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<tr>
<th>NCCN Guidelines Panel Members</th>
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<tr>
<td>Genetic/Familial High-Risk Assessment: Breast and Ovarian</td>
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<tr>
<th>Name</th>
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| NCCN Guidelines Panel Disclosures |

* *Discussion Writing Committee Member * |
† Medical oncology 
∆ Cancer/Medical genetics 
♂ Internal medicine 
‡ Hematology/Hematology oncology 
Ω Gynecologic oncology/Gynecology 
¶ Breast surgical oncology 
& Public health and preventive medicine 
￥ Patient advocacy
Clinical Trials: NCCN believes that the best management for any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.

To find clinical trials online at NCCN Member Institutions, click here: nccn.org/clinical_trials/physician.html.

NCCN Categories of Evidence and Consensus: All recommendations are category 2A unless otherwise specified.

See NCCN Categories of Evidence and Consensus.

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Global
- “Epithelial” ovarian cancer was replaced by “invasive” ovarian cancer.
- Footnotes throughout the Guidelines related to risk assessment and counseling were moved to a new page titled, “Principles of Cancer Risk Assessment and Counseling.” (BR/OV-A). A new footnote was added throughout to reference this new page, “For further details regarding the nuances of genetic counseling and testing, see BR/OV-A.”

Breast and Ovarian Cancer Genetic Assessment:

BR/OV-1
- First column, heading revised, “An affected individual with a cancer diagnosis meeting any with one or more of the following”
  - 3rd bullet was modified, “Triple negative (ER-, PR-, HER2-) breast cancer ≤60 y.”
- Second column, heading revised, “An unaffected individual with no personal history of cancer but with a family history of any one or more of the following”
  - 4th bullet was modified, “≥1 invasive ovarian cancer primary from the same side of family.”
- Both personal and family history columns
  - 1st bullet was revised, “A known mutation in a breast cancer susceptibility gene within the family.”
  - Bullet was revised, “The following bullet was revised, ≥1 family member on same side of family with a combination of breast cancer and ≥1 of the following Personal and/or family history of three or more of the following (especially if early onset): pancreatic cancer, prostate cancer (Gleason score ≥7); sarcoma, adrenocortical carcinoma, brain tumors, endometrial cancer, leukemia/lymphoma; thyroid cancer, kidney cancer, dermatologic manifestations and/or macrocephaly, hamartomatous polyps of GI tract; diffuse gastric cancer (can include multiple primaries in same individual).”
- Footnote e was modified, “For the purposes of these guidelines, includes fallopian tube and primary peritoneal cancers are included. BRCA-related ovarian cancers are associated with epithelial non-mucinous histology. Other cancer genetic syndromes may be associated with mucinous ovarian cancer. Non-epithelial ovarian cancer may be associated with PJS and possibly other cancer syndromes.” Also for footnote e on HBOC-1.

BR/OV-2
- Detailed family history
  - 1st bullet was modified, “Expanded pedigree, particularly around affected individuals...”
- Detailed medical and surgical history
  - 1st bullet was modified, “Any personal cancer history (eg, age, type, histology, laterality).”
  - 4th bullet was modified by adding, “Hormone or oral contraceptive use.”
  - 5th bullet was modified by adding, “Previous breast biopsies and pathology results.”

HBOC-1
- Two statements were moved from under family history and listed above the criteria.
  - Meeting one or more of these criteria warrants further personalized risk assessment, genetic counseling, and often genetic testing and management.
  - Testing of unaffected individuals should only be considered when an appropriate affected family member is unavailable for testing.
  - First bullet was revised, “Individual from a family with a known deleterious BRCA1/BRCA2 mutation or other cancer susceptibility gene.”
  - Second bullet, Personal history of breast cancer + one or more of the following
    - 2nd sub-bullet, the 1st tertiary bullet was clarified, “An additional breast primary.”
  - Bullets regarding prostate and pancreatic cancer were separated and revised.
    - 5th bullet was revised as, “Personal history of prostate cancer (Gleason score ≥7) at any age with ≥2 1 close blood relative with breast (≤50 y) and/or invasive ovarian and/or pancreatic or prostate cancer (Gleason score ≥7) at any age.”
  - 6th bullet was revised as, “Personal history of pancreatic cancer at any age with ≥2 close blood relative with breast (≤50 y) and/or invasive ovarian and/or pancreatic cancer at any age.”
  - 7th bullet was revised as, “Personal history of pancreatic cancer, and Ashkenazi Jewish ancestry, only one additional affected relative is needed.”
  - Footnote f was revised by adding, “Comprehensive genetic testing full sequencing may be considered if ancestry also includes non-Ashkenazi Jewish...”

HBOC-2
- Footnote g was modified by adding, “Additional testing may be indicated if there is also a significant family history of cancer on the side of the family without the known mutation.”
- Footnote i was modified by adding, “If no mutation found, consider testing another family member with next highest likelihood of having a mutation and/or other hereditary breast/ovarian cancer syndromes...” Similar footnote was revised on LIFR-2 and COWD-2.

Continued on next page
NCCN Guidelines Version 1.2015 Updates
Genetic/Familial High-Risk Assessment: Breast and Ovarian

HBOC-A 1 of 2

- HBOC syndrome management for women
  - 3rd bullet, Breast screening
    - 1st sub-bullet was revised, “Age 25–29 y, annual breast MRI screening (preferred) or mammogram if MRI is unavailable or individualize based on earliest age of onset in family history if breast cancer diagnosis under age 25 is present.”
    - New sub-bullet was added, “For women with a BRCA mutation who are treated for breast cancer, screening of remaining breast tissue with annual mammography and breast MRI should continue.”
  - 5th bullet was revised, “Recommend risk-reducing salpingo-oophorectomy (ideally in consultation with a gynecologist oncologist), ideally typically between 35 and 40 y, and upon completion of childbearing, or individualized based on earliest age of onset of ovarian cancer in the family. See Risk-Reducing Salpingo-Oophorectomy (RRSO) Protocol in NCCN Guidelines for Ovarian Cancer-Principles of Surgery.”
    - New sub-bullet was added, “Salpingectomy is not the standard of care and is discouraged outside a clinical trial. The concern for risk-reducing salpingectomy alone is that women are still at risk for developing ovarian cancer. In addition, in premenopausal women, oophorectomy reduces the risk of developing breast cancer by 50%.” with a corresponding reference.
  - 7th bullet was revised, “For those patients who have not elected risk-reducing salpingo-oophorectomy, consider concurrent transvaginal ultrasound (preferably day 1–10 of menstrual cycle in premenopausal women) + CA-125 (preferably after day 5 of menstrual cycle in premenopausal women) every 6 mo starting at age 30–35 y before the earliest age of first diagnosis of ovarian cancer in the family; while there may be circumstances where clinicians find screening helpful, data do not support routine ovarian screening. Transvaginal ultrasound for ovarian cancer has not been shown to be sufficiently sensitive or specific as to support a positive recommendation, but may be considered at the clinician’s discretion starting at age 30–35 y. Serum CA-125 is an additional ovarian screening test with caveats similar to transvaginal ultrasound.”

- 8th bullet was revised, “Consider chemoprevention risk reduction agents as options for breast and ovarian cancer...”

- Footnotes
  - Footnote was removed: “There are data that show that annual transvaginal ultrasound and CA-125 are not effective strategies for screening for ovarian cancer in high-risk women. There are limited data regarding the effectiveness of a six-month screening interval. Thus, until such data are available it is reasonable to consider this approach in high-risk women, especially in the context of a clinical research setting.”
  - Footnote text was removed except for reference to See Discussion.

HBOC-A 2 of 2

- HBOC syndrome management for men
  - 2nd bullet was revised, “Clinical breast exam, every 6–12 mo, starting at age 35 y.”
  - Bullet was removed, “Consider baseline mammogram at age 40 y; annual mammogram if gynecomastia or parenchymal/glandular breast density on baseline study.”

- Reproductive options
  - 2nd bullet was revised from “For BRCA2 mutation carriers, risk of a rare (recessive) Fanconi anemia/brain tumor phenotype in offspring should be discussed if both partners carry a BRCA2 mutation” to “BRCA2 gene mutations may be associated with the rare autosomal recessive condition, Fanconi anemia. Thus, for this gene, consideration would be given to carrier testing the partner for mutations in the same gene if it would inform reproductive decision-making and/or risk assessment and management.”

Continued on next page
Updates in Version 1.2015 of the NCCN Guidelines for Genetic/Familial High-Risk Assessment: Breast and Ovarian from Version 2.2014 include:

**Li-Fraumeni Syndrome:**

**LIFR-1**
- Fourth bullet was revised, “Individual with breast cancer ≤35 y, TP53 testing can be ordered alone, concurrently with BRCA1/2 testing and/or other gene testing or as a follow-up test after negative BRCA1/2 testing.”
- List of cancers associated with LFS was removed from the page since it is included in the criteria.

**LIFR-A**
- Breast cancer risk for women
  - 3rd bullet, Breast screening
    ◊ 1st sub-bullet was revised, “Age 20–29 y, annual breast MRI screening (preferred) or mammogram if MRI is unavailable or individualized based on earliest age of onset in family.”
    ◊ 4th sub-bullet was added, “For women with a TP53 mutation who are treated for breast cancer, screening of remaining breast tissue should continue.”
- Other cancer risks
  - 4th bullet was modified: “Consider colonoscopy every 2–5 y starting no later than at 25 y or 5 y before the earliest known colon cancer in the family (whichever comes first).”
  - Bullet was removed, “Discuss option to participate in novel screening approaches using technologies, such as whole-body MRI, abdominal ultrasound, and brain MRI” and replaced with two bullets,
    ◊ Perform annual whole body MRI (rapid non-contrast exams per ACRIN model).
    ◊ The brain may be examined as part of whole body MRI or as a separate exam
  - 6th bullet was added, “Perform annual dermatologic examination.”
  - Footnote 4 was removed, “A surveillance study has been published that utilizes these screening approaches (Villani A, Tabori U, Schiffman J, et al. Lancet Oncol 2011;12:559-567). See Discussion” and replaced with “Whole body MRI is being evaluated in multiple international trials. Other components of screening are being evaluated in protocols, including regular blood screening for hematologic malignancies, and biochemical screening.”

**Cowden Syndrome/PTEN Hamartoma Tumor Syndrome:**

**COWD-1**
- Minor criteria
  - “Intellectual disability” replaced “mental retardation.”
- Footnotes
  - Footnote d was added, “Current evidence does not support testing for succinate dehydrogenase (SDH) gene mutations in patients with PHTS. (Am J Hum Genet 2011;88:674-675).”
  - Footnote g was added, “Multiple polyp types are often seen in patients with PHTS, and less commonly may include adenomas, hyperplastic polyps, and other histologies.”

**COWD-A**
- Cowden syndrome/PHTS management for women
  - 2nd bullet was modified, “Clinical breast exam, every 6–12 mo, starting at age 25 y or 5–10 y before the earliest known breast cancer in the family (whichever comes first).”
  - 3rd bullet was revised by clarifying as
    ◊ **Breast Screening**
      - 1st sub-bullet was revised, “Annual mammography and breast MRI screening starting at age 30–35 y or 5–10 y before individualized based on the earliest known breast cancer age of onset in the family (whichever comes first).”
      - 2nd sub-bullet was added, “Age >75 y, management should be considered on an individual basis.”
      - 3rd sub-bullet was added, “For women with a **PTEN** mutation who are treated for breast cancer, screening of remaining breast tissue with annual mammography and breast MRI should continue.”
- Cowden syndrome/PHTS management for men and women
  - 2nd bullet was modified, “Annual thyroid ultrasound starting at time of PHTS diagnosis age 18 y or 5–10 y before the earliest known thyroid cancer in the family, whichever is earlier.”
  - 3rd bullet was revised, “Colonoscopy, starting at age 35 y unless symptomatic or close relative with colon cancer under age 40 y. Colonoscopy should be done every 5 y or more frequently if patient is symptomatic or polyps found.”

**Continued on next page**
Updates in Version 1.2015 of the NCCN Guidelines for Genetic/Familial High-Risk Assessment: Breast and Ovarian from Version 2.2014 include:

**Multi-Gene Testing**

**GENE-1**
- This section was extensively revised and text will be included in the corresponding discussion.

**ADDIT-2**
- A new table was added, “Breast and Ovarian Management Based on Genetic Test Results.”
CRITERIA FOR FURTHER GENETIC RISK EVALUATION

An individual with a cancer diagnosis meeting any of the following:

- A known mutation in a cancer susceptibility gene within the family
- Early-age-onset breast cancer
- Triple negative (ER-, PR-, HER2-) breast cancer ≤60 y
- Two breast cancer primaries in a single individual
- Breast cancer at any age, and
  †≥1 close blood relative with breast cancer ≤50 y, or
  †≥1 close blood relative with invasive ovarian cancer at any age, or
  †≥2 close blood relatives with breast cancer and/or pancreatic cancer at any age, or
- From a population at increased risk
- Personal and/or family history of three or more of the following (especially if early onset): pancreatic cancer, prostate cancer (Gleason score ≥7), sarcoma, adrenocortical carcinoma, brain tumors, endometrial cancer; thyroid cancer, kidney cancer, dermatologic manifestations and/or macrocephaly, hamartomatous polyps of gastrointestinal (GI) tract; diffuse gastric cancer (can include multiple primary cancers in same individual)
- Invasive ovarian cancer
- Male breast cancer

An individual with no personal history of cancer but with a family history of any of the following:

- A known mutation in a cancer susceptibility gene within the family
- ≥2 breast cancer primaries in a single individual
- ≥2 individuals with breast cancer primaries on the same side of family
- ≥1 invasive ovarian cancer primary
- First- or second-degree relative with breast cancer ≤45 y
- Personal and/or family history of three or more of the following (especially if early onset): pancreatic cancer, prostate cancer (Gleason score ≥7), sarcoma, adrenocortical carcinoma, brain tumors, endometrial cancer; thyroid cancer, kidney cancer, dermatologic manifestations and/or macrocephaly, hamartomatous polyps of GI tract; diffuse gastric cancer (can include multiple primary cancers in same individual)
- Male breast cancer

†For populations at increased risk, requirements for inclusion may be modified (eg, individuals of Ashkenazi Jewish descent with breast or ovarian or pancreatic cancer at any age).

For dermatologic manifestations, see COWD-1.

For hamartomatous colon polyps in conjunction with breast cancer and hyperpigmented macules of the lips and oral mucosa, STK11 testing should be considered. See NCCN Guidelines for Genetic/Familial High-Risk Assessment: Colorectal—Peutz-Jeghers syndrome. Melanoma has been reported in some HBOC families.

For lobular breast cancer with a family history of diffuse gastric cancer, CDH1 gene testing should be considered.

†For further details regarding the nuances of genetic counseling and testing, see BR/OV-A.

See Assessment (BR/OV-2)
**Gene Testing**

- **See Targeted Testing Criteria for**
  - Hereditary Breast/Ovarian Syndrome (HBOC-1)
  - Li-Fraumeni Syndrome (LIFR-1)
  - Cowden Syndrome/PHTS (COWD-1)

- **See Multi-Gene Testing (GENE-1)**

**Assessment**

**Patient Needs and Concerns:**
- Knowledge of genetic testing for cancer risk, including benefits, risks, and limitations
- Goals for cancer family risk assessment

**Detailed Family History:**
- Expanded pedigree, particularly around affected individuals, to include first-, second-, and third-degree relatives (parents, siblings, children, grandparents, aunts, uncles, nieces, nephews, grandchildren, half-siblings, great-grandparents, great-aunts, great-uncles, great-grandchildren, and first cousins) *(See BR/OV-B)*
- Types of cancer, bilaterality, age at diagnosis
- History of chemoprevention and/or risk-reducing surgery
- Medical record documentation as needed, particularly pathology reports of primary cancers

**Detailed Medical and Surgical History:**
- Any personal cancer history (eg, age, histology, laterality)
- Carcinogen exposure (eg, history of radiation therapy)
- Reproductive history
- Hormone or oral contraceptive use
- Previous breast biopsies and pathology results
- History of salpingo-oophorectomy

**Focused Physical Exam (Conducted by Qualified Clinician):**
- Breast/ovarian
- Cowden syndrome/PHTS specific:
  - Dermatologic, including oral mucosa
  - Head circumference
  - Thyroid (enlarged or nodular on palpation)

\(^k\)For Cowden syndrome dermatologic manifestations, see COWD-1 and for PJS dermatologic manifestations, see NCCN Guidelines for Genetic/Familial High-Risk Assessment: Colorectal.

\(^l\)In some cases, multi-gene testing may be a preferable way to begin testing over the single-gene testing process.

**Note:** All recommendations are category 2A unless otherwise indicated.

Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.
PRINCIPLES OF CANCER RISK ASSESSMENT AND COUNSELING

- Cancer risk assessment and genetic counseling is highly recommended when genetic testing is offered (i.e., pre-test counseling) and after results are disclosed (i.e., post-test counseling).\(^1\)\(^-\)\(^5\) A genetic counselor, medical geneticist, oncologist, surgeon, oncology nurse or other health professional with expertise and experience in cancer genetics should be involved early in the counseling of patients.

- Pre-test counseling includes:
  - Collection of a comprehensive family history
    - Note that when assessing family history, close blood relatives include first-, second-, and third-degree relatives on each side of the family (See BR/OV-B)
  - Evaluation of a patient’s cancer risk
  - Generating a differential diagnosis and educating the patient on inheritance patterns, penetrance, variable expressivity, and the possibility of genetic heterogeneity

- Post-test counseling includes discussions of:
  - Results along with their significance and impact and recommended medical management options
  - Informing and testing at-risk family members
  - Available resources such as disease specific support groups and research studies.

Genetic Testing Considerations

- Testing should be considered in appropriate high risk individuals where it will impact the medical management of the tested individual and/or their at-risk family members. It should be, performed in a setting in which it can be adequately interpreted, and impact the medical management of the tested individual and/or their at-risk family members.\(^1\)

- The probability of mutation detection associated with these criteria will vary based on family structure. Individuals with unknown or limited family history/structure, such as fewer than 2 female first- or second-degree relatives having lived beyond age 45 in either lineage, may have an underestimated probability of familial mutation detection. The estimated likelihood of mutation detection may be very low in families with a large number of unaffected female relatives.

- Patients who have received an allogenic bone marrow transplant should not have molecular genetic testing via blood or buccal samples due to unreliable test results from contamination by donor DNA. If available, DNA should be extracted from a fibroblast culture. If this source of DNA is not possible, buccal samples can be considered, subject to the risk of donor DNA contamination.

- Comprehensive genetic testing includes full sequencing and testing for large genomic rearrangements.

- Genetic testing for adult onset diseases (e.g., \textit{BRCA1/2}) in children <18 y is generally not recommended.\(^6\)

\(^1\)\(^-\)\(^5\) Note: All recommendations are category 2A unless otherwise indicated.

Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.
PRINCIPLES OF CANCER RISK ASSESSMENT AND COUNSELING

Genetic Testing Approach

• If more than one family member is affected with cancers highly associated with a particular inherited cancer susceptibility syndrome, consider testing first a family member with youngest age at diagnosis, bilateral disease, multiple primary cancers, or other cancers associated with the syndrome, or most closely related to the proband/patient. If there are no living family members with cancer that is a cardinal feature of the syndrome in question, consider testing first- or second-degree family members affected with other cancers thought to be related to the gene in question (eg, prostate, pancreas, melanoma with BRCA1/2).

