Breast cancer

Breast cancer is a disease in which certain cells in the breast become abnormal and multiply uncontrollably to form a tumor. Although breast cancer is much more common in women, this form of cancer can also develop in men. In both women and men, the most common form of breast cancer begins in cells lining the milk ducts (ductal cancer). In women, cancer can also develop in the glands that produce milk (lobular cancer). Most men have little or no lobular tissue, so lobular cancer in men is very rare.

In its early stages, breast cancer usually does not cause pain and may exhibit no noticeable symptoms. As the cancer progresses, signs and symptoms can include a lump or thickening in or near the breast; a change in the size or shape of the breast; nipple discharge, tenderness, or retraction (turning inward); and skin irritation, dimpling, or scaliness. However, these changes can occur as part of many different conditions. Having one or more of these symptoms does not mean that a person definitely has breast cancer.

In some cases, cancerous tumors can invade surrounding tissue and spread to other parts of the body. If breast cancer spreads, cancerous cells most often appear in the bones, liver, lungs, or brain. Tumors that begin at one site and then spread to other areas of the body are called metastatic cancers.

A small percentage of all breast cancers cluster in families. These cancers are described as hereditary and are associated with inherited gene mutations. Hereditary breast cancers tend to develop earlier in life than noninherited (sporadic) cases, and new (primary) tumors are more likely to develop in both breasts.

Frequency

Breast cancer is the second most commonly diagnosed cancer in women. (Only skin cancer is more common.) About one in eight women in the United States will develop invasive breast cancer in her lifetime. Researchers estimate that more than 230,000 new cases of invasive breast cancer will be diagnosed in U.S. women in 2015.

Male breast cancer represents less than 1 percent of all breast cancer diagnoses. Scientists estimate that about 2,300 new cases of breast cancer will be diagnosed in men in 2015.

Particular gene mutations associated with breast cancer are more common among certain geographic or ethnic groups, such as people of Ashkenazi (central or eastern European) Jewish heritage and people of Norwegian, Icelandic, or Dutch ancestry.
Causes

Cancers occur when a buildup of mutations in critical genes—those that control cell growth and division or repair damaged DNA—allow cells to grow and divide uncontrollably to form a tumor. In most cases of breast cancer, these genetic changes are acquired during a person's lifetime and are present only in certain cells in the breast. These changes, which are called somatic mutations, are not inherited. Somatic mutations in many different genes have been found in breast cancer cells. Less commonly, gene mutations present in essentially all of the body's cells increase the risk of developing breast cancer. These genetic changes, which are classified as germline mutations, are usually inherited from a parent. In people with germline mutations, changes in other genes, together with environmental and lifestyle factors, also influence whether a person will develop breast cancer.

Some breast cancers that cluster in families are associated with inherited mutations in particular genes, such as \textit{BRCA1} or \textit{BRCA2}. These genes are described as "high penetrance" because they are associated with a high risk of developing breast cancer, ovarian cancer, and several other types of cancer in women who have mutations. Men with mutations in these genes also have an increased risk of developing several forms of cancer, including breast cancer. The proteins produced from the \textit{BRCA1} and \textit{BRCA2} genes are involved in fixing damaged DNA, which helps to maintain the stability of a cell's genetic information. They are described as tumor suppressors because they help keep cells from growing and dividing too fast or in an uncontrolled way. Mutations in these genes impair DNA repair, allowing potentially damaging mutations to persist in DNA. As these defects accumulate, they can trigger cells to grow and divide without control or order to form a tumor.

A significantly increased risk of breast cancer is also a feature of several rare genetic syndromes. These include Cowden syndrome, which is most often caused by mutations in the \textit{PTEN} gene; hereditary diffuse gastric cancer, which results from mutations in the \textit{CDH1} gene; Li-Fraumeni syndrome, which is usually caused by mutations in the \textit{TP53} gene; and Peutz-Jeghers syndrome, which typically results from mutations in the \textit{STK11} gene. The proteins produced from these genes act as tumor suppressors. Mutations in any of these genes can allow cells to grow and divide unchecked, leading to the development of a cancerous tumor. Like \textit{BRCA1} and \textit{BRCA2}, these genes are considered "high penetrance" because mutations greatly increase a person's chance of developing cancer. In addition to breast cancer, mutations in these genes increase the risk of several other types of cancer over a person's lifetime. Some of the conditions also include other signs and symptoms, such as the growth of noncancerous (benign) tumors.

