



Instructions: The accurate interpretation and reporting of genetic results is contingent upon the reason for testing, clinical information, family history, and ancestry. To help provide the best possible service, supply the information requested below and **send this paperwork with the specimen or return by fax to Mayo Clinic Laboratories, Attn: Genetic Counselors at 507-284-1759. Phone: 800-533-1710 / International clients: +1-507-266-5700 or email MLIINT@mayo.edu**

Patient Information (required)

Patient Name (Last, First Middle)		Birth Date (mm-dd-yyyy)
Sex Assigned at Birth <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Unknown <input type="checkbox"/> Choose not to disclose	Legal/Administrative Sex <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Nonbinary	

Referring Healthcare Professional Information

Requesting Healthcare Professional Name (Last, First)	Phone	Fax*
Genetic Counselor Name (Last, First)	Phone	Fax*

*Fax number given must be from a fax machine that complies with applicable HIPAA regulations.

Reason for Testing

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Clinical Information

Age of onset: _____	Exposure to aminoglycoside antibiotics (eg, gentamicin, tobramycin, amikacin): <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown
Temporal bone abnormalities: <input type="checkbox"/> Yes <input type="checkbox"/> No	If "Yes," specify: _____
Type of hearing loss; check all that apply:	
<input type="checkbox"/> Sensorineural	<input type="checkbox"/> Conductive
<input type="checkbox"/> Stable	<input type="checkbox"/> Progressive
<input type="checkbox"/> Bilateral	<input type="checkbox"/> Unilateral
<input type="checkbox"/> Auditory neuropathy/dyssynchrony	<input type="checkbox"/> Mixed
<input type="checkbox"/> Fluctuating	<input type="checkbox"/> Unknown
Syndrome(s) suspected: <input type="checkbox"/> Yes <input type="checkbox"/> No	If "Yes," specify: _____
Other clinical features; list all relevant clinical symptoms, attach clinic note:	
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Audiogram; describe results, attach audiogram:	
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Family History

Pedigree; draw pedigree below or attach pedigree:		Pedigree Key <input type="radio"/> Female <input checked="" type="radio"/> <input checked="" type="radio"/> Affected <input type="radio"/> Male <input checked="" type="radio"/> <input checked="" type="radio"/> Carrier
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Paternal ancestry: _____	Maternal ancestry: _____	Consanguinity: <input type="checkbox"/> Yes <input type="checkbox"/> No
Are other relatives known to be affected?	<input type="checkbox"/> Yes <input type="checkbox"/> No	If "Yes," indicate relationship to patient: _____
Are other relatives known to be carriers?	<input type="checkbox"/> Yes <input type="checkbox"/> No	If "Yes," indicate relationship to patient: _____
Have other relatives had molecular genetic testing?	<input type="checkbox"/> Yes <input type="checkbox"/> No	If "Yes," complete the information below:
Gene: _____		
Name of individual tested (Last, First Middle): _____		
Birth date of individual tested (mm-dd-yyyy): _____		
Mutations/Variants: _____		
Laboratory at which testing was performed: _____		