



Patient Information Sheet

Instructions: The accurate interpretation and reporting of the 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.**

Patient Information

Patient Name <i>(First, Middle, Last)</i>		Birth Date <i>(Month DD, YYYY)</i>	Sex <input type="checkbox"/> Male <input type="checkbox"/> Female
Referring Physician Name	Phone Number	Fax Number	
Other Contact	Phone Number	Fax Number	

*Fax number given must be from a fax machine that complies with applicable HIPAA regulation.

Study Purpose

Study Purpose: <input type="checkbox"/> Diagnostic <input type="checkbox"/> Presymptomatic
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HHT Known Mutation (if applicable)

If 89391 ENG Gene Known Mutation, or 89393 ACVRL1 Gene Known Mutation is ordered, the following information must be provided or testing cannot be completed: Known familial mutation to be analyzed: Proband's relationship to patient:

HHT Deletion/Duplication Detection (if applicable)

If 89587 ENG and ACVRL1, Large Del/Dup is ordered for familial mutation testing, the following information must be provided or testing cannot be completed: Familial deletion or duplication: Proband's relationship to patient:
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Clinical History (check all that apply)

Pertinent Clinical and Laboratory History	
Telangiectasia(s)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Location and/or number: <input type="checkbox"/> Lips _____ <input type="checkbox"/> Oral Cavity _____ <input type="checkbox"/> Fingers _____ <input type="checkbox"/> Nose _____ <input type="checkbox"/> GI mucosa _____	
Nosebleed(s)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Frequency _____	
Visceral AVM(s)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Location and/or number: <input type="checkbox"/> Hepatic _____ <input type="checkbox"/> Cerebral _____ <input type="checkbox"/> Pulmonary _____ <input type="checkbox"/> Spinal _____ <input type="checkbox"/> Gastrointestinal _____	
Juvenile polyps? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown Other relevant clinical information (surgeries, etc.):	
Ethnic Background - <i>Ethnic background is necessary to provide appropriate interpretation of test results.</i>	
<input type="checkbox"/> Northern European Caucasian <input type="checkbox"/> Mixed European Caucasian <input type="checkbox"/> Southern European Caucasian If Caucasian, indicate countries of origin: _____	
<input type="checkbox"/> African American <input type="checkbox"/> Hispanic <input type="checkbox"/> Asian <input type="checkbox"/> French Canadian <input type="checkbox"/> African American <input type="checkbox"/> Other (specify): _____	
Family History	
Are other relatives known to be affected? <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, indicate their relationship to the patient.
Have other relatives had molecular genetic testing for HHT? <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, please indicate the performing laboratory and attach a copy of the genetic test lab report if available:
If the relative was tested at Mayo Clinic, include the name of the family member:	
Please include a detailed pedigree, if available.	