

**Virant Diagnostics, Inc.**

11002 Veirs Mill Rd, Suite 404
 Wheaton, MD 20902
 Phone: (877) 888-2973
 Fax: (888) 314-3456
 CLIA #: 21D2184276
 www.virantdx.com

Angioedema Complement Assays and Molecular Diagnosis Requisition Form

Place Barcode Label Here

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 Family history consistent with HAE type? <input type="checkbox"/> Yes <input type="checkbox"/> No Clinical history consistent with HAE? <input type="checkbox"/> Yes <input type="checkbox"/> No
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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"/> L29.8: Pruritus
<input type="checkbox"/> L29.8: Pruritus	<input type="checkbox"/> L29.8: Pruritus	<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.
- For whole exome sequencing, I acknowledge that I have read and understood the detailed consent information on:
www.virantdx.com/testing-solutions/genetics-solution/.

Patient Signature: _____ Date: ___ / ___ / ___

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: ___ / ___ / ___

TEST MENU**ANGIOEDEMA COMPLEMENT AND OTHER ASSAYS**

Specimen Type: Whole blood (Sodium citrate, light blue top tube)

Specimen Process and Shipment Requirements: 3 cc of whole blood must be centrifuged, and the plasma must be transferred to a plastic transport tube. The tube must be kept in the refrigerator at 4 °C and then shipped to the laboratory with a cold pack AND SAMPLES FOR GENETIC TESTING IF ORDERED.

<input type="checkbox"/> Comprehensive C1 INH function (C1s, Factor XII, and Plasma Kallikrein)	<input type="checkbox"/> Chromogenic C1 INH function
<input type="checkbox"/> Complement C1q (Currently unavailable)	<input type="checkbox"/> Complement C4 (Currently unavailable)
<input type="checkbox"/> Bradykinin 1-5 (Currently only for research)	

SEE REVERSE FOR MOLECULAR TESTING INFORMATION**FOR LABORATORY USE ONLY**

Accession #:	Patient ID:
Client/Physician ID:	Date Received: ___ / ___ / ___ Time Received: ___ : ___ <input type="checkbox"/> AM <input type="checkbox"/> PM

MOLECULAR DIAGNOSTIC TESTING			
Specimen Type: <input type="checkbox"/> Whole blood (EDTA, purple top tube) <input type="checkbox"/> Buccal swab <input type="checkbox"/> Saliva			
Specimen Requirements: For whole blood, 3 cc must be sent to the laboratory the day of collection WITH SAMPLES FOR COMPLEMENT TESTING IF ORDERED.			
C1 INH Deficiency	Exon(s)	Region Sequenced	
<input type="checkbox"/> SERPING1	exons 2-8	coding ± 10bp	
nC1INH Gene	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
<input type="checkbox"/> KNG1 gene	exon 10	c.1136T>A	p.Met379Lys
		c.1720C>G	p.Pro574Ala
<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
	exon 25	c.2576delG	p.Gly859Glufs*8
<input type="checkbox"/> HS3ST6 gene	exon 2	c.430A>T	p.Thr144Ser
<input type="checkbox"/> nC1INH Angioedema Panel (all 6 nC1INH genes)			
<input type="checkbox"/> Angioedema Panel (all 6 nC1INH genes plus SERPING1 gene)			
<input type="checkbox"/> Comprehensive Angioedema Panel (77 genes involved in the complement, coagulation, and tissue kallikrein pathways)		Please visit www.virantdx.com/testing-solutions/genetics-solution/ for more information.	
<input type="checkbox"/> Whole Exome Sequencing (22,000 genes)		Please visit www.virantdx.com/testing-solutions/genetics-solution/ for more information.	
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.):
FOR LABORATORY USE ONLY			
Accession #:		Patient ID:	
Client/Physician ID:	Date Received: ___/___/___	Time Received: ___:___	<input type="checkbox"/> AM <input type="checkbox"/> PM

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

To place a test order online, please visit our laboratory information system's client portal at: virant.stratusdx.net/