myriad | Hereditary Cancer Test Request Form

- Please submit both pages of this form
- Make sure information is complete and legible

FOR LAB USE

SPECIMEN COLLECTION DATE (REQUIRED)
__/__/____ (MM/DD/YYYY)

NOTE: Affix Patient Identifier Label to Specimen Tube

At the time of specimen collection: ☐ Hospital Inpatient (>24 hour stay) Discharge date: __/__/____ (MM/DD/YYYY) ☐ Hospital Outpatient ☐ Non-Hospital Patient

1. Patient Information (Complete information required)

Name (last)	Name (first)	(m.i.)	Gender ☐ Male ☐ Female	Birthdate (MM/DD/YYYY)	Patient ID #

Email	Cell phone	Daytime phone
	- -	- -

Address	City	State	Zip

2. Ordering Provider Information (Only name and HCP Account # required unless you're a new customer or HCP # is unknown)

Name (last)	Name (first)	Myriad HCP Account #	Degree	NPI #

Address	City	State	Zip

Office Contact Name	Phone	Fax	Email
	- -	- -	

3. Send Results To (Optional - additional clinician can be listed to receive test status updates and the patient's copy of the test results)

Name (last)	Name (first)	Myriad HCP Account #	Degree	NPI #

Address	City	State	Zip

Office Contact Name	Phone	Fax	Email
	- -	- -	

4. Test Requested (For test descriptions see reverse)

Tests ordered will be processed and billed based on payer criteria.

Myriad myRisk® Hereditary Cancer Panel Testing

HEREDITARY BREAST AND OVARIAN CANCER SYNDROME CRITERIA:

☐ Integrated BRAC*Analysis*® with Myriad myRisk® Hereditary Cancer Update Test

FOR PATIENTS OF ASHKENAZI JEWISH ANCESTRY:

☐ Multisite 3 BRAC*Analysis*®

☐ REFLEX to Integrated BRAC*Analysis*® with Myriad myRisk® Hereditary Cancer Update Test if the Multisite 3 is negative

LYNCH SYNDROME OR *MYH*-ASSOCIATED POLYPOSIS (MAP) CRITERIA:

☐ COLARIS®PLUS with Myriad myRisk® Hereditary Cancer Update Test

FAMILIAL POLYPOSIS SYNDROME CRITERIA:

☐ COLARIS *AP*®PLUS with Myriad myRisk® Hereditary Cancer Update Test

FOR PATIENTS PREVIOUSLY TESTED AT MYRIAD:

☐ Myriad myRisk® Update Test (Available to patients previously tested negative with BRAC*Analysis*®, COLARIS®, and/or COLARIS *AP*®. BART and/or *PMS2* testing will be included in the test order unless previously performed or restricted by payor criteria.)

RISK ANALYSIS OPTIONS (to be excluded on report, see reverse for details):

☐ riskScore® is not appropriate for this patient

☐ Tyrer-Cuzick and riskScore® are not appropriate for this patient

Single-Syndrome Testing

HEREDITARY BREAST AND OVARIAN CANCER SYNDROME CRITERIA:

☐ Integrated BRAC*Analysis*®

FOR PATIENTS OF ASHKENAZI JEWISH ANCESTRY:

☐ Multisite 3 BRAC*Analysis*®

☐ REFLEX to Integrated BRAC*Analysis*® if the Multisite 3 is negative

LYNCH SYNDROME OR *MYH*-ASSOCIATED POLYPOSIS (MAP) CRITERIA:

☐ COLARIS®PLUS

FAMILIAL POLYPOSIS SYNDROME CRITERIA:

☐ COLARIS *AP*®PLUS

Additional Tests*

☐ **Single Site Testing:**

Specify Gene: _____

_____

Mutation: _____

_____

Relationship: My patient is the _____

(e.g. maternal aunt) of the known mutation carrier. **Required: Include a copy of the known mutation carrier's report.**

☐ **Other:** (e.g. single gene analysis)

_____

5. Confirmation of Informed Consent & Statement of Medical Necessity

I affirm each of the following: I have provided genetic testing information to the patient and the patient has consented to genetic testing. This test is medically necessary for the diagnosis of a disease or syndrome. The results will be used in the patient's medical management and treatment decisions. The person listed as the Ordering Physician is authorized by law to order the test(s) requested herein.