• Testing for unaffected family members when no affected member is available should be considered. Significant limitations of interpreting test results should be discussed.

• If no mutation is found, consider other hereditary cancer syndromes. For additional information on other genetic mutations associated with breast/ovarian cancer risk for which genetic testing is clinically available, see ADDIT-1 and ADDIT-2.

• Testing family members for a variant of unknown significance should not be used for clinical purposes. Consider a referral to research studies that aim to define the functional impact of variants.

Risk to relatives

• Advise about possible inherited cancer risk to relatives, options for risk assessment, and management.

• Recommend genetic counseling and consideration of genetic testing for at-risk relatives.

Reproductive options

• For patients of reproductive age, advise about options for prenatal diagnosis and assisted reproduction including pre-implantation genetic diagnosis. Discussion should include known risks, limitations, and benefits of these technologies. See Discussion for details.

• BRCA2 gene mutations may be associated with the rare autosomal recessive condition, Fanconi anemia. Thus, for this gene, consideration would be given to carrier testing the partner for mutations in the same gene if it would inform reproductive decision-making and/or risk assessment and management.


6. Committee on Bioethics; Committee on Genetics, and American College of Medical Genetics and; Genomic Social; Ethical; Legal Issues Committee. Ethical and policy issues in genetic testing and screening of children. Pediatrics 2013;131:620-622.

Note: All recommendations are category 2A unless otherwise indicated.

Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.
NCCN Guidelines Version 1.2015
Breast and/or Ovarian Cancer Genetic Assessment

PEDIGREE: FIRST-, SECOND-, AND THIRD-DEGREE RELATIVES OF PROBAND

- First-degree relatives: parents, siblings, and children;
- Second-degree relatives: grandparents, aunts, uncles, nieces, nephews, grandchildren, and half-siblings;
- Third-degree relatives: great-grandparents, great-aunts, great-uncles, great-grandchildren, and first cousins.

Note: All recommendations are category 2A unless otherwise indicated.
Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.
HEREDITARY BREAST AND/OR OVARIAN CANCER SYNDROME TESTING CRITERIA

Meeting one or more of these criteria warrants further personalized risk assessment, genetic counseling, and often genetic testing and management. Testing of unaffected individuals should only be considered when an appropriate affected family member is unavailable for testing.

- Individual from a family with a known deleterious BRCA1/BRCA2 mutation or other cancer susceptibility gene
- Personal history of breast cancer + one or more of the following:
  - Diagnosed ≤45 y
  - Diagnosed ≤50 y with:
    - An additional breast cancer primary
    - ≥1 close blood relative with breast cancer at any age
    - An unknown or limited family history
  - Diagnosed ≤60 y with:
    - A triple negative breast cancer
  - Diagnosed at any age with:
    - ≥1 close blood relative with breast cancer diagnosed ≤50 y
    - ≥2 close blood relatives with breast cancer at any age
    - ≥1 close blood relative with invasive ovarian cancer
    - ≥2 close blood relatives with pancreatic cancer and/or prostate cancer (Gleason score ≥7) at any age
    - A close male blood relative with breast cancer
  - For an individual of ethnicity associated with higher mutation frequency (eg, Ashkenazi Jewish) no additional family history may be required
- Personal history of invasive ovarian cancer
- Personal history of male breast cancer

For further details regarding the nuances of genetic counseling and testing, see BR/OV-A.

For the purposes of these guidelines, invasive and ductal carcinoma in situ breast cancers should be included.

Two breast cancer primaries includes bilateral (contralateral) disease or two or more clearly separate ipsilateral primary tumors either synchronously or asynchronously.

Close blood relatives include first-, second-, and third-degree relatives on same side of family. (See BR/OV-B)

Includes fallopian tube and primary peritoneal cancers. BRCA-related ovarian cancers are associated with epithelial non-mucinous histology. Other cancer genetic syndromes may be associated with mucinous ovarian cancer. Non-epithelial ovarian cancer may be associated with PJS and possibly other cancer syndromes. Ovarian/fallopian tube/primary peritoneal cancers are component tumors of Lynch syndrome; be attentive for clinical evidence of this syndrome. See NCCN Guidelines for Genetic/Familial High-Risk Assessment: Colorectal.

Testing for Ashkenazi Jewish founder-specific mutation(s) should be performed first. Comprehensive genetic testing may be considered if ancestry also includes non-Ashkenazi Jewish relatives or if other HBOC criteria are met. Founder mutations exist in other populations.
# NCCN Guidelines Version 1.2015
## Hereditary Breast and/or Ovarian Cancer Syndrome

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<th>HBOC FOLLOW-UP</th>
<th>FAMILY STATUS</th>
<th>GENETIC TESTING&lt;sup&gt;a&lt;/sup&gt;</th>
<th>TEST OUTCOME&lt;sup&gt;a&lt;/sup&gt;</th>
<th>SCREENING RECOMMENDATION</th>
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<tr>
<td>HBOC testing criteria met</td>
<td>Risk assessment and counseling:&lt;sup&gt;a&lt;/sup&gt; - Psychosocial assessment and support - Risk counseling - Education - Discussion of genetic testing - Informed consent</td>
<td>Deleterious familial BRCA1/BRCA2 mutation known</td>
<td>Recommend BRCA1/BRCA2 testing for specific familial mutation&lt;sup&gt;g&lt;/sup&gt;</td>
<td>See HBOC Syndrome Management (HBOC-A)</td>
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<td>No known familial BRCA1/BRCA2 mutation</td>
<td>Consider comprehensive BRCA1/BRCA2 testing of patient or if unaffected, test family member with highest likelihood of a mutation&lt;sup&gt;h&lt;/sup&gt; or</td>
<td>See HBOC Syndrome Management (HBOC-A)</td>
</tr>
<tr>
<td></td>
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<td></td>
<td>Mutation found</td>
<td>Offer research and individualized recommendations according to personal and family history</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Not tested</td>
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<td></td>
<td>No mutation found&lt;sup&gt;i&lt;/sup&gt;</td>
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<td></td>
<td>Variant of unknown significance found (uninformative)&lt;sup&gt;i&lt;/sup&gt;</td>
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<td></td>
<td>Consider multi-gene testing, if appropriate</td>
<td>See Multi-Gene Testing (GENE-1)</td>
</tr>
</tbody>
</table>

<sup>a</sup>For further details regarding the nuances of genetic counseling and testing, see BR/OV-A.

<sup>g</sup>If of Ashkenazi Jewish descent, in addition to the specific familial mutation, test for all three founder mutations. Additional testing may be indicated if there is also a significant family history of cancer on the side of the family without the known mutation.

<sup>h</sup>For both affected and unaffected individuals of Ashkenazi Jewish descent with no known familial mutation, first test for the three common mutations. Then, if negative for the three mutations and ancestry also includes non-Ashkenazi Jewish relatives or other HBOC criteria are met, consider comprehensive genetic testing. For both affected and unaffected individuals who are non-Ashkenazi Jewish and who have no known familial mutation, comprehensive genetic testing is the approach, if done.

<sup>i</sup>If no mutation found, consider testing another family member with next highest likelihood of having a mutation and/or other hereditary breast/ovarian cancer syndromes such as Li-Fraumeni (LIFR-1) and/or Cowden syndrome (COWD-1) or multi-gene testing (GENE-1). For additional information on other genetic mutations associated with breast/ovarian cancer risk for which genetic testing is clinically available, see ADDIT-1.

Note: All recommendations are category 2A unless otherwise indicated.

Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.
WOMEN

• Breast awareness starting at age 18 y.
• Clinical breast exam, every 6–12 mo, starting at age 25 y.
• Breast screening
  ▪ Age 25–29 y, annual breast MRI screening (preferred) or mammogram if MRI is unavailable or individualized based on family history if a breast cancer diagnosis before age 25 is present.
  ▪ Age 30–75 y, annual mammogram and breast MRI screening.
  ▪ Age >75 y, management should be considered on an individual basis.
• For women with a BRCA mutation who are treated for breast cancer, screening of remaining breast tissue with annual mammography and breast MRI should continue.
• Discuss option of risk-reducing mastectomy
  ▪ Counseling may include a discussion regarding degree of protection, reconstruction options, and risks.
  ▪ Recommend risk-reducing salpingo-oophorectomy (ideally in consultation with a gynecologist oncologist), typically between 35 and 40 y, and upon completion of child bearing. See Risk-Reducing Salpingo-Oophorectomy (RRSO) Protocol in NCCN Guidelines for Ovarian Cancer- Principles of Surgery.
  ▪ Counseling includes a discussion of reproductive desires, extent of cancer risk, degree of protection for breast and ovarian cancer, management of menopausal symptoms, possible short-term hormone replacement therapy to a recommended maximum age of natural menopause, and related medical issues.
  ▪ Salpingectomy alone is not the standard of care and is discouraged outside a clinical trial. The concern for risk-reducing salpingectomy alone is that women are still at risk for developing ovarian cancer. In addition, in premenopausal women, oophorectomy reduces the risk of developing breast cancer by 50%.
• Address psychosocial, social, and quality-of-life aspects of undergoing risk-reducing mastectomy and/or salpingo-oophorectomy.
• For those patients who have not elected risk-reducing salpingo-oophorectomy, while there may be circumstances where clinicians find screening helpful, data do not support routine ovarian screening. Transvaginal ultrasound for ovarian cancer has not been shown to be sufficiently sensitive or specific as to support a positive recommendation, but may be considered at the clinician's discretion starting at age 30–35 y. Serum CA-125 is an additional ovarian screening test with caveats similar to transvaginal ultrasound.
• Consider risk reduction agents as options for breast and ovarian cancer, including discussing risks and benefits (See Discussion for details). (See NCCN Guidelines for Breast Cancer Risk Reduction).
• Consider investigational imaging and screening studies, when available (eg, novel imaging technologies, more frequent screening intervals) in the context of a clinical trial.

1Women should be familiar with their breasts and promptly report changes to their health care provider. Periodic, consistent breast self exam (BSE) may facilitate breast self awareness. Premenopausal women may find BSE most informative when performed at the end of menses.
2Randomized trials comparing clinical breast exam versus no screening have not been performed. Rationale for recommending clinical breast exam every 6–12 mo is the concern for interval breast cancers.
4High-quality breast MRI limitations include having: a need for a dedicated breast coil, the ability to perform biopsy under MRI guidance, experienced radiologists in breast MRI, and regional availability. Breast MRI is performed preferably days 7–15 of menstrual cycle for premenopausal women.
5Given the high rate of occult neoplasms, special attention should be given to sampling and pathologic review of the ovaries and fallopian tubes. (See Discussion for details.) See the College of American Pathologists, Protocol for the Examination of Specimens from Patients with Carcinoma of the Ovary. See NCCN Guidelines for Ovarian Cancer for treatment of findings.
HBOC SYNDROME MANAGEMENT (2 of 2)

MEN
• Breast self-exam training and education starting at age 35 y
• Clinical breast exam, every 12 mo, starting at age 35 y
• Starting at age 40 y:
  ‣ Recommend prostate cancer screening for BRCA2 carriers
  ‣ Consider prostate cancer screening for BRCA1 carriers

MEN AND WOMEN
• Education regarding signs and symptoms of cancer(s), especially those associated with BRCA gene mutations.
• No specific screening guidelines exist for pancreatic cancer and melanoma, but screening may be individualized based on cancers observed in the family.8

RISK TO RELATIVES
• Advise about possible inherited cancer risk to relatives, options for risk assessment, and management.
• Recommend genetic counseling and consideration of genetic testing for at-risk relatives.

REPRODUCTIVE OPTIONS
• For patients of reproductive age, advise about options for prenatal diagnosis and assisted reproduction including pre-implantation genetic diagnosis. Discussion should include known risks, limitations, and benefits of these technologies. See Discussion for details.
• BRCA2 gene mutations may be associated with the rare autosomal recessive condition, Fanconi anemia. Thus, for this gene, consideration would be given to carrier testing the partner for mutations in the same gene if it would inform reproductive decision-making and/or risk assessment and management.9

7There are only limited data to support breast imaging in men.
8Consider full-body skin and eye exam for melanoma and investigational protocols for pancreatic cancer.
**LI-FRAUMENI SYNDROME TESTING CRITERIA**

- Individual from a family with a known *TP53* mutation
- Classic Li-Fraumeni syndrome (LFS) criteria:
  - Combination of an individual diagnosed age <45 y with a sarcoma
    - AND
    - A first-degree relative diagnosed age <45 y with cancer
    - AND
    - An additional first- or second-degree relative in the same lineage with cancer diagnosed age <45 y, or a sarcoma at any age
- Chompret criteria:
  - Individual with a tumor from LFS tumor spectrum (eg, soft tissue sarcoma, osteosarcoma, brain tumor, breast cancer, adrenocortical carcinoma, leukemia, lung bronchoalveolar cancer) before 46 years of age, AND at least one first- or second-degree relative with any of the aforementioned cancers (other than breast cancer if the proband has breast cancer) before the age of 56 years or with multiple primaries at any age
    - OR
  - Individual with multiple tumors (except multiple breast tumors), two of which belong to LFS tumor spectrum with the initial cancer occurring before the age of 46 years
    - OR
  - Individual with adrenocortical carcinoma or choroid plexus carcinoma at any age of onset, regardless of the family history
- Early-age-onset breast cancer:
  - Individual with breast cancer ≤35 y, *TP53* testing can be ordered alone, concurrently with *BRCA1/2* testing and/or other gene testing or as a follow-up test after negative *BRCA1/2* testing

**FOLLOW-UP**

- LFS testing criteria met
  - See Follow-up (LIFR-2)

- LFS testing criteria not met
  - Individualized recommendations according to personal and family history

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*aFor further details regarding the nuances of genetic counseling and testing, see BR/OV-A.


*cTo date, there have been no reports of Ewing sarcoma, GIST, desmoid tumor, or angiosarcoma in *TP53* mutation carriers.


### Li-Fraumeni Syndrome

#### Clinical Trials
- NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.

#### Genetics
- **Li-Fraumeni Syndrome (LIFR-2)**
- **Youngest age at diagnosis, bilateral disease, multiple primaries, or sarcoma at age <45 y.**
- **If no mutation is found, consider testing another family member with next highest likelihood of having a mutation and/or other hereditary breast cancer syndromes such as HBOC (HBOC-1) and/or Cowden syndrome (COWD-1) or multi-gene testing (GENE-1).** For additional information on other genetic mutations associated with breast/ovarian cancer risk for which genetic testing is clinically available, see ADDIT-1.

#### Follow-Up

- **Risk assessment and counseling.**
  - Psychosocial assessment and support
  - Risk counseling
  - Education
  - Discussion of genetic testing
  - Informed consent

#### Family Status
- **Deleterious familial TP53 mutation known**
  - Consider TP53 testing for specific familial mutation (category 2A for adults; category 2B for children)

- **No known familial TP53 mutation**
  - Consider comprehensive TP53 testing of patient or, if unaffected, test family member with highest likelihood of a mutation
    - **Mutation found**
      - **Not tested**
        - **No mutation found**
          - **Variant of unknown significance found (uninformative)**
            - **See Multi-Gene Testing (GENE-1)**
        - **See Li-Fraumeni Syndrome Management (LIFR-A)**
      - **See Li-Fraumeni Syndrome Management (LIFR-A)**
    - **See Multi-Gene Testing (GENE-1)**
  - Consider multi-gene testing, if appropriate

#### Genetic Testing
- **Consider TP53 testing for specific familial mutation (category 2A for adults; category 2B for children)**

#### Test Outcome
- **Positive for familial TP53 mutation**
- **TP53 testing not performed**
- **Negative for familial TP53 mutation**

#### Screening Recommendation
- **See Li-Fraumeni Syndrome Management (LIFR-A)**
- **Cancer screening as per NCCN Screening Guidelines**
- **Offer research and individualized recommendations according to personal and family history**

---

**Note:** All recommendations are category 2A unless otherwise indicated.
LI-FRAUMENI SYNDROME MANAGEMENT

BREAST CANCER RISK FOR WOMEN
• Breast awareness\(^1\) starting at age 18 y.
• Clinical breast exam, every 6–12 mo, starting at age 20–25 y or 5–10 y before the earliest known breast cancer in the family (whichever comes first).
• Breast screening\(^2\)
  ‣ Age 20–29 y, annual breast MRI\(^3\) screening (preferred) or mammogram if MRI is unavailable
  ‣ Age 30–75 y, annual mammogram and breast MRI\(^3\) screening
  ‣ Age >75 y, management should be considered on an individual basis.
  ‣ For women with a TP53 mutation who are treated for breast cancer, screening of remaining breast tissue with annual mammography and breast MRI should continue.
• Discuss option of risk-reducing mastectomy and counsel regarding degree of protection, degree of cancer risk, and reconstruction options.
• Address psychosocial, social, and quality-of-life aspects of undergoing risk-reducing mastectomy.

OTHER CANCER RISKS
• Address limitations of screening for many cancers associated with LFS. Because of the remarkable risk of additional primary neoplasms, screening may be considered for cancer survivors with LFS and a good prognosis from their prior tumor(s).
• Pediatricians should be apprised of the risk of childhood cancers in affected families.
• Annual comprehensive physical exam with high index of suspicion for rare cancers and second malignancies in cancer survivors: include neurologic examination.
• Therapeutic RT for cancer should be avoided when possible.
• Consider colonoscopy every 2-5 y starting at 25 y or 5 y before the earliest known colon cancer in the family (whichever comes first).
• Perform annual dermatologic examination.
• Perform annual whole body MRI (rapid non-contrast exams per ACRIN model).\(^4\)
• The brain may be examined as part of whole body MRI or as a separate exam.\(^4\)
• Provide additional surveillance based on family history of cancer.
• Provide education regarding signs and symptoms of cancer.

\(^1\)Women should be familiar with their breasts and promptly report changes to their health care provider. Periodic, consistent breast self exam (BSE) may facilitate breast self awareness. Premenopausal women may find BSE most informative when performed at the end of menses.
\(^2\)The appropriateness of imaging modalities and scheduling is still under study.
\(^3\)High-quality breast MRI limitations include having: a need for a dedicated breast coil, the ability to perform biopsy under MRI guidance, experienced radiologists in breast MRI, and regional availability. Breast MRI is performed preferably days 7–15 of menstrual cycle for premenopausal women.
\(^4\)Whole body MRI is being evaluated in multiple international trials. Other components of screening are being evaluated in protocols, including regular blood screening for hematologic malignancies, and biochemical screening.
LI-FRAUMENI SYNDROME MANAGEMENT

REPRODUCTIVE OPTIONS
• For patients of reproductive age, advise about options for prenatal diagnosis and assisted reproduction including pre-implantation genetic diagnosis. Discussion should include known risks, limitations, and benefits of these technologies. See Discussion for details.

RISK TO RELATIVES
• Advise about possible inherited cancer risk to relatives, options for risk assessment, and management.
• Recommend genetic counseling and consideration of genetic testing for at-risk relatives.

