

Specific Site Analysis Test Requisition Form - Page 1 of 2 (Known Familial Alteration Analysis)

COMPLETE ENTIRE FORM TO AVOID DELAYS

PATIENT INFORMATION	То	submit an	order via email, please s	send the	e comp	oleted test requis	ition fo	orm to	info@ambrygen.com	
Name (Last, First, MI)			<u>, </u>	Sex at	Birth	Date of Birth (MM/DD,	YY)	MRN		
				□F [□М					
Ethnicity: Asian Ashkenazi Jewish Black/African American White French Canadian/Cajun Hispanic/Latino Mediterranean Middle Eastern Native American Pacific Islander Portuguese Unknown Other:										
Address			City				State		Zip	
									·	
Di		F 11					ъ (I D:II:		
Phone		Email						ed Billing		
							∐Insu	rance L	Self-pay □Institutional	
SPECIMEN INFORMATION*	(For phlebotomy service, select a	all services you a	re requesting)							
Type(s) ☐ Blood (EDTA preferred)	☐ Saliva ☐ Cord Blood** ☐ D	NA, Source:	☐ Other**:	☐ Per	rsonal hi	story of allogenic bone	marrow	or perip	heral stem cell transplant*	
Collection Date	Specimen ID [☐ Send saliva I	kit to patient	N	Лedical F	Record #				
PRENATAL SAMPLES ONLY	k*									
FRENATAL SAMFLES ONET										
Sample type: Direct CVS Cultured CVS Cultured amnio POC Cultured POC Gestational age at sample collection							ample collection			
* Blood/saliva from patients with a history of allogenic bone marrow or stem cell transplant cannot be used for genetic testing. Blood/saliva from patients with active hematological disease is not										
recommended. An alternative specimen may be needed. See ambrygen.com/specimen-requirements for details. ** Fetal specimens, cord blood and POC will have maternal cell contamination studies added for a charge. Maternal and fetal specimen required. Please see bottom of page 2 for Maternal Cell Contamination sample submission test codes.										
Phlebotomy Services Request: Phle	ebotomy draw Send blood kit	to patient \(\subseteq \)	Send saliva kit to patient							
As the patient's clinician, I am unaware of any potential for complication or difficulty in drawing blood for the listed patient(s). I understand that the phlebotomist has full authority to refuse to draw any patient if the safety of the phlebotomist and/or patient(s) are in question.										
INDICATION(S) FOR TESTIN	NG									
ICD-10 code(s):			Testing could aid in syste	mic therap	py and/o	or surgical decision-m	aking for	my affe	cted patient ☐ Yes ☐ No	
ORDERING LICENSED PROV	IDER/SENDING FACILIT	「Y (Each listed	person will receive a copy of	the report	t)					
Facility Name (Facility Code)	Address		City		<u>'</u>	State / Country 2	Zip		Phone	
Ordering Licensed Provider Name (Li	ast, First)(Code)	NPI#	Phone		Fax/	'Email				
ADDITIONAL RESULTS RECI	DIENTS									
Genetic Counselor or Other Medical		ode)	Phone/Fax/Email							
Genetic counselor of other Medical	Trovider Hume (East, First) (C	ouc)	Thomey raxy Email							
Genetic Counselor or Other Medical	Provider Name (Last, First) (C	ode)	Phone/Fax/Email							
CONFIDMATION OF INFORME	D CONCENT DDE TECT C	ENIETIC COLL	NCELING AND MEDICAL	NECECC	·ITV 50	D CENETIC TECT				
The undersigned person (or represer	·		·					nationt h	ass diven appropriate	
consent.l confirm that testing is med										
genetic counseling services by a third applies to the attached letter of mediapplies.		the patient's in	surance provider. Furthermore	, all inforn	nation o	n this TRF is true to th	e best o	f my kno	owledge. My signature	
applies to the attached letter of med	ical flecessity.									
Signature Required for Processing	Medical Professional Sig	nature:					D	ate:		
■ INSURANCE BILLING (Inclu	ide copy of both sides of insura	ance card)				INSTITUTIONAL	BILLI	NG		
Patient Relation to Policy Holder?	Name and DOB of				Fa	icility Name	☐ Send	d invoice	to facility address above	
☐ Self ☐ Spouse ☐ Child	Policy Holder (if not self)		LIMO			1.1				
Insurance Company	Policy #		HMO Auth #		A	ddress				
Out of Pocket:	<u> </u>		•			ontact Name				
In many cases, we will start testing immediately (may vary by test or insurance provider). We will attempt to contact the patient if the estimated out-of-pocket costs are > USD \$100.										
Special Billing Notes:	•					none Number		E-mail/	'Fax	
Special billing ractes.										
						PATIENT PAYM	ENT		(Payable to Ambry Genetics) t Card (Call 949-900-5795)	
Patient Acknowledgement: I acknowledge										
(Ambry), authorize Ambry to release medical information concerning my testing to my insurer, to be my designated representative for purposes of appealing any denial of benefits as needed and to request additional medical records for this purpose. I understand that I am financially responsible for any amounts not covered by my insurer and responsible for sending Ambry money received from my health insurance company.										
For patient payment by credit card: I hereby authorize Ambry Genetics Corporation to bill my credit card as indicated above. In order to expedite consideration for eligibility for Ambry's Patient Assistance Program,										
please provide the total annual gross hou					y the liste	ed income:	I auth	orize Am	bry Genetics Corporation to	
verify the above information for the sole property.	purpose or assessing financial nee	u, including the r	ignit to seek supporting document	ation.						
☐ I am a New York resident and I give Ambry Genetics permission to store my sample for longer than 60 days. NOTE: If left blank, consent is interpreted as "NO".										
Signature Required For Insurance/	<u> </u>			,					Date:	



Patient Name: [DOB:
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Specific Site Analysis Test Requisition Form - Page 2 of 2

SPECIFIC SITE ANALYSIS (5555)										
Positive control: ☐ Sent ☐ To be sent ☐ Not available ☐ Available at Ambry, accession #:										
The following will be requested when ordering known mutation analysis for a mutation identified in an outside laboratory:		ALTERATION TO BE TESTED								
		Gene 1	,	Alteration 1						
1. Proband report (mandatory)										
2. Positive control (recommended; required for prenatal testing)		Gene 2	,	Alteration 2						
ACMG guidelines, CAP and CLIA regulatory provisions recommend use of a positive cor										
to provide evidence of amplification when interrogating a specific sequence alteration. It recommended that individuals for a known genotype for the locus tested be included as		Gene 3	,	Alteration 3						
control to ensure assay performance.		Gene 4	,	Alteration 4						
PATIENT CLINICAL INFORMATION										
☐ Healthy ☐ Affected/Symptomatic, age at diagnosis:										
Please list relevant clinical findings with ICD-10 codes:										
PREVIOUS TEST HISTORY (Please include copy of test results if performed at another laboratory)										
			Testing Lab							
Previously Detected Alteration(s) Gene		e Name		resulting Edus						
Patient previously tested at Ambry?										
Name	Date of Birth (MM/DD/YY) Relation									
FOR PRENATAL SPECIMENS, POC OR CORD BLOOD: MATERNAL CELL CONTAMINATION ANALYSIS REQUIRED										
Both test codes required for fetal specimens										
□ 1260 MCC for fetal specimen or cord blood □ 1262 MCC Reference for maternal blood sample (No Charge)										