

Patient name	Patient DOB (MM/DD/YYYY)	Age	Gender
Healthcare provider		Today's date (MM/DD/YYYY)	

## PERSONAL AND FAMILY HISTORY OF CANCER

Please include: yourself, parents, siblings, children, grandparents, grandchildren, aunts, uncles, nephews, nieces, half siblings, first cousins, great grandparents, and great grandchildren. Please be as thorough and accurate as possible.

☐ Adopted/unknown family history

CANCER	YOU Age of diagnosis	PARENTS/SIBLINGS/ CHILDREN	Age of diagnosis	RELATIVES ON YOUR MOTHER'S SIDE	Age of diagnosis	RELATIVES ON YOUR FATHER'S SIDE	Age of diagnosis
<input type="checkbox"/> Y <input type="checkbox"/> N EXAMPLE: Breast Cancer	44	—	—	Grandmother Aunt	47 51	Cousin	54
<input type="checkbox"/> Y <input type="checkbox"/> N BREAST CANCER							
<input type="checkbox"/> Y <input type="checkbox"/> N OVARIAN CANCER (Peritoneal/fallopian tube)							
<input type="checkbox"/> Y <input type="checkbox"/> N UTERINE/ENDOMETRIAL CANCER							
<input type="checkbox"/> Y <input type="checkbox"/> N PROSTATE CANCER							
<input type="checkbox"/> Y <input type="checkbox"/> N COLON/RECTAL CANCER							
<input type="checkbox"/> Y <input type="checkbox"/> N PANCREATIC CANCER							
<input type="checkbox"/> Y <input type="checkbox"/> N OTHER CANCER(S) (Specify cancer type)							
<input type="checkbox"/> Y <input type="checkbox"/> N Are you of Ashkenazi Jewish descent? (Jewish with ancestors from Central or Eastern Europe)							
<input type="checkbox"/> Y <input type="checkbox"/> N Have you or anyone in your family had genetic testing for a hereditary cancer syndrome? (Please describe and include a copy of result if possible)							

## HEREDITARY CANCER FEATURES

Please complete this section with your healthcare provider

YOUR PERSONAL HISTORY	YOUR FAMILY HISTORY
<b>HEREDITARY BREAST CANCER SYNDROMES*</b> <ul style="list-style-type: none"> <li><input type="checkbox"/> Breast cancer diagnosed at or before age 50</li> <li><input type="checkbox"/> Two primary occurrences of breast cancer</li> <li><input type="checkbox"/> Male breast cancer</li> <li><input type="checkbox"/> Triple negative breast cancer diagnosed at or before age 60</li> <li><input type="checkbox"/> Ovarian cancer</li> <li><input type="checkbox"/> Pancreatic cancer</li> <li><input type="checkbox"/> Metastatic or intraductal/criform prostate cancer</li> <li><input type="checkbox"/> Ashkenazi Jewish ancestry, regardless of personal history of cancer</li> </ul>	<b>HEREDITARY BREAST CANCER SYNDROMES</b> <ul style="list-style-type: none"> <li><input type="checkbox"/> Relative with breast cancer at or before age 50</li> <li><input type="checkbox"/> Male relative with breast cancer</li> <li><input type="checkbox"/> Relative with ovarian cancer</li> <li><input type="checkbox"/> Relative with pancreatic cancer</li> <li><input type="checkbox"/> Relative with metastatic or intraductal/criform prostate cancer</li> <li><input type="checkbox"/> Three or more relatives with breast and/or prostate cancer</li> <li><input type="checkbox"/> A previously identified pathogenic variant ("mutation") in the family</li> <li><input type="checkbox"/> Ashkenazi Jewish ancestry, regardless of family history of cancer</li> </ul>
<b>HEREDITARY COLON CANCER SYNDROMES</b> <ul style="list-style-type: none"> <li><input type="checkbox"/> Colorectal cancer before age 50</li> <li><input type="checkbox"/> Endometrial/uterine cancer before age 50</li> <li><input type="checkbox"/> Tumor with mismatch repair (MMR) deficiency<sup>†</sup></li> <li><input type="checkbox"/> Two or more Lynch syndrome cancers<sup>‡</sup></li> <li><input type="checkbox"/> One Lynch syndrome cancer and one or more relatives with a Lynch syndrome cancer</li> </ul>	<b>HEREDITARY COLON CANCER SYNDROMES</b> <ul style="list-style-type: none"> <li><input type="checkbox"/> At least one first-degree relative with colon or endometrial cancer before age 50</li> <li><input type="checkbox"/> At least one first-degree relative with more than one Lynch syndrome cancer</li> <li><input type="checkbox"/> Two or more relatives with a Lynch syndrome cancer,<sup>‡</sup> at least one before age 50</li> <li><input type="checkbox"/> Three or more relatives with a Lynch syndrome cancer</li> <li><input type="checkbox"/> A previously identified pathogenic variant ("mutation") in the family</li> </ul>

\* Including: Breast (female and male), ovarian, pancreatic, prostate cancer

<sup>†</sup> Via PCR, NGS, or IHC. Screening for MMR deficiency is recommended for all colorectal and endometrial cancer tumors and should be considered for other Lynch syndrome cancers.

<sup>‡</sup> Including: Colon, endometrial/uterine, gastric/stomach, ovarian, ureter/renal pelvis, biliary tract, small bowel, pancreas, and brain cancer, as well as sebaceous adenomas

## CANCER RISK ASSESSMENT REVIEW

To be completed after discussion with healthcare provider

If any of the boxes above are checked, this history has features that may indicate a hereditary cancer syndrome and warrants consideration of genetic testing.

Patient's signature	Date (MM/DD/YYYY)
Healthcare provider's signature	Date (MM/DD/YYYY)

For office use only:

Patient offered hereditary cancer genetic testing?

☐ YES

☐ NO

Follow-up appointment scheduled:

☐ YES

☐ NO

☐ ACCEPTED

☐ DECLINED

Date of next appointment