Note: All recommendations are category 2A unless otherwise indicated.
Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.
# NCCN Guidelines Version 1.2015
## Cowden Syndrome/PHTS

### COWDEN SYNDROME/PTEN HAMARTOMA TUMOR SYNDROME TESTING CRITERIA

<table>
<thead>
<tr>
<th>Major criteria:</th>
<th>Minor criteria:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Breast cancer</td>
<td>• Thyroid structural lesions (eg, adenoma, nodule(s), goiter)</td>
</tr>
<tr>
<td>• Endometrial cancer</td>
<td>• Renal cell carcinoma</td>
</tr>
<tr>
<td>• Follicular thyroid cancer</td>
<td>• Single GI hamartoma or ganglioneuroma</td>
</tr>
<tr>
<td>• Multiple GI hamartomas or ganglioneuromas</td>
<td>• Testicular lipomatosis</td>
</tr>
<tr>
<td>• Macrocephaly (megealocephaly) (ie, ≥97%, 58 cm in adult women, 60 cm in adult men)</td>
<td>• Vascular anomalies (including multiple intracranial developmental venous anomalies)</td>
</tr>
<tr>
<td>• Macular pigmentation of glans penis</td>
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<tr>
<td>• Mucocutaneous lesions</td>
<td></td>
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<tr>
<td>‣ One biopsy-proven trichilemmoma</td>
<td>• Autism spectrum disorder</td>
</tr>
<tr>
<td>‣ Multiple palmoplantar keratoses</td>
<td>• Colon cancer</td>
</tr>
<tr>
<td>‣ Multifocal or extensive oral mucosal papillomatosis</td>
<td>• ≥3 esophageal glycogenic acanthoses</td>
</tr>
<tr>
<td>‣ Multiple cutaneous facial papules (often verrucous)</td>
<td>• Lipomas</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>At-risk individual with a relative with a clinical diagnosis of CS/PHTS or BRRS for whom testing has not been performed</th>
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</thead>
<tbody>
<tr>
<td>The at-risk individual must have the following:</td>
</tr>
<tr>
<td>♦ Any one major criterion or</td>
</tr>
<tr>
<td>♦ Two minor criteria</td>
</tr>
</tbody>
</table>

### FOLLOW-UP

- **CS/PHTS testing criteria met**
  - Individualized recommendations according to personal and family history

- **CS/PHTS testing criteria not met**
  - See Follow-up (COWD-2)

---

**Note:** All recommendations are category 2A unless otherwise indicated. Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.

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For further details regarding the nuances of genetic counseling and testing, see BR/OV-A.

These are testing criteria; clinical diagnostic criteria can be found on COWD-3.

If two criteria involve the same structure/organ/tissue, both may be included as criteria.

Current evidence does not support testing for succinate dehydrogenase (SDH) gene mutations in patients with PHTS. (Am J Hum Genet 2011;88:674-675).


If an individual has two or more major criteria, such as breast cancer and non-medullary thyroid cancer, but does not have macrocephaly, one of the major criteria may be included as one of the three minor criteria to meet testing criteria.

Multiple polypl types are often seen in patients with PHTS, and less commonly may include adenomas, hyperplastic polyplps, and other histologies.


The literature available on mucocutaneous lesions is not adequate to accurately specify the number or extent of mucocutaneous lesions required to be a major criterion for CS/PHTS. Clinical judgment should be used.

Insufficient evidence exists in the literature to include fibrocystic disease of the breast, fibromas, and uterine fibroids as diagnostic criteria.
# Cowden Syndrome/PHTS

## COWDEN SYNDROME FOLLOW-UP

<table>
<thead>
<tr>
<th>Family Status</th>
<th>Genetic Testing</th>
<th>Test Outcome</th>
<th>Screening Recommendation</th>
</tr>
</thead>
</table>
| Risk assessment and counseling:  
- Psychosocial assessment and support  
- Risk counseling  
- Education  
- Discussion of genetic testing  
- Informed consent | Deleterious familial PTEN mutation known | Consider PTEN testing for specific familial mutation | Positive for familial PTEN mutation | See Cowden Syndrome/PHTS Management (COWD-A) |
| No known familial PTEN mutation | Consider comprehensive PTEN testing of patient or, if unaffected, test family member with highest likelihood of a mutation or Consider multi-gene testing, if appropriate | Mutation found | Meets CS/PHTS diagnostic criteria (see COWD-3) | See Cowden Syndrome/PHTS Management (COWD-A) |
| Mutation found | Not tested | No mutation found | Does not meet CS/PHTS diagnostic criteria (see COWD-3) | Offer research and individualized recommendations according to personal and family history |
| Variant of unknown significance found (uninformative)| | | |

\[^a\] For further details regarding the nuances of genetic counseling and testing, see BR/OV-A.

\[^k\] If no mutation is found, consider testing another family member with next highest likelihood of having a mutation and/or other hereditary breast cancer syndromes such as HBOC (HBOC-1) and/or Li-Fraumeni syndrome (LIFR-1) or multi-gene testing (GENE-1). For additional information on other genetic mutations associated with breast/ovarian cancer risk for which genetic testing is clinically available, see ADDIT-1.
REvised PTEn Hamartoma tumor syndrome clinical diagnostic criteria

MAJOR CRITERIA:
- Breast cancer
- Endometrial cancer (epithelial)
- Thyroid cancer (follicular)
- GI hamartomas (including ganglioneuromas, but excluding hyperplastic polyps; ≥3)
- Lhermitte-Duclos disease (adult)
- Macrocephaly (≥97 percentile: 58 cm for females, 60 cm for males)
- Macular pigmentation of the glans penis
- Multiple mucocutaneous lesions (any of the following):
  - Multiple trichilemmomas (≥3, at least one biopsy proven)
  - Acral keratoses (≥3 palmoplantar keratotic pits and/or acral hyperkeratotic papules)
  - Mucocutaneous neuromas (≥3)
  - Oral papillomas (particularly on tongue and gingiva), multiple (≥3)
  - OR biopsy proven OR dermatologist diagnosed

MINOR CRITERIA:
- Autism spectrum disorder
- Colon cancer
- Esophageal glycogenic acanthoses (≥3)
- Lipomas (≥3)
- Intellectual disability (ie, IQ ≤75)
- Renal cell carcinoma
- Testicular lipomatosis
- Thyroid cancer (papillary or follicular variant of papillary)
- Thyroid structural lesions (eg, adenoma, multinodular goiter)
- Vascular anomalies (including multiple intracranial developmental venous anomalies)

Operational diagnosis in an individual (either of the following):
1. Three or more major criteria, but one must include macrocephaly, Lhermitte-Duclos disease, or GI hamartomas; or
2. Two major and three minor criteria.

Operational diagnosis in a family where one individual meets revised PTEn hamartoma tumor syndrome clinical diagnostic criteria or has a PTEn mutation:
1. Any two major criteria with or without minor criteria; or
2. One major and two minor criteria; or
3. Three minor criteria.

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Note: All recommendations are category 2A unless otherwise indicated.
Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.
COWDEN SYNDROME/PHTS MANAGEMENT

WOMEN
• Breast awareness starting at age 18 y.
• Clinical breast exam, every 6–12 mo, starting at age 25 y or 5–10 y before the earliest known breast cancer in the family (whichever comes first).
• Breast screening
  ▶ Annual mammography and breast MRI screening starting at age 30–35 y or 5–10 y before the earliest known breast cancer in the family (whichever comes first).\(^2,3\)
  ▶ Age >75 y, management should be considered on an individual basis.
  ▶ For women with a PTEN mutation who are treated for breast cancer, screening of remaining breast tissue with annual mammography and breast MRI should continue.
• For endometrial cancer screening,\(^4\) encourage patient education and prompt response to symptoms (eg, abnormal bleeding). Consider annual random endometrial biopsies and/or ultrasound beginning at age 30–35 y.
• Discuss option of risk-reducing mastectomy and hysterectomy and counsel regarding degree of protection, extent of cancer risk, and reconstruction options.
• Address psychosocial, social, and quality-of-life aspects of undergoing risk-reducing mastectomy and/or hysterectomy.

MEN AND WOMEN
• Annual comprehensive physical exam starting at age 18 y or 5 y before the youngest age of diagnosis of a component cancer in the family (whichever comes first), with particular attention to thyroid exam.
• Annual thyroid ultrasound starting at time of PHTS diagnosis
• Colonoscopy, starting at age 35 y unless symptomatic or close relative with colon cancer under age 40 y. Colonoscopy should be done every 5 y or more frequently if patient is symptomatic or polyps found.
• Consider renal ultrasound starting at age 40 y, then every 1–2 y
• Dermatologic management may be indicated for some patients
• Consider psychomotor assessment in children at diagnosis and brain MRI if there are symptoms.
• Education regarding the signs and symptoms of cancer.

Continued on next page

\(^1\)Women should be familiar with their breasts and promptly report changes to their health care provider. Periodic, consistent breast self exam (BSE) may facilitate breast self awareness. Premenopausal women may find BSE most informative when performed at the end of menses.
\(^2\)The appropriateness of imaging modalities and scheduling is still under study.
\(^3\)High-quality breast MRI limitations include having: a need for a dedicated breast coil, the ability to perform biopsy under MRI guidance by experienced radiologists in breast MRI, and regional availability. Breast MRI is preferably preformed on days 7–15 of a menstrual cycle for premenopausal women.
\(^4\)There are limited data regarding the lifetime risk of endometrial cancer in CS/PHTS. Surveillance screening and surgical intervention should be on an individual basis.
\(^5\)Oophorectomy is not indicted for CS/PHTS alone but may be indicated for other reasons.
RISK TO RELATIVES
• Advise about possible inherited cancer risk to relatives, options for risk assessment, and management.
• Recommend genetic counseling and consideration of genetic testing for at-risk relatives.

REPRODUCTIVE OPTIONS
• For women of reproductive age, advise about options for prenatal diagnosis and assisted reproduction including pre-implantation genetic diagnosis. Discussion should include known risks, limitations, and benefits of these technologies. See Discussion for details.
MULTI-GENE TESTING

Overview of multi-gene testing

• The recent introduction of multi-gene testing for hereditary forms of cancer has rapidly altered the clinical approach to testing at-risk patients and their families. Based on next-generation sequencing technology, these tests simultaneously analyze a set of genes that are associated with a specific family cancer phenotype or multiple phenotypes.

• Patients who have a personal or family history suggestive of a single inherited cancer syndrome are most appropriately managed by genetic testing for that specific syndrome. When more than one gene can explain an inherited cancer syndrome, than multi-gene testing, may be more efficient and/or cost-effective.

• There is also a role for multi-gene testing in individuals who have tested negative (indeterminate) for a single syndrome, but whose personal or family history remains strongly suggestive of an inherited susceptibility.

• As commercially available tests differ in the specific genes analyzed (as well as classification of variants and many other factors), choosing the specific laboratory and test panel is important.

• Multi-gene testing can include “intermediate” penetrant (moderate-risk) genes. For many of these genes, there are limited data on the degree of cancer risk and there are no clear guidelines on risk management for carriers of mutations. Not all genes included on available multi-gene tests are necessarily clinically actionable. As is the case with high-risk genes, it is possible that the risks associated with moderate-risk genes may not be entirely due to that gene alone, but may be influenced by gene/gene or gene/environment interactions. Therefore, it may be difficult to use a known mutation alone to assign risk for relatives. In many cases the information from testing for moderate penetrance genes does not change risk management compared to that based on family history alone.

• There is an increased likelihood of finding variants of unknown significance when testing for mutations in multiple genes.

• It is for these and other reasons that multigene testing are ideally offered in the context of professional genetic expertise for pre- and post-test counseling.

References (GENE-2)

See Breast and Ovarian Management Based on Genetic Test Results (ADDIT-2)
MULTI-GENE TESTING

REFERENCES

EXAMPLES OF ADDITIONAL GENETIC MUTATIONS ASSOCIATED WITH BREAST/OVARIAN CANCER RISK

- Hereditary Diffuse Gastric Cancer Syndrome (See NCCN Guidelines for Gastric Cancer)
  - CDH1 gene
  - Diffuse gastric cancer — 67%–83% risk
  - Lobular cancer of the breast — 39%–52% risk

- Peutz-Jeghers Syndrome (See NCCN Guidelines for Genetic/Familial High-Risk Assessment: Colorectal for more information)
  - STK11/LKB1 gene
  - Breast cancer — 44%–50% risk
  - Ovarian cancer — 18%–21% risk (ovarian sex cord tumors are the most common)

- Lynch Syndrome (See NCCN Guidelines for Genetic/Familial High-Risk Assessment: Colorectal for more information)
  - Mismatch Repair (MMR) genes — MLH1, MSH2, MSH6, PMS2
  - EPCAM gene deletion
  - Ovarian cancer — 9% risk
  - Breast cancer — conflicting data regarding increased risks
# NCCN Guidelines Version 1.2015
## Genetic/Familial High-Risk Assessment: Breast and Ovarian

## BREAST AND OVARIAN MANAGEMENT BASED ON GENETIC TEST RESULTS

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Recommend MRI&lt;sup&gt;c&lt;/sup&gt; (&lt;20% risk of breast cancer)&lt;sup&gt;d&lt;/sup&gt;)</th>
<th>Recommend RRSO</th>
<th>Discuss Option of RRM</th>
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<tbody>
<tr>
<td>ATM</td>
<td>BRCA1</td>
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<sup>a</sup>Other genes may be included in multi-gene testing.

<sup>b</sup>Intervention may still be warranted based on family history or other clinical factors.

<sup>c</sup>See NCCN Guidelines for Breast Cancer Screening and Diagnosis.

<sup>d</sup>May be modified based on family history or specific gene mutation.

<sup>e</sup>See NCCN Guidelines for Genetic/Familial High-Risk Assessment: Colorectal.

Note: All recommendations are category 2A unless otherwise indicated.

Clinical Trials: NCCN believes that the best management of any cancer patient is in a clinical trial. Participation in clinical trials is especially encouraged.
NCCN Guidelines Version 1.2015
Genetic/Familial High-Risk Assessment: Breast and Ovarian

Discussion

This discussion is being updated to correspond with the newly updated algorithm. Last updated 08/08/13

NCCN Categories of Evidence and Consensus

Category 1: Based upon high-level evidence, there is uniform NCCN consensus that the intervention is appropriate.

Category 2A: Based upon lower-level evidence, there is uniform NCCN consensus that the intervention is appropriate.

Category 2B: Based upon lower-level evidence, there is NCCN consensus that the intervention is appropriate.

Category 3: Based upon any level of evidence, there is major NCCN disagreement that the intervention is appropriate.

All recommendations are category 2A unless otherwise noted.

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Overview

All cancers develop as a result of mutations in certain genes, such as those involved in the regulation of cell growth and/or DNA repair, although not all of these mutations are inherited from a parent. For example, sporadic mutations can occur in somatic/tumor cells only, and de novo mutations can occur for the first time in a germ cell (i.e., egg or sperm) or in the fertilized egg itself during early embryogenesis. However, family studies have long documented an increased risk of several forms of cancer among first-degree relatives (i.e., parents, siblings, and children) and second-degree relatives (i.e., grandparents, aunts or uncles, grandchildren, and nieces or nephews) of affected individuals. These individuals may have an increased susceptibility to cancer as the result of one or more gene mutations present in parental germline cells; cancers developing in these individuals may be classified as hereditary or familial cancers.

Hereditary cancers are often characterized by mutations associated with a high probability of cancer development (i.e., a high penetrance genotype), vertical transmission through either mother or father, and an association with other types of tumors. They often have an early age of onset, and exhibit an autosomal dominant inheritance pattern (i.e., occur when the individual has a mutation in only one copy of a gene). Familial cancers share some but not all features of hereditary cancers. For example, although familial breast cancers occur in a given family more frequently than in the general population, they generally do not exhibit the inheritance patterns or onset age consistent with hereditary cancers. Familial cancers may be associated with chance clustering of sporadic cancer cases within families, genetic variation in lower penetrance genes, a shared environment, or combinations of these factors.

Assessment of an individual’s risk of familial or hereditary cancer is based on a thorough evaluation of the family history. With respect to hereditary cancers, advances in molecular genetics have identified a number of genes associated with inherited susceptibility to breast and/or ovarian cancers (e.g., BRCA1, BRCA2, PTEN, TP53, CDH1) and provided a means of characterizing the specific gene mutation or mutations present in certain individuals and families exhibiting an increased risk of cancer. The field of cancer genetics has implications for all aspects of cancer management of individuals with hereditary or familial cancers, including prevention, screening, and treatment.

The NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®) for Genetic/Familial High-Risk Assessment: Breast and Ovarian were developed with an acute awareness of the preliminary nature of much of our knowledge regarding the clinical application of the rapidly emerging field of molecular genetics, and with an appreciation for the need for flexibility when applying these guidelines to individual families. Furthermore, it should be emphasized that these guidelines were not developed as a substitute for professional genetic counseling. Rather, they are intended to serve as a resource for healthcare providers to identify individuals who may benefit from cancer risk assessment and genetic counseling, to provide genetic counselors with an updated tool for the assessment of individual breast cancer and ovarian cancer risk and to guide decisions related to genetic testing, and to facilitate a multidisciplinary approach in the management of individuals at increased risk of hereditary breast and/or ovarian cancer. Although cancers other than breast and ovarian cancers are associated with these hereditary syndromes, the main focus of this NCCN Guidelines is on the management of breast and ovarian cancer risk in these individuals. During the last few years, a number of genetic aberrations that may contribute to increased risks for development of
breast and/or ovarian cancers have been identified. The current NCCN Guidelines for Genetic/Familial High-Risk Assessment: Breast and Ovarian focus specifically on assessment of mutations in the genes BRCA1/BRCA2, TP53 and PTEN, and recommended approaches to genetic testing/counseling and management strategies in individuals with these genetic mutations.

A glossary of genetic terms is included in Table 1 for reference.

**Hereditary Breast or Breast/Ovarian Cancer Syndromes**

Breast cancer is the most prevalent type of cancer in women in the U.S. and the second leading cause of cancer death in women. In the U.S., approximately 234,580 new cases of breast cancer and 40,030 deaths are estimated in 2013 (estimated figures include both genders). Up to 10% of breast cancers are due to specific mutations in single genes that are passed down in a family. Specific patterns of hereditary breast/ovarian cancers are linked to mutations in the BRCA1 or BRCA2 genes. In addition, two very rare hereditary cancer syndromes exhibiting an increased risk of breast cancer are Li-Fraumeni syndrome and Cowden syndrome, which are related to germline mutations in the TP53 and PTEN genes, respectively. Similar to the BRCA 1/2 genes, the TP53 and PTEN genes encode for proteins involved in processes related to tumor suppression, such as DNA repair and cell cycle regulation. Hereditary diffuse gastric cancer (HDGC) is another rare hereditary syndrome that is also associated with development of lobular breast cancer. This syndrome arises from mutation(s) in the CDH1 (cadherin 1, type 1, E-cadherin [epithelial]) gene which encodes for a tumor suppressor gene product. In an analysis of 4 predominantly gastric cancer pedigrees from Newfoundland with a specific CDH1 mutation, the cumulative risk of female lobular breast cancer by the age of 75 was estimated to be as high as 52%.

