



**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, supply the information requested below and **send paperwork with the specimen.**

**Patient Information**

Patient Name <i>(Last, First, Middle)</i>	Birth Date <i>(mm-dd-yyyy)</i>	Sex <input type="checkbox"/> Male <input type="checkbox"/> Female
Referring Provider Name <i>(Last, First)</i>	Phone	Fax*
Other Contact Name <i>(Last, First)</i>	Phone	Fax*

\*Fax number provided must be from a fax machine that complies with applicable HIPAA regulations.

**Purpose of Study** Check all that apply.

**Clinical Status:**  Symptomatic  Asymptomatic  
**Study Purpose:**  Diagnostic  Presymptomatic

**Note:** If testing for a previously identified familial SDHB, SDHC, or SDHD mutation or variant is desired, order test FMTT / Familial Mutation, Targeted Testing and provide documentation of the familial mutation or variant to the laboratory by attaching a copy of the genetic test lab report and filling in the familial mutation or variant below.

- Mutation or variant to be detected: \_\_\_\_\_
- Proband's relationship to the patient: \_\_\_\_\_

**Pertinent Clinical and Laboratory History** Check all that apply.

Paragangliomas  Yes  No If yes, number and location: \_\_\_\_\_  
Pheochromocytomas  Yes  No If yes, unilateral or bilateral:  Unilateral  Bilateral  
Renal Cell Carcinoma  Yes  No Other tumors  Yes  No If yes, specify: \_\_\_\_\_  
Hypertension  Yes  No Headaches  Yes  No Profuse sweating  Yes  No  
Palpitations  Yes  No Anxiety  Yes  No

Supporting Biochemical Test Results (list)

Other Relevant Clinical Information (surgeries, malignancy, etc.)

**Ethnic Background** Ethnic background is necessary to provide appropriate interpretation of test results.

European Caucasian  African American  Hispanic  Asian  Other, specify: \_\_\_\_\_  
Indicate countries of origin: \_\_\_\_\_

**Family History** Include a detailed pedigree, if available.

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

Have other relatives had molecular genetic testing for SDHB, SDHC, or SDHD?  Yes  No  
If yes, which gene: \_\_\_\_\_  
If yes, 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: \_\_\_\_\_