

 **Hereditary Breast & Ovarian Cancer:** Family history clinical decision support tool for risk assessment and management

**Average risk** screening with mammography is recommended for all women between the ages of 40 and 74, every 2-3 years, following a discussion of benefits and harms, as part of shared decision making. [[Canadian Taskforce for Preventative Health Care, 2024\(draft\)](#)]

Those with **additional** breast cancer **risk factors** (e.g. extremely dense breast tissue (ACR D), family history of breast/ovarian cancer) could be considered for modified breast cancer screening e.g. supplemental ultrasound, MRI, or mammography beginning at a younger age or arranged more frequently. [Check your local breast cancer screening program for more.](#)

For those in Ontario, find the high-risk [Ontario Breast Screening Program requisition here.](#)

Individuals who meet one or more of the criteria below may be eligible for **high-risk** breast cancer screening with MRI, genetic counselling and/or genetic testing. The eligibility for high-risk screening and/or genetic testing is determined following a genetic assessment.

**Risk assessment:**

Consider referral for a genetics assessment if your patient has a personal and/or family history of at least one of the following:

- Breast cancer at a young age ( $\leq 45$  years)
- Breast cancer  $\leq 50$  years **and** a limited family structure (e.g. adoption, few close relatives)
- Breast cancer  $\leq 50$  years of age **and** a second primary breast cancer
- Triple negative breast cancer diagnosed  $\leq 60$  years of age
- Male breast cancer at any age
- Ovarian cancer at any age (excluding borderline, pure clear cell, mucinous and low malignant potential tumours)
- Multiple primaries in the same individual e.g. breast and ovarian cancer
- Breast **or** ovarian cancer at any age **AND**  $\geq 1$  close relative(s)\* diagnosed with a related cancer, including breast (one diagnosis should be  $< 50y$ ), ovarian, pancreatic or high-risk prostate cancer<sup>+</sup>

\*Closely related individuals typically refers to first- and second-degree blood relatives on the same side of the family (e.g. parent, sibling, offspring, grandparent, aunt/uncle)

<sup>+</sup>Confirmed with evidence of one or more of the following features: T3 (or higher) staging, Grade Group 4 or 5 (Gleason Score 8 to 10), lymph node involvement, PSA  $\geq 20$

- Breast **or** ovarian cancer at any age **AND**  $\geq 2$  close relatives diagnosed with breast cancer or prostate cancer at any age

**Ashkenazi Jewish ancestry AND:**

- Personal history of breast cancer, colorectal cancer and/or GI polyposis at any age
- Family history on the Ashkenazi Jewish side, that includes  $\geq 1$  first- or second-degree relative(s) with breast cancer, ovarian cancer, gastrointestinal polyposis at any age, or colorectal cancer diagnosed  $< 60$  years
- A blood relative with a known pathogenic/likely pathogenic genetic variant in a cancer susceptibility gene *Report or documentation containing specific gene and variant should ideally be sent with referral.*
- An established risk model calculating a  $\geq 5\%$  chance to carry a *pathogenic/likely pathogenic* variant in a gene associated with HBOC (e.g. *BRCA1, BRCA2, CHEK2, PALB2, ATM*) using family and personal history

**Resources**



[GECKO point of care tool](#) for Hereditary Breast and Ovarian cancer

[Provincial breast cancer screening programs](#)

Find the contact information for your local [Genetics Centre here](#)

More on breast density at [Dense Breast Canada.org](#)