

# Clinical Lab Request

## Genetics

Ship specimens to:  
 UW MEDICAL CENTER  
 LABORATORY MEDICINE - GENETICS LAB  
 1959 NE PACIFIC ST, ROOM NW220  
 SEATTLE, WA 98195-7110

LAB ACC. #	LOGGED BY:
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Genetics Lab (206) 598-6429 (Clinical questions)  
 Genetics Lab genelab@uw.edu (Clinical questions)  
 Laboratory Fax (206) 598-0304  
 Reference Services (206) 520-4600 (Specimen transport)

Complete a requisition for each individual tested

PATIENT NUMBER		Test Information: http://depts.washington.edu/labweb/genetics	
PATIENT NAME		REASONS FOR TESTING:	
M <input type="checkbox"/> F <input type="checkbox"/>		<input type="checkbox"/> Carrier Detection	
DATE OF BIRTH		<input type="checkbox"/> Diagnostic / Rule out	
SPECIMEN COLLECTION DATE AND TIME		<input type="checkbox"/> Mutation Typing	
SENDER SPECIMEN NUMBER		<input type="checkbox"/> Presymptomatic	
		<input type="checkbox"/> Other (Specify) _____	
Ethnic Background:		ICD Code: <b>REQUIRED</b>	
ORDERING PROVIDER		PATIENT BILLING ADDRESS	
NPI #		STREET	
PHONE:		CITY	
FAX:		STATE	
EMAIL (PRIMARY CONTACT):		ZIP	
CONTACT PERSON:		TELEPHONE	
PHONE:		SUBSCRIBER NAME	
FAX:		GROUP #	
Send Reports To (mailing address):		<input type="checkbox"/> Premera Blue Cross <input type="checkbox"/> Regence <input type="checkbox"/> DSHS (attach current coupon)	
<b>REQUIRED</b>		<input type="checkbox"/> Medicare (answer required question below) Is this either a hospital outpatient or inpatient? Yes      No (see reverse side for additional information)	
FAX		Other Insurance Name/Address	
Fax Results? <input type="checkbox"/> Yes <input type="checkbox"/> No			

