 2017;317(23):2402-2416. doi:10.1001/jama.2017.7112. **7.** Referenced with permission from the NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®) for Genetic/Familial High Risk Assessment: Breast, Ovarian and Pancreatic V.1.2021. © National Comprehensive Cancer Network, Inc. 2020. All rights reserved. Accessed September 2020. To view the most recent and complete version of the guideline, go online to NCCN.org. NCCN makes no warranties of any kind whatsoever regarding their content, use or application and disclaims any responsibility for their application or use in any way. **8.** Referenced with permission from the NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®) for Breast Cancer V.6.2020. © National Comprehensive Cancer Network, Inc. 2020. All rights reserved. Accessed September 2020. To view the most recent and complete version of the guideline, go online to NCCN.org. NCCN makes no warranties of any kind whatsoever regarding their content, use or application and disclaims any responsibility for their application or use in any way. **9.** National Cancer Institute. Genetics of breast and gynecologic cancers (PDQ®) - health professional version. <https://www.cancer.gov/types/breast/hp/breast-ovarian-genetics-pdq/>. Accessed September 2020. **10.** O'Neill SC, Mays D, Patenaude AF. Women's concerns about the emotional impact of awareness of heritable breast cancer risk and its implications for their children. *J Community Genet.* 2015;6:55-62. **11.** Gallo AM, Angst DB, Knapp KA. Disclosure of genetic information within families. *Am J Nurs.* 2009;109(4):65-69. **12.** National Cancer Institute. BRCA mutations: Cancer risk and genetic testing. <https://www.cancer.gov/about-cancer/causes-prevention/genetics/brca-fact-sheet>. Accessed September 2020. **13.** Senter L, O'Connor M, Oriyo F, Sweet K, Toland AE. Linking distant relatives with BRCA gene mutations: potential for cost savings. *Clin Genet.* 2014;85(1):54-58. **14.** American College of Surgeons. Cancer program standards 2012: ensuring patient-centered care. Version 1.2. <https://www.facs.org/-/media/files/quality%20programs/cancer/coc/programstandards2012.ashx>. Accessed September 2020. **15.** American Joint Committee on Cancer. Breast cancer staging. <http://cancerstaging.org/references-tools/quickreferences/documents/breastmedium.pdf>. Accessed April 6, 2017. **16.** Early Stage Breast Cancer, Union for International Cancer Control 2014 Review of Cancer Medicines on the WHO List of Essential Medicines https://www.who.int/selection_medicines/committees/expert/20/applications/EarlyStageBreast.pdf?ua=1#:~:text=Executive%20Summary,distant%20metastatic%20disease%20are%20not. Accessed September 2020.