



FAILURE TO COMPLETE MAY DELAY RESULTS

Patient's Last Name		First	Middle	Birth date (required)	Sex
Outside Patient Number	Outside Specimen Number		Send Report To:		
Ordering Provider			Address:		
Provider Phone Number	DIAGNOSIS / ICD CODE:		Phone:	Fax:	

IMPORTANT INFORMATION REGARDING BILLING AND MEDICAL NECESSITY ON BACK

FAX ADDITIONAL RESULTS TO: NAME (please print): _____ FAX #: _____	HEALTHCARE PROFESSIONAL TO CALL FOR INFO/ABNORMAL RESULTS: NAME (please print): _____ PHONE #: _____
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SPECIMEN INFORMATION

Date collected: ____ / ____ / ____ Specimen Type: EDTA Cultured Skin Fibroblasts Banked DNA @ SCH, date: _____
 Time collected: _____ ACD Extracted DNA Other: _____

Also acceptable for DNA banking
 Cultured Amniocytes
 Cultured CVS

PATIENT/FAMILY HISTORY REQUIRED FOR ALL INDIVIDUALS TESTED

PEDIGREE DIAGRAM (ONLY ONE NEEDED PER FAMILY)

REASON FOR STUDY: Diagnostic (affected) Diagnostic (not affected)
 Carrier Testing (affected family member) Carrier Testing (no family history)
CLINICAL FINDINGS, PREVIOUS RELEVANT GENETICS TEST RESULT(S), FAMILY HISTORY: _____

PATIENT PREGNANT? NO YES LMP: _____ EDC: _____
ETHNICITY (check all that apply): African American Ashkenazi Jewish
 Asian European Caucasian Hispanic Native American Indian/Alaskan
 Pacific Islander Other: _____

Blank area for Pedigree Diagram.

DNA ANALYSIS

DIABETES TESTING:
 MODY Sequencing Panel (13 genes)
 Neonatal Diabetes Panel (22 genes)
 Congenital Hyperinsulinism Panel (10 genes)
Individual Diabetes Gene Sequencing (please indicate gene):
 ABCC8 GCK HNF1A HNF4A INS KCNJ11

DNA BANK DNA Banking **ONLY** (Separate Consent Form Required)
FRAGX DNA Fragile X DNA Analysis (FMR1 only)
GALT DNA Galactosemia (GALT) Panel (8 mutations)
GAUCHE DNA Gaucher Disease (GBA) Panel (11 mutations)
GAUCHSEQ Gaucher Disease (GBA) Sequencing Analysis

HEARING LOSS TESTING:
 Connexin 26/30 (GJB2/GJB6) DNA Analysis**
 Connexin 30 (GJB6) Sequencing
 Pendred (SLC26A4) Sequencing

LCHAD SEQ LCHAD/TFP (HADHA) Sequencing Analysis
LCHAD SEQ LCHAD/TFP (HADHB) Sequencing Analysis
LCHAD SEQ LCHAD/TFP (HADHA + HADHB) Sequencing
 Simultaneous Sequential (please indicate order)

LIPA SEQ Lysosomal Acid Lipase (LIPA) Sequencing (Wolman Disease/CESD)
MCAD SEQ MCAD (ACADM) Sequencing Analysis
MD DNA Duchenne/Becker Muscular Dystrophy (DMD) Deletion/Duplication***
POLG1 SEQ Polymerase Gamma (POLG) Sequencing Analysis
POLG2 SEQ Polymerase Gamma 2 (POLG2) Sequencing Analysis
POMPE SEQ Pompe (GAA) Sequencing Analysis
PWS-AS Prader-Willi/Angelman DNA Methylation study (one or both)

AGXT SEQ Primary Hyperoxaluria Type 1 (AGXT) Sequencing Analysis
ALDH7A1 SEQ Pyridoxine-Dependent Seizures (ALDH7A1) Sequencing Analysis

