

**FAILURE TO COMPLETE MAY DELAY RESULTS**

Patient's Last Name	First	Middle	Birth Date (Required)	Legal Sex	Gender Identity
Outside Patient Number	Outside Specimen Number	Send Report To:			
Ordering Provider			Address:		
Provider Phone#/Email	Diagnosis/ICD10 Code (Required)	Phone:	Fax:		

**IMPORTANT INFORMATION REGARDING BILLING AND MEDICAL NECESSITY ON LAST PAGE**

<b>CLINICIAN TO CONTACT FOR INFO/ABNORMAL RESULTS:</b>			<b>FAX ADDITIONAL RESULTS TO:</b>		
NAME:	EMAIL:	PHONE #:	NAME:	FAX #:	

**SPECIMEN INFORMATION**
**ALL SPECIMENS MUST BE LABELED WITH A MINIMUM OF TWO UNIQUE IDENTIFIERS**

Date collected: \_\_\_\_\_ Time collected: \_\_\_\_\_

<input type="checkbox"/> Blood <input type="checkbox"/> Cord Blood <input type="checkbox"/> Extracted gDNA from blood <input type="checkbox"/> EDTA <input type="checkbox"/> ACD	<input type="checkbox"/> Saliva (OrageneDx OGD-575/675 only)	<input type="checkbox"/> Fresh Tissue Tissue source (Exact Anatomical Site): _____
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**FOR SCH LAB USE:**  Room Temp  Refrig  Frozen

Container(s) received:  EDTA  NaHep  Sterile cup/cryovial  DNA tube  Saliva  Other: \_\_\_\_\_

**PATIENT/FAMILY HISTORY REQUIRED - ATTACH RELEVANT CLINIC NOTES**

Clinical information provided will aid in interpretation, decrease testing delays and improve reporting.

**REASON FOR STUDY:**  Diagnostic (affected)  Diagnostic (not affected)  Carrier Testing (affected family member)  Carrier Testing (no family history)

**CLINICAL FINDINGS, FAMILY HISTORY:** \_\_\_\_\_

**RELEVANT PREVIOUS GENETIC TEST RESULT(S):** \_\_\_\_\_

**PATIENT PREGNANT?**  No  Yes, estimated due date: \_\_\_\_\_

**MOLECULAR ANALYSIS**

 Test information, specimen and shipping requirements & gene lists available at: <http://seattlechildrenslab.testcatalog.org>

Testing performed by next-generation sequencing (NGS) analyzed to identify both sequence and copy number variants.

Test code	CHILDHOOD INTERSTITIAL LUNG DISEASE (ChILD) Panel	Test code	IMMUNODEFICIENCY (ImmuneSeq) Panels
LAB1825	<input type="checkbox"/> ChILD Expanded Seq Panel	LAB3799	<input type="checkbox"/> ImmuneSeq Autoimmune Lymphoproliferative Syndrome (ALPS)
<b>CRANIOSYNOSTOSIS Panels</b>		LAB3800	<input type="checkbox"/> ImmuneSeq Familial Hemophagocytic Lymphohistiocytosis (FHLH)
LAB1835	<input type="checkbox"/> Craniosynostosis Focused Seq Panel	LAB3904	<input type="checkbox"/> ImmuneSeq Primary Ciliary Dyskinesia (PCD) Panel
LAB1835	<input type="checkbox"/> Craniosynostosis Expanded Seq Panel	LAB3798	<input type="checkbox"/> ImmuneSeq Severe Combined Immunodeficiency (SCID)
<b>DIABETES Panels</b>		LAB3801	<input type="checkbox"/> ImmuneSeq (VEO-IBD)/Early Onset Enteropathy
LAB2943	<input type="checkbox"/> Congenital Hyperinsulinism Seq Panel	LAB3797	<input type="checkbox"/> ImmuneSeq Expanded Panel
LAB1884	<input type="checkbox"/> Maturity Onset Diabetes of the Young Seq Panel	<b>TARGETED GENE ANALYSIS from SCH Panels</b>	
LAB1885	<input type="checkbox"/> Neonatal Diabetes Seq Panel	Target gene(s) must be specified: _____	
<b>DIFFERENCES IN SEX DEVELOPMENT (DSD) Panel</b>		LAB3617	<input type="checkbox"/> Targeted Gene Sequencing by NGS
LAB1840	<input type="checkbox"/> Differences in Sex Dev Seq Panel	LAB3616	<input type="checkbox"/> Targeted Gene Deletion/Duplication by Array
<b>HEREDITARY HEMORRHAGIC TELANGIECTASIA Panel</b>		<b>MOLECULAR FAMILY FOLLOW-UP STUDY*</b>	
LAB1856	<input type="checkbox"/> Hereditary Hemorrhagic Telangiectasia Seq Panel	*Targeted testing is available <u>only</u> for family follow-up of individuals previously tested at Seattle Children's Hospital Genetics Lab. All below fields are <u>required</u> .	
<b>INTESTINAL PSEUDO-OBSTRUCTION Panel</b>		LAB1915	<input type="checkbox"/> Targeted Gene Variant:
LAB1866	<input type="checkbox"/> Intestinal Pseudo-Obstruction Seq Panel	Gene: _____	
<b>RETT/ANGELMAN SYNDROME Panel</b>		Variant(s): _____	
LAB1898	<input type="checkbox"/> Rett/Angelman Syndrome Seq Panel	Proband Name: _____	
		Relationship to Proband: _____	

