

THIS IS NOT A TEST REQUEST FORM.
The information below is required to perform *BMP9*-related telangiectasia syndrome testing.
Please fill out this form and submit it with the test request form or electronic packing list.

PATIENT HISTORY FOR *BMP9*-RELATED TELANGIECTASIA SYNDROME TESTING

Patient Name _____ **Date of Birth** ____ / ____ / ____ **Gender** F M

Physician _____ **Physician Phone** (____) _____ **Practice Specialty** _____

Genetic Counselor _____ **Counselor Phone** (____) _____

Patient's Ethnicity (check all that apply)

- | | | | |
|---|---|--|--------------------------------------|
| <input type="checkbox"/> African American | <input type="checkbox"/> Ashkenazi Jewish | <input type="checkbox"/> Asian | <input type="checkbox"/> Caucasian |
| <input type="checkbox"/> Hispanic | <input type="checkbox"/> Middle Eastern | <input type="checkbox"/> Native American | <input type="checkbox"/> Other _____ |

Does the patient have SYMPTOMS? No Yes; please check all that apply

- Telangiectasia (locations and numbers): _____
- Nosebleeds (frequency): _____
- AVM(s) (locations): _____
- Other: _____

Does the patient have a FAMILY HISTORY of telangiectases or nosebleeds? No Yes Unknown

If yes, attach a PEDIGREE or specify the relatives RELATIONSHIP to the patient. List their symptoms:

Has DNA testing been performed for these family member(s)? No Yes Unknown

Please attach a copy of the relative's DNA laboratory result. (REQUIRED for familial mutation testing).

Has the patient undergone previous DNA testing? No Yes Unknown

If yes, please describe test(s) and results: _____

Circle the test you intend to order

2010015 Telangiectasia Syndrome (*BMP9/GDF2*) Sequencing – Clinical sensitivity ~1%. Order for individuals with negative *ACVRL1*, *ENG* and *SMAD4* testing who have multiple cutaneous telangiectasia accompanied by nosebleeds.

2009337 Hereditary Hemorrhagic Telangiectasia (HHT) Panel, Sequencing and Deletion/Duplication, 5 Genes (*ACVRL1*, *BMP9*, *ENG*, *RASAI*, *SMAD4*) – Preferred initial test to confirm a diagnosis of a telangiectasia/AVM disorder.

2001961 Familial Mutation, Targeted Sequencing- Tests for a *BMP9* sequence change identified in a family member; a copy of a relative's DNA laboratory result is **REQUIRED**.

For questions, contact an ARUP genetic counselor at (800) 242-2787, ext. 2141

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