



Virant Diagnostics, Inc.

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 CAP #: 954036801, CLIA #: 21D2184276
www.virantdx.com

Genetics Test Requisition Form

Place Barcode Label Here

Specimen Information

Collection Date: ___/___/___
 Collection Time: ___:___ AM PM

PATIENT INFORMATION

Last Name:	First Name:	Date of Birth:
Email:	Phone:	Gender: <input type="checkbox"/> M <input type="checkbox"/> F
Address:	City: State:	Zip:

FAMILY HISTORY

Sample Pedigree: <input type="checkbox"/> Proband <input type="checkbox"/> Mother <input type="checkbox"/> Father <input type="checkbox"/> Siblings <input type="checkbox"/> Other:	HAE Type: <input type="checkbox"/> Type I <input type="checkbox"/> Type II <input type="checkbox"/> Unknown <input type="checkbox"/> NA Family history consistent with HAE type? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> No family history <input type="checkbox"/> NA Clinical history consistent with HAE? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> NA
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DIAGNOSIS CODES

<input type="checkbox"/> D84.1: Defects in the complement system	<input type="checkbox"/> D82.8: HAE with normal C1-INH	<input type="checkbox"/> T78.3XXA: Angioedema
<input type="checkbox"/> T78.2XXA: Anaphylaxis	<input type="checkbox"/> L29.8: Pruritus	<input type="checkbox"/> D89.9: Disorder of immune mechanism
<input type="checkbox"/> D84.9: Immunodeficiency, unspecified	<input type="checkbox"/> Other:	

PATIENT CONSENT/AUTHORIZATION

- I hereby authorize the release of medical information related to this service for submission of personal reports to my healthcare providers and insurance carriers.
- In addition, I agree to assume responsibility for payment of charges for laboratory services that are not covered by my healthcare insurer.
- I acknowledge that I have read and understood the Genetics Informed Consent Form at <https://virantdx.com/wp-content/uploads/2022/12/Genetics-Informed-Consent-Form-221212.pdf>.
- For family relative(s) providing sample(s) for Carrier Testing or **Whole Exome Sequencing**, patient consent/authorization has been completed on the Genetics Informed Consent Form.

Signature: _____ **Date:** ___/___/___

Printed Name:	Relationship: <input type="checkbox"/> Self <input type="checkbox"/> Parent <input type="checkbox"/> Legal Guardian <input type="checkbox"/> Durable Power of Attorney for Health Care	
Patient information (if not same as above):	First Name:	Date of Birth:
Last Name:		

INSURANCE AND PAYMENT INFORMATION

Bill Insurance (Attach copy of insurance card, front and back) Bill Client Bill Patient (Cash/Check/Credit Card) Other:

Primary Plan Name:	Policy Holder Name:
Policy #:	Group #:
Secondary Plan Name:	Policy Holder Name:
Policy #:	Group #:

HEALTHCARE PROVIDER INFORMATION

Provider Name:	NPI:
Organization:	Phone:
Address:	City: State: Zip:

Provider Authorization Signature: _____ **Date:** ___/___/___

SPECIMEN REQUIREMENTS

Specimen Type: Whole blood (EDTA, purple top tube) Saliva Buccal swab (Currently for research use ONLY)
 Specimen must be whole blood for the **Angioedema Panel** and **Whole Exome Sequencing**.

Specimen Process and Shipment Requirements:

- For **whole blood**, at least 3 cc must be sent to the laboratory on the day of collection at ambient temperature for overnight delivery. Whole blood samples must be stored at 4°C until shipment. Please contact us at (877) 888-2973 or info@virantdx.com to obtain overnight shipping labels.
- For **saliva**, collect and store samples according to the collection kit manufacturer's instructions. Saliva samples must be sent to the laboratory on the day of collection at ambient temperature for overnight delivery. Please contact us at (877) 888-2973 or info@virantdx.com to obtain overnight shipping labels.

PLEASE SEE THE NEXT PAGE FOR THE GENETIC TEST MENU

GENETIC TESTS			
C1INH Deficiency	Exon(s)	Region Sequenced	
<input type="checkbox"/> SERPING1 gene	exons 2-8	coding ± 10bp	
nC1INH Genes	Exon(s)	Screen for Known HAE Variant(s)	
<input type="checkbox"/> F12 gene	exon 9	c.983C>A	p.Thr328Lys
		c.983C>G	p.Thr328Arg
		c.971_1018 +24del72	
		c.892_909dup	
<input type="checkbox"/> KNG1 gene	exon 10	c.1136T>A	p.Met379Lys
<input type="checkbox"/> ANGPT1 gene	exon 2	c.807G>T	p.Ala119Ser
<input type="checkbox"/> PLG gene	exon 9	c.988A>G	p.Lys330Glu
	exon 18	c.2183T>A	p.Val728Glu
<input type="checkbox"/> MYOF gene	exon 7	c.651G>T	p.Arg217Ser
<input type="checkbox"/> HS3ST6 gene	exon 2	c.430A>T	p.Thr144Ser
<input type="checkbox"/> nC1INH Hereditary Angioedema (HAE) Panel (all 6 nC1INH genes)			
<input type="checkbox"/> Hereditary Angioedema (HAE) Panel (all 6 nC1INH genes and SERPING1 gene)			
<input type="checkbox"/> Angioedema Panel (77 genes involved in the complement, coagulation, and tissue kallikrein pathways) Specimen must be whole blood. Please visit https://virantdx.com/testing-solutions/genetic-testing/comprehensive-angioedema-gene-panel/ for more information.			
<input type="checkbox"/> Whole Exome Sequencing (22,000 genes) Specimen must be whole blood. Must provide clinical information: https://virantdx.com/wp-content/uploads/2022/12/Genetics-Clinical-Information-Form-221212.pdf . Providing maternal and paternal (or other family relative) samples in addition to the proband sample is helpful for result interpretation.			
<input type="checkbox"/> MCAS Panel (Alpha-tryptasemia Copy Number Variation and KIT D816V Mutation Hotspot) (Currently for research use ONLY)			
<input type="checkbox"/> Alpha-tryptasemia Copy Number Variation (Currently for research use ONLY)			
<input type="checkbox"/> KIT D816V Mutation Hotspot (Currently for research use ONLY)			
<input type="checkbox"/> Primary Immunodeficiency Panel (Currently for research use ONLY)			
TARGETED VARIANT TESTING			
<input type="checkbox"/> Known Familial Variant(s)			
<input type="checkbox"/> Confirmation of Variant Identified in Research Lab			
Proband Name:		Relationship to Proband:	
Variant Information:			
Gene:	Transcript (NM#):	Coding Position (c.):	Protein (p.):

Please visit our webpage at <https://virantdx.com/testing-solutions/genetic-testing/> and contact us at angioedemalab@virantdx.com or (877) 888-2973 for any inquiries.