

# HEREDITARY CANCER TEST REQUISITION

PATIENT INFO	
NAME	PATIENT ID NUMBER
DATE OF BIRTH (MM/DD/YYYY)	SEX <input type="checkbox"/> Female <input type="checkbox"/> Male
ETHNICITY <input type="checkbox"/> Asian <input type="checkbox"/> French Canadian/Cajun <input type="checkbox"/> Jewish, non-Ashkenazi <input type="checkbox"/> Native American <input type="checkbox"/> African American <input type="checkbox"/> Hispanic <input type="checkbox"/> Middle Eastern <input type="checkbox"/> Caucasian <input type="checkbox"/> Jewish, Ashkenazi <input type="checkbox"/> Other/Mixed/Unknown _____	
TELEPHONE NUMBER	E-MAIL
ADDRESS / CITY / STATE / ZIP	

CLINICIAN INFO	
PROGENITY ACCOUNT NUMBER	
ORDERING CLINICIAN NAME / NPI#	
CLINIC NAME	
TELEPHONE NUMBER	FAX NUMBER
ADDRESS / CITY / STATE / ZIP	

**ACKNOWLEDGEMENT:** I authorize the laboratory to provide to my health plan the information on this form and other information provided by my healthcare provider if necessary for reimbursement. I understand that the laboratory may seek prior authorization for testing from my health plan on my behalf. I also authorize all benefits of the plan to be payable directly to the laboratory, and I agree to remit to the laboratory any payment for these services made directly to me. I understand that the laboratory may be an out-of-network provider for my health plan and that I am responsible for all amounts not reimbursed by my health plan. I hereby designate the laboratory as my Authorized Representative, as provided under ERISA, 29 C.F.R. § 2560.5031(b)(4), and/or as my Attorney in Fact, for the purpose of pursuing administrative appeals to which I am entitled and, if the laboratory deems it appropriate, any legal and/or equitable claims that I could bring against my health plan, and/or its fiduciaries, and/or its administrators, with respect to their handling or resolution of my insurance claim.

I authorize the laboratory to retain and use my de-identified specimen and test data (where all information that could link me to the specimen or data has been removed) for research and/or to help develop new products or services, in compliance with applicable laws.

I do not authorize the laboratory to retain and use my de-identified specimen and test data as described above.

If signature is present but box is not checked, consent is implied. All leftover specimens from New York State will be destroyed within 60 days.

PATIENT SIGNATURE	DATE (MM/DD/YY)
<b>REQUIRED</b> X	
PATIENT NAME	

**ACKNOWLEDGEMENT:** I hereby confirm that information has been provided to the patient about the test(s) to be performed and the patient has given consent as required under applicable laws and regulations for the test(s) to be performed. The test(s) to be performed are medically necessary and the results will be used for medical management and treatment decision purposes for this patient. I confirm that the person listed as the Ordering Clinician is authorized by law to order the test(s) requested herein.

ORDERING CLINICIAN SIGNATURE	DATE (MM/DD/YY)
<b>REQUIRED</b> X	
GENETIC COUNSELOR NAME / NPI#	
TELEPHONE NUMBER	E-MAIL
FAX NUMBER	
ADDITIONAL REPORTS TO: CLINICIAN NAME / NPI#	
FAX NUMBER	

BILLING INFO		
<input type="checkbox"/> <b>BILL INSURANCE</b> (Attach legible front and back copy of insurance cards)		
NAME OF INSURED	INSURANCE COMPANY	
MEMBER ID	GROUP #	PRE-AUTHORIZATION #, IF OBTAINED
<input type="checkbox"/> <b>BILL PATIENT</b> Please call us to review payment options.		
<input type="checkbox"/> <b>BILL ORDERING CLINICIAN</b>		

**Collection Requirement:** 1 x 4 lavender-top EDTA tube **QR** 1 x 0 ORAcollect® buccal swab **QR** 1 x M 15 mL mouthwash

