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THIS IS NOT A TEST REQUEST FORM. Please complete and submit with the test request form or electronic packing list.

PATIENT HISTORY FOR JUVENILE POLYPOSIS SYNDROME (JPS) / HEREDITARY HEMORRHAGIC TELANGIECTASIA (HHT) TESTING

Patient Name:	Date of Birth:	Sex: ☐ Female ☐ Male
Ordering Provider:	Provider's Phone:	
Practice Specialty:		
Genetic Counselor.	Counselor Phone:	
Patient's Ethnicity/Ancestry (check all that apply)		
☐ African American/Black ☐ Asian ☐ Hispanic	☐ White ☐ Other: _	
List country of origin (if known):		
Does the patient have <u>symptoms</u> ?	□ No □	Yes (check all that apply and describe)
☐ Cerebral arteriovenous malformation (AVM)		
☐ GI carcinoma (location:		; age of diagnosis:)
☐ Hepatic AVM		
☐ Juvenile polyposis (number of polyps:; location(s):_		; age of onset:)
☐ Pulmonary AVM		
☐ Spinal AVM		
☐ Spontaneous, recurrent nosebleeds (frequency:)
☐ Telangiectasia (location(s):)
☐ Other symptom(s):		
Has the patient had an allogeneic bone marrow or umbilical	cord blood transplant	□ No □ Yes □ Unknown
Is there any relevant family history?	······································	□ No □ Yes □ Unknown
If yes, specify:	D JPS	☐ JP/HHT ☐ Neither ☐ Unknown
Attach a pedigree or specify the relative's $\underline{\text{relationship}}$ to the	patient. List their <u>sympton</u>	ns and age of onset:
Has DNA testing been performed for the family member(s)?. If yes, attach a copy of the relative's DNA laboratory result (E) Check the test you intend to order.		
☐ 2004992 Juvenile Polyposis (BMPR1A) Sequencing and D	el/Dup: Preferred test for o	confirmation of a clinical diagnosis of
or predictive testing for JPS. Clinical sensitivity		
□ 2009337 Hereditary Hemorrhagic Telangiectasia (HHT) Pa	nnel, Sequencing and Del/[Oup: Clinical sensitivity SMAD4: ~28%
for JPS, unknown for JP/HHT. ☐ 2001961 Familial Mutation, Targeted Sequencing: Tests f	or sequence	
variant(s) previously identified in a family memb		
lab result is REQUIRED.		
☐ 3003144 Deletion/Duplication Analysis by MPLA: Tests fo		Master Label
deletion/duplication previously identified in a fa	mily member; a copy	
of a relative's lab report is REQUIRED.		
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For questions, contact an ARUP gene	tic counselor at 800-242-2	2787 ext. 2141.