



MYRIAD®

MYRIAD GENETIC LABORATORIES, INC.
A CLIA Certified Laboratory
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COLARIS®

A Test for Lynch Syndrome (HNPCC) and MYH-Associated Polyposis (MAP)

Test Request Form and Statement of Medical Necessity
TO AVOID DELAYS PLEASE COMPLETE ENTIRE FORM

NOTE: Affix Bar Code Label to Specimen Tube

SPECIMEN COLLECTION DATE (REQUIRED)

05/10/2012

ORDERING PHYSICIAN and SEND RESULTS TO (IF OTHER THAN ORDERING PHYSICIAN)
Physician, Bob, MD
3456789012
Shirley Assistant
222-222-2222
222-222-2223

PATIENT INFORMATION (COMPLETE INFORMATION REQUIRED FOR INSURANCE COVERAGE)
Doe, Jane T
00000
FEMALE
09/10/1954
123 Generic Street
This City
ST
45678
222-000-0000
janedoe@email.com

ANCESTRY AND CLINICAL HISTORY
WESTERN/NORTHERN EUROPE
ASHKENAZI
CENTRAL/EASTERN EUROPE
LATIN AMERICAN/CARIBBEAN
AFRICA
ASIA
NEAR EAST/MIDDLE EAST
NATIVE AMERICAN
OTHER

PATIENT PERSONAL HISTORY OF CANCER (Check all that apply)
ICD-9 Code(s)/Dx: 153.0, V16.40
Colorectal Invasive*
Mucinous
Signet Ring
Medullary Growth Pattern
Tumor Infiltrating Lymphocytes
Crohn's-like Lymphocytic Reaction
Adenomatous Polyps
Endometrial/Uterine
Other
Bone Marrow Transplant Recipient

FAMILY HISTORY OF CANCER (Please Indicate Relationship, Maternal or Paternal, Site of Cancer or Adenoma Number, Age at Diagnosis)
Mother
Aunt
Colon
Endometrial
56
48

TESTS REQUESTED (FOR DETAILED INFORMATION ABOUT THESE TESTS, PLEASE SEE BACK OF THIS FORM)

INITIAL TESTING FOR LYNCH SYNDROME (HNPCC) AND MYH-ASSOCIATED POLYPOSIS (MAP)

COLARIS - Analysis of MLH1, MSH2, MSH6, PMS2, MYH, and EPCAM (PMS2 and MYH will be reported independently. PMS2 will be billed separately, and coverage may vary based on payor criteria.)

OTHER TESTS

SINGLE SITE TESTING (For family of known mutation carriers)
Specify Gene: and Mutation:
Relationship: My patient is the (eg, maternal aunt) of the known mutation carrier. Include a copy of the known mutation carrier's report, if available.
GENE-SPECIFIC TESTING (Please specify):
OTHER:

INFORMED CONSENT AND STATEMENT OF MEDICAL NECESSITY

I have supplied information to the patient regarding genetic testing and the patient has agreed for genetic testing to be performed. I further confirm that this test is medically necessary for the diagnosis or detection of a disease, illness, impairment, symptom, syndrome or disorder, and that the information will be used in the medical management and treatment decisions for the patient. I confirm that the person listed in the Ordering Physician space above is authorized by law to order the test(s) requested.
Bob Physician, MD
05/10/2013

BILLING/PAYMENT INFORMATION

OPTION 1: PLEASE BILL MY INSURANCE (Option 1 requires patient signature and enlarged copy of both sides of insurance card(s). If two cards are submitted, indicate which is primary)

Name of Policy Holder: JANE DOE
DOB: 09/10/1954
Insurance ID #/SSN: ABC-123
Patient Relation to Policy Holder: Self
I hereby represent that I am covered by insurance and authorize Myriad Genetic Laboratories, Inc. (MGL) to furnish my designated insurance carrier, health plan, or third party administrator (collectively "Plan") the information on this form and other information provided by my healthcare provider necessary for reimbursement. I authorize Plan benefits to be payable to me and that MGL will contact me prior to test start ONLY if my total financial responsibility will exceed \$375 (for any reason, including co-insurance and deductibles, and other services). If requested, 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. I understand that I may not alter any printed text in this paragraph.
Jane Doe
05/10/2013

REMINDER: INCLUDE A COPY OF BOTH SIDES OF YOUR INSURANCE CARD(S)

OPTION 2: PATIENT PAYMENT (Please call Customer Service for questions regarding test prices)

Please bill my credit card (all major credit cards accepted) in the amount of \$ Card # Exp. Date:
Cardholder Name (please print): Cardholder Signature:
Personal check, cashiers check, or money order enclosed, payable to Myriad Genetic Laboratories, Inc.

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 offers the following tests for Lynch syndrome, also known as Hereditary Nonpolyposis Colorectal Cancer (HNPCC), and *MYH*-Associated Polyposis (MAP):

COLARIS®:

Sequence and large rearrangement analysis of *MLH1*, *MSH2*, *MSH6*, *MYH*, and *PMS2*,* and large rearrangement analysis of *EPCAM*

NOTE: *MYH* and *PMS2* will be reported independently. *PMS2* will be billed separately, and coverage may vary based on payor criteria.

Single Site Lynch Syndrome Testing:

Mutation-specific analysis for individuals with a known *MLH1*, *MSH2*, *MSH6*, *PMS2*, *MYH*, or *EPCAM* mutation in the family

Gene-Specific Lynch Syndrome Testing and MAP Analysis:

MLH1 Analysis – Sequence and large rearrangement analysis of *MLH1*

MSH2 Analysis – Sequence and large rearrangement analysis of *MSH2* including *EPCAM* rearrangements

MSH6 Analysis – Sequence and large rearrangement analysis of *MSH6*

PMS2 Analysis – Sequence and large rearrangement analysis of *PMS2*

MYH Analysis – Sequence and large rearrangement analysis of *MYH*

**PMS2* analysis will be run as a reflex test following a negative result for *MLH1*, *MSH2* and *MSH6* for submissions to Medicare and certain other payors.

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