Furthermore, germline CDH1 mutations may be associated with lobular breast cancer in the absence of diffuse gastric cancer.

These hereditary syndromes share several features beyond elevation of breast cancer risk. These syndromes arise from germline gene mutations that are not within sex-linked genes; hence, the mutations can be inherited from either parent. The syndromes are associated with breast cancer onset at an early age and development of other types of cancer, and exhibit an autosomal dominant inheritance pattern (see Table 1). Offspring of an individual with one of these hereditary syndromes have a 50% chance of inheriting the mutation. In addition, individuals with these hereditary syndromes share increased risks of multiple cases of early onset disease as well as bilateral disease. The gene mutations associated with these hereditary syndromes are considered to be highly penetrant, although a subsequent alteration in the second copy of the gene without the hereditary mutation is believed to be necessary for the initiation of cancer development (i.e., 2-hit hypothesis). In addition, the manifestations (i.e., expression) of these hereditary syndromes are often variable in individuals within a single family (e.g., age of onset, tumor site, and number of primary tumors). The risk of developing cancer in individuals with one of these hereditary syndromes depends upon numerous variables including the gender and age of the individual.

**Hereditary Breast/Ovarian Cancer Syndrome**

The overall prevalence of disease-related mutations in BRCA1 and BRCA2 genes has been estimated as 1 in 300 and 1 in 800, respectively. Currently, hundreds of unique mutations have been identified in both BRCA1 and BRCA2 genes. However, a number of founder effects (see Table 1) have been observed in certain populations, wherein the same mutation has been found in multiple,
unrelated families and can be traced back to a common ancestor. Among the Ashkenazi Jewish population, for example, the frequency of 187delAG and 5385insC mutations in BRCA1 and the 6174delT mutation in BRCA2 approximates 1 in 40.\textsuperscript{6,22} Certain founder mutations have also been identified in other populations.\textsuperscript{20,23-28} It has been estimated that over 90% of early onset cancers in families with both breast and ovarian cancers are caused by mutation(s) in the BRCA1 or BRCA2 genes.\textsuperscript{29} Hence, the degree of clinical suspicion for a BRCA mutation in a single individual with both breast and ovarian cancer or someone with a family history of both breast and ovarian cancer should be very high.

Both the BRCA1 and BRCA2 genes encode for proteins involved in tumor suppression. The BRCA1 gene is located on chromosome 17 and is believed to be involved in both DNA repair and in the regulation of cell-cycle checkpoints in response to DNA damage. However, the molecular mechanism through which BRCA1 functions to preserve genomic stability remains unclear.\textsuperscript{30} The BRCA2 gene, located on chromosome 13, is involved in repair of replication-mediated double-strand DNA breaks.\textsuperscript{31,32}

Mutations in the BRCA1 or BRCA2 genes can be highly penetrant (for definition, see Table 1) although the probability of cancer development in carriers of BRCA1 or BRCA2 mutations is variable, even within families with the same mutation.\textsuperscript{33-35} Estimates of penetrance range from 41% to 90% lifetime risk for breast cancer, with an increased risk of contralateral breast cancer.\textsuperscript{36-42} In addition, female carriers of these genes have an estimated 8% to 62% lifetime risk for ovarian cancer, depending upon the population studied.\textsuperscript{37,54,38-44} In a meta-analysis (2007) of published data that evaluated BRCA1 and BRCA2 penetrance, estimates for mean cumulative risks of breast cancer and ovarian cancer by age 70 years for BRCA1 mutation carriers were 57% and 40%, respectively.\textsuperscript{38} The corresponding estimates for BRCA2 mutation carriers were 49% and 18%, respectively. In a recent prospective analysis of risk estimates from individuals with BRCA1 and BRCA2 mutations in the UK (N=1887), estimates for mean cumulative risks of breast cancer and ovarian cancer by age 70 years for BRCA1 mutation carriers were 60% and 59%, respectively.\textsuperscript{41} The corresponding estimates for BRCA2 mutation carriers were 55% and 16.5%, respectively. Among the patients diagnosed with unilateral breast cancer (n=651), the mean cumulative risks for contralateral breast cancer by age 70 years were estimated to be 83% for BRCA1 carriers and 62% for BRCA2 carriers.\textsuperscript{41} At present, it is unclear whether penetrance is related to the specific mutation identified in a family or whether additional factors, either genetic or environmental, affect disease expression. It is generally accepted, however, that carriers of mutations in BRCA1 or BRCA2 genes have an excessive risk for both breast and ovarian cancer that warrants consideration of more intensive screening and preventive strategies.

Some histopathologic features have been reported to occur more frequently in breast cancers characterized by a BRCA1/2 mutation. For example, several studies have shown that BRCA1 breast cancer is more likely to be characterized as ER-, PR-negative, and HER2-negative (i.e., “triple negative”).\textsuperscript{45-50} Studies have reported BRCA1 mutations in 9% to 28% of patients with triple-negative breast cancer.\textsuperscript{50-56} In addition, it appears that among patients with triple-negative disease, BRCA mutation carriers were diagnosed at a younger age compared with non-carriers.\textsuperscript{53,57} A recent study in a large cohort of patients with triple-negative breast cancer (N=403) reported a median age of diagnosis of 39 years among carriers of BRCA1 mutations (n=65).\textsuperscript{52} Patients in this population-based study were unselected for family history or age. Among the group of patients with early onset (age
at diagnosis <40 years) triple-negative breast cancer (n=106), the incidence of BRCA1 mutations was 36%; among those diagnosed before age 50 years (n=208), the incidence was 27%. For patients with triple-negative breast cancer with a family history of breast and/or ovarian cancer (n=105), BRCA1 mutations were found in 48%.

An increased incidence of BRCA mutations was reported in triple-negative breast cancer cases from at-risk populations. Among Ashkenazi Jewish women with breast cancer unselected for family history (N=451), triple-negative disease was observed in 14% and BRCA founder mutations were found in 11% of patients. Among the subgroup with triple-negative breast cancer (n=65), the incidence of BRCA mutations was 39% (BRCA1 mutation in 30%; BRCA2 mutation in 9%). Although many of the mutation studies in triple-negative breast cancer have reported on the association with BRCA1 mutations, several reports have also suggested the role of BRCA2 mutations in triple-negative breast cancer. The incidence of BRCA2 mutations range from 4% to 17% in studies of triple-negative breast cancer cases unselected for age or family history.

An increased frequency of other malignancies has been reported in families with mutations in the BRCA1 or BRCA2 gene. Germline BRCA1 and BRCA2 mutations have been associated with an increased risk of prostate cancer in numerous reports. In particular, BRCA2 mutations have been associated with 2- to 6-fold increase in risk of prostate cancer, while increased risks were not observed for BRCA1 mutation carriers in some studies. Prostate cancer with germline BRCA mutations appear to have a more aggressive phenotype (e.g., more frequently associated with Gleason score ≥8) than tumors from non-carrier patients. A recent study in a large cohort of patients with prostate cancer from Spain (N=2019) showed that the group of patients with BRCA mutations had significantly higher rates of aggressive prostate cancer (Gleason score ≥8), nodal involvement and distant metastasis compared with non-carriers. Moreover, cause-specific survival outcome was significantly poorer in BRCA mutation carriers compared with non-carriers (median survival 8.6 years vs. 15.7 years; P=0.015). Subgroup analysis by mutation type showed poor outcomes in patients with BRCA2 mutations (n=61); the role of BRCA1 mutations was not well defined, possibly due to the small patient size (n=18) and limited follow-up in this subgroup. Prostate cancer in patients with BRCA2 mutations has also been associated with a higher histologic grade in other studies.

In addition, analyses of data obtained from cancer registries and treatment center databases showed that BRCA2 mutation carriers with prostate cancer had more aggressive or rapidly progressive disease, and significantly decreased survival compared with patients who were BRCA1 mutation carriers or non-carriers. In a study of patients with prostate cancer from a population-based cancer registry in Iceland (N=596), patients with BRCA2 mutations had significantly decreased median survival compared with non-carriers (having wild type BRCA2) patients (2 years vs. 12 years; P<0.001). Moreover, in a study of patients with prostate cancer using data obtained from cancer center databases (N=301), patients with BRCA2 mutations had significantly decreased median survival compared with patients with BRCA1 mutations (4 years vs. 8 years; P<0.01). BRCA2 mutation carriers have also been reported to have a higher risk of pancreatic cancer and melanoma. Both BRCA1 and BRCA2 mutations have been associated with increased propensity for developing pancreatic cancer. In an analysis of samples taken from patients with familial pancreatic cancer (kindreds in which ≥3 family members had pancreatic cancer, at least 2 of which were first-degree relatives), BRCA2 mutations were detected in 17% of patient samples. Among the Ashkenazi Jewish population, BRCA2 mutations were detected in 48%.
mutations have been identified in about 4% of patients with pancreatic cancer.\textsuperscript{74,79}

Some data related to the risk of cancers in this population at some sites other than the breast/ovary are contradictory.\textsuperscript{80} For example, it has been suggested that the increased risk of endometrial cancer observed in some \textit{BRCA1} or \textit{BRCA2} mutation carriers is mainly due to the use of tamoxifen therapy by these women rather than the presence of a gene mutation.\textsuperscript{81}

Germline mutations in \textit{BRCA1} and \textit{BRCA2} are responsible for 5% to 10% of epithelial ovarian cancers (i.e., ovarian cancer developing on the surface of the ovary).\textsuperscript{82} Increased risks of cancers of the fallopian tube and primary peritoneal cancer are also observed in this population. In the setting of an invasive ovarian cancer diagnosis, as many as 15% of unselected individuals will have a germline \textit{BRCA1} or \textit{BRCA2} mutation.\textsuperscript{42,83} However, it has been reported that about half of families showing a genetic predisposition to ovarian cancer do not have identifiable mutations in \textit{BRCA1}/2 genes.\textsuperscript{84} Hence, other gene mutations predisposing to ovarian cancer are likely to exist.\textsuperscript{85} Of note, ovarian cancer is a component tumor of Lynch syndrome which is associated with germline mutations in mismatch repair genes.\textsuperscript{86} Interestingly, results from a prospective study suggest that women from families at increased risk of hereditary breast cancer without site-specific \textit{BRCA} mutations are not at increased risk for ovarian cancer, although these results may have been confounded by the ethnic characteristics and size of the study population.\textsuperscript{87}

It is interesting to note that several recent studies have reported more favorable survival outcomes among \textit{BRCA1}/2 mutation carrier patients with ovarian cancer compared with non-carrier patients.\textsuperscript{88-93} In a case-control study of patients with epithelial ovarian cancer (N=66), patients with \textit{BRCA1}/2 mutations had improved outcomes compared with patients with non-hereditary ovarian cancer, including significantly longer median survival from time of diagnosis (101 months vs. 35 months; \textit{P}<0.002).\textsuperscript{92} In a large case-control study of Jewish patients with epithelial invasive ovarian cancer (N=779), patients with \textit{BRCA1}/2 mutations had significantly longer median survival compared with non-carrier patients (54 months vs. 38 months; \textit{P}=0.002).\textsuperscript{91} Results from a recent pooled analysis from 26 observational studies that included invasive epithelial ovarian cancer cases from \textit{BRCA1}/2 mutation carriers (n=1213) and non-carriers (n=2666) showed favorable survival outcomes for patients with \textit{BRCA1}/2 mutations.\textsuperscript{89} The 5-year survival rate for non-carriers, \textit{BRCA1} carriers and \textit{BRCA2} carriers was 36%, 44%, and 52%, respectively. The survival advantage compared with non-carriers was significant for both the \textit{BRCA1} carriers (hazard ratio=0.78; 95% CI, 0.68-0.89; \textit{P} <0.001) and \textit{BRCA2} mutation carriers (hazard ratio=0.61; 95% CI, 0.50-0.76; \textit{P} <0.001).\textsuperscript{89} In a recent population-based case-control study of women with invasive epithelial (nonmucinous) ovarian cancer (N=1001) from the Australian Ovarian Cancer Study Group, \textit{BRCA1}/2 mutation carriers had improved survival outcomes compared with non-carriers in terms of median progression-free survival (20 months vs. 16 months; not statistically significant) and median survival (62 months vs. 55.5 months; \textit{P}=0.031).\textsuperscript{88} Moreover, \textit{BRCA} mutation carriers appeared to be more responsive to cytotoxic chemotherapy (regardless of class of agent) compared with non-carrier patients. Outcomes appeared to be most favorable for \textit{BRCA2} mutation carriers; in the subgroup of patients with \textit{BRCA2} mutations (n=53), the median survival was 70 months.\textsuperscript{88} In an observational study of patients with high-grade serous ovarian cancer (N=316), patients with \textit{BRCA2} mutations had significantly favorable survival outcomes (hazard ratio=0.33; 95% CI, 0.16–0.69; \textit{P}=0.003; 5-year rate: 61% vs. 25%) and progression-free survival (hazard ratio=0.40; 95% CI, 0.22–0.74;
P=0.004; 3-year rate: 44% vs. 16%) compared with non-carrier patients (having wild type BRCA). Additionally, BRCA2 mutations were associated with significantly higher response rates (compared with non-carriers or with BRCA1 mutation carriers) to primary chemotherapy. In contrast, BRCA1 mutations were not associated with prognosis or improved chemotherapy response.

The histology of ovarian cancers in carriers of a BRCA1 or BRCA2 mutation is more likely to be characterized as serous adenocarcinoma and high grade compared with ovarian cancers in non-mutation carriers, although endometrioid and clear cell ovarian cancers have also been reported in the former population. In studies of women with BRCA1/2 mutations who underwent risk reduction salpingo-oophorectomy (RRSO), occult gynecological carcinomas were identified in 4.5%-9% of cases based on rigorous pathological examinations of the ovaries and fallopian tubes. Tubal intraepithelial carcinoma (TIC) is thought to represent an early precursor lesion for serous ovarian cancers, and TIC (with or without other lesions) was detected in 5% to 8% of cases from patients with BRCA1/2 mutations who underwent RRSO. The fimbriae or distal tube was reported to be the predominant site of origin for these early malignancies found in patients with BRCA1/2 mutations. Although TIC appeared to present more frequently among BRCA1/2 mutation carriers compared with non-carriers undergoing RRSO, TIC has also been documented among patients with serous carcinomas unselected for family history or BRCA mutation status. Because TIC was identified in individuals who underwent surgery for risk reduction (for BRCA1/2 mutation carriers) or other gynecological indications, the incidence and significance of these early lesions within the general population is unclear. Hence, at the present time, there is no justifiable role for BRCA testing for cases based solely on the finding of TIC during pathology evaluation for gynecological indications.

Male carriers of a BRCA gene mutation also have a greater risk for cancer susceptibility. In one study of 26 high-risk families with at least one case of male breast cancer, 77% demonstrated a BRCA2 mutation. Among male patients with breast cancer who were not selected on the basis of family history, 4% to 14% tested positive for a germline BRCA2 mutation. In a recent series of male breast cancer cases (N=115; primarily from cancer registry data), BRCA2 mutations were detected in 16% of cases; the incidence of BRCA2 mutations was 40% among patients selected for family history of breast cancer and 13% among those unselected for family history. For males with a BRCA2 mutation, the cumulative lifetime risk of breast cancer has been estimated at 7% to 8%. In contrast, for men without such a mutation, the lifetime risk of breast cancer has been estimated at approximately 0.1% (1 in 1,000). The NCCN panel recommends that individuals from a family with a known deleterious BRCA1 or BRCA2 mutation be considered for testing (see Guidelines section on HBOC Syndrome Testing Criteria).

In individuals from a family without a known deleterious BRCA mutation, testing should be considered for those individuals who meet the testing criteria discussed below. In evaluating risks based on family history factors, the maternal and paternal sides should be considered independently. For the testing criteria mentioned below, “close relatives” pertain to first-, second- or third-degree blood relatives on the same side (either maternal or paternal side) of the family. Individuals with a limited family history (e.g., having fewer than 2 first- or second-degree relatives or female relatives surviving beyond 45 years of age on either the maternal or paternal side) may have an underestimated probability...
of a familial gene mutation. The panel recommends that patients with a personal history of breast cancer in addition to one or more of the following criteria be considered for BRCA1/BRCA2 testing:

- Diagnosed at age 45 years or younger;
- Having 2 breast primaries (bilateral tumors or 2 or more clearly separate ipsilateral tumors, occurring synchronously or asynchronously) with the first breast cancer diagnosed at age 50 years or younger;
- Diagnosed at 50 years or younger with 1 or more close relative with breast cancer at any age (or with a limited family history);
- Diagnosed with triple-negative breast cancer at age 60 years or younger;
- Diagnosed at any age with 1 or more close relative with breast cancer diagnosed at age 50 years or younger;
- Diagnosed at any age with 2 or more close relatives with breast cancer at any age; diagnosed at any age with 1 or more close relative with epithelial ovarian cancer diagnosed at any age;
- Diagnosed at any age with 2 or more close relatives with pancreatic cancer or aggressive prostate cancer (Gleason score ≥7) at any age; or
- Having a close male relative with breast cancer at any age.

In unaffected individuals with a family history only (i.e., no personal history of breast or ovarian cancer), significant limitations of interpreting test results should be discussed prior to any testing. Moreover, testing of unaffected individuals should only be considered when an appropriate affected family member is unavailable for testing. Clinical judgement should be used to evaluate each unaffected individual for his/her likelihood of carrying the mutation based on factors such as the unaffected individual's current age and the age of unaffected female relatives who link the individual with an affected close relative.

In patients with a personal history of breast cancer and with an ethnic background associated with higher mutation frequency (e.g., Ashkenazi Jewish heritage), no additional family history may be needed to meet testing criteria. In addition, the NCCN panel recommends testing for patients with a personal history of the following:

- Epithelial ovarian cancer diagnosed at any age;
- Male breast cancer diagnosed at any age; or
- Pancreatic cancer or aggressive prostate cancer (Gleason score ≥7) diagnosed at any age, with 2 or more close relatives with breast cancer and/or ovarian cancer and/or pancreatic or aggressive prostate cancer diagnosed at any age.

Li-Fraumeni Syndrome

Li-Fraumeni syndrome (LFS) is a rare hereditary cancer syndrome associated with germline TP53 gene mutations.\(^\text{13}\) It has been estimated to be involved in only about 1% of hereditary breast cancer cases,\(^\text{112}\) although results from a recent study suggest that germline TP53 gene mutations may be more common than previously believed.\(^\text{113}\) The tumor suppressor gene, TP53, is located on chromosome 17,\(^\text{114,115}\) and the protein product of the TP53 gene (i.e., p53) is located in the cell nucleus and binds directly to DNA. It has been called the “guardian of the genome” and plays important roles in controlling the cell cycle and apoptosis.\(^\text{114-116}\) Germline mutations in the TP53 gene have been observed in over 50% (and in over 70% in some studies) of families
meeting the classic definition of LFS (see Guidelines section on Li-
Fraumeni Syndrome Testing Criteria). Additional studies are
needed to investigate the possibility of other gene mutations in families
meeting these criteria not carrying germline TP53 mutations.