Test information, specimen and shipping requirements & gene lists available at: <http://seattlechildrenslab.testcatalog.org>

Test code OTHER MOLECULAR ANALYSIS		Test code BIOCHEMICAL GENETICS MOLECULAR ANALYSIS	
LAB1808	<input type="checkbox"/> 22q11.2 Deletion/Duplication by MLPA (22q11.2 DS, DiGeorge, VCFS)	LAB3912	<input type="checkbox"/> Galactosemia (GALT) Sequencing
LAB1847	<input type="checkbox"/> Fragile X DNA (FMR1)	LAB1850	<input type="checkbox"/> Gaucher Disease Panel (11 variants)
LAB1894	<input type="checkbox"/> Prader-Willi/Angelman Methylation & CNV Analysis	LAB1850	<input type="checkbox"/> Gaucher DNA Sequencing (GBA)
LAB1912	<input type="checkbox"/> Spinal Muscular Atrophy <u>Diagnostic</u> (SMN1 & SMN2 Copy Number)	LAB1870	<input type="checkbox"/> LCHAD/TFP (HADHA) Sequencing
LAB1911	<input type="checkbox"/> Spinal Muscular Atrophy <u>Carrier</u> Test (SMN1 copy number)	LAB1870	<input type="checkbox"/> LCHAD/TFP (HADHB) Sequencing
		LAB1872	<input type="checkbox"/> Lysosomal Acid Lipase (LIPA) Sequencing (Wolman Disease/CESD)
		LAB1877	<input type="checkbox"/> MCAD (ACADM) Sequencing
		LAB1889	<input type="checkbox"/> Polymerase Gamma (POLG) Sequencing
		LAB1891	<input type="checkbox"/> Polymerase Gamma 2 (POLG2) Sequencing
		LAB1892	<input type="checkbox"/> Pompe (GAA) Sequencing
		LAB1862	<input type="checkbox"/> Primary Hyperoxaluria Type 1 (AGXT) Sequencing
		LAB1811	<input type="checkbox"/> Pyridoxine-Dependent Seizures (ALDH7A1) Sequencing
		LAB3530	<input type="checkbox"/> Tyrosinemia type 1 (FAH) Panel (6 variants)
		LAB1921	<input type="checkbox"/> VLCAD (ACADVL) Sequencing
		LAB1926	<input type="checkbox"/> Wilson Disease (ATP7B) Sequencing
Test code ARRAY ANALYSIS		CYTO/ARRAY FAMILY FOLLOW-UP STUDY***	
LAB1803	<input type="checkbox"/> Chromosomal SNP Microarray	<p>***Targeted testing is available <u>only</u> for family follow-up of individuals previously tested at Seattle Children's Hospital Genetics Lab.</p> <p>Proband control specimen may be required. Consult with laboratory genetic counselors prior to placing order if uncertain.</p> <p><b>Proband Name (Required):</b> _____</p> <p><b>Relationship to Proband (Required):</b> _____</p> <p>Test (indicated on Proband report):</p>	
Test code CYTOGENETIC ANALYSIS		<p>LAB3622 <input type="checkbox"/> Family Study Karyotype</p> <p>LAB3621 <input type="checkbox"/> Family Study FISH</p> <p>LAB3625 <input type="checkbox"/> Family Study qPCR</p> <p>LAB3623 <input type="checkbox"/> Family Study Chromosomal SNP Microarray</p> <p>LAB3682 <input type="checkbox"/> Proband Control specimen for:            Family Member Name (Required): _____            Relationship to Family Member (Required): _____            Check test that will be performed on family member:</p> <p><input type="checkbox"/> Family Study Karyotype</p> <p><input type="checkbox"/> Family Study FISH</p> <p><input type="checkbox"/> Family Study qPCR</p> <p><input type="checkbox"/> Family Study Chromosomal SNP Microarray</p>	
LAB1797	<input type="checkbox"/> Constitutional Karyotype - Mosaic Study (30 cells)**		
LAB1797	<input type="checkbox"/> Constitutional Karyotype - Routine (20 cells)**		
LAB1797	<input type="checkbox"/> Constitutional Karyotype - Limited Study (5 cells)		
LAB1800	<input type="checkbox"/> Constitutional FISH (Fluorescence <i>in situ</i> hybridization) for SRY Sex Determination		
LAB1804	<input type="checkbox"/> Fibroblast Culture		
LAB1804	<input type="checkbox"/> Fibroblast Culture and Storage (Cryopreservation)		
<p>** <input type="checkbox"/> Preliminary karyotype notification requested (24-72 hours).</p> <ul style="list-style-type: none"> <li>• <u>Only</u> available for infants &lt;2 weeks of age.</li> <li>• Minimum sample volume: 1.5mL.</li> <li>• Additional charge applies.</li> </ul> <p>Contact Name (Required): _____</p> <p>Contact Number (Required): _____</p>			

**See Page 3 for Billing Information**

## 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