



## Form 1b: HHT Genetic Test Requisition

### LAB USE ONLY DO NOT FILL OUT

Date received: Y \_\_\_\_\_ M \_\_\_\_\_ D \_\_\_\_\_

Specimen type: \_\_\_\_\_

Condition: \_\_\_\_\_

MRN: \_\_\_\_\_ Tech: \_\_\_\_\_

#### Patient

Last name: \_\_\_\_\_

First name: \_\_\_\_\_

Date of birth: Y \_\_\_\_\_ M \_\_\_\_\_ D \_\_\_\_\_

Gender:  Male  Female

Pregnant:  Yes  No

Expected delivery date: Y \_\_\_\_\_ M \_\_\_\_\_ D \_\_\_\_\_

#### Patient History

Confirmed clinical diagnosis

Suspected clinical diagnosis  Unaffected

Symptoms:

PAVM  Rare nose bleeds  Liver shunts

CAVM  Frequent nose bleeds  Stroke

Telangiectasia  Other: \_\_\_\_\_

#### Family History

Isolated case  Positive family history

Family previously tested:  Yes  No

Mutation identified:  Yes  No

*If mutation identified at lab other than Impact Genetics please provide report.*

Proband name (first person in a family to be studied):  
\_\_\_\_\_

Mutation: \_\_\_\_\_

#### Relationship To Proband

Proband  Parent of proband

Brother or sister of proband  Child of proband

Other: \_\_\_\_\_

#### Specimen Information

Sample:

Blood sample for DNA

Blood sample for RNA (at Impact Genetics' request)

DNA from blood  DNA from tumor

Fresh tumor  Frozen tumor

Buccal swab (only for preapproved familial mutation confirmation, contact lab directly before submitting)

Other: \_\_\_\_\_

Pre-natal:

Cord blood  CVS  Cultured amniocytes

Direct amniotic fluid  DNA extracted from CVS

DNA extracted from amniocytes

Date of collection: Y \_\_\_\_\_ M \_\_\_\_\_ D \_\_\_\_\_

Time of collection: HH:MM (24hr) \_\_\_\_\_

Sample to test maternal cell contamination

#### Referring Specialist

Name: \_\_\_\_\_

Specialty: \_\_\_\_\_

Contact: \_\_\_\_\_

Telephone: \_\_\_\_\_ Fax: \_\_\_\_\_

Email: \_\_\_\_\_

Signature: \_\_\_\_\_

Institution: \_\_\_\_\_

Address: \_\_\_\_\_  
\_\_\_\_\_

City: \_\_\_\_\_ Prov/State: \_\_\_\_\_

Postal code: \_\_\_\_\_ Country: \_\_\_\_\_

Additional copies to: \_\_\_\_\_

Email: \_\_\_\_\_ Fax: \_\_\_\_\_

#### Pedigree

*Please complete the following page for detailed family history information if available.*

#### Billing

*Impact Genetics tests ordered through LabCorp test menu. USA only.*

a) Client Bill

Provide details: \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

b) Patient Pay

*Complete LabCorp Financial Responsibility Form.*

c) Third party insurance (USA only)

*Complete Form 1c: U.S. Insurance Information and to expedite testing, complete LabCorp Financial Responsibility Form.*

1100 Bennett Road - Unit 4, Bowmanville, ON L1C 3K5

t: 647-478-4902 or 1-877-624-9769 f: 905-697-9786

e: info@impactgenetics.com *Please ensure to use secure email*



## HHT Genetic Test Requisition: Family History Information

Proband name: \_\_\_\_\_ Date of birth: Y \_\_\_\_\_ M \_\_\_\_\_ D \_\_\_\_\_

Enter family members starting with the proband's parents and then brothers/sisters and biological children.  
Use additional sheets as needed to provide information on additional affected family members.

### Name of Family Member Relationship to Proband

### Symptoms

Please check all applicable boxes

Name: \_\_\_\_\_

Relationship: \_\_\_\_\_

Date of birth: Y \_\_\_\_\_ M \_\_\_\_\_ D \_\_\_\_\_

Carries HHT mutation?

Yes  No  Unknown

PAVM

CAVM

Rare nose bleeds

Frequent nose bleeds

GI bleeding

Telangiacteses

Liver shunts

Stroke

Unaffected

Other (list below):  
\_\_\_\_\_

Name: \_\_\_\_\_

Relationship: \_\_\_\_\_

Date of birth: Y \_\_\_\_\_ M \_\_\_\_\_ D \_\_\_\_\_

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