

**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:	PHONE #:	NAME:	FAX #:
EMAIL:		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"/> Saliva (OrageneDx OGD-575/675 only)	<input type="checkbox"/> Fresh Tissue
<input type="checkbox"/> Cord Blood	Tissue source (Exact Anatomical Site): _____	
<input type="checkbox"/> Extracted gDNA from blood		
<input type="checkbox"/> EDTA <input type="checkbox"/> ACD		

**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.

<b>Test code CHILDHOOD INTERSTITIAL LUNG DISEASE (ChILD) Panel</b> LAB1825 <input type="checkbox"/> ChILD Expanded Seq Panel <b>DIABETES Panels</b> LAB2943 <input type="checkbox"/> Congenital Hyperinsulinism Seq Panel LAB1884 <input type="checkbox"/> Maturity Onset Diabetes of the Young Seq Panel LAB1885 <input type="checkbox"/> Neonatal Diabetes Seq Panel <b>HEREDITARY HEMORRHAGIC TELANGIECTASIA Panel</b> LAB1856 <input type="checkbox"/> Hereditary Hemorrhagic Telangiectasia Seq Panel <b>IMMUNODEFICIENCY Panels</b> LAB3799 <input type="checkbox"/> Autoimmune Lymphoproliferative Syndrome (ALPS) Panel LAB3800 <input type="checkbox"/> Familial Hemophagocytic Lymphohistiocytosis (FHLH) Panel LAB3904 <input type="checkbox"/> Primary Ciliary Dyskinesia (PCD) Panel LAB3798 <input type="checkbox"/> Severe Combined Immunodeficiency (SCID) Panel LAB3801 <input type="checkbox"/> Very Early Onset Inflammatory Bowel Disease (VEO-IBD) Panel LAB3797 <input type="checkbox"/> ImmuneSeq Inborn Errors of Immunity Panel <b>INTESTINAL PSEUDO-OBSTRUCTION Panel</b> LAB1866 <input type="checkbox"/> Intestinal Pseudo-Obstruction Seq Panel	<b>Test code CRANIOSYNOSTOSIS Panels</b> LAB1835 <input type="checkbox"/> Craniosynostosis Focused Seq Panel LAB1835 <input type="checkbox"/> Craniosynostosis Expanded Seq Panel <b>DIFFERENCES IN SEX DEVELOPMENT (DSD) Panel</b> LAB1840 <input type="checkbox"/> Differences in Sex Dev Seq Panel <b>TARGETED GENE ANALYSIS from SCH Panels</b> Target gene(s) must be specified: _____ LAB3617 <input type="checkbox"/> Targeted Gene Sequencing by NGS LAB3616 <input type="checkbox"/> Targeted Gene Deletion/Duplication by Array <b>MOLECULAR FAMILY FOLLOW-UP STUDY**</b> <b>**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> LAB1915 <input type="checkbox"/> Targeted Gene Variant: Gene: _____ Variant(s): _____ Proband Name: _____ Relationship to Proband: _____
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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
<p>LAB1808 <input type="checkbox"/> 22q11.2 Deletion/Duplication by MLPA (22q11.2 DS, DiGeorge, VCFS)</p> <p>LAB1847 <input type="checkbox"/> Fragile X DNA (FMR1)</p> <p>LAB1894 <input type="checkbox"/> Prader-Willi/Angelman Methylation &amp; CNV Analysis</p> <p>LAB1912 <input type="checkbox"/> Spinal Muscular Atrophy <u>Diagnostic</u> (SMN1 &amp; SMN2 Copy Number)</p> <p>LAB1911 <input type="checkbox"/> Spinal Muscular Atrophy <u>Carrier</u> Test (SMN1 copy number)</p>	<p>LAB3912 <input type="checkbox"/> Galactosemia (GALT) Sequencing</p> <p>LAB1877 <input type="checkbox"/> MCAD (ACADM) Sequencing</p> <p>LAB3941 <input type="checkbox"/> Ornithine Transcarbamylase (OTC) Deficiency Sequencing</p> <p>LAB1892 <input type="checkbox"/> Pompe (GAA) Sequencing</p> <p>LAB1921 <input type="checkbox"/> VLCAD (ACADVL) Sequencing</p> <p>LAB1926 <input type="checkbox"/> Wilson Disease (ATP7B) Sequencing</p>
Test code ARRAY ANALYSIS	CYTO/ARRAY FAMILY FOLLOW-UP STUDY***
<p>LAB1803 <input type="checkbox"/> Chromosomal SNP Microarray</p>	<p>***Targeted testing is available <u>only</u> for family follow-up of individuals previously tested at Seattle Children's Hospital Genetics Lab.</p>
Test code CYTOGENETIC ANALYSIS	<p>Proband control specimen may be required. Consult with laboratory genetic counselors prior to placing order if uncertain.</p>
<p>LAB1797 <input type="checkbox"/> Constitutional karyotype - Mosaic Study (30 cells)</p> <p>LAB1797 <input type="checkbox"/> Constitutional karyotype - Routine (20 cells)</p> <p>LAB1797 <input type="checkbox"/> Constitutional karyotype - Limited Study (5 cells)</p> <p>LAB1800 <input type="checkbox"/> Constitutional FISH (Fluorescence <i>in situ</i> hybridization) for SRY Sex Determination</p> <p>LAB1804 <input type="checkbox"/> Fibroblast Culture</p> <p>LAB1804 <input type="checkbox"/> Fibroblast Culture and Storage (Cryopreservation)</p>	<p><b>Proband Name (Required):</b> _____</p> <p><b>Relationship to Proband (Required):</b> _____</p> <p>Test (indicated on Proband report):</p> <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>
<div style="border: 1px solid black; padding: 5px;"> <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> </div>	<p>LAB3682 <input type="checkbox"/> Proband Control specimen for:</p> <p>Family Member Name (Required): _____</p> <p>Relationship to Family Member (Required): _____</p> <p>Check test that will be performed <u>on family member</u>:</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>

## 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