**SPECIMEN INFO:** Date Collected (MM/DD/YY) Time Collected  AM  PM Collected by: \_\_\_\_\_

TESTS REQUESTED (SEE DESCRIPTIONS ON REVERSE)
<input type="checkbox"/> <b>Riscover™ Comprehensive</b> Analysis of 31 genes related to breast, ovarian, endometrial, colorectal, and other cancers.
<input type="checkbox"/> <b>Riscover™ BRCA</b> Analysis of 2 genes related to hereditary breast, ovarian, and other cancers. <input type="checkbox"/> Limit to <b>Riscover™ BRCA Ashkenazi Jewish Three-Site Analysis</b>
<input type="checkbox"/> <b>Riscover™ Lynch Syndrome</b> Analysis of 5 genes related to hereditary colorectal, endometrial, and other cancers.
<input type="checkbox"/> <b>Known Family Variant</b> Gene _____ Variant _____ <i>A copy of the affected relative's result report is required.</i> If the relative was tested at Progenity, please provide: Full name: _____ Date of birth: _____ Relationship to patient: _____ <input type="checkbox"/> Check if this testing is for the Variant Classification Program (only the familial variant of uncertain significance will be tested)
<input type="checkbox"/> <b>Other:</b> _____

PATIENT CLINICAL HISTORY
<b>REQUIRED</b> Has patient ever been diagnosed with cancer? <input type="checkbox"/> <b>Yes</b> – complete the section below and attach relevant records. <input type="checkbox"/> <b>No</b>
<input type="checkbox"/> <b>BREAST CANCER</b> <input type="checkbox"/> left <input type="checkbox"/> right <input type="checkbox"/> bilateral Age at dx: _____ <input type="checkbox"/> pre-menopausal <input type="checkbox"/> post-menopausal Pathology: <input type="checkbox"/> ductal invasive [Z85.3] <input type="checkbox"/> DCIS [Z86.000] <input type="checkbox"/> lobular invasive [Z85.3] <input type="checkbox"/> other: _____ Triple negative? (ER-, PR-, Her2/neu-) <input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> <b>OVARIAN CANCER</b> [Z85.43] (female patients) Age at dx: _____ Pathology: _____
<input type="checkbox"/> <b>UTERINE / ENDOMETRIAL CANCER</b> [Z85.42] (female patients) Age at dx: _____ Pathology: _____ Was tumor screening performed? <input type="checkbox"/> Yes <input type="checkbox"/> No <i>If "Yes," attach results.</i>
<input type="checkbox"/> <b>PROSTATE CANCER</b> [Z85.46] (male patients) Age at dx: _____ Gleason score: _____
<input type="checkbox"/> <b>COLORECTAL CANCER</b> Location: <input type="checkbox"/> colon [Z85.030] <input type="checkbox"/> rectum [Z85.040] Age at dx: _____ Pathology: _____ Was tumor screening performed? <input type="checkbox"/> Yes <input type="checkbox"/> No <i>If "Yes," attach results.</i>
<input type="checkbox"/> <b>GI POLYPS</b> [Z86.010] Age at dx: _____ Number of polyps: _____ Pathology: <input type="checkbox"/> adenomatous <input type="checkbox"/> other type: _____
<input type="checkbox"/> <b>HEMATOLOGIC CANCER</b> Age at dx: _____ Pathology: _____ Was bone marrow or stem cell transplant performed? <input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> <b>OTHER CANCER</b> Age at dx: _____ Type: _____
Is patient adopted with no family history? <input type="checkbox"/> Yes <input type="checkbox"/> No
<b>Additional information:</b>

REQUISITION CHECKLIST
<b>DID YOU INCLUDE?</b> <input type="checkbox"/> Riscover™ Cancer Compass family history form (both sides) <b>REQUIRED</b> <input type="checkbox"/> ICD-10 diagnosis code(s) <b>REQUIRED</b> <input type="checkbox"/> Patient's insurance card (legible front and back copy) <b>REQUIRED</b> <input type="checkbox"/> Patient clinical history (if personal history of cancer) <input type="checkbox"/> Genetic counseling visit notes (if performed) <input type="checkbox"/> Results for any previous testing <input type="checkbox"/> Previous genetic testing on patient <input type="checkbox"/> Previous genetic testing on patient's family members <input type="checkbox"/> Tumor testing results, such as MSI/IHC for colorectal or uterine cancer

ADDITIONAL CLINICAL INFO
<b>PREVIOUS CANCER GENETIC TESTING</b> <input type="checkbox"/> No previous testing <input type="checkbox"/> Testing performed: (Attach results if available) gene(s) tested: _____ results: _____
<b>GENETIC COUNSELING PERFORMED?</b> <input type="checkbox"/> Yes <input type="checkbox"/> No Date of GC visit: _____ / _____ / _____ <i>Please attach GC visit note.</i>
<b>OTHER RELEVANT CLINICAL HISTORY</b>

**REQUIRED:** ICD-10 Code(s) *Commonly used codes are noted above and additional codes are listed on the reverse.*

PATIENT NAME	PATIENT NAME	PATIENT NAME	PATIENT NAME
DOB	DOB	DOB	DOB

## RISCOVER™ PANEL DESCRIPTIONS

### Riscover™ Comprehensive

Analysis of *APC, ATM, BARD1, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN2A, CHEK2, EPCAM, MEN1, MLH1, MSH2, MSH6, MUTYH, NBN, NF1, PALB2, PMS2, POLD1, POLE, PTEN, RAD51C, RAD51D, RET, SMAD4, STK11, TP53, VHL*

### Riscover™ BRCA

Sequence and deletion/duplication analysis for *BRCA1* and *BRCA2*

### Riscover™ BRCA Ashkenazi Jewish Three-Site Analysis

Targeted analysis of three mutations: *BRCA1* c.68\_69delAG; *BRCA1* c.5266dupC; *BRCA2* c.5946delT

### Riscover™ Lynch Syndrome

Sequence and deletion/duplication analysis for *MLH1, MSH2, MSH6*, and *PMS2*; Deletion/duplication analysis only for *EPCAM*

## ICD-10 CODES

### PERSONAL HISTORY OF CANCER

Personal history of malignant carcinoid tumor of large intestine .....	Z85.030
Personal history of other malignant neoplasm of large intestine .....	Z85.038
Personal history of malignant carcinoid tumor of rectum .....	Z85.040
Personal history of other malignant neoplasm of rectum, rectosigmoid junction, and anus.....	Z85.048
Personal history of malignant neoplasm of pancreas.....	Z85.07
Personal history of malignant neoplasm of breast.....	Z85.3
Personal history of malignant neoplasm of other parts of uterus.....	Z85.42
Personal history of malignant neoplasm of ovary.....	Z85.43
Personal history of malignant neoplasm of prostate.....	Z85.46
Personal history of in-situ neoplasm of breast.....	Z86.000
Personal history of colonic polyps.....	Z86.010

### FAMILY HISTORY OF CANCER

Family history of malignant neoplasm of breast.....	Z80.3
Family history of malignant neoplasm of ovary .....	Z80.41
Family history of malignant neoplasm of prostate .....	Z80.42
Family history of malignant neoplasm of other organs or systems.....	Z80.8
Family history of malignant neoplasm, unspecified.....	Z80.9
Family history of colonic polyps .....	Z83.71
Family history of carrier of genetic disease.....	Z84.81

### LIMITED OR NO FAMILY HISTORY

Genetic susceptibility to malignant neoplasm of breast.....	Z15.01
Genetic susceptibility to malignant neoplasm of endometrium .....	Z15.04
Genetic susceptibility to malignant neoplasm of ovary .....	Z15.02
Genetic susceptibility to malignant neoplasm of prostate .....	Z15.03
Genetic susceptibility to other malignant neoplasm .....	Z15.09