

Clinical Cancer Genetics Clinical Referral Guidelines

Breast Center 2011 Update

Strongly encourage referral:

Patients with any of the following:

- Breast cancer diagnosed ≤ 50 years of age
- Breast cancer diagnosed at any age, AND one or more of the following:
 - Personal history of ovarian cancer
 - Family history* of ovarian cancer
 - Family history of breast cancer diagnosed ≤ 50 years of age
 - Family history of male breast cancer
 - Family history of pancreas cancer
 - Personal or family history of follicular thyroid cancer, endometrial cancer, and/or dermatologic manifestations of Cowden syndrome, germline PTEN mutation
 - Personal or family history of sarcoma, adrenocortical cancer, brain tumors, leukemia/lymphoma, manifestations of Li-Fraumeni Syndrome, germline p53 mutation
 - Ashkenazi Jewish ancestry
- Male with a personal history of breast cancer
- Personal or family history of a known mutation (BRCA1/2, p53, PTEN) for a cancerpredisposition syndrome

Consider referral:

Patients with any of the following:

- Bilateral cancer (or two breast primaries) diagnosed at any age
- Breast cancer diagnosed at any age AND a family history of ≥ 2 relatives diagnosed with breast cancer at any age
- No personal history of cancer, but a family member who meets any of the above (Strongly encourage referral) criteria
- Patients with previous genetic test results (whether negative or variant of uncertain significance)

*Family history should be all on the same side of the family, i.e. either maternal *or* paternal and includes first, second and third degree relatives.

*Breast cancer includes invasive and ductal carcinoma in situ (DCIS)

Adapted from the NCCN Practice Guidelines in Oncology, v.1.2010