Mutations in dozens of other genes have been studied as possible risk factors for breast cancer. These genes are described as "low penetrance" or "moderate penetrance" because changes in each of these genes appear to make only a small or moderate contribution to overall breast cancer risk. Some of these genes provide instructions for making proteins that interact with the proteins produced from the
BRCA1 or BRCA2 genes. Others act through different pathways. Researchers suspect that the combined influence of variations in these genes may significantly impact a person's risk of developing breast cancer.

In many families, the genetic changes associated with hereditary breast cancer are unknown. Identifying additional genetic risk factors for breast cancer is an active area of medical research.

In addition to genetic changes, researchers have identified many personal and environmental factors that contribute to a person's risk of developing breast cancer. These factors include gender, age, ethnic background, a history of previous breast cancer, certain changes in breast tissue, and hormonal and reproductive factors. A history of breast cancer in closely related family members is also an important risk factor, particularly if the cancer occurred in early adulthood.

**Inheritance Pattern**

Most cases of breast cancer are not caused by inherited genetic factors. These cancers are associated with somatic mutations in breast cells that are acquired during a person's lifetime, and they do not cluster in families.

In hereditary breast cancer, the way that cancer risk is inherited depends on the gene involved. For example, mutations in the BRCA1 and BRCA2 genes are inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to increase a person's chance of developing cancer. Although breast cancer is more common in women than in men, the mutated gene can be inherited from either the mother or the father.

In the other syndromes discussed above, the gene mutations that increase cancer risk also have an autosomal dominant pattern of inheritance. It is important to note that people inherit an increased likelihood of developing cancer, not the disease itself. Not all people who inherit mutations in these genes will ultimately develop cancer.

In many cases of breast cancer that clusters in families, the genetic basis for the disease and the mechanism of inheritance are unclear.

**Other Names for This Condition**

- breast cancer, familial
- breast carcinoma
- cancer of breast
- malignant neoplasm of breast
- malignant tumor of breast
- mammary cancer
Diagnosis & Management

Genetic Testing Information

- What is genetic testing?
  
- American Society of Breast Surgeons: Consensus Guideline on Genetic Testing for Hereditary Breast Cancer

- Genetic Testing Registry: Breast cancer, familial male

- Genetic Testing Registry: Breast cancer, susceptibility to

- Genetic Testing Registry: Familial cancer of breast

Research Studies from ClinicalTrials.gov

- ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22breast+cancer%22

Other Diagnosis and Management Resources

- GeneReview: BRCA1- and BRCA2-Associated Hereditary Breast and Ovarian Cancer
  https://www.ncbi.nlm.nih.gov/books/NBK1247

- GeneReview: Hereditary Diffuse Gastric Cancer
  https://www.ncbi.nlm.nih.gov/books/NBK1139

- GeneReview: Li-Fraumeni Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1311

- GeneReview: Peutz-Jeghers Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1266

- GeneReview: PTEN Hamartoma Tumor Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1488

- Genomics Education Programme (UK): Hereditary Breast and Ovarian Cancer

- National Cancer Institute: Breast Cancer Risk Assessment Tool
  https://bcrisktool.cancer.gov/

- National Cancer Institute: Breast Cancer Treatment (PDQ®)–Patient Version
  https://www.cancer.gov/types/breast/patient/breast-treatment-pdq
• National Cancer Institute: Genetic Testing for BRCA1 and BRCA2: It's Your Choice
• National Cancer Institute: Genetic Testing for Hereditary Cancer Syndromes

Additional Information & Resources

Health Information from MedlinePlus

• Encyclopedia: Breast Cancer
  https://medlineplus.gov/ency/article/000913.htm
• Health Topic: Breast Cancer
  https://medlineplus.gov/breastcancer.html
• Health Topic: Male Breast Cancer
  https://medlineplus.gov/malebreastcancer.html

Genetic and Rare Diseases Information Center

• Familial breast cancer
  https://rarediseases.info.nih.gov/diseases/10415/familial-breast-cancer
• Metaplastic carcinoma of the breast

