

Patient ID SA00062769	Patient Name TESTINGRNV, HHTM NORM	Birth Date 1980-01-21	Gender F	Age 33
Order Number SA00062769	Client Order Number SA00062769	Ordering Physician Client, Client	Report Notes	
Account Information C7028846 DLMP Rochester		Collected 24 Sep 2013 08:57		

ENG and ACVRL1, Large Del/Dup

ENG-ACVRL1, Large Del/Dup Result

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The familial ENG deletion of Exon 3–8 duplication was not detected in this individual.

ENG-ACVRL1, Large Del/Dup Interp

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The ENG deletion of Exon 3–8 duplication was previously identified in a family member with features of a hereditary hemorrhagic telangiectasia (HHT). Since this mutation was not detected in this individual, this suggests that this patient is at no greater risk than someone in the general population for having HHT.

ADDITIONAL INFORMATION

Multiplex-ligation dependent probe amplification (MLPA) was used to test for the presence of large genomic alterations in the ENG and ACVRL1 genes (GenBank numbers NM_001114753.1 and NM_000020.2, respectively). This PCR-based method utilizes probes for all 15 exons of the ENG gene, and all 10 exons of the ACVRL1 gene.

A genetic consultation may be of benefit.

CAUTIONS:

Rare polymorphisms exist that could lead to false negative or positive results. If results obtained do not match the clinical findings, additional testing should be considered.

Test results should be interpreted in the context of clinical findings, family history, and other laboratory data. Misinterpretation of results may occur if the information provided is inaccurate or incomplete.

If the patient has had an allogeneic blood or marrow transplant or a recent (i.e. less than 6 weeks from time of sample collection) heterologous blood transfusion these results may be inaccurate due to the presence of donor DNA.

ENG-ACVRL1, Large Del/Dup Rev by

MCR

Jamie Brufat

Received: 25 Sep 2013 14:01

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Performing Site Legend

Code	Laboratory	Address
MCR	Mayo Clinic Dept. of Lab Med and Pathology	200 First Street SW, Rochester, MN 55905