

# **Test Definition: VHLZZ**

Von Hippel Lindau Syndrome, VHL, Full Gene Analysis, Varies

Reporting Title: VHL Full Gene Analysis

Performing Location: Rochester

#### **Ordering Guidance:**

For patients suspected of having hereditary erythrocytosis or polycythemia, order HEMP / Hereditary Erythrocytosis Mutations, Whole Blood.

For a comprehensive hereditary cancer panel that includes the VHL gene, consider one of the following tests:

- -ENDCP / Hereditary Endocrine Cancer Panel, Varies
- -HPGLP / Hereditary Paraganglioma/Pheochromocytoma Panel, Varies
- -RENCP / Hereditary Renal Cancer Panel, Varies

Testing for VHL gene as part of a customized panel is available. For more information see CGPH / Custom Gene Panel, Hereditary, Next-Generation Sequencing, Varies.

Targeted testing for familial variants (also called site-specific or known mutations testing) is available for this gene. For more information see FMTT / Familial Variant, Targeted Testing, Varies. To obtain more information about this testing option, call 800-533-1710.

### **Shipping Instructions:**

Specimen preferred to arrive within 96 hours of collection.

### **Specimen Requirements:**

 $\textbf{Patient Preparation:} \ \textbf{A previous bone marrow transplant from an allogenic donor will interfere with testing.} \ \textbf{Call}$ 

800-533-1710 for instructions for testing patients who have received a bone marrow transplant.

Specimen Type: Whole blood

Container/Tube:

Preferred: Lavender top (EDTA) or yellow top (ACD)

Acceptable: Any anticoagulant Specimen Volume: 3 mL Collection Instructions:

- 1. Invert several times to mix blood.
- 2. Send whole blood specimen in original tube. Do not aliquot.

Specimen Stability Information: Ambient (preferred) 4 days/Refrigerated

### Forms:

- 1. **New York Clients-Informed consent is required.** Document on the request form or electronic order that a copy is on file. The following documents are available:
- -Informed Consent for Genetic Testing (T576)
- -Informed Consent for Genetic Testing (Spanish) (T826)
- 2. Molecular Genetics: Inherited Cancer Syndromes Patient Information Sheet (T519)
- 3. If not ordering electronically, complete, print, and send a Oncology Test Request (T729) with the specimen.

Specimen Type	Temperature	Time	Special Container
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# **Test Definition: VHLZZ**

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Varies Varies
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## **Result Codes:**

Result ID	Reporting Name	Туре	Unit	LOINC®
614875	Test Description	Alphanumeric		62364-5
614876	Specimen	Alphanumeric		31208-2
614877	Source	Alphanumeric		31208-2
614878	Result Summary	Alphanumeric		50397-9
614879	Result	Alphanumeric		82939-0
614880	Interpretation	Alphanumeric		69047-9
614881	Resources	Alphanumeric		99622-3
614882	Additional Information	Alphanumeric		48767-8
614883	Method	Alphanumeric		85069-3
614884	Genes Analyzed	Alphanumeric		48018-6
614885	Disclaimer	Alphanumeric		62364-5
614886	Released By	Alphanumeric		18771-6

LOINC® and CPT codes are provided by the performing laboratory.

## **Supplemental Report:**

Supplemental

### **CPT Code Information:**

81404

# **Reference Values:**

An interpretive report will be provided.