LFS, a highly penetrant cancer syndrome associated with a high life-
time risk of cancer, is characterized by a wide spectrum of neoplasms
occurring at a young age. It is associated with soft-tissue sarcomas,
osteosarcomas (although Ewing’s sarcoma is less likely to be
associated with LFS), premenopausal breast cancer, acute leukemia,
and cancer of the colon, adrenal cortex, and brain tumors. Sarcoma, breast cancer, adrenocortical tumors and certain brain tumors
have been referred to as the “core” cancers of LFS since they account
for the majority of cancers observed in individuals with germline
mutations in the TP53 gene, and in one study, at least one of these
cancers was found in one or more members of all families with a
germline TP53 gene mutation. Interestingly, recent retrospective
studies have reported a very high frequency of HER2-positive breast
tumors (67%–83% of evaluated breast tumors) among patients with
germline TP53 mutations, which suggest that amplification of HER2
may arise in conjunction with TP53 mutations. This association
between HER2-positive breast cancer and germline TP53 mutations
warrants further investigation, as such patients may potentially benefit
from chemoprevention therapies that incorporate HER2-targeted
agents.

Individuals with LFS often present with certain cancers (e.g., soft-tissue
sarcomas, brain tumors, and adrenocortical carcinomas) in early
childhood, and have an increased risk of developing multiple primary
cancers during their lifetimes. Results of a segregation analysis of
data collected on the family histories of 159 patients with childhood soft
tissue sarcoma showed carriers of germline TP53 mutations to have
estimated cancer risks of approximately 60% and 95% by age 45 and
70 years, respectively. Although similar cancer risks are observed in
men and women with LFS when gender-specific cancers are not
considered, female breast cancer is commonly associated with the
syndrome. It is important to mention that estimations of cancer risks
associated with LFS are limited to at least some degree by selection
bias since dramatically affected kindreds are more likely to be identified
and become the subject of further study.

A number of different sets of criteria have been used to help identify
individuals with LFS. For the purposes of the NCCN Guidelines, 2 sets
of these criteria are used to facilitate the identification of individuals who
are candidates for TP53 gene mutation testing.

Classic LFS criteria, based on a study by Li and Fraumeni involving 24
LFS kindreds, include the following: member of a kindred with a known
TP53 mutation; combination of an individual diagnosed at age 45 years
or younger with a sarcoma, and a first-degree relative diagnosed with
cancer at age 45 years or younger, and an additional first- or second-
degree relative in the same lineage with cancer diagnosed at age
younger than 45 years or a sarcoma at any age (see Guidelines section
on Li-Fraumeni Syndrome Testing Criteria). Classic LFS criteria have
been estimated to have a high positive predictive value (estimated at
56%) as well as a high specificity, although the sensitivity is relatively
low (estimated at 40%). Thus, it is not uncommon for individuals with
patterns of cancer outside of these criteria to be carriers of germline
TP53 mutations. Classic LFS criteria make up one set of criteria
included in the Guidelines to guide selection of individuals for TP53
gene mutation testing (see Guidelines section on Li-Fraumeni
Syndrome Testing Criteria).
Other groups have broadened the classic LFS criteria to facilitate identification of individuals with LFS. One set of these less strict criteria proposed by Birch and colleagues shares many of the features of classic LFS criteria, although a larger range of cancers are included. Uncommonly, individuals with de novo germline TP53 mutations (no mutation in either biological parent) have been identified. These cases would not be identified as TP53 testing candidates based upon classic LFS criteria due to requirement of a family history. This issue is circumvented, in part, by the criteria for TP53 testing proposed by Chompret and colleagues, which recommends testing for patients with multiple primary tumors of at least 2 “core” tumor types (i.e., sarcoma, breast cancer, adrenocortical carcinoma, brain tumors) diagnosed at age <36 years or patients with adrenocortical carcinoma diagnosed at any age, regardless of family history (see Guidelines section on Li-Fraumeni Syndrome Testing Criteria). The Chompret criteria have an estimated positive predictive value of 20% to 35% and when incorporated as part of TP53 testing criteria in conjunction with classic LFS criteria, have been shown to improve the sensitivity to 95% (i.e., the Chompret criteria added to classic LFS criteria detected 95% of patients with TP53 mutations). The Chompret criteria are the second set of criteria included in the NCCN Guidelines. Although not part of the original published criteria set forth by Chompret et al., the panel recommends adding lung bronchoalveolar cancer and leukemia as one of the core tumor types (for inclusion in criterion 1 and 2 of the Chompret criteria) and also recommends testing individuals with choroid plexus carcinoma (i.e., updated Chompret criteria) was recently proposed by Tinat et al and is supported by the NCCN Guidelines panel. The panel also supports the broader age cut-offs proposed by Tinat et al, based upon a study in a large number of families, which detected germline TP53 mutations in affected individuals with later tumor onsets.

Women with early-onset breast cancer (age of diagnosis ≤35 years), with or without family history of core tumor types, are another group for whom TP53 gene mutation testing may be considered. Several recent studies have investigated the likelihood of a germline TP53 mutation in this population. In a study of TP53 mutations evaluated at a single reference laboratory, Gonzalez et al. found that all women younger than 30 years of age with breast cancer who had a first- or second-degree relative with at least one of the core cancer types (n=5), had germline TP53 mutations. In a recent analysis of data of patients with early-onset breast cancer (age of diagnosis <30 years) tested for TP53 mutation at a single institution (N=28), 6 patients (33%) were found to have TP53 mutations. Among the patients who were tested, a TP53 mutation was found in approximately 8% who did not meet traditional LFS criteria for testing. In another recent study in patients with BRCA1/BRCA2 mutation-negative early-onset breast cancer (age of diagnosis ≤35 years) tested for TP53 mutation at a single institution (N=83), approximately 5% were found to have TP53 mutations. Deleterious TP53 mutations were identified in 3 of 4 patients (75%) with a family history of at least 2 LFS-associated tumors (breast cancer, bone or soft tissue sarcoma, brain tumors or adrenocortical cancer) and in 1 of 17 patients (6%) with a family history of breast cancer only. Among women <30 years of age with breast cancer and without a family history, the incidence of TP53 mutations has been reported at 3% to 8%. Other studies have found an even lower incidence of
germline TP53 gene mutations in this population. For example, Bougeard et al reported that only 0.7% of unselected women with breast cancer before age 33 were carriers of a germline TP53 mutation. Furthermore, Ginsburg and colleagues found no germline TP53 mutations in 95 unselected women with early-onset breast cancer who previously tested negative for BRCA mutations.

Finally, a member of a family with a known TP53 mutation is considered to be at sufficient risk to warrant gene mutation testing, even in the absence of any other risk factors. Individuals not meeting testing criteria should be followed according to recommendations tailored to his/her personal cancer history and family history.

**Cowden Syndrome**

Cowden syndrome, a rare hereditary cancer syndrome, was first described in 1963 and named after the Cowden family, the first family documented with signs of the disease. The incidence of Cowden syndrome has been reported to be 1 in 200,000, although it is likely to be underestimated due to difficulties associated with making a clinical diagnosis of the disease. Cowden syndrome is an autosomal dominant disorder associated with germline mutations in the PTEN ("phosphatase and tensin homologue deleted on chromosome 10") tumor suppressor gene located on chromosome 10q23; the gene is thought to be involved in cell cycle arrest and apoptosis, and other cell survival pathways. It is considered to be part of the spectrum of PTEN hamartoma tumor syndromes (PHTS), which also includes Bannayan-Riley-Ruvalcaba syndrome (BRRS), Proteus syndrome, and Proteus-like syndrome. Additional clinical syndromes related to germline mutations in PTEN include Lhermitte-Duclos disease and autism spectrum disorders with macrocephaly, both of which have been associated with Cowden syndrome. The estimated penetrance of PTEN mutation is high, at approximately 80%. Hamartomas, a common manifestation of these syndromes, are benign tumors resulting from an overgrowth of normal tissue.

Cowden syndrome is associated with multiple hamartomatous and/or cancerous lesions in various organs and tissues, including the skin, mucous membranes, breast, thyroid, endometrium and brain. This syndrome is the only PHTS disorder associated with a documented predisposition to malignancies; hence, it is included in these Guidelines. However, it has been suggested that patients with other PTHS diagnoses associated with PTEN mutations should be assumed to have Cowden-associated cancer risks. In a study of patients meeting diagnostic criteria for Cowden syndrome (N=211; identified from published literature and records from a single institution), the cumulative lifetime risk of any cancer was 89%. PTEN mutations had been identified in 97 of 105 patients (92%) who underwent testing. The cumulative lifetime cancer risks for all evaluable patients (n=210) were 81% for female breast cancer, 21% for thyroid cancer, 19% for endometrial cancer, 15% for renal cancer, and 16% for colorectal cancer. In a recent prospective study that evaluated genotype-phenotype associations between PTEN mutations and cancer risks, a large number of patients meeting modified (relaxed) International Cowden Consortium criteria (N=3399) were enrolled and tested for PTEN mutations. Deleterious germline mutations in PTEN were identified in 368 patients (11%). Calculation of age-adjusted standardized incidence ratios (SIRs) using cancer incidence data from the SEER database showed elevated SIRs among individuals with PTEN mutations for breast cancer (25), thyroid cancer (51), endometrial cancer (43), colorectal cancer (10), renal cancer (31), and melanoma (8.5). The estimated cumulative lifetime cancer risks were 85% for breast, 35% for thyroid, 28% for endometrial, 9% for colorectal, 34% for
renal and 6% for melanoma.\(^{151}\) In another recent study in individuals with PHTS found to have deleterious germline \(PTEN\) \(PTEN\) mutations (\(N=154;\) detailed information available in \(n=146\)), age- and gender-adjusted SIRs were elevated for female breast cancer (39), endometrial cancer (49), female thyroid cancer (43), male thyroid cancer (199.5), female melanoma (28), and male melanoma (39).\(^{152}\) The cumulative lifetime risks in these individuals were 77% for female breast cancer and 38% for thyroid cancer. The cumulative lifetimes risk for any cancer was 85% overall, and women with PHTS were found to have a 2-fold greater cancer risk compared with men with PHTS.\(^{152}\)

Women diagnosed with Cowden syndrome have a high risk of benign fibrocystic breast disease and their lifetime risk of breast cancer has been estimated at 25% to 50% with an average age of 38 to 46 years at diagnosis.\(^{12,149,153}\) Recent studies (as discussed above) have reported a higher cumulative lifetime risk of breast cancer (77–85%) in individuals with Cowden syndrome or \(PTEN\) \(PTEN\) mutations.\(^{150-152}\) There have been only 2 cases of breast cancer reported in men with Cowden syndrome.\(^{12}\)

Thyroid disease, including benign multinodular goiter, adenomatous nodules, and follicular adenomas have been reported to occur in up to approximately 70% of individuals with Cowden syndrome\(^{154}\) and the lifetime risk of thyroid cancer (follicular or papillary) has been estimated at 3% to 10%.\(^{12,155}\) A higher cumulative lifetime risk of thyroid cancer (21–38%) was reported in several recent studies in individuals with Cowden syndrome or \(PTEN\) \(PTEN\) mutations (as discussed earlier).\(^{150-152}\) As in many other hereditary cancer syndromes, affected individuals are more likely to develop bilateral and multifocal cancer in paired organs.\(^{148}\)

Although not well defined, women with Cowden syndrome may have a 5% to 10% risk of endometrial cancer,\(^{12,156}\) and an increased risk of uterine fibroids. Recent studies showed a higher lifetime risk of endometrial cancer (19–28%) in women with Cowden syndrome or \(PTEN\) \(PTEN\) mutations.\(^{150,151}\) As discussed earlier, increased lifetime risks for colorectal cancer (9–16%), renal cancer (15–34%) and melanoma (6%) were also reported recently in individuals with Cowden syndrome or \(PTEN\) \(PTEN\) mutations.\(^{150,151}\) In addition, brain tumors and vascular malformations affecting any organ are occasionally seen in individuals with Cowden syndrome, although the risks for developing these conditions are not well defined.\(^{12}\) It is important to note, however, that most of the data on the frequencies of the clinical features of Cowden syndrome are from compilations of case reports of relatively young individuals who may have subsequently developed additional signs of the disease (i.e., new cancerous lesions), and these data are also likely to be confounded by selection bias.\(^{12}\) Furthermore, a considerable number of these studies were published prior to the establishment in 1996 of the International Cowden Consortium operational diagnostic criteria for the syndrome which were based on published data and the expert opinion of individuals representing a group of centers mainly in North America and Europe.\(^{12,157}\)

Classic features of Cowden syndrome include mucocutaneous papillomatous papules, palmoplantar keratoses, and trichilemmomas (i.e., benign tumors derived from the outer root sheath epithelium of a hair follicle).\(^{12,158}\) Most individuals with Cowden syndrome exhibit characteristic mucocutaneous lesions by their twenties, and such lesions have been reported to occur in 99% of individuals with Cowden syndrome, a syndrome showing nearly complete penetrance.\(^{82,145}\) The presence of 2 or more trichilemmomas has been reported to be pathognomonic for Cowden syndrome.\(^{159,160}\) However, since most of this evidence is from the older literature, it is possible that the association between these 2 entities is somewhat overestimated.\(^{12}\) There are reports of individuals with a solitary trichilemmoma who do not have Cowden syndrome.\(^{159,160}\) Nevertheless, due to the strong association
between these lesions and Cowden syndrome and the difficulty in clinically distinguishing between a trichilemmoma and another mucocutaneous lesion, it is important that a diagnosis of trichilemmoma is histologically confirmed.

It has historically been reported that about 40% individuals with Cowden syndrome have gastrointestinal polyps (often colonic), although more recent data suggest that this risk may be 80% or higher. Indeed, a recent analysis of PTEN mutation carriers reported gastrointestinal polyps in 93% of patients. Most of the polyps are hamartomatous, although ganglioneuromas (i.e., rare, benign peripheral nervous system tumors) have also been reported to occur. However, early-onset (age <50 years) colorectal cancer has been reported in 13% of patients with PTEN mutation-associated Cowden syndrome, suggesting that routine colonoscopy may be warranted in this population.

Adult Lhermitte-Duclos disease (LDD) and autism spectrum disorder characterized by macrocephaly are strongly associated with Cowden syndrome. A rare, slow growing, benign hamartomatous lesion of the brain, LDD is a dysplastic gangliocytoma of the cerebellum. In a study of individuals meeting the diagnostic criteria for Cowden syndrome, the cumulative lifetime risk of LDD was reported to be 32%. The preponderance of evidence supports a strong association between adult-onset LDD and the presence of a PTEN gene mutation, although exceptions have been reported. In addition, there is a relatively large body of evidence to support that 10% to 20% of individuals with autism spectrum disorder and macrocephaly carry germline PTEN mutations. Macrocephaly (defined as head circumference greater than the 97th percentile) is a common finding in patients with Cowden syndrome. It has been estimated that approximately 80% of individuals with this syndrome will exhibit this clinical finding.

The BRRS variant of PHTS has been characterized by the presence of multiple lipomas, gastrointestinal hamartomatous polyps, macrocephaly, hemangiomas, developmental delay, and in males, pigmented macules on the glans penis, although formal diagnostic criteria have not been established for this syndrome. PTEN gene mutations testing in individuals characterized with BRRS have been reported in approximately 60% of these patients. Further, in another study, 10% of patients with BRRS for whom a PTEN gene mutation test was negative were shown to be carriers of large PTEN gene deletions.

The PTEN mutation frequency in individuals meeting International Cowden Consortium criteria for Cowden syndrome has been estimated at about 80%. However, a recent evaluation of data based on samples analyzed at a single academic pathology laboratory (N=802 evaluable) reported a much lower frequency (34%) of PTEN mutations among individuals meeting diagnostic criteria for Cowden syndrome. The authors concluded that the current Consortium diagnostic criteria are not as sensitive in identifying individuals with PTEN mutations as previously estimated. The International Cowden Consortium criteria have been updated several times since 1996 and they have largely served as the basis for the list of PTEN mutation testing criteria included in the NCCN Guidelines. On the basis of literature reports and expert consensus, the panel has recently revised both the list of criteria associated with this genetic syndrome as well as the combinations of criteria that establish which individuals are candidates for PTEN gene mutation testing (see Guidelines section on Cowden Syndrome Testing Criteria). Similar to earlier versions, criteria are grouped into 3 general categories. A patient is considered for PTEN gene mutation testing based on whether he/she meets certain criteria or combinations of criteria from these 3 categories. The first criteria category includes individuals meeting diagnostic criteria for Cowden
syndrome\textsuperscript{172}; or a personal history of BRRS, adult LDD, autism spectrum disorder with macrocephaly, or 2 or more biopsy proven trichilemmomas. Any individual presenting with one or more of these diagnoses warrants \textit{PTEN} testing. Previously, some of the criteria from this group have sometimes been referred to as “pathognomonic” although it is unlikely that any of these conditions can stand alone as a definitive diagnostic criterion of Cowden syndrome. Another criterion which can be considered to be sufficient to warrant \textit{PTEN} gene mutation testing is a family history which includes the presence of a known deleterious \textit{PTEN} mutation.

The next category of criterion represents “major” features associated with Cowden syndrome.\textsuperscript{143,146,172,174} The major criteria include the presence of breast cancer, macrocephaly (i.e., megalocephaly),\textsuperscript{169} endometrial cancer, follicular thyroid cancer, multiple gastrointestinal hamartomas or ganglioneuromas, macular pigmentation of glans penis, and certain mucocutaneous lesions that are often observed in patients with Cowden syndrome (i.e., one biopsy proven trichilemmoma, multiple palmoplantar keratoses, multiple or extensive oral mucosal papillomatosis, multiple cutaneous facial papules). With respect to decisions related to the presence of mucocutaneous lesions, the panel did not consider the available literature to be adequate to accurately specify the number or extent of these lesions required for the condition to be defined as a major criterion for Cowden syndrome, and clinical judgment is needed when evaluating such lesions. An individual exhibiting 2 or more major criteria where one of these is macrocephaly meets the testing threshold. An individual with 3 or more major criteria (without macrocephaly) also meets the threshold for \textit{PTEN} testing. In addition, individuals exhibiting 1 major criterion with 3 or more minor criteria (discussed below) also meet the testing threshold; if an individual exhibits 2 or more major criteria (e.g., breast cancer and follicular thyroid cancer) but does not have macrocephaly, then one of the major criteria may be included as one of the 3 minor criteria to meet the testing threshold.

The final category of criteria represents features with a “minor” association with Cowden syndrome.\textsuperscript{143,146,172,174} These include autism spectrum disorder (without macrocephaly), colon cancer, esophageal glycogenic acanthosis (3 or more), lipomas, mental retardation, papillary or follicular variant of papillary thyroid cancer, thyroid structural lesions other than follicular thyroid cancer (e.g., adenoma, nodules, goiter), renal cell carcinoma, a single gastrointestinal hamartoma or ganglioneuroma, testicular lipomatosis, or vascular anomalies (including multiple intracranial developmental venous anomalies). The panel felt that evidence from the literature was insufficient to include fibrocystic breast disease, fibromas or uterine fibroids as part of the testing criteria. An individual would need to exhibit 4 or more minor criteria or as discussed above, 3 or more minor and one major criterion to meet testing criteria (see Guidelines section on Cowden Syndrome Testing Criteria and the Discussion section below on Risk Assessment, Counseling, and Management: Cowden Syndrome).

Lastly, an at-risk individual (first-degree relative of an affected individual) with one or more major criterion or 2 or more minor criteria, along with a relative diagnosed with Cowden syndrome or BRRS (for whom testing has not been performed), would also meet the threshold for \textit{PTEN} testing. Individuals not meeting testing criteria should be followed according to recommendations tailored to his/her personal cancer history and family history.

Other Genetic Lesions Associated with Breast/Ovarian Cancer

Although the highly penetrant \textit{BRCA1} and \textit{BRCA2} gene mutations (together with genomic rearrangements in \textit{BRCA} and other high-
penetrance mutations such as \( TP53 \) or \( PTEN \) mutations discussed above) are thought to account for a large proportion of familial breast cancers, other breast and/or ovarian cancer susceptibility genes have been identified. For instance, germline mutations in \( CDH1 \) are associated with hereditary diffuse gastric cancer and lobular breast cancer, and studies have reported a cumulative lifetime risk of breast cancer of 39% to 52% among women who carry \( CDH1 \) mutations.\(^ {15,175} \)

Germline mutations in \( STK11 \) is associated with Peutz-Jeghers syndrome, an autosomal dominant disorder characterized by gastrointestinal polyps, mucocutaneous pigmentation, and elevated risk of gastrointestinal cancers as well as breast or ovarian cancers. Further information on Peutz-Jeghers syndrome can be found in the NCCN Guidelines for Genetic/Familial High-Risk Assessment: Colorectal. Other breast and/or ovarian cancer susceptibility genes with lower or moderate penetrance have been identified in recent years, and include mutations in \( CHEK2, PALB2, BRIP1, RAD51C \), among others.\(^ {176-179} \) In a study of breast cancer patients in the U.S. with strong family history of breast or ovarian cancer but who tested negative for \( BRCA1 \) or \( BRCA2 \) mutations, 12% were found to have large genomic rearrangements (deletion or duplication) in \( BRCA \), and 5% had \( CHEK2 \) mutations.\(^ {178} \) Deleterious \( CHEK2 \) mutations have been reported to occur with a higher frequency in Northern and Eastern European countries compared with North America.\(^ {176,177,180,181} \) The cumulative lifetime risk of breast cancer in women with \( CHEK2 \) mutations and familial breast cancer has been estimated to range from approximately 28% to 37%.\(^ {182,183} \) In the NCCN Guidelines for Genetic/Familial High-Risk Assessment: Breast and Ovarian, the panel specifically focuses on assessment of known high-penetrance mutations (i.e., \( BRCA1, BRCA2, TP53 \) and \( PTEN \)) and recommendations for genetic testing, counseling and management strategies in individuals with these mutations. A comprehensive review of other lower or moderate-penetrance susceptibility genes is beyond the scope of the Guidelines, but has been reviewed in a recent publication.\(^ {176} \)

With recent advances in genomic sequencing technologies, it is now possible to test for multiple breast and/or ovarian cancer susceptibility genes in parallel using multigene or multiplex panels.\(^ {176,179,184} \) Although multigene sequencing approaches may be resource efficient in terms of time and costs, several issues must be addressed before multiplex testing panels can be incorporated as part of standard clinical practice. At the present time, no consensus exists on recommendations for optimal management or surveillance approaches for carriers of lower or moderate penetrance genes, and no data are available to address cancer risk assessments in individuals who are found to carry multiple gene mutations with moderate penetrance.\(^ {184} \) Importantly, additional genetic counseling approaches must be vetted and developed in order to adequately address the limitations and implications associated with interpretation of multiplex testing results.

**Initial Risk Assessment**

For a patient concerned about or suspected of having a hereditary propensity to breast and/or ovarian cancer, an initial risk evaluation should be performed in order to determine if a formal risk assessment should be undertaken (see Guidelines section on Criteria for Further Genetic Risk Evaluation). The first step in this preliminary assessment is a broad and flexible evaluation of the personal and family history of the individual with respect to breast and/or ovarian cancer.\(^ {185,186} \) The magnitude of the risk increases with the number of affected relatives in the family, the closeness of the relationship, and is affected by the age at which the affected relative was diagnosed.\(^ {187,188} \) The younger the age at diagnosis, the more likely it is that a genetic component is present. When assessing a family history for a hereditary pattern, the equal...
likelihood of paternal or maternal transmission of a gene that predisposes to breast cancer must also be kept in mind.

If an individual or a close family member of that individual meets any one of the criteria presented in the NCCN Guidelines (see Guidelines section on Criteria for Further Genetic Risk Evaluation), that individual may be at increased risk for breast and/or ovarian cancer, and a referral for genetic assessment is recommended. The maternal and paternal sides of the family should be considered independently for familial patterns of cancer.

For individuals potentially meeting established criteria for one or more of the hereditary cancer syndromes, genetic testing should be considered along with appropriate pre-test counseling. A genetic counselor and/or a medical geneticist should be involved in this process. Those not meeting criteria for testing who are still considered at increased risk of familial breast cancer are also likely to benefit from appropriate risk-reduction strategies (e.g., a change in the frequency of, or modalities used for, breast cancer screening). The panel recommends that these individuals follow recommendations in the NCCN Guidelines for Breast Cancer Screening and Diagnosis.

**Formal Risk Assessment and Genetic Counseling**

**Risk Assessment**

Cancer genetic risk assessment and genetic counseling is a multi-step process of identifying and counseling individuals at risk for familial or hereditary cancer.

Cancer genetic risk assessment involves use of pedigree analysis with available risk assessment models to determine whether a family history is suggestive of sporadic, familial, or hereditary cancer. Risk assessment includes both an evaluation of an individual’s absolute risk of breast and/or ovarian cancer as well as an estimation of the likelihood that the individual has a heritable genetic mutation in his/her family. Genetic risk assessment is a dynamic process and can change if additional relatives are diagnosed with cancer.

Statistical models based on personal and family history characteristics have been developed to estimate a person’s interval and lifetime risks of developing breast cancer. For example, the Claus tables may be useful in providing breast cancer risk estimates for white women without a known cancer-associated gene mutation who have one or two first- or second-degree female relatives with breast cancer. In addition, decision models developed to estimate the likelihood that a $BRCA1/2$ mutation is present include BRCAPRO$^{190,191}$ and the Breast and Ovarian Analysis of Disease Incidence and Carrier Estimation Algorithm (BOADICEA)$^{190}$. A lifetime risk of breast cancer of 20% to 25% or greater as assessed by models based largely on family history has been used in some guidelines to identify a woman as being at high risk of breast cancer. For example, this risk threshold was used in updates to the American Cancer Society (ACS) guidelines on breast screening which incorporates magnetic resonance imaging (MRI).$^{192,193}$

First-degree relatives of individuals with a known deleterious gene mutation in $BRCA1/2$, $TP53$ or $PTEN$ genes are considered to have a 50% risk of carrying that mutation.

**Evaluation of Patient’s Needs and Concerns**

The first step in evaluating a individual’s risk for hereditary breast cancer is to assess her/his concerns and reasons for seeking counseling and to guarantee that her/his personal needs and priorities will be addressed in the counseling process. Several studies have documented a highly exaggerated perception of risk among women with a family history of breast cancer who seek cancer risk counseling.$^{194}$
This is a situation that can interfere with the adoption of appropriate health behaviors. In addition, the patient’s knowledge about the benefits, risks, and limitations of genetic testing should be assessed as well as the patient's goals. A positive, supportive interaction with the counseling team is an important determinant of ultimate satisfaction with the counseling process and of adherence to recommended health behaviors.

**Detailed Family History**
A detailed family history is the cornerstone of effective genetic counseling. An examination of family history involves development of an expanded pedigree collected beginning with the health of the proband (index case) and proceeding outward to include first-, second-, and third-degree relatives on both the maternal and paternal sides. Standardized pedigree nomenclature should be used. Unaffected family members, both living and deceased, are also included, as their histories also provide information about the magnitude of genetic risk.

Information collected includes cancer diagnoses by primary site, age at diagnosis, bilaterality (when appropriate), and current age or age at death. Whenever possible, cancer diagnoses in the family are verified by obtaining medical records, pathology reports, or death certificates. This is particularly important in the case of a report of an “abdominal” cancer in a female relative—a situation in which cancers of the cervix, uterus, ovary, and/or colon is often confused. It is also important to know the ancestry/ethnicity of the individual.

Other medical conditions that may be associated with or predispose an individual to breast and/or ovarian cancer should also be noted. Family history data are then graphically represented on a pedigree that follows standard nomenclature to illustrate family relationships and disease information. Factors that limit the informativeness of the pedigree are small family size, a small number of individuals of the susceptible gender for sex-limited cancers, reduced penetrance, early deaths in family members (which precludes the possibility that they will develop adult diseases), prophylactic surgeries that remove an organ from subsequent risk of cancer (e.g., hysterectomy for uterine fibroids in which the ovaries are also removed), adoptions, and inaccurate or incomplete information on family members.

A recent prospective registry study of 306 women diagnosed with breast cancer at < 50 years of age, who had no first- or second-degree relatives with breast or ovarian cancer, showed that those individuals with a limited family history (defined as fewer than 2 first- or second-degree female relatives or fewer than 2 female relatives surviving beyond age 45 years in either lineage) may have an underestimated probability of a *BRCA1/2* gene mutation based on models dependent on family history.

**Medical and Surgical History**
The collection of a detailed medical and surgical history from the proband allows the counselor to estimate the contribution of other risk factors that may interact with or modify family history to determine the risk of breast cancer. A history of previous breast biopsies, especially those in which the pathology revealed atypical hyperplasia or lobular carcinoma in situ (LCIS), is associated with an increased risk of breast cancer. Pathologic verification of these diagnoses is encouraged. History of salpingo-oophorectomy and potential exposure to carcinogens (e.g., radiation therapy) should also be included in the patient’s assessment. When taking the medical history, the clinician should also be alert to the physical manifestations of Cowden syndrome, especially skin conditions (see below under Focused Physical Examination).
Reproductive variables are important determinants of risk for both breast and ovarian cancer, suggesting a significant contribution of hormones to the etiology of these cancers. This possible link is supported by the increased breast cancer risk seen among women who have had prolonged exposure to exogenous estrogens and progestins and the reduction in risk for ovarian cancer observed among women who report using oral contraceptives.

### Focused Physical Examination

A physical examination performed by a physician or nurse should be part of the risk assessment. Particular attention should be paid to organs/areas of the body known to be affected in individuals with specific hereditary breast and/or ovarian syndromes. For example, certain patterns of mucocutaneous manifestations are associated with Cowden syndrome, as discussed earlier; a focused physical examination for Cowden syndrome should include a comprehensive dermatologic examination (including oral mucosa), evaluation of head circumference (to determine presence of macrocephaly) and palpation of the thyroid (see section above on Cowden Syndrome).

### Genetic Counseling

Genetic counseling is a critical component of the cancer risk assessment process. Counseling for hereditary breast and/or ovarian cancer uses a broad approach to place genetic risk in the context of other related risk factors, thereby customizing counseling to the experiences of the individual. The purpose of cancer genetic counseling is to educate individuals about the genetic, biological, and environmental factors related to the individual’s cancer diagnosis and/or risk of disease to help them derive personal meaning from cancer genetic information, to and empower them to make educated, informed decisions about genetic testing, cancer screening, and cancer prevention. Individuals need to understand the relevant genetic, medical, and psychosocial information and be able to integrate this information before they can make an informed decision. The presentation of information is most effective when tailored to the age and education of the person undergoing counseling, and that individual’s personal exposure to the disease, level of risk, and social environment.

Pre-test counseling is an essential element of the genetic counseling process in the event that genetic testing for a gene mutation associated with a hereditary cancer syndrome is under consideration. The foundation of pre-test genetic counseling is based on the principle of informed consent. Pre-test counseling should include a discussion of why the test is being offered and how test results may impact medical management, cancer risks associated with the gene mutation in question, the significance of possible test results (see section on Genetic Testing, below), the likelihood of a positive result, technical aspects and accuracy of the test, economic considerations, risks of genetic discrimination, psychosocial aspects, confidentiality issues, as well as other topics. A discussion of confidentiality issues should include an explanation of the federal Genetic Information Nondiscrimination Act (GINA) enacted in 2008 which prohibits health insurers and employers from discrimination on the basis of genetic test results.

Post-test counseling must also be performed and includes disclosure of results, a discussion of the significance of the results, an assessment of the impact of the results on the emotional state of the individual, a discussion of the impact of the results on the medical management of the individual, and how and where the patient will be followed. In addition, identification of a gene mutation associated with a hereditary predisposition to breast and/or ovarian cancer in an individual necessitates a discussion of possible inherited cancer risk to relatives.
and the importance of informing family members about test results. It may also be appropriate to offer genetic testing to both parents of an individual who tests positive for one of these gene mutations (i.e., BRCA1/2, PTEN, TP53) when the lineage is in question.

**Genetic Testing**

The selection of appropriate candidates for genetic testing is based on the personal and familial characteristics that determine the individual’s prior probability of being a mutation carrier, and on the psychosocial degree of readiness of the person to receive genetic test results. The potential benefits, limitations, and risks of genetic testing are also important considerations in the decision-making process. Many women feel that they are already doing everything they can to minimize their risk of developing breast cancer, and others fear the emotional toll of finding out that they are a mutation carrier, especially if they have children who would be at risk of inheriting the mutation. For those who choose not to proceed with testing, the counseling team tailors recommendations for primary and secondary prevention based on the individual’s personal and family history.

In the statement on Genetic Testing for Cancer Susceptibility from the American Society of Clinical Oncology (ASCO) updated in 2003, genetic testing is recommended when there is: (i) a personal or family history suggesting genetic cancer susceptibility (ii) the test can be adequately interpreted and (iii) the results will aid in the diagnosis or influence the medical or surgical management of the patient or family members at hereditary risk of cancer. These recommendations were reiterated in the latest 2010 ASCO update on Genetic and Genomic Testing for Cancer Susceptibility with respect to testing individuals for gene mutations known to cause hereditary breast and/or ovarian cancer(s).

As part of pre-test counseling, the counselor reviews the distinctions between true-positive, true-negative, indeterminate (or uninformative), and inconclusive (or variants of unknown significance) test results (see Table 2), as well as the technical limitations of the testing process. A clear distinction is made between the probability of being a mutation carrier and the probability of developing cancer. The probabilistic nature of genetic test results and the potential implications for other family members must also be discussed. Individuals who have received allogeneic hematopoietic stem cell transplantation (HSCT) should not have molecular genetic testing performed on blood samples, as these blood cells would represent donor-derived DNA. In such cases, DNA of the individual being tested should be extracted from a fibroblast culture, if available. If this is not possible, buccal cells may be considered as an alternative source for DNA; however, a study has reported that over time, buccal epithelial cells are replaced by donor-derived cells in allogeneic HSCT recipients. Therefore, genetic testing using buccal swab samples may be limited given this known risk of donor DNA contamination.

The genetic testing strategy is greatly facilitated when a deleterious mutation has already been identified in another family member. In that case, the genetic testing laboratory can limit the search for mutations in additional family members to the same location in the gene. In most cases, an individual testing negative for a known familial gene mutation predisposing to breast cancer can be followed with routine breast screening. Individuals who meet testing criteria but do not undergo gene testing should be followed as if a gene mutation (i.e., BRCA, PTEN, or TP53 gene mutation) is present, if they have a close family member who is a known carrier of the deleterious mutation.

For the majority of families in whom mutation status is unknown, it is best to consider testing an affected family member first, especially a
family member with early-onset disease, bilateral disease, or multiple primaries, because that individual has the highest likelihood for a positive test result. Unless the affected individual is a member of an ethnic group for which particular founder gene mutations are known, comprehensive genetic testing (i.e., full sequencing of the genes and detection of large gene rearrangements) should be performed.

For individuals with family histories consistent with a pattern of hereditary breast and/or ovarian cancer on both the maternal and paternal sides, the possibility of a second deleterious mutation in the family should be considered, and full sequencing may be indicated.

In the situation of an unaffected individual with a family history only, the testing of the unaffected individual (or of unaffected family members) should only be considered when no affected family member is available for testing. In such cases, the unaffected individual or unaffected close relative with the highest likelihood of testing positive for the mutation should be tested. A negative test result in such cases, however, is considered indeterminate (see Table 2) and does not provide the same level of information as when there is a known deleterious mutation in the family. Thus, one should be mindful that when testing unaffected individuals (in the absence of having tested affected family members), significant limitations may exist in interpreting the test results.

In the case of hereditary breast/ovarian cancer (i.e., BRCA mutation), if no family member with breast or ovarian cancer is living, consideration can be given to testing first- or second-degree family members affected with cancers thought to be related to the deleterious mutation in question (e.g., prostate or pancreatic cancer).

Another counseling dilemma is posed by the finding of a variant or mutation of unknown significance (see Table 2), a mutation that may actually represent a benign polymorphism unrelated to an increased breast cancer risk or may indicate an increased breast cancer risk. The individual must be counseled in such a situation, because additional information about that specific mutation will be needed before its significance can be understood. These patients should be considered for referral to research studies that aim to define the functional impact of the gene variant.

Finally, it is important to mention that certain large genomic rearrangements are not detectable by a primary sequencing assay, thereby necessitating supplementary testing, in some cases. For example, there are tests that detect rare, large cancer-associated rearrangements of DNA in the BRCA1 and BRCA2 genes that are otherwise not detected by direct sequencing of the BRCA1/2 genes. Therefore, the NCCN Guidelines panel emphasizes the need for comprehensive testing, which encompasses full BRCA1/2 sequencing and detection of large gene rearrangements.

Risk Assessment, Counseling, and Management: Hereditary Breast/Ovarian Cancer Syndrome

Detailed in the NCCN Guidelines is a set of specific risk assessment criteria which form part of the decision-making process in evaluating whether an individual suspected of being carriers of a BRCA1/2 mutation should be considered for genetic testing (see Guidelines section on Hereditary Breast and/or Ovarian Cancer Syndrome Testing Criteria). Following risk assessment and counseling, genetic testing should be considered for individuals for whom hereditary breast/ovarian cancer syndrome testing criteria are met. Testing is generally not recommended in children under the age of 18 years.

Individuals from a family with a known deleterious BRCA1 or BRCA2 mutation should be tested for the specific familial mutation. For
individuals from a family without a known \textit{BRCA1} or \textit{BRCA2} mutation (and who meet testing criteria), genetic testing should be comprehensive, including full sequencing of \textit{BRCA1} and \textit{BRCA2}, and testing for large genomic rearrangements. Individuals from a family with a known deleterious \textit{BRCA1}/\textit{BRCA2} mutation who test positive for the familial mutation, or for whom \textit{BRCA1}/\textit{BRCA2} mutation testing is not performed, should follow the screening recommendations outlined under the Guidelines section on HBOC Syndrome Management (and discussed below). Those who test negative for \textit{BRCA1}/\textit{BRCA2} mutations can be considered for risk assessment/genetic testing for other hereditary breast/ovarian cancer syndromes such as Li-Fraumeni syndrome or Cowden syndrome, if testing criteria for these syndromes are met. For individuals who have not been tested or for those in whom variants of unknown significance are found (uninformative testing results), participation in a research program or individualized recommendations based on personal history and family history should be offered.

Counseling issues specific for both female and male carriers of a \textit{BRCA1}/2 mutation include the increased incidence of pancreatic cancer and melanoma. In addition, the risks to family members of individuals with a known \textit{BRCA1}/2 gene mutation (see Discussion sections on Risk Assessment and Genetic Testing) should also be discussed as well as the importance of genetic counseling for these individuals. Counseling issues pertaining specifically to male breast cancer have also been described, and include an increased risk of prostate cancer in male carriers of a \textit{BRCA1}/2 mutation.\cite{214-216}

Recommendations for the medical management of hereditary breast/ovarian cancer syndrome are based on an appreciation of the early onset of disease, the increased risk of ovarian cancer, and the risk for male breast cancer in \textit{BRCA1}/2 carriers. An individual with a known deleterious \textit{BRCA1}/2 mutation in a close family member who does not undergo gene testing should be followed according to the same screening/management guidelines as a carrier of a \textit{BRCA1}/2 mutation. An individual from a family with a known deleterious \textit{BRCA1}/2 mutation
who tests negative for the familial mutation should be followed according to the recommendations in the NCCN Guidelines for Breast Cancer Screening and Diagnosis. In situations where an individual (or family member) from a family with no known familial BRCA1/2 mutations undergoes genetic testing, and no mutation is found, testing for other hereditary breast syndromes should be considered if testing criteria are met (see sections on Li-Fraumeni Syndrome Testing Criteria and Cowden Syndrome Testing Criteria).

**Screening Recommendations**

The emphasis on initiating screening considerably earlier than standard recommendations is a reflection of the early age of onset seen in hereditary breast/ovarian cancer.\(^{217}\) For a woman who is a carrier of a BRCA1/2 mutation, training in breast awareness with regular monthly practice should begin at age 18 years, and semiannual clinical breast examinations should begin at age 25 years. The woman should have annual mammograms and breast MRI screening (to be performed on day 7-15 of menstrual cycle for premenopausal women) beginning at age 25 years or on an individualized timetable based on the earliest age of cancer onset in family members.\(^{192,217-220}\)

Mammography has served as the standard screening modality for detection of breast cancer during the last few decades. False-negative mammography results have been correlated with factors such as presence of BRCA1/2 mutation and high breast tissue density,\(^{221-224}\) both of which may occur more frequently among younger women. Rapidly growing or aggressive breast tumors—also more common among younger women—have also been associated with decreased sensitivity of mammographic screening methods.\(^{221,225}\)

Prospective studies on comparative surveillance modalities in women at high risk for familial breast cancer (i.e., confirmed BRCA1/BRCA2 mutation or suspected mutation based on family history) have consistently reported higher sensitivity of MRI screening (77–94%) compared with mammography (33–59%) in detecting breast cancers; false-positive rates were higher with MRI in some reports, resulting in a slightly lower or similar specificity with MRI screening (81–98%) compared with mammography (92–100%).\(^{217-219,226-228}\) The sensitivity with ultrasound screening (33–65%) appeared similar to that of mammography in this high-risk population.\(^{217,226-228}\) In a recent prospective screening trial (conducted from 1997–2009) that evaluated the performance of annual MRI and mammography in women (age 25–65 years; N=496) with confirmed BRCA1/BRCA2 mutation, sensitivity with MRI was significantly higher compared with mammography during the entire study period (86% vs. 19%; \(P<0.0001\)).\(^{229}\) Sensitivity with MRI was higher during the early years (1997–2002; 74% vs. 35%) as well as the later years of the study (2003–2009; 94% vs. 9%). Factors such as age, mutation type or invasiveness of the tumor did not significantly influence the relative sensitivity of the 2 screening modalities. Importantly, the large majority (97%) of cancers detected by MRI screening were early stage tumors.\(^{229}\) Among previously unaffected women diagnosed with invasive breast cancer during the study (n=28), 1 patient had died due to the cancer and 3 additional patients died due to other causes; the annual breast cancer-specific mortality rate was 0.5%. At a median follow up of 8 years from diagnosis, none of the surviving patients (n=24) has developed distant recurrence.\(^{229}\)

All of the studies discussed above evaluated a screening strategy that was conducted on an annual basis, and many of the studies included individuals without confirmed BRCA1/BRCA2 mutation status. A recent retrospective study evaluated a different screening interval, using alternating mammography and MRI screening every 6 months in women with confirmed BRCA1/BRCA2 mutation (N=73).\(^{230}\) After a median follow up of 2 years, 13 breast cancers were detected among 11 women; 12 of the tumors were
detected by MRI screening but not by mammography obtained 6 months earlier. The sensitivity and specificity with MRI screening was 92% and 87%, respectively.230

The optimal surveillance approach in women at high risk for familial breast cancer remains uncertain, especially for women between the ages of 25 and 30 years. Although earlier studies have reported an unlikely association between radiation exposure from mammography and increased risk of breast cancer in carriers of \textit{BRCA1}/\textit{BRCA2} mutation,231,232, a recent report from a large cohort study suggested an increased risk in women exposed to radiation at a young age.233 A retrospective cohort study (from the GENE-RAD-RISK study) showed that exposure to diagnostic radiation (including mammography) prior to age 30 years was associated with increased risk of breast cancer in women with \textit{BRCA1}/\textit{BRCA2} mutation (N=1993).233 Thus, one of the potential benefits of incorporating MRI modalities into surveillance strategies may include minimizing the radiation risks associated with mammography, in addition to the higher sensitivity of MRI screening in detecting tumors. The use of MRI, however, may potentially be associated with higher false-positive results and higher costs relative to mammography. The appropriate imaging modalities and surveillance intervals are still under investigation. In a recent report based on a computer simulation model that evaluated different annual screening strategies in \textit{BRCA1}/\textit{BRCA2} mutation carriers, a screening approach that included annual MRI starting at age 25 years combined with alternating digital mammography/MRI starting at age 30 years was shown to be the most effective strategy when radiation risks, life expectancy and false-positive rates were considered.234 Future prospective trials are needed to evaluate the different surveillance strategies in individuals at high risk for familial breast cancer. Annual MRI as an adjunct to screening mammogram and clinical breast examination for women aged 25 years or older with a genetic predisposition for breast cancer is supported by guidelines from the ACS.192

Post-test counseling in women with confirmed \textit{BRCA1}/\textit{BRCA2} mutation (or highly suspected of having the mutation based on presence of known deleterious mutation in the family) includes discussion of risk-reducing mastectomy and/or salpingo-oophorectomy. Counseling for these risk-reducing surgeries may include discussion of extent of cancer risk reduction/protection, risks associated with surgeries, reconstructive options, management of menopausal symptoms, and discussion of reproductive desires. It is important to address the psychosocial and quality-of-life aspects of undergoing risk-reducing surgical procedures.

For women who have not elected ovarian cancer risk-reducing surgery, concurrent transvaginal ultrasound and CA-125 determination should be considered every 6 months, starting at age 30 years or 5 to 10 years earlier than the earliest age of first diagnosis of ovarian cancer in the family, for the early detection of ovarian cancer (see Guidelines section on HBOC Syndrome Management). Although there are retrospective data indicating that annual ovarian screening using transvaginal ultrasound and measurement of serum CA-125 levels is neither an effective strategy for the early detection of ovarian tumors nor a reasonable substitute for a bilateral risk-reduction salpingo-oophorectomy,235,236 the data are limited regarding the effectiveness of these screening interventions when used every 6 months. Investigational imaging and screening studies may be considered for this population.

Men testing positive for a \textit{BRCA1}/2 mutation should have a semiannual clinical breast examination, and undergo training in breast self-examination with regular monthly practice starting at age 35 years.
Baseline mammography should be considered at age 40 years, followed by annual screening with mammography for those men with gynecomastia or parenchymal/glandular breast density on baseline study. In addition, screening for prostate cancer starting at age 40 years should be considered. Involvement in population screening guidelines for prostate cancer is recommended. For both men and women testing positive for a BRCA1/2 mutation, a full body skin exam for melanoma screening and investigational protocols for pancreatic cancer screening should be considered. Although no specific screening guidelines exist for these tumor types, individualized screening approaches may be provided according to personal or family history of cancer.

Risk Reduction Surgery

**Bilateral Total Mastectomy**

Retrospective analyses with median follow-up periods of 13-14 years have indicated that bilateral risk reduction mastectomy (RRM) decreased the risk of developing breast cancer by at least 90% in moderate- and high-risk women and in known BRCA1/2 mutation carriers. Results from smaller prospective studies with shorter follow-up periods have provided support for concluding that RRM provides a high degree of protection against breast cancer in women with a BRCA1/2 mutation.

The NCCN Guidelines panel supports discussion of the option of RRM for women on a case-by-case basis. Counseling regarding the degree of protection offered by such surgery and the degree of cancer risk should be provided.

It is important that the potential psychosocial effects of RRM are addressed, although these effects have not been well studied. Multidisciplinary consultations are recommended prior to surgery and should include the discussions of the risks and benefits of surgery, and surgical breast reconstruction options. Immediate breast reconstruction is an option for many women following RRM, and early consultation with a reconstructive surgeon is recommended for those considering either immediate or delayed breast reconstruction.

**Bilateral Salpingo-oophorectomy**

Women with a BRCA1/2 mutation are at increased risk for both breast and ovarian cancers (including fallopian tube cancer and primary peritoneal cancer). Although the risk of ovarian cancer is generally considered to be lower than the risk of breast cancer in a BRCA1/2 mutation carrier, the absence of reliable methods of early detection and the poor prognosis associated with advanced ovarian cancer have lent support for the performance of bilateral risk reduction salpingo-oophorectomy (RRSO) after completion of childbearing in these women. In the studies of Rebbeck et al, the mean age at diagnosis of ovarian cancer was 50.8 years for BRCA1/2 carriers.

The effectiveness of RRSO in reducing the risk of ovarian cancer in carriers of a BRCA1/2 mutation has been demonstrated in a number of studies. For example, results of a meta-analysis involving 10 studies of BRCA1/2 mutation carriers showed an approximately 80% reduction in the risk of ovarian or fallopian cancer following RRSO. In a large prospective study of women who carried deleterious BRCA1 or BRCA2 mutations (N=1079), RRSO significantly reduced the risk of BRCA1-associated gynecological tumors (including ovarian, fallopian tube or primary peritoneal cancers) by 85% compared with observation, during a 3-year follow-up period (hazard ratio=0.15; 95% CI, 0.04-0.56; P=0.005). However, a 1%-4.3% residual risk of a primary peritoneal carcinoma has been reported in some studies. RRSO is also reported to reduce the risk of breast cancer in carriers of a BRCA1/2 mutation by approximately 50%.
control international study by Eisen et al., a 56% (odds ratio=0.44; 95% CI, 0.29-0.66; \(P\leq0.001\)) and a 43% (odds ratio=0.57; 95% CI, 0.28-1.15; \(P=0.11\)) breast cancer risk reduction (adjusted for oral contraceptive use and parity) was reported following RRSO in carriers of a \textit{BRCA1} and a \textit{BRCA2} mutation, respectively.\(^{252}\) Hazard ratios of 0.47 (95% CI, 0.29-0.77)\(^{246}\) and 0.30 (95% CI, 0.11-0.84; \(P=0.022\))\(^{250}\) were reported in two other studies comparing breast cancer risk in women with a \textit{BRCA1/2} mutation who had undergone RRSO with carriers of these mutations who opted for surveillance only. These studies are further supported by a recent meta-analysis which found similar reductions in breast cancer risk of approximately 50% for \textit{BRCA1} and \textit{BRCA2} mutation carriers following RRSO,\(^{247}\) although results of a prospective cohort study suggest that RRSO may be associated with a greater reduction in breast cancer risk for \textit{BRCA2} mutation carriers compared with \textit{BRCA1} mutation carriers.\(^{248}\)

Reductions in breast cancer risk for carriers of a \textit{BRCA1/2} mutation undergoing RRSO may be associated with decreased hormonal exposure following surgical removal of the ovaries. Greater reductions in breast cancer risk were observed in women with a \textit{BRCA1} mutation who had a RRSO at age 40 years or younger (odds ratio=0.36, 95% CI, 0.20-0.64) relative to \textit{BRCA1} carriers aged 41-50 years who had this procedure (odds ratio=0.50, 95% CI, 0.27-0.92).\(^{253}\) A nonsignificant reduction in breast cancer risk was found for women aged 51 or older although only a small number of women were included in this group.\(^{252}\) However, results from Rebbeck et al also suggest that RRSO after age 50 is not associated with a substantial decrease in breast cancer risk.\(^{251}\) Due to the limited data, an optimal age for RRSO is difficult to specify.

The NCCN Guidelines panel recommends RRSO for women with a known \textit{BRCA1/2} mutation, ideally between ages 35 and 40 years and upon completion of child bearing or at an individualized age based on earliest age of ovarian cancer diagnosed in the family. Peritoneal washings should be performed at surgery, and pathologic assessment should include fine sectioning of the ovaries and fallopian tubes.\(^{100,101}\)

Other topics which should be addressed with respect to RRSO include the increased risk of osteoporosis and cardiovascular disease associated with premature menopause, as well as the potential effects of possible cognitive changes, accelerated bone loss, and vasomotor symptoms on quality of life.

It has been reported that short-term hormone replacement therapy (HRT) in women undergoing RRSO does not negate the reduction in breast cancer risk associated with the surgery.\(^{254}\) In addition, results of a recent case-control study of \textit{BRCA1} mutation carriers showed no association between use of HRT and increased breast cancer risk in postmenopausal \textit{BRCA1} mutation carriers.\(^{255}\) However, caution should be used when considering use of HRT in mutation carriers following RRSO, given the limitations inherent in nonrandomized studies.\(^{256,257}\)

**Chemoprevention**

The use of selective estrogen receptor modulators (i.e., tamoxifen, raloxifene) has been proven to reduce the risk of invasive breast cancer in postmenopausal women considered at high risk for developing breast cancer.\(^{258-263}\) However, only limited data are available on the specific use of these agents in patients with \textit{BRCA} mutations. As previously discussed, patients with \textit{BRCA} mutations who are diagnosed with breast cancer have elevated risks for developing contralateral breast tumors. In one of the largest prospective series of \textit{BRCA} mutation carriers evaluated, the mean cumulative lifetime risks for contralateral breast cancer were estimated to be 83% for \textit{BRCA1} carriers and 62%
Patients with BRCA mutations who have intact contralateral breast tissue (and who do not undergo oophorectomy or receive chemoprevention) have an estimated 40% risk of contralateral breast cancer at 10 years. Case control studies from the Hereditary Breast Cancer Clinical Study Group reported that the use of tamoxifen protected against contralateral breast cancer with an odds ratio of 0.38 (95% CI, 0.19–0.74) to 0.50 (95% CI, 0.30–0.85) among BRCA1 mutation carriers and 0.42 (95% CI, 0.17–1.02) to 0.63 (95% CI, 0.20–1.50) among BRCA2 carriers. This translates to an approximately 45% to 60% reduction in risk of contralateral tumors among BRCA mutation carriers with breast cancer. The data were not consistent with regards to the protective effects of tamoxifen in the subset of BRCA mutation carriers who also underwent oophorectomy. In addition, no data were available on the estrogen receptor status of the tumors. An evaluation of the subset of healthy individuals with a BRCA1/2 mutation in the Breast Cancer Prevent Trial revealed that breast cancer risk was reduced by 62% in those with a BRCA2 mutation receiving tamoxifen relative to placebo (risk ratio=0.38; 95% CI, 0.06–1.56). However, tamoxifen use was not associated with a reduction in breast cancer risk in those with a BRCA1 mutation. These findings may be related to the greater likelihood for development of estrogen receptor-negative tumors in BRCA1 mutation carriers relative to BRCA2 mutation carriers. However, this analysis was limited by the very small number of individuals with a BRCA1/2 mutation (n=19; 7% of study population). Recently, common single-nucleotide polymorphisms were identified in genes (ZNF423 and CTSO genes) that are involved in estrogen-dependent regulation of BRCA1 expression. These gene variants were associated with alterations in breast cancer risk during treatment with selective estrogen receptor modulators, and may eventually pave the way for predicting the likelihood of benefit with these chemopreventive approaches in individual patients.

With respect to the evidence regarding the effect of oral contraceptives on cancer risks in women with known BRCA1/2 gene mutations, case-control studies have demonstrated that oral contraceptives reduced the risk of ovarian cancer by 45%-50% in BRCA1 mutation carriers and by 60% in BRCA2 mutation carriers; moreover, risks appeared to decrease with longer duration of oral contraceptive use. In a recent meta-analysis conducted in a large number of BRCA1/2 mutation carriers with (n=1503) and without (n=6315) ovarian cancer, use of oral contraceptives significantly reduced the risk of ovarian cancer by approximately 50% for both the BRCA1 mutation carriers (summary relative risk [SRR]=0.51; 95% CI, 0.30–0.85) and BRCA2 mutation carriers (SRR=0.52; 95% CI, 0.31–0.87).

Studies on the effect of oral contraceptive use on breast cancer risk among BRCA1/2 mutation carriers have reported conflicting data. In one case-control study, use of oral contraceptives was associated with a modest but statistically significant increase in breast cancer risk among BRCA1 mutation carriers (odds ratio=1.20; 95% CI, 1.02-1.40), but not among BRCA2 mutation carriers. Among BRCA1 mutation carriers, breast cancer risks with oral contraceptives were significantly associated with ≥5 years of oral contraceptive use (odds ratio=1.33; 95% CI,1.11–1.60), breast cancer diagnosed before age 40 (odds ratio=1.38; 95% CI,1.11–1.72), and use of oral contraceptives before 1975 (odds ratio=1.42; 95% CI, 1.17–1.75). In another case-control study, oral contraceptive used for at least 1 year was not significantly associated with breast cancer risks in either BRCA1 or BRCA2 mutation carriers. However, among BRCA2 mutation carriers, use of oral contraceptives for at least 5 years was associated with a significantly increased risk for breast cancer (odds ratio=2.06; 95% CI, 1.08-3.94); results were similar when only the cases with oral contraceptives use on or after 1975 were considered. Other case-control studies have
reported no significant associations with oral contraceptives use (especially with the use of low-dose formulations after 1975) and risks for breast cancer in \textit{BRCA1/2} mutation carriers.\textsuperscript{274,275} In fact, in one study, the use of low-dose oral contraceptives for at least 1 year was associated with significantly decreased risks for breast cancer among \textit{BRCA1} mutation carriers (odds ratio=0.22; 95% CI, 0.10–0.49; \(P<0.001\)), though not for \textit{BRCA2} mutation carriers.\textsuperscript{275} Differences in the study design employed by these case-control studies make it difficult to compare outcomes between studies, and likely accounts for the conflicting results. The study design might have differed with regards to factors such as the criteria for defining the “control” population for the study (e.g., non-\textit{BRCA1/2} mutation carriers vs. mutation carriers without a cancer diagnosis), consideration of family history of breast or ovarian cancer, baseline demographics of the population studied (e.g., nationality, ethnicity, geographic region, age groups), age of onset of breast cancer, and formulations or duration of oral contraceptives used. In a meta-analysis conducted in a large number of \textit{BRCA1/2} mutation carriers with (n=2855) and without (n=2954) breast cancer, use of oral contraceptives was not found to be significantly associated with breast cancer risks in either the \textit{BRCA1} mutation carriers (SRR=1.09; 95% CI, 0.77–1.54) or the \textit{BRCA2} mutation carriers (SRR=1.15; 95% CI, 0.61–2.18).\textsuperscript{271}

Reproductive Options

The outcomes of genetic testing can have profound impact on family planning decisions for individuals of reproductive age who are found to be carriers of \textit{BRCA1/2} mutations. For example, in cases where both partners carry a \textit{BRCA2} mutation, there may be a high risk for the offspring to develop a rare Fanconi anemia/brain tumor phenotype (recessive disorder).\textsuperscript{215} Counseling for reproductive options such as prenatal diagnosis, preimplantation genetic diagnosis (PGD) and assisted reproduction may therefore be warranted for couples expressing concern over the \textit{BRCA} mutation carrier status of their future offspring. Such counseling should include a comprehensive discussion of the potential risks, benefits, and limitations of reproductive options.

Prenatal diagnosis involves postimplantation genetic analysis of an early embryo, utilizing chorionic villi or amniotic fluid cell samples; genetic testing is typically conducted between week 12 and week 16 of gestation, and testing results may potentially lead to a couple’s decision to terminate pregnancy.\textsuperscript{216,276} During the past 2 decades, PGD has emerged as an alternative method of genetic testing in early embryos. PGD involves the testing of 1 or 2 cells from embryos in very early stages of development (i.e., 6 to 8 cells) after in vitro fertilization (IVF). This procedure allows for the selection of unaffected embryos to be transferred to the uterus,\textsuperscript{216,276} and may, therefore, offer the advantage of avoiding potential termination of pregnancy. However, procedures such as PGD are not without limitations as it may still require a confirmatory prenatal diagnosis depending upon a couple’s medical needs or requests. Moreover, the PGD process requires the use of IVF regardless of the fertility status of the couple (i.e., also applies to couples without infertility issues), and IVF may not always lead to a successful pregnancy. Lastly, the technology or expertise may not be readily available in a couple’s geographical location. Various factors, both medical and personal, must be weighed in the decision to utilize prenatal diagnosis or PGD. Medical considerations may include factors such as the age of onset of the hereditary cancer, penetrance, severity or associated morbidity and mortality of the cancer, and availability of effective cancer risk reduction methods or effective treatments.\textsuperscript{216,276} Although the use of prenatal diagnosis or PGD is relatively well established for severe hereditary disorders with very high penetrance,
their use in conditions associated with lower penetrance (e.g., hereditary breast or ovarian cancer syndrome) remains somewhat controversial from both an ethical and regulatory standpoint. Personal considerations for the decision to utilize prenatal diagnosis or PGD may include individual ethical beliefs, value systems, cultural and religious beliefs, as well as social and economic factors. Based on results from surveys administered to women at high risk for hereditary breast or ovarian cancer, 50% to 75% of respondents felt that PGD was an acceptable option for high-risk individuals, yet only about 14% to 33% would consider undergoing PGD themselves. A survey in high-risk men (N=228; carriers of BRCA mutation; or having a partner or first-degree relative with BRCA mutation) showed that 80% of these men were unaware of PGD; after being informed of the definition of PGD, 34% indicated that they would consider the option of using PGD. Importantly, these surveys suggested that the majority of high-risk women and men have little or no knowledge of PGD, highlighting the need for better awareness and education regarding potential reproductive options. Successful births have been reported with the use of PGD and IVF in BRCA1/2 mutation carriers, but data in the published literature are still very limited. In addition, data pertaining to long-term safety or outcomes of PGD and assisted reproduction in BRCA mutation carriers are not yet available.

Risk Assessment, Counseling, and Management: Li-Fraumeni Syndrome

The approach to families with other hereditary breast cancer syndromes, such as LFS, reflects that of hereditary breast/ovarian cancer in many ways. However, there are some syndrome-specific differences with regard to assessment and management. In the case of LFS, there are multiple associated cancers, both pediatric and adult, that should be reflected in the expanded pedigree (see Guidelines section on Li-Fraumeni Syndrome Testing Criteria). Cancers associated with LFS include but are not limited to premenopausal breast cancer, bone and soft tissue sarcomas, acute leukemia, brain tumor, adrenocortical carcinoma, unusually early onset of other adenocarcinomas, or other childhood cancers. Verification of these sometimes very rare cancers is particularly important.

Following risk assessment and counseling, genetic testing should be considered in individuals for whom testing criteria are met. This recommendation is category 2A for adults and 2B for children. The NCCN Guidelines panel also suggests consideration of TP53 mutation testing in those with early onset breast cancer (≤35 years of age) for whom BRCA1/2 testing result is negative, especially if there is a family history of LFS related cancers. The NCCN Guidelines panel recommends comprehensive testing, which should include full sequencing and analysis of gene deletion/duplication. In the absence of additional family history, early breast cancer alone is associated with a low likelihood of mutation identification. Individuals who have tested positive for a TP53 mutation may have greater distress than anticipated, so provisions for supportive interventions should be provided. An individual with a known deleterious TP53 mutation in a close family member who does not undergo testing should be followed according to the same recommendations as a carrier of a TP53 mutation (see Guidelines section on Li-Fraumeni Syndrome Management). In situations where an individual (or family member) from a family with no known familial TP53 mutation undergoes genetic testing, and no mutation is found, testing for other hereditary breast syndromes should be considered if testing criteria are met (see sections on Hereditary Breast and/or Ovarian Cancer Syndrome Testing Criteria and Cowden Syndrome Testing Criteria). As previously discussed in the BRCA1/BRCA2 testing section above, testing of unaffected individuals...
should only be considered when an appropriate affected family member is not available for testing; importantly, the significant limitations of interpreting testing results for an unaffected individual should be discussed prior to testing.

Management of LFS should address the limitations of screening for the many cancers associated with this syndrome. For those at risk for breast cancer, training and education in breast self-examination should start at age 18 years, with the patient performing regular self-examination on a monthly basis. For members of families with LFS, it is recommended that breast cancer surveillance by clinical breast examination, every 6 to 12 months, begin between the ages of 20 and 25 years (or 5 to 10 years before the earliest known breast cancer in the family, whichever is earlier) because of the very early age of breast cancer onset seen in these families. Annual mammograms and breast MRI screening should begin at ages 20 to 25 years or be individualized, based on earliest age of onset in the family. For relatively young patients (age 20 to 30 years), only an annual breast MRI screening may be considered (given the theoretical risks of radiation exposure with mammography in these patients) based on the physician’s discretion. Although there are no data regarding risk reduction surgery in women with LFS, options for risk reducing mastectomy should be discussed on a case-by-case basis (see Discussion section on Bilateral Total Mastectomy for HBOC). Counseling for risk-reducing surgeries may include discussion of extent of cancer risk reduction/protection, risks associated with surgeries, and reconstructive options. It is also important to address the psychosocial and quality-of-life aspects of undergoing risk-reducing surgical procedures.

Many of the other cancers associated with germline mutations in TP53 do not lend themselves to early detection. Thus, additional recommendations are general and include annual comprehensive physical examinations starting at age 20 to 25 years among family members who have survived one cancer when there is a high index of suspicion for second malignancies (Guidelines section on Li-Fraumeni Syndrome Management). Clinicians should address screening limitations for other cancers associated with LFS. The option to participate in clinical trials evaluating novel screening approaches using technologies such as whole-body MRI, abdominal ultrasound and brain MRI should also be discussed if such trials are available. Colonoscopy should be considered every 2 to 5 years, starting at no later than 25 years. Education regarding signs and symptoms of cancer is important. Patients should be advised about the risk to relatives, and genetic counseling for relatives is recommended. Annual physical examination is recommended for cancer survivors with a high index of suspicion for rare cancers and second malignancies. Pediatricians should be made aware of the risk of childhood cancers in affected families. For couples expressing the desire that their offspring not carry a familial TP53 mutation, options for prenatal diagnosis should be discussed (for discussion on known risks, limitations, and benefits of such technologies, see section above on Reproductive Options under Risk Assessment, Counseling, and Management: Hereditary Breast and/or Ovarian Cancer Syndrome).

A recent prospective observational study incorporated a clinical surveillance protocol for asymptomatic TP53 mutation carriers from eight families affected by LFS. In this study, 18 of the 33 asymptomatic mutation carriers agreed to undergo surveillance while the remainder of the carriers did not. The surveillance protocol included both biochemical methods and imaging techniques, such as annual brain MRI for brain tumor surveillance (both children and adults); annual rapid total-body MRI (both children and adults) and ultrasound of abdomen and pelvis every 6 months (for adults only) for soft
tissue/bone sarcoma surveillance; colonoscopy every 2 years beginning at age 40 years (or 10 years before earliest known colon cancer in the family); ultrasound of abdomen and pelvis every 3-4 months, complete urinalysis every 3-4 months, blood test every 4 months for adrenocortical carcinoma surveillance (children only); and complete blood counts and blood tests every 4 months for leukemia/lymphoma surveillance (both children and adults); for surveillance of breast cancers, the protocol was similar to the NCCN Guidelines for LFS management.\(^2\)\(^8\) Using this surveillance protocol, asymptomatic tumors were detected in 7 of the patients; after a median follow-up time of 24 months, all 7 of these carriers were alive. Ten individuals in the non-surveillance group developed high-grade, advanced stage tumors; only 2 of these individuals were alive at the end of follow up. The 3-year overall survival rate was significantly higher for the surveillance group compared with the non-surveillance group (100% vs. 21%; \(P=0.016\)).\(^2\)\(^8\) Although this was a small study in a limited number of patients, the clinical surveillance protocol employed was feasible and detected asymptomatic tumors in about 40% of individuals with \(TP53\) mutations. The protocol may represent an emerging option for surveillance/management of at-risk individuals from families with LFS; further evaluation of this protocol is warranted.

Only very limited data exists on the use of prenatal diagnostics/genetic testing for \(TP53\) mutations in families with LFS.\(^2\)\(^8\)\(^5\)\(^6\) Counseling for reproductive options such as prenatal diagnosis, preimplantation genetic diagnosis (PGD) and assisted reproduction may be warranted for couples expressing concern over the mutation carrier status of their future offspring. Such counseling should include a comprehensive discussion of the potential risks, benefits, and limitations of reproductive options. For general discussions on the topic of reproductive options and counseling considerations, see the Discussion section above on Reproductive Options under Risk Assessment, Counseling, and Management: Hereditary Breast/Ovarian Cancer Syndrome.

### Risk Assessment, Counseling, and Management: Cowden Syndrome

The assessment of individuals suspected of having Cowden syndrome incorporates both a history of the benign and malignant conditions associated with the syndrome and a targeted physical examination, including the skin and oral mucosa, breast, and thyroid gland (see Guidelines section on Cowden Syndrome Testing Criteria). The NCCN Guidelines panel has recently revised both the list of criteria associated with this genetic syndrome as well as the combinations of criteria that establish which individuals are candidates for \(PTEN\) gene mutation testing (see Guidelines section on Cowden Syndrome Testing Criteria and Discussion section on Cowden Syndrome). These criteria are recommended to assess the need for further risk assessment and genetic testing, but are not intended to serve as clinical diagnostic criteria. Following risk assessment and counseling, genetic testing should be considered in individuals for whom testing criteria are met. The NCCN Guidelines panel recommends comprehensive testing, which should include full sequencing, gene deletion/duplication analysis, and promoter analysis. Unlike the “pathognomonic” criteria, none of the individual major or minor criteria are considered by the NCCN Guidelines panel to be sufficient to warrant genetic testing in the absence of other clinical evidence of Cowden syndrome. However, the panel recommends genetic testing in an individual exhibiting 2 or more major criteria when one is macrocephaly, 3 or more major criteria when one is not macrocephaly, one major criterion along with 3 or more minor criteria, or in someone meeting specifications for 4 minor criteria. Furthermore, any of the major criteria can be classified as a minor criterion for the purpose of meeting the threshold required for genetic
testing if 2 or more major criteria are present in a single individual but the individual does not have macrocephaly. The testing threshold is lower for an individual considered to be “at risk” (e.g., a first-degree relative of an individual and/or proband with a clinical diagnosis of Cowden syndrome or BRRS for whom genetic testing has not been performed). In this case, any one major criterion or 2 minor criteria are considered to be sufficient for genetic testing to be recommended. Recommendations for individuals not meeting these testing criteria should be individualized according to personal and family history.

An individual with a known deleterious PTEN mutation in a close family member who does not undergo gene testing should be followed according to the same guideline as a carrier of a PTEN mutation (see Guidelines section on Cowden Syndrome Management). In situations where an individual (or family member) from a family with no known familial PTEN mutation undergoes genetic testing, and no mutation is found, testing for other hereditary breast syndromes should be considered if testing criteria are met (see sections on Hereditary Breast and/or Ovarian Cancer Syndrome Testing Criteria and Li-Fraumeni Syndrome Testing Criteria).

Current medical management recommendations for individuals with Cowden syndrome focus on primary and secondary prevention options for breast cancer and on annual physical examinations, starting at age 18 years (or 5 years before the youngest age of diagnosis of a component cancer in the family) to detect skin changes and to monitor the thyroid gland for abnormalities. A baseline thyroid ultrasound should be performed at age 18 years and considered annually thereafter for both men and women with Cowden syndrome. Annual dermatological examination should also be considered. In addition, colonoscopy should be considered starting at age 35 years, performed every 5 to 10 years or more frequently in cases where the patient is symptomatic or polyps are found. Education regarding the signs and symptoms of cancer is important; patients should also be advised about the risk to relatives, and genetic counseling is recommended for at-risk relatives.

Women should begin regular monthly breast self-examinations at age 18 years and have a semiannual clinical breast examination, beginning at age 25 years or 5-10 years earlier than the earliest known breast cancer in the family. Women should also have an annual mammogram and breast MRI screening starting at ages 30-35 years, or 5 to 10 years earlier than the earliest known breast cancer in the family. Although there are no data regarding risk reduction surgery in women with Cowden syndrome, the option of risk-reduction mastectomy and hysterectomy should be discussed on a case-by-case basis (see Discussion section on Bilateral Total Mastectomy). Oophorectomy is not indicated for Cowden syndrome alone, but may be indicated for other reasons. Counseling for risk-reducing surgeries may include discussion of extent of cancer risk reduction/protection, risks associated with surgeries, reconstructive options and reproductive desires. It is also important to address the psychosocial and quality-of-life aspects of undergoing risk-reducing surgical procedures. The panel recommends patient education regarding the symptoms of endometrial cancer including the necessity of a prompt response to such symptoms. Women diagnosed with Cowden syndrome should consider participation in a clinical trial to determine the effectiveness and necessity of endometrial cancer screening. No published data exists on the use of prenatal diagnostics/genetic testing for PTEN mutations in families with Cowden syndrome. However, for couples expressing the desire that their offspring not carry a familial PTEN mutation, options for prenatal diagnosis, preimplantation genetic diagnosis (PGD) and assisted reproduction can be discussed. Such counseling should include a comprehensive discussion of the potential risks, benefits, and
limitations of reproductive options. For general discussions on the topic of reproductive options and counseling considerations, see the Discussion section above on Reproductive Options, under Risk Assessment, Counseling, and Management: Hereditary Breast/Ovarian Cancer Syndrome.
Table 1. Glossary of relevant genetic terms (from the National Cancer Institute [NCI])

<table>
<thead>
<tr>
<th>Term</th>
<th>Definition</th>
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<tbody>
<tr>
<td><strong>Autosomal dominant</strong></td>
<td>Autosomal dominant inheritance refers to genetic conditions that occur when a mutation is present in one copy of a given gene (i.e., the person is heterozygous).</td>
</tr>
<tr>
<td><strong>Autosomal recessive</strong></td>
<td>Autosomal recessive inheritance refers to genetic conditions that occur only when mutations are present in both copies of a given gene (i.e., the person is homozygous for a mutation, or carries two different mutations of the same gene, a state referred to as compound heterozygosity).</td>
</tr>
<tr>
<td><strong>de novo mutation</strong></td>
<td>An alteration in a gene that is present for the first time in one family member as a result of a mutation in a germ cell (egg or sperm) of one of the parents, or a mutation that arises in the fertilized egg itself during early embryogenesis. Also called new mutation.</td>
</tr>
<tr>
<td><strong>Familial</strong></td>
<td>A phenotype or trait that occurs with greater frequency in a given family than in the general population; familial traits may have a genetic and/or nongenetic etiology.</td>
</tr>
<tr>
<td><strong>Family history</strong></td>
<td>The genetic relationships within a family combined with the medical history of individual family members. When represented in diagram form using standardized symbols and terminology, it is usually referred to as a pedigree or family tree.</td>
</tr>
<tr>
<td><strong>Founder effect</strong></td>
<td>A gene mutation observed with high frequency in a population founded by a small ancestral group that was once geographically or culturally isolated, in which one or more of the founders was a carrier of the mutant gene.</td>
</tr>
<tr>
<td><strong>Germline</strong></td>
<td>The cells from which eggs or sperm (i.e., gametes) are derived.</td>
</tr>
<tr>
<td><strong>Kindred</strong></td>
<td>An extended family.</td>
</tr>
<tr>
<td><strong>Pedigree</strong></td>
<td>A graphic illustration of family history.</td>
</tr>
<tr>
<td><strong>Penetrance</strong></td>
<td>A characteristic of a genotype; it refers to the likelihood that a clinical condition will occur when a particular genotype is present.</td>
</tr>
<tr>
<td><strong>Proband</strong></td>
<td>The individual through whom a family with a genetic disorder is ascertained. In males this is called a propositus, and in females it is called a proposita.</td>
</tr>
<tr>
<td><strong>Sporadic cancer</strong></td>
<td>This term has two meanings. It is sometimes used to differentiate cancers occurring in people who do not have a germline mutation that confers increased susceptibility to cancer from cancers occurring in people who are known to carry a mutation. Cancer developing in people who do not carry a high-risk mutation is referred to as sporadic cancer. The distinction is not absolute, because genetic background may influence the likelihood of cancer even in the absence of a specific predisposing mutation. Alternatively, sporadic is also sometimes used to describe cancer occurring in individuals without a family history of cancer.</td>
</tr>
</tbody>
</table>
# Table 2. Genetic test results to determine the presence of a cancer-predisposing gene

<table>
<thead>
<tr>
<th><strong>Result</strong></th>
<th><strong>Description</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>True-positive</strong></td>
<td>The person is a carrier of an alteration in a known cancer-predisposing gene.</td>
</tr>
<tr>
<td><strong>True-negative</strong></td>
<td>The person is not a carrier of a known cancer-predisposing gene that has been positively identified in another family member.</td>
</tr>
<tr>
<td><strong>Indeterminate (Uninformative)</strong></td>
<td>The person is not a carrier of a known cancer-predisposing gene, and the carrier status of other family members is either also negative or unknown.</td>
</tr>
<tr>
<td><strong>Inconclusive (Variants of unknown significance)</strong></td>
<td>The person is a carrier of an alteration in a gene that currently has no known significance.</td>
</tr>
</tbody>
</table>
References


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