

## II UPDATE SWISS GUIDELINES FOR GENETIC COUNSELLING AND TESTING FOR PREDISPOSITION TO CANCER 2024

<b>WOMEN WITH A PERSONAL HISTORY OF BREAST CANCER OR DCIS AND ONE OF THE FOLLOWING</b>	
Age at diagnosis $\leq 40$ y (any case) or $\leq 50$ y at oncogeneticist's discretion	
Triple negative (ER, PR and HER2 negative) BC $\leq 60$ y or older at oncogeneticist's discretion	
Bilateral BC or second separate primary	<ul style="list-style-type: none"> <li>• if the first cancer was diagnosed <math>\leq 50</math> y</li> <li>• with <math>\geq 1</math> close relative with BC (if only one relative affected, then age at diagnosis <math>\leq 50</math> y)</li> </ul>
Age at diagnosis $\leq 50$ y	<ul style="list-style-type: none"> <li>• with 1 close relative with BC <math>\leq 50</math> y</li> <li>• limited family history</li> </ul>
Diagnosed at any age	<ul style="list-style-type: none"> <li>• with <math>\geq 2</math> close relatives with BC or Prostate CA</li> <li>• a close male relative with BC</li> <li>• with <math>\geq 1</math> close relative with epithelial OC, Pancreatic CA or metastatic or high-risk Prostate CA (see section Prostate CA below)</li> <li>• Ashkenazi Jewish ancestry (see section Ashkenazi Jewish ancestry below)</li> </ul>
Treatment indications	<ul style="list-style-type: none"> <li>• PARP inhibitors for BC in the metastatic setting</li> <li>• PARP inhibitors for high-risk HER2-negative BC in the adjuvant setting</li> </ul>
<b>MEN WITH A PERSONAL HISTORY OF BREAST CANCER</b>	
<b>HEREDITARY PREDISPOSITION TO OVARIAN CANCER (including fallopian tube or peritoneal CA)</b>	
<ul style="list-style-type: none"> <li>• Personal history of epithelial OC (including fallopian tube or peritoneal CA) at any age</li> <li>• Unaffected with OC with a first- or second-degree relative with epithelial OC (including fallopian tube or peritoneal CA) at any age</li> </ul>	
<b>ASHKENAZI JEWISH HERITAGE</b>	
Search for the 3 founder BRCA1 and BRCA2 P/LP variants may be considered regardless of personal or family history	
<b>RISK ACCORDING TO CALCULATIONS OF RISK MODELS</b>	
Individuals affected or unaffected with BC or OC not meeting the criteria above with a probability $> 5\%$ of a BRCA1/2 P/LP variant based on prior probability models (eg, Tyrer-Cuzick, CanRisk)	
<b>PANCREATIC CANCER</b>	
Exocrine pancreatic CA (adenocarcinoma) at any age	
Unaffected individuals with	<ul style="list-style-type: none"> <li>• 1 first-degree relative with <math>\geq 1</math> or more first- or second-degree relatives with pancreatic CA</li> <li>• <math>\geq 3</math> individuals with pancreatic CA (same side of the family)</li> </ul>
<b>PROSTATE CANCER</b>	
<ul style="list-style-type: none"> <li>• Metastatic Prostate CA at any age</li> <li>• High-risk localised or locally advanced Prostate CA (PSA <math>&gt;20</math>ng/mL or ISUP Grade Group 4 or 5 or <math>\geq</math> cT2c or cN1) irrespective of the family history</li> </ul>	
<b>FAMILY HISTORY ONLY</b>	
Testing of an unaffected or affected individual not meeting criteria above when an appropriate affected family member is unavailable for testing with $\geq 1$ close relative with BC, OC, Pancreatic CA and/or Prostate CA fulfilling one of the above criteria	

### CARRIER TESTING

Testing of an individual from a family with a known P/LP variant in a gene conferring high or moderate risk for CA

### TUMOR PATHOGENIC VARIANT

Germline confirmation of a P/LP variant of a gene conferring a high or moderate risk for BC, OC, Pancreatic CA, Prostate CA, Renal cell CA, Urothelial CA, Paraganglioma, Pheochromocytoma and Neuroendocrine tumors detected by tumor profiling on any tumor type

### FURTHER RECOMMENDATIONS

- Hereditary Renal cell CA, Urothelial CA, Paraganglioma, Pheochromocytoma and Neuroendocrine tumors: See current version of the NCCN Guidelines
- Patient should preferably be referred to experts/centers with solid experience in these issues

#### Abbreviations:

BC, breast cancer; CA, cancer; DCIS, ductal carcinoma in situ; ER, estrogen receptor; HER2, human epidermal growth factor receptor 2; LP, likely pathogenic; P, pathogenic; PR, progesterone receptor; OC, ovarian cancer; y, years

#### Definitions:

- Ashkenazi Jewish founder P/LP variants: BRCA1: c.68\_69delAG, c.5266dupC; BRCA2: c.5946delT
- Ashkenazi Jewish heritage: At least one parent or grandparent of Ashkenazi Jewish ancestry
- Close relative: First- or second-degree relative on the same side of the family. First-degree relatives: Mother/father, sister/brother, daughter/son. Second-degree relatives: Grandparents, aunt/uncle, niece/nephew, grandchildren
- Limited family history:  $\leq 2$  female close relatives having lived beyond age 45 y in either lineage