

LABORATORY OF GENETICS AND GENOMICS

For local courier service and/or inquiries, please contact 513-636-4474 • Fax: 513-636-4373 www.cincinnatichildrens.org/moleculargenetics • Email: moleculargenetics@cchmc.org

Mailing Address:

3333 Burnet Avenue, Room R1042 Cincinnati, OH 45229

CUSTOM GENE SEQUENCING OR DELETION/DUPLICATION ASSAY REQUISITION

All Information Must Be Completed Before Sample Can Be Processed					
PATIENT INFORMATION	ETHNIC/RACIAL BACKGROUND (Choose All)				
Patient Name:,,,	□ European American (White) □ African-American (Black) □ Native American or Alaskan □ Asian-American □ Pacific Islander □ Ashkenazi Jewish ancestry □ Latino-Hispanic □				
Home Phone: MR# Date of Birth// Gender: □ Male □ Female BILLING INFORMATION (Choc	(specify country/region of origin) ☐ Other				
□ REFERRING INSTITUTION Institution:	Insurance can only be billed if requested at the time of service. Policy Holder Name:				
SAMPLE/SPECIMEN INFORMATION SPECIMEN TYPE: Amniotic fluid Blood Bone marrow	REFERRING PHYSICIAN Physician Name (print):				
□ Cord blood □ CVS □ Cytobrushes □ DNA □ Saliva □ Tissue (specify):	Address:				

 $\hfill\square$ Patient signed completed ABN

Medical Necessity Regulations: At the government's request, the Molecular Genetics Laboratories would like to remind all physicians that when ordering tests that will be paid under federal health care programs, including Medicare and Medicaid programs, that these programs will pay only for those tests the relevant program deems to be (1) included as covered services, (2) reasonable, (3) medically necessary for the treatment and diagnosis of the patient, and (4) not for screening purposes.

6	Cincinnati Children's
	changing the outcome together

Patient Name:	Date of Birth:	

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Reason for Testing:

☐ Mutation detection in suspected affected patient

☐ Carrier testing

☐ Pre-symptomatic diagnosis of at-risk relative

☐ Prenatal testing (by previous arrangement only)

Please call 513-636-4474 to discuss any prenatal testing with a genetic counselor prior to shipment.

PEDIGREE OR FAMILY HISTORY

Parental Consanguinity \square Y \square N

TEST(S) REQUESTED

CUSTOM GENE SEQUENCING

Gene(s) to be sequenced (specify): ___

Only genes with clear published functional relationship to rare diseases are accepted.

Suspected syndrome/ condition: _

Please choose one of the following:

☐ Full gene(s) sequencing

 \square Full gene(s) sequencing with reflex to deletion and duplication analysis, if indicated (please see list of genes available for del/dup at

www.cincinnatichildrens.org/deldup)

☐ Familial mutation analysis

Proband's name: _ Proband's DOB: __

Proband's mutation: ____

Patient's relation to proband: __

If testing was not performed at Cincinnati Children's, please include proband's report and at least 100ng of proband's DNA to use as a positive control.

DELETION AND DUPLICATION ASSAY

Gene(s) to be analyzed (specify): ___

Please see list of available genes at: www.cincinnatichildrens.org/deldup

Suspected syndrome/ condition: _

Please choose one of the following:

- ☐ Deletion and duplication analysis of gene(s) specified above
- \square Deletion and duplication analysis of gene(s) specified above with reflex to sequencing, if indicated
- $\hfill\square$ Analysis of gene(s) specified above from previously analyzed deletion and duplication
- ☐ Familial deletion analysis

Proband's name: ___

Proband's DOB: ___

Proband's mutation: ____

Patient's relation to proband: ___

If testing was not performed at Cincinnati Children's, please include proband's report and at least 100ng of proband's DNA to use as a positive control.

CLINICAL HISTORY

Symptoms:	Medical procedures:
Laboratory tests and results:	Medical imaging tests and results:
Previous genetic tests and results:	Other non-genetic diagnostics tests and results: