

Molecular Genetics-Inherited Cancer Syndromes Patient Information Sheet (Supply T519)



*Molecular Genetics – Inherited Cancer Syndromes
Patient Information Sheet*

The accurate interpretation and reporting of genetic results is contingent upon the reason for referral, clinical information, ethnic background, and family history. To help provide the best possible service, please supply the information requested below and **send paperwork with the specimen or return by fax to Laboratory Genetics 507-284-0670 (phone 507-538-2996).**

Patient Name (Last, First, Middle)	Sex <input type="checkbox"/> Male <input type="checkbox"/> Female	Birth Date (Month DD, YYYY)
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Check box(es) below and complete information to indicate who should receive reports.

<input type="checkbox"/> Requesting Physician	Phone	Fax
<input type="checkbox"/> Genetic Counselor	Phone	Fax
MML Account Number		

Reason for Testing NOTE: It is recommended to first document a mutation in an affected family member prior to performing a pre-symptomatic testing.

<input type="checkbox"/> Patient has a definite diagnosis of:	<input type="checkbox"/> FAP <input type="checkbox"/> HNPCC <input type="checkbox"/> MEN2 <input type="checkbox"/> MYH	Age when diagnosed _____
<input type="checkbox"/> Patient has a possible diagnosis of:	<input type="checkbox"/> FAP <input type="checkbox"/> HNPCC <input type="checkbox"/> MEN2 <input type="checkbox"/> MYH	
<input type="checkbox"/> Patient is asymptomatic but has a family history of:	<input type="checkbox"/> FAP <input type="checkbox"/> HNPCC <input type="checkbox"/> MEN2 <input type="checkbox"/> MYH	

If more than one test is ordered, should all tests be performed at the same time? Yes No
If No, indicate preferred order of testing:

Family History

Are other relatives known to be affected? Yes No If yes, indicate their relationship to the patient:

Are other relatives known to be a carrier? Yes No If yes, indicate their relationship to the patient:

Have other relatives had molecular genetic testing? Yes No If yes, indicate the results (specific mutation(s) identified and the laboratory at which testing was performed):

If the relative was tested at the Mayo Clinic, include the name of the family member:

Please include a detailed pedigree, if available.

FAP (Familial Adenomatous Polyposis) and MYH (MYH Gene Analysis for Multiple Adenoma) Clinical Information

NUMBER OF POLYPS <input type="checkbox"/> Not screened <input type="checkbox"/> Screen negative (0 polyps) <input type="checkbox"/> 1-5 <input type="checkbox"/> 6-20 <input type="checkbox"/> 21-50 <input type="checkbox"/> 51-100 <input type="checkbox"/> 101-500 <input type="checkbox"/> >500 Has histopathology confirmed that polyps are adenomatous? <input type="checkbox"/> Yes <input type="checkbox"/> No	OTHER CLINICAL MANIFESTATIONS CHRPE <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Desmoid tumors <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Epidermoid cysts <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Osteomas <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Duodenal or gastric polyps <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Colon Cancer <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown List location of Colon Cancer: Other - (describe)
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HNPCC (Hereditary Nonpolyposis Colorectal Cancer)

NUMBER OF POLYPS <input type="checkbox"/> Not screened <input type="checkbox"/> Screen negative (0 polyps) <input type="checkbox"/> List number of polyps present: Has MSI and/or IHC been performed? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, please indicate results:	OTHER CLINICAL MANIFESTATIONS Endometrial Cancer <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Gastric Cancer <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Upper Tract Urothelial Cancer <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Brain Tumor <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Colon Cancer <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown List location of Colon Cancer: Other - (describe)
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MEN2 (Multiple Endocrine Neoplasia Type 2)

CLINICAL SYMPTOMS (in this individual) <input type="checkbox"/> Medullary Thyroid Cancer <input type="checkbox"/> Ganglioneuromas	<input type="checkbox"/> Elevated Calcitonin <input type="checkbox"/> Hyperparathyroidism	<input type="checkbox"/> Pheochromocytoma <input type="checkbox"/> Other: _____
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