SIGN HERE: Medical Professional (required to process form) X _____ Date: __/__/____ (MM/DD/YYYY)

(Signature date is the specimen collection date if a different date is not provided above)

6. Billing/Payment Information

☐ **OPTION 1: BILL INSURANCE** (Please attach copy of authorization/referral)

Name of Policy Holder: _____ DOB: __/__/____ (MM/DD/YYYY)

Insurance ID#: _____ Patient Relation to Policy Holder: ☐ Self ☐ Spouse ☐ Child ☐ Other Authorization/Referral: _____

SIGN HERE: Patient/Responsible Party **I AGREE TO THE BILLING TERMS ON REVERSE** X _____ DATE: __/__/____ (MM/DD/YYYY)

Reminder: Include a copy of BOTH SIDES of your insurance card(s). If you submit more than one card, indicate which is primary.

I understand that Myriad will contact me if I will be financially responsible for any non-covered service. To be considered for the Myriad Financial Assistance Program, please provide the following information: Annual household income $ _____,_____. Number of family members in household _____.

☐ **OPTION 2: PATIENT PAYMENT** (Please call Customer Service for questions regarding test prices or for credit card payment)

☐ **OPTION 3: OTHER BILLING** (To establish an account, submit billing information with this form)

☐ Bill our institutional account #: _____ or established research project code #: _____ or Authorization/Voucher #: _____

MYRIAD GENETIC LABORATORIES, INC. A CLIA Certified Laboratory
320 Wakara Way • Salt Lake City, UT 84108 / **(800) 469-7423 • Fax (801) 584-3615** • myriad.com

PP-0718-MR1

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Testing for Myriad myRisk® Hereditary Cancer

IMPORTANT INFORMATION FOR PATIENT*

BILLING TERMS: I represent that I am covered by insurance and authorize Myriad Genetic Laboratories, Inc. (MGL) to give my designated insurance carrier, health plan, or third party administrator (collectively "Plan") the relevant health information necessary for reimbursement. I authorize Plan benefits to be payable to MGL. I understand MGL will contact me if I will be financially responsible for any non-covered service. I agree to assist MGL in resolving insurance claim issues and if I don't assist, I may be responsible for the full test cost. I permit a copy of this authorization to be used in place of the original.

NON-DISCRIMINATION: Federal law (GINA) and laws in most states prohibit discrimination regarding employment eligibility, health benefits, or health insurance premiums based solely on genetic information. Myriad Genetic Laboratories, Inc. (Myriad) complies with applicable Federal civil rights laws and does not discriminate on the basis of race, color, national origin, age, disability, or sex.

*Translation of Billing Terms are available in Mandarin and Spanish at MyriadPromise.com. Myriad also provides free language services to people whose primary language is not English through qualified interpreters. If you need these services, contact Customer Service at 800-469-7423,

AFFORDABILITY: Myriad Promise™

- The majority of appropriate patients pay $0
- Myriad will work with your insurance provider to help you get the appropriate coverage
- If you encounter ANY financial hardship associated with your bill, Myriad will work with you toward your complete satisfaction
- For more information please refer to the billing information at MyriadPromise.com

TEST DESCRIPTIONS (For a full list of tests offered, visit www.myriadpro.com)

Integrated BRAC*Analysis*®: Analysis of *BRCA1* and *BRCA2* for susceptibility to Hereditary Breast and Ovarian Cancer syndrome.

Integrated BRAC*Analysis*® with Myriad myRisk® Hereditary Cancer Update Test: Analysis of *BRCA1* and *BRCA2* for susceptibility to Hereditary Breast and Ovarian Cancer syndrome with additional genes associated with hereditary cancer risk (see table below). May also include riskScore® breast cancer analysis. See below details for inclusion criteria.

Multisite 3 BRAC*Analysis*®: Three-mutation *BRCA1* and *BRCA2* analysis for individuals of Ashkenazi Jewish ancestry: *BRCA1* c.68_69del (p.Glu23Valfs*17) (aka *BRCA1* 185delAG, 187delAG); *BRCA1* c.5266dupC (p.Gln1756Profs*74) (aka *BRCA1* 5382insC, 5385insC); *BRCA2* c.5946del (p.Ser1982Argfs*22) (aka *BRCA2* 6174delT).

COLARIS®PLUS: Analysis of *MLH1, MSH2, MSH6, PMS2, MUTYH,* and *EPCAM* for susceptibility to Lynch syndrome (HNPCC) and *MYH*-Associated Polyposis (MAP).

COLARIS®PLUS with Myriad myRisk® Hereditary Cancer Update Test: Analysis of *MLH1, MSH2, MSH6, PMS2, MUTYH,* and *EPCAM* for susceptibility to Lynch syndrome (HNPCC) and *MYH*-Associated Polyposis (MAP) with additional genes associated with hereditary cancer risk (see table below). May also include riskScore® breast cancer analysis. See below details for inclusion criteria.

COLARIS *AP*®PLUS: Analysis of *APC* for susceptibility to FAP/AFAP.

