

COMPLETE TO SUBMIT FAMILY MEMBERS FOR EXOME ORDERS

All family member specimens must be received within 4 weeks of order to be included in analysis

To submit an order via email, please send the completed test requisition form to info@ambrygen.com

COLLECTION DATE (REQUIRED)
<p>If date of collection is not provided, three calendar days before specimen receipt will be used (for specimens stored longer than 30 days, the day of archive retrieval will be used as the date of service)</p>

PATIENT INFORMATION			
Legal Name (Last, First, MI)	Date of Birth (MM/DD/YY)	Sex Assigned at Birth <input type="checkbox"/> F <input type="checkbox"/> M	Gender (optional) <input type="checkbox"/> Man <input type="checkbox"/> Woman <input type="checkbox"/> Nonbinary <input type="checkbox"/> Self-described
Genetic Ancestry: <input type="checkbox"/> Ashkenazi Jewish <input type="checkbox"/> Asian <input type="checkbox"/> Black/African American <input type="checkbox"/> French Canadian/Cajun <input type="checkbox"/> Hispanic/Latino <input type="checkbox"/> Mediterranean <input type="checkbox"/> Middle Eastern <input type="checkbox"/> Native American <input type="checkbox"/> Pacific Islander <input type="checkbox"/> Portuguese <input type="checkbox"/> White <input type="checkbox"/> Unknown <input type="checkbox"/> Other:			MRN
Address		City	State Zip
Mobile #	Email		Preferred Billing <input type="checkbox"/> Insurance <input type="checkbox"/> Self-pay <input type="checkbox"/> Institutional

SPECIMEN INFORMATION (Please see ambrygen.com/specimen-requirements for details)	
<input type="checkbox"/> Personal history of allogenic bone marrow or peripheral stem cell transplant	<input type="checkbox"/> Current diagnosis of heme malignancy, Type:
Specimen ID	Medical Record #
Collection Assistance: <input type="checkbox"/> Phlebotomy draw** <input type="checkbox"/> Insurance preverification first <input type="checkbox"/> Send saliva kit to patient <input type="checkbox"/> Send buccal kit to patient <small>**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.</small>	

CLINICAL INFORMATION
Is family member affected with the same phenotype as the proband? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Partially <input type="checkbox"/> Possibly, describe: _____

TEST MENU	
<input type="checkbox"/> Family member for ExomeNext® (no charge) <input type="checkbox"/> Family member for ExomeNext-Rapid® (no charge) <input type="checkbox"/> Other _____ (Test Code/Test Name)	Proband Name: _____ Relationship to proband: _____

SECONDARY FINDINGS
Secondary findings results are only available for each family member who is fully sequenced as part of the duo/trio. Check "decline" to opt-out of the ACMG Recommended List of secondary findings. If left unchecked, secondary findings will be reported for this family member. <input type="checkbox"/> Decline

ORDERING PHYSICIAN/SENDING FACILITY (Each listed person will receive a copy of the report)										
<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <th style="width: 30%;">Facility Name (Facility Code)</th> <th style="width: 30%;">Address</th> <th style="width: 15%;">City</th> <th style="width: 15%;">State /Country</th> <th style="width: 10%;">Zip</th> <th style="width: 10%;">Phone</th> </tr> </table> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <th style="width: 40%;">Ordering Licensed Provider Name (Last, First)(Code)</th> <th style="width: 15%;">NPI#</th> <th style="width: 15%;">Phone</th> <th style="width: 30%;">Fax/Email</th> </tr> </table>	Facility Name (Facility Code)	Address	City	State /Country	Zip	Phone	Ordering Licensed Provider Name (Last, First)(Code)	NPI#	Phone	Fax/Email
Facility Name (Facility Code)	Address	City	State /Country	Zip	Phone					
Ordering Licensed Provider Name (Last, First)(Code)	NPI#	Phone	Fax/Email							

ADDITIONAL RESULTS RECIPIENTS		
<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <th style="width: 70%;">Genetic Counselor or Other Medical Provider Name (Last, First) (Code)</th> <th style="width: 30%;">Phone/Fax/Email</th> </tr> </table>	Genetic Counselor or Other Medical Provider Name (Last, First) (Code)	Phone/Fax/Email
Genetic Counselor or Other Medical Provider Name (Last, First) (Code)	Phone/Fax/Email	

CONFIRMATION OF INFORMED CONSENT, PRE-TEST GENETIC COUNSELING, AND MEDICAL NECESSITY FOR GENETIC TESTING

I confirm that the genetic test ordered is medically appropriate. All information on this TRF is true to the best of my knowledge. I also confirm that the patient has consented to proceed with genetic testing, including the transfer and processing of their sample and personal/sensitive information in the United States. I agree to allow Ambry Genetics to facilitate the provision of pre-test genetic counseling services by a third-party service, as required by the patient's insurance provider.

<i>Signature Required for Processing</i> Medical Professional Signature: _____	Date: _____
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Family Member Acknowledgement: I affirm that the medical professional listed above has offered genetic counseling and has reviewed with me the exome sequencing process prior to testing, and I would like to proceed with test processing. I understand that the exome testing is being performed in order to assist analysis for my family member (proband), that a primary report will only be generated for the proband, and that it may be possible to infer information about my results based on the proband's report.

I agree to be contacted regarding future research studies for which I may be a candidate. Any future research projects will be subject to a separate informed consent process and participation is voluntary. Learn more about Ambry's privacy practices at <https://www.ambrygen.com/legal/notice-of-privacy-practices>.

For NY Residents:

I understand that New York State law requires Ambry Genetics to destroy my sample at the end of the testing process or not more than sixty days after the sample was taken. By checking this box, I agree that Ambry Genetics will instead retain my sample for at least 6 months after the testing above has been completed, and may (a) retain and use samples and health information for an indefinite period of time in accordance with applicable law; and (b) de-identify such samples and information and use and share the resulting de-identified samples and information in accordance with applicable law.

If family member signature is not completed below, the medical professional listed above affirms the family member has given consent for genetic testing to be performed and the signed consent form is on file.

Family Member/Guardian Signature: _____	Date: _____
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For all exome orders, Ambry includes testing for co-segregation analysis (aka: family testing for candidate alterations) if samples are sent before testing begins.

Supplemental Information

Electronic ordering is also available via AmbryPort®. Please visit <https://portal.ambrygen.com/login> to log in to the online portal.

Specimen Requirements

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.

Buccal swab samples from patients with a history of allogenic bone marrow or stem cell transplant should not be used for genetic testing. For these patients, an alternative specimen (e.g. cultured fibroblasts) is required. Testing on buccal swab samples from patients with active hematological disease is not recommended. An alternative specimen (e.g. cultured fibroblasts) is recommended. Please see ambrygen.com/specimen-requirements for details.