Additional NIH Resources

• National Cancer Institute: Breast Cancer Treatment (PDQ®)–Patient Version
  https://www.cancer.gov/types/breast/patient/breast-treatment-pdq
• National Cancer Institute: Genetics of Breast and Gynecologic Cancers (PDQ®)–Health Professional Version
• National Human Genome Research Institute: Learning About Breast Cancer
  https://www.genome.gov/Genetic-Disorders/Breast-Cancer
• National Institute of Environmental Health Sciences: Breast Cancer Risk and Environmental Factors
Educational Resources

- Cancer.Net: Breast Cancer in Men
  https://www.cancer.net/cancer-types/breast-cancer-men

- Centers for Disease Control and Prevention: Does Breast or Ovarian Cancer Run In Your Family?
  https://www.cdc.gov/Features/HereditaryCancer/

- Centre for Genetics Education

- Genetic Science Learning Center, University of Utah
  https://learn.genetics.utah.edu/content/disorders/multifactorial/

- MalaCards: breast cancer
  https://www.malacards.org/card/breast_cancer

- Merck Manual Consumer Version
  https://www.merckmanuals.com/home/women-s-health-issues/breast-disorders/breast-cancer

- Orphanet: Hereditary breast cancer
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=227535

- The Sister Study: A Study of the Environmental and Genetic Risk Factors for Breast Cancer
  https://sisterstudy.niehs.nih.gov/English/index1.htm

- U.S. Preventive Services Task Force: Risk Assessment, Genetic Counseling, and Genetic Testing for BRCA-related Cancer in Women
  https://www.uspreventiveservicestaskforce.org/Home/GetFileByID/1030

Patient Support and Advocacy Resources

- American Cancer Society: Breast Cancer

- Breast Cancer Resources Directory
  http://www.cancerindex.org/clinks3.htm

- Breastcancer.org
  https://www.breastcancer.org/

- CancerCare
  https://www.cancercare.org/

- Centers for Disease Control and Prevention: National Breast and Cervical Cancer Early Detection Program
  https://www.cdc.gov/cancer/nbccedp/
• FORCE (Facing Our Risk of Cancer Empowered)
https://www.facingourrisk.org/index.php

• Living Beyond Breast Cancer
https://www.lbbc.org

• National Coalition for Cancer Survivorship
https://www.canceradvocacy.org/

• Susan G. Komen
https://ww5.komen.org/

• Susan G. Komen: Breast Cancer in Men

Clinical Information from GeneReviews
• BRCA1- and BRCA2-Associated Hereditary Breast and Ovarian Cancer
https://www.ncbi.nlm.nih.gov/books/NBK1247

• Hereditary Diffuse Gastric Cancer
https://www.ncbi.nlm.nih.gov/books/NBK1139

• Li-Fraumeni Syndrome
https://www.ncbi.nlm.nih.gov/books/NBK1311

• Peutz-Jeghers Syndrome
https://www.ncbi.nlm.nih.gov/books/NBK1266

• PTEN Hamartoma Tumor Syndrome
https://www.ncbi.nlm.nih.gov/books/NBK1488

Scientific Articles on PubMed
• PubMed
https://www.ncbi.nlm.nih.gov/pubmed?term=%28Breast+Neoplasms%5BMAJR%5D%29+AND+%28%28breast+cancer%5BTI%5D%29+AND+%28hereditary%5BTIAB%5D%29+AND+review%5Bpt%5D+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1800+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM
• BREAST CANCER
http://omim.org/entry/114480

• BREAST-OVARIAN CANCER, FAMILIAL, SUSCEPTIBILITY TO, 1
http://omim.org/entry/604370

• BREAST-OVARIAN CANCER, FAMILIAL, SUSCEPTIBILITY TO, 2
http://omim.org/entry/612555
Medical Genetics Database from MedGen

- Breast cancer, familial male  
- Hereditary Breast Carcinoma  

Sources for This Summary

- American Cancer Society: Cancer Facts and Statistics  
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/23586058
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3618918/
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25467110
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/23747889
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/24116874
- National Cancer Institute: Genetics of Breast and Gynecologic Cancers (PDQ®)–Health Professional Version  
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22957996
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301425
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25605744
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4478970/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25186627

Reprinted from Genetics Home Reference:

Reviewed: May 2015
Published: September 10, 2019

Lister Hill National Center for Biomedical Communications
U.S. National Library of Medicine
National Institutes of Health
Department of Health & Human Services