COLARIS *AP*®PLUS with Myriad myRisk® Hereditary Cancer Update Test: Analysis of *APC* for susceptibility to FAP/AFAP with additional genes associated with hereditary cancer risk. May also include riskScore® breast cancer analysis. See below details for inclusion criteria.

Single Site Testing: Analysis of single, familial mutation.

Myriad myRisk® Update Test: Analysis of 29 hereditary cancer genes for patients who previously tested negative for BRAC*Analysis*®, COLARIS®, and/or COLARIS *AP*®. BART and/or *PMS2* testing will be included in the test order unless previously performed or restricted by payor criteria. May also include riskScore® breast cancer analysis. See below details for inclusion criteria.

Certain payers do not cover genetic testing when Single Nucleotide Polymorphisms (SNPs) are a component of the test. For payers who do not reimburse for a hereditary cancer test due to SNP analysis inclusion, Myriad will report the myRisk Hereditary Cancer Test without SNPs and these patients will not receive a SNP based riskScore®. Please visit www.myriadpro.com/payeroptout to determine if your patient's payer does not reimburse for hereditary cancer genetic testing with SNP analysis.

Genes & Associated Cancers*	Br	Ov	Co	En	Me	Pa	Ga	Pr
BRCA1	●	●				●		●
BRCA2	●	●			●	●		●
*MLH1, MSH2, MSH6, PMS2, EPCAM***		●	●	●		●	●	●
APC			●			●	●	
MUTYH			●					
CDK4, CDKN2A (p16INK4a, p14ARF)					●	●		
TP53	●	●	●	●	●	●	●	●
PTEN	●		●	●	●			
STK11	●	●	●	●		●	●	
CDH1	●		●				●	
BMPR1A, SMAD4			●			●	●	
PALB2, ATM	●					●		
CHEK2	●		●					
NBN	●							●
BARD1	●							
BRIP1		●						
RAD51C, RAD51D		●						
POLD1, POLE			●					
GREM1			●					
HOXB13								●

Br: Breast / **Ov**: Ovarian / **Co**: Colorectal / **En**: Endometrial / **Me**: Melanoma / **Pa**: Pancreatic / **Ga**: Gastric / **Pr**: Prostate

*Additional risks may be associated with each gene/syndrome. **Large rearrangement only.

Turnaround Time:

- The majority of Myriad myRisk® results are completed within 14 days
- We will notify you in the unusual event results take longer than 21 days

Myriad myRisk® Report includes:

- myRisk Genetic Result
- riskScore® Result
  - Personalized breast cancer risk assessment based on an analysis of biomarkers combined with patient clinical and family history data
- myRisk Management Tool
  - Guideline based (NCCN, CAPS, Amsterdam, and others) cancer management for both positive and negative results
  - Includes a Tyrer-Cuzick breast cancer risk estimate

Completing the Test Request Form:

- Please include:
  - Age, cancer diagnosis, ancestry, gender and cancer family history

The myRisk Management Tool and riskScore may not be reported without an accurate and specific personal and family history included on the Patient Cancer Family History in Sections 7 - 11.

riskScore® is calculated for women under age 85, of solely White / Non-Hispanic and/or Ashkenazi Jewish ancestry, without a personal history of breast cancer, LCIS, hyperplasia, atypical hyperplasia, or a breast biopsy with unknown results. riskScore® is not calculated if a woman or blood relative is known to carry a mutation in a breast cancer risk gene. riskScore® and Tyrer-Cuzick model will not be calculated if provider indicates that they are not appropriate for the patient by selecting the check box in Section 4. Not all data collected on the TRF is incorporated into Tyrer-Cuzick or riskScore® calculations. Some fields may be used for anonymized, internal validation studies only.

- The genes associated with Myriad myRisk® Hereditary Cancer Panel are subject to change. To ensure you have a current version of the TRF and the genes included with the Myriad myRisk panel please visit **www.myriadpro.com/documents-and-forms/test-request-forms** and **www.myriadpro.com/myrisk/why-myriad-myrisk/gene-selection**.
- For additional information visit MySupport360.com and MyriadPro.com

7. Patient Information (Make sure information is the same as entered on page 1)

Name (last)	(first)	(m.i.)	Birthdate (MM/DD/YYYY)
			/ /

8. Ancestry (riskScore® is currently only validated and provided for patients of solely White/Non-Hispanic and/or Ashkenazi Jewish ancestry)

Select all that apply:
☐ Ashkenazi Jewish ☐ Asian ☐ Black / African ☐ Hispanic / Latino ☐ Middle Eastern ☐ Native American ☐ Pacific Islander ☐ White / Non-Hispanic