Check Test Requested	Genetics/Hereditary	Specimen Submitted
<b>Tumor/Somatic</b> <input type="checkbox"/> BRCA 1/2 - Complete Analysis (Tissue) <b>BRCA12</b> <input type="checkbox"/> ALK FISH 671 <input type="checkbox"/> ALK Resistance <b>ALKHSP</b> <input type="checkbox"/> ROS1 FISH 423 <input type="checkbox"/> EGFR Mutations <b>EGFRHS</b> <input type="checkbox"/> EGFR T790M <b>T790</b> <input type="checkbox"/> PDL-1 - IHC (attach separate PhenoPath Req) <input type="checkbox"/> BRAF Mutations <b>BRAF</b> <input type="checkbox"/> BRAF Glioma <b>BRAF</b> <input type="checkbox"/> HRAS Mutations <b>HRASHS</b> <input type="checkbox"/> NRAS Mutations <b>NRAS</b> <input type="checkbox"/> KRAS Mutations <b>KRAS</b> <input type="checkbox"/> IDH Mutations <b>IDH</b> <input type="checkbox"/> KIT Mutations, Melanoma <b>KITML</b> <input type="checkbox"/> KIT & PDGFRA Mutations, GIST <b>KITG</b> <input type="checkbox"/> Microsatellite Instability DNA Test <b>MSI</b> <input type="checkbox"/> Mismatch Repair IHC 669 <input type="checkbox"/> MLH1 methylation <b>MLH1MT</b> <input type="checkbox"/> UW-Oncoplex Cancer Gene Panel <b>OPX</b> Indicate prior molecular results in clinical history <input type="checkbox"/> UW-Oncoplex Single Gene <b>OPG</b>	<input type="checkbox"/> Alpha Thalassemia (deletions) <b>ATHAL</b> <input type="checkbox"/> Alpha Hemoglobin DNA Sequence <b>HASEQ</b> <input type="checkbox"/> Beta Hemoglobin DNA Sequence <b>HBSEQ</b> <input type="checkbox"/> Cystic Fibrosis <input type="checkbox"/> 5T Allele (CBAVD) <b>CFDNA</b> <input type="checkbox"/> Prothrombin 20210A (Factor II) <b>PRODS</b> <input type="checkbox"/> Factor V Leiden (APC Resistance) <b>F5DNA</b> <input type="checkbox"/> Fragile X DNA Test <b>FX</b> <input type="checkbox"/> Hemochromatosis (HFE) <b>HEMDNA</b> <input type="checkbox"/> Huntington Disease <b>HDTEST</b> <input type="checkbox"/> Maternal Cell Contamination <b>MCC</b> <input type="checkbox"/> Spinocerebellar Ataxia Panel (Types 1, 2, 3, 6, 7) <b>SCAPN</b> <input type="checkbox"/> Spinocerebellar Ataxia, single test Specify Type: _____ <b>SCA 1, 2, 3, 6, 7</b> <input type="checkbox"/> Other: _____	<input type="checkbox"/> BRCA 1/2 - Complete Analysis <b>BRCA12</b> <input type="checkbox"/> BRCA 1/2 - Ashkenazi Jewish 3-site <b>BRCAAJ</b> <input type="checkbox"/> BROCA - Cancer Risk Panel <b>BROCA</b> <input type="checkbox"/> ColoSeq - Lynch and Polyposis Panel <b>COSEQ</b> <input type="checkbox"/> ColoSeq Tumor - Full Panel <input type="checkbox"/> ColoSeq Tumor - Single Gene(s) _____ SPS: Do NOT log CSQTS/CSQTP <input type="checkbox"/> Normal Control (blood) <b>DNAPRP</b> (Required for ColoSeq Tumor tests) <input type="checkbox"/> ColoSeq - Polyposis (APC and MUTYH) <b>COSEQP</b> <input type="checkbox"/> Epileptic Encephalopathy Panel <b>EPIPX</b> <input type="checkbox"/> Megalencephaly Panel <b>MEGPX</b> <input type="checkbox"/> Immunoplex Panel (specify full or subpanel) <b>IMD</b> <input type="checkbox"/> Full <input type="checkbox"/> SCID <input type="checkbox"/> Agamma <input type="checkbox"/> FHLH <input type="checkbox"/> MarrowSeq Panel <b>MRW</b> <input type="checkbox"/> Single Gene Analysis (specify gene) <b>SGN</b> <input type="checkbox"/> Known Mutation* (specify gene & mutation) <b>KMU</b>
		<input type="checkbox"/> Amniocytes Fresh tissues are typically cultured prior to testing <input type="checkbox"/> Chorionic Villi Fresh tissues are typically cultured prior to testing <input type="checkbox"/> Blood (5 mL EDTA /Lavender Top) <input type="checkbox"/> Blood (BCT-1) Required for cell free DNA <input type="checkbox"/> DNA (source _____) <input type="checkbox"/> Solid Tissue, FFPE <input type="checkbox"/> Solid Tissue, fresh Contact lab for acceptable types <input type="checkbox"/> Solid Tissue, other <input type="checkbox"/> Other: _____
		<b>Medical Necessity Information</b> When ordering tests for which Medicare reimbursement will be sought, physicians should only order tests which are medically necessary for diagnosis or treatment of the patient. You should be aware that Medicare generally does not cover routine screening tests, and will only pay for tests that are covered by the program and are reasonable and necessary to treat or diagnose the patient.

Pedigree / Family History/ Clinical History and Prior Molecular Results (Please attach notes and/or pedigree)

#### MEDICARE BILLING INFORMATION

Medicare billing policy prevents us from submitting a Medicare claim for laboratory testing referred to us on hospital inpatients or hospital outpatients. For these samples, we will bill the sending location.

#### CMS MEDICAL NECESSITY INFORMATION

It is our policy to provide health care providers with the ability to order only those lab tests medically necessary for the individual patient and to ensure that the convenience of ordering standard panels and custom profiles does not impact this ability. While we recognize the value of this convenience, indiscriminate use of panels and profiles can lead to ordering tests that are not medically necessary. Therefore, all tests offered in our panels and profiles can be ordered individually as well. If a component test is not listed individually on the request form, it may be written in the "OTHER REQUESTS" box. We encourage you to order individual tests or a less inclusive profile when not all of the tests included in the panel or profile are medically necessary for the individual patient.