

GENETIC TESTING PROGRAM REQUISITION FORM

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Promotion/Contract Code:

CLIA# 99D2092375, CAP# 9257331

See test codes and detailed descriptions on tests on blueprintgenetics.com

REQUIRED FIELDS ARE MARKED WITH AN ASTERISK (*)

*TEST CODE

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TEST INFORMATION

* Test Name:

All tests include analysis of both small exonic and splice site variations, and large deletions and insertions.

Sample type ⁱⁱ⁾ : <input type="checkbox"/> Blood <input type="checkbox"/> Saliva <input type="checkbox"/> DNA, source:	Sample Collection Date:
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ii) Please note that this information affects interpretation for mitochondrial DNA testing.
 More information about sample requirements on blueprintgenetics.com/sample-requirements.

PROGRAM-SPECIFIC INFORMATION

<p>Eligibility Criteria</p> <p>Eligibility for this program is a current or prior clinical diagnosis of auditory neuropathy, or a medical history consistent with auditory neuropathy. Auditory neuropathy (AN) is a hearing disorder characterized by an absent or abnormal auditory brainstem response (ABR) with preservation of otoacoustic emissions (OAEs) or cochlear microphonics (CMs).</p> <p>Question 1:</p> <p>1a. OAE or CM present with absent or abnormal ABR: Yes No</p> <p>1b. OAE or CM previously present with absent or abnormal ABR: Yes No</p> <p>Question 2:</p> <p>Patient does not have evidence of syndromic medical history: Yes No</p>	<p>Genetic Counseling</p> <p>Select one of the options</p> <p style="text-align: center;">My institute will provide genetic counseling for the patient</p> <p style="text-align: center;">I request for my patient post-test genetic counseling service offered by the genetic testing program</p>
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ORDERING HEALTH CARE PROFESSIONAL INFORMATION

* Name and Full Address:	* Institution:	
	* Email:	
	* NPI# (REQUIRED, US only):	
	Phone:	Fax:

Delivery of results Mail Fax Nucleus Results will always be available on our online reporting system at nucleus.blueprintgenetics.com.

SHARE RESULTS WITH

Name:		Role/Title:	
Email:		Street Address:	
City:	State:	Zip/Post Code:	Country:
Phone:	Fax:	Mail Results	Fax Results Nucleus

Results can be shared within the same hospital on our ordering portal, Nucleus.

PATIENT INFORMATION

* First Name:		* Last Name:	
* DOB: Year / Month / Day		MRN/SSN:	
Street Address:			
City:	State:	Zip/Post Code:	Country:
Phone:	Email:		

PATIENT HISTORY

* Sex: Male <input type="checkbox"/> Female <input type="checkbox"/> Unknown/uncertain <input type="checkbox"/>	Ethnicity:	ICD-10 Codes:
* Has the Patient Received a Hematopoietic Stem Cell Transplantation? Yes No		
* Has the Patient Received Granulocyte Transfusions in the Past Two Weeks? Yes No		
* Describe the Relevant Clinical Findings Supporting the Diagnosis (attach possible supportive material such as ECG):		
Family History (attach pedigree if available):	Previous Genetic Testing Results:	

VARIANT SPECIFIC TESTING INFORMATION

REQUIRED FIELDS ARE MARKED WITH AN ASTERISK (*)

Please select either Familial Variant Testing or Targeted Variant Testing

Familial Variant Testing Select this test when you want to test your patient for a variant that has been found in one of their relatives.	Targeted Variant Testing Select this test for founder mutation testing, confirmation of research results or clarifications of variant interpretation from another laboratory.								
* Index Patient's Order ID: * Index Patient's Subject ID:	* Indication for testing: Confirmation of research results Clarifying interpretation Founder/common mutation Other: Study								
* Indication for testing <table border="0"> <tr> <td>Diagnostic</td> <td>Predictive</td> <td>Carrier</td> <td>Segregation</td> </tr> <tr> <td colspan="4">Other: Study</td> </tr> </table>		Diagnostic	Predictive	Carrier	Segregation	Other: Study			
Diagnostic		Predictive	Carrier	Segregation					
Other: Study									
* Complete the following sentence to explain the relationship between the person being tested and the index patient. The person being tested is the index patient's: (e.g. son, daughter, brother, sister, mother, father)									

VARIANTS TO BE TESTED

	*Gene: (e.g. LMNA)	*Transcript: (e.g. NM_170707.3)	*cDNA change: (e.g. c.4375C>T or c.612_615del)	*Protein change: (e.g. Arg190Gln)
*Variant 1:				
Variant 2:				
Variant 3:				

*** BILLING INFORMATION**

INSTITUTIONAL BILLING	
Facility Name:	
Street Address:	City:
	State:
	Zip/Post Code:
Country:	Contact Person:
Phone:	Email:

GENERAL TERMS

By placing the order the Customer accepts Blueprint Genetics' General Terms. Blueprint Genetics reserves the right to amend its General Terms, of which the latest version shall always be applied. The latest version can be found at <https://blueprintgenetics.com/general-terms/>

ORDERING HEALTH CARE PROFESSIONAL SIGNATURE

I have discussed and obtained the Genetic Testing Program Informed Consent with the patient or their legal guardian and obtained any other consent from the patient that is required under the laws of my country/Province and local laws. I will maintain a record of the consent(s) and promptly notify Blueprint Genetics in writing of any changes, including revocation, of a consent. I certify that the patient is eligible and suitable for genetic testing services and/or genetic counseling I am ordering under the program. I agree I will not bill the patient or their insurance for the genetic testing services and/or genetic counseling offered as part of this program. I authorize Blueprint Genetics, and others working with Blueprint Genetics to contact me by mail, email or phone to inform me about ongoing clinical trials, clinical studies, services and products that might be relevant to the patient. I understand that I may revoke this authorization by contacting Blueprint Genetics Client Services.	
* Signature:	* Date:
* Name:	

HEALTH CARE PROFESSIONAL SIGNATURE REQUIRED FOR PROCESSING