RETT/ATYPICAL RETT SYNDROME TESTING:
 Rett Panel** (MECP2 Seq, MECP2 Del/Dup***, CDKL5 Seq, FOXP1 Seq)
 MECP2 Panel** (MECP2 Sequencing with reflex to Deletion/Duplication***)
 MECP2 Sequencing
 MECP2 Deletion/Duplication***
 CDKL5 Sequencing
 FOXP1 Sequencing

SMA Spinal Muscular Atrophy (SMN1) (Diagnostic)
SMACARRIER Spinal Muscular Atrophy (SMN1) Carrier Testing***
TYR DNA Tyrosinemia type 1 (FAH) Panel (6 mutations)
VLCAD SEQ VLCAD (ACADVL) Sequencing Analysis
ATP7B SEQ Wilson Disease (ATP7B) Sequencing Analysis

KNOWN VARIANT ANALYSIS (Information Below is Required for this Testing)
 1 Variant 2 Variants 3 Variants
GENE: _____
VARIANT(S)/MUTATION(S): _____
NAME OF FAMILY MEMBER PREVIOUSLY TESTED AT SEATTLE CHILDREN'S LAB*: _____
RELATIONSHIP TO FAMILY MEMBER PREVIOUSLY TESTED: _____

*If relative was not tested at Seattle Children's Laboratory, please contact the Laboratory Genetic Counselors (206-987-5400) for approval of testing prior to submitting samples.

** Panel tests are performed sequentially in the order in which they are listed
 *** DNA extracted at other laboratories is not acceptable for SMA Carrier, DMD & MECP2 del/dup testing

BILLING INFORMATION

PHYSICIAN NOTIFICATION: Only tests that you believe are appropriate for patient care should be ordered. Medicare/Medicaid will pay only for tests that are medically necessary for the diagnosis and treatment of the patient, rather than for screening purposes.

BILLING NOTIFICATION: All samples will be billed to the referring institution unless complete billing and diagnosis information is provided on this form. Contact Seattle Children's Laboratory Client Services for additional assistance (206) 987-2617.

BILL TO:

- Referring Institution (Preferred)** - Provide billing address or stamp institution's information.
(Institutional billing will be done for all patients with Medicare except for established Seattle Children's patients.)

Billing Address:	Billing Contact Name:
Billing Contact Phone/Fax:	Billing Contact Email:

- Primary Insurance** (Attach copy of card.) **Medicaid** (Only Alaska, Idaho, Montana and Washington Medicaid are accepted.)

Patient Address		
Guarantor Name	DOB	Relationship to Patient
Guarantor Address (if different from patient's)		
Guarantor Phone (if different from patient's)	Employer	
Primary Care Physician	Phone Number	
Insurance Company/Medical Coverage		
Claims Address	Phone Number	
Policy Number	Group Number	
Subscriber	Sex	Subscriber's DOB

- Secondary Insurance** (Attach copy of card.) **Medicaid** (Only Alaska, Idaho, Montana and Washington Medicaid are accepted.)

Insurance Company/Medical Coverage		
Claims Address	Phone Number	
Policy Number	Group Number	
Subscriber	Sex	Subscriber's DOB

- Self Pay**- First, call Lab Client Services for pricing. Then, provide credit card information below or enclose a check with the sample.

Patient Address		
Guarantor Name	DOB	Relationship to Patient
Guarantor Address (if different from patient's)		
Guarantor Phone (if different from patient's)		
Name on Credit Card	Payment Amount	CVN
Card Number	Card Type	Expiration

Please visit our test catalog at <http://seattlechildrenslab.testcatalog.org> for testing information or call:

Lab Genetic Counselors (206) 987-5400 Lab Client Services (206) 987-2617 Molecular Genetics Lab (206) 987-3872



Ship to: LABORATORY
4800 Sand Point Way NE, M/S: OC.8.720
SEATTLE, WA 98105