9. Patient Personal History of Cancer & Other Clinical Information (Select all that apply)

☐ **No personal history of cancer**

Patient has been diagnosed with:	Age at Diagnosis	Patient is Currently Being Treated	Pathology / Other Info
☐ **Breast Cancer** ☐ **Left** ☐ **Right**		☐	☐ Ductal Invasive ☐ Lobular Invasive ☐ DCIS ☐ Bilateral ☐ Triple-Negative (ER-, PR-, HER2-) ☐ Metastatic: HER2 Status: ☐ + ☐ - Previous Chemotherapy: ☐ Yes ☐ No If ER/PR+, previous Endocrine Therapy: ☐ Yes ☐ No ☐ N/A or inappropriate
☐ **Endometrial / Uterine Cancer**		☐	☐ Tumor MSI-High or IHC Abnormal - Result: ☐ Tumor not available for MSI or IHC testing
☐ **Ovarian Cancer**		☐	☐ Non-epithelial
☐ **Prostate Cancer**		☐	Gleason Score: ☐ Metastatic
☐ **Colon / Rectal Cancer**		☐	Type: ☐ Mucinous ☐ Signet Ring ☐ Medullary Growth Pattern ☐ Tumor Infiltrating Lymphocytes ☐ Crohn's-like Lymphocytic Reaction ☐ Patient's tumor is MSI-High or IHC Abnormal - Result: ☐ Tumor not available for MSI or IHC testing
☐ **Colon / Rectal Adenomas**		☐	Cumulative Adenomatous Polyp #: ☐ 1 ☐ 2-5 ☐ 6-9 ☐ 10-19 ☐ 20-99 ☐ 100+
☐ **Hematologic Cancer**		☐	
☐ **Other Cancer**		☐	Type
☐ **Other Cancer**		☐	Type

Check if applicable to patient:
- % on one of the Lynch Syndrome Risk Models (PREMM$_5$, MMRpro, or MMRpredict)
- ☐ Bone Marrow Transplant Recipient Type: ☐ Autologous ☐ Allogeneic (If allogeneic please call 800-469-7423 x3850)
- ☐ Blood Transfusion Recipient Type: ☐ Whole blood ☐ Packed red blood cells Date: / / (MM/DD/YYYY)

10. Family History of Cancer

Provide complete and specific information to ensure proper insurance reimbursement, determine cancer risk estimates, and optimize medical management recommendations.

☐ **No Known Family History of Cancer**

☐ **Limited Family Structure** Limited family history available such as fewer than two female 1st or 2nd degree maternal or paternal relatives having lived beyond age 45

Relationship to Patient	Maternal (mother's side)	Paternal (father's side)	Cancer Site or Polyp Type (add # for colon/rectal adenomas)	Age at Each Diagnosis
	☐	☐		
	☐	☐		
	☐	☐		
	☐	☐		
	☐	☐		
	☐	☐		
	☐	☐		

11. Breast Cancer Risk Model Information

Only complete for female patients NEVER diagnosed with breast cancer.

Patient information:

Height - ft: in: **Weight** (lbs):

Age at time of first menstrual period:

Is patient: ☐ Pre-menopausal ☐ Peri-menopausal ☐ Post-menopausal: **Age of onset:**

Has this patient had a live birth?: ☐ No ☐ Yes: **patient's age at first child's birth:**

Has patient ever used Hormone Replacement Therapy? ☐ No ☐ Yes

If Yes, Treatment Type: ☐ Combined ☐ Estrogen only ☐ Progesterone only

If Yes, is patient a: ☐ **Current User:** Started ____ years ago Intended use for ____ more years ☐ **Past User:** Stopped ____ years ago

INFORMATION about PATIENT'S FEMALE RELATIVES:

Number of daughters:

Number of sisters:

Number of maternal aunts (mother's sisters):

Number of paternal aunts (father's sisters):

OTHER INFORMATION:

Mammographic Density:

Has the patient had her breast density assessed? ☐ No ☐ Yes

If yes, complete one of the following for the most recent assessment:

☐ **Volpara® Volumetric Density:** ____.____ %

☐ **VAS Percentage Density:** ____.____ %

☐ **BI-RADS® ATLAS Density (Select one of the following):**

☐ Almost entirely fatty ☐ Heterogeneously dense ☐ Scattered fibroglandular density ☐ Extremely dense ☐ Unknown

NOTE: Risk associated with mammographic density is not incorporated into riskScore (v.1), nor Tyrer-Cuzick (v.7) calculations provided on the clinical report.

Please indicate if the patient has had a breast biopsy showing one or more of the following results: ☐ N/A (No biopsy or none of the listed results)

☐ Hyperplasia ☐ Atypical Hyperplasia ☐ LCIS ☐ Biopsy with unknown or pending results