

Test Request Form

A test for the polyposis syndromes
(FAP, AFAP, MAP)

page 1 of 2

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

Note: Affix patient identifier label to specimen tube

Specimen collection date (required)

(mm/dd/yyyy)

FOR LAB USE

At the time of specimen collection: Non-hospital patient Hospital outpatient Hospital inpatient (>24 hour stay) Discharge date: (mm/dd/yyyy)

1. Patient information (Complete information required)

Legal name (last)	Legal name (first)	(m.i.)	Sex at birth <input type="checkbox"/> M <input type="checkbox"/> F	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)

Testing for Lynch syndrome (HNPCC) and MYH-associated polyposis (MAP)
<input type="checkbox"/> COLARIS AP [®] PLUS – Analysis of APC and MYH
Other tests
<input type="checkbox"/> Single Site testing – Mutation-specific analysis for individuals with known mutations in their family. Specify gene: _____ Specify variant (mutation): _____ Relationship: My patient is the _____ (e.g., maternal aunt) of the known mutation carrier. Required: Include a copy of the known mutation carriers report.
<input type="checkbox"/> Other:

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. I authorize Myriad to assist my patients in obtaining pre-test genetic counseling services if required by the patient's insurance provider (see reverse). The person listed as the ordering provider is authorized by law to order the test(s) requested herein.

Sign here: **Medical professional** (required to process form) X

(Signature date is the specimen collection date if a different date is not provided here) Date: _____ (mm/dd/yyyy)

6. Billing/payment information

Option 1: Bill insurance (Please attach copy of authorization/referral)

Name of policy holder: _____
DOB: _____ (mm/dd/yyyy)

Name of insurance: _____
Insurance ID#: _____
Authorization/referral: _____

Patient relation to policy holder: Self Spouse Child Other

Sign here: **Patient/responsible party** I agree to the billing terms on reverse. X

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: Uninsured (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 #: _____



Testing for COLARIS AP[®]PLUS

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. By agreeing to testing I also authorize Myriad to obtain a consumer credit report on me from a consumer reporting agency selected by Myriad. I understand and agree that Myriad may use my consumer credit report to confirm whether my income qualifies me for financial assistance. I further understand that this is not a credit application and will not impact my credit score. 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.

[^]Translation of Billing Terms are available in Mandarin and Spanish at www.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
- Myriad is committed to provide patients with access to accurate and affordable genetic results
- For more information please refer to the billing information at www.myriadpromise.com

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.

Sex assigned at birth is a label given to an individual at birth, typically "male" or "female".

A legal name identifies a person for legal and administrative purposes. It is recorded on a birth certificate, marriage certificate, or other government issued document that records a name change.

COLARIS AP[®]

A genetic test for adenomatous polyposis syndromes (including familial adenomatous polyposis (FAP), attenuated FAP, and MYH-associated polyposis (MAP))

COLARIS AP[®]PLUS :

Full sequence and large rearrangement analysis of *APC* and *MYH*

Single Site Polyposis Testing:

Mutation-specific analysis for individuals with a known *APC* or *MYH* mutation(s) in the family

Gene-Specific COLARIS AP[®]:

APC gene analysis – Full sequence and large rearrangement analysis of *APC*

MYH gene analysis – Full sequence and large rearrangement analysis of *MYH*

NOTE: If COLARIS[®]PLUS and COLARIS AP[®]PLUS are ordered on the same patient, *MYH* analysis will only be performed once.

Authorization of referral to genetic counseling

In signing section 5 of the test request form, you hereby authorize Myriad to assist your patient in obtaining genetic counseling from a third-party service. The specific process will vary by third-party counseling service but in most situations the genetic counselor will be added as the healthcare provider receiving a copy of the patient's results, and also be allowed to change the test order should there be a clinical or payer-related reason to do so. You authorize the genetic counselor to facilitate the completion of any test requisition forms and/or submit any prior authorization, if necessary, on your behalf and identifying you as the ordering provider in any such forms by including your name and NPI.

Special instructions (if applicable): *Please note: some options may not be possible if an alternate is required by the patient's insurance or if the patient requests otherwise.

- Expedite genetic counseling for immediate management decision
- Maintain my test as ordered
- Allow me to review results with my patient prior to their follow-up counseling session
- Other: _____

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

Legal name (last)	Legal name (first)	(m.i.)	Birthdate (mm/dd/yyyy)
-------------------	--------------------	--------	------------------------

8. 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)

Patient has never been diagnosed with cancer

Patient has been diagnosed with:	Age at diagnosis	Patient is currently being treated	Pathology / other info
<input type="checkbox"/> Colon cancer <input type="checkbox"/> Rectal cancer		<input type="checkbox"/>	Type: <input type="checkbox"/> Mucinous <input type="checkbox"/> Signet ring <input type="checkbox"/> Medullary growth pattern <input type="checkbox"/> Tumor infiltrating lymphocytes <input type="checkbox"/> Crohn's-like lymphocytic reaction <input type="checkbox"/> Patient's tumor is MSI-high or IHC abnormal - result: _____ <input type="checkbox"/> Tumor not available for MSI or IHC testing
<input type="checkbox"/> Colon adenomas <input type="checkbox"/> Rectal adenomas		<input type="checkbox"/>	Cumulative adenomatous polyp #: <input type="checkbox"/> 1 <input type="checkbox"/> 2-5 <input type="checkbox"/> 6-9 <input type="checkbox"/> 10-19 <input type="checkbox"/> 20-99 <input type="checkbox"/> 100+
<input type="checkbox"/> Endometrial cancer - not sarcoma		<input type="checkbox"/>	<input type="checkbox"/> Tumor MSI-high or IHC abnormal - result: _____ <input type="checkbox"/> Tumor not available for MSI or IHC testing
<input type="checkbox"/> Prostate cancer		<input type="checkbox"/>	Gleason score: _____ <input type="checkbox"/> Metastatic (includes distant metastasis and regional bed/nodes) <input type="checkbox"/> NCCN high / very high risk
<input type="checkbox"/> Hematologic cancer		<input type="checkbox"/>	
<input type="checkbox"/> Other cancer		<input type="checkbox"/>	Type: _____
<input type="checkbox"/> ICD-10 code(s)/Dx:			


Check if applicable to patient: _____ % on one of the Lynch syndrome risk models (PREMM₅, MMRpro, or MMRpredict)

Bone marrow transplant recipient Type: Autologous Allogeneic (If allogeneic please call 800-469-7423 x3850)

Blood transfusion recipient within 28 days of sample collection Type: Whole blood Packed red blood cells

Blood transfusion recipient within 12 months of sample collection 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

Relationship to patient	Maternal (mother's side)	Paternal (father's side)	Cancer site, Gleason score, or polyp type (if colon/rectal adenomas, include total number)	Age at each diagnosis	Unavailable for testing	If relative has <u>not</u> been tested, why?		
						Relative is deceased	Patient has no contact with relative	Relative declines testing
	<input type="checkbox"/>	<input type="checkbox"/>			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	<input type="checkbox"/>	<input type="checkbox"/>			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	<input type="checkbox"/>	<input type="checkbox"/>			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	<input type="checkbox"/>	<input type="checkbox"/>			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	<input type="checkbox"/>	<input type="checkbox"/>			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	<input type="checkbox"/>	<input type="checkbox"/>			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	<input type="checkbox"/>	<input type="checkbox"/>			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

