



Colorado Genetics Laboratory  
www.coloradogeneticslab.com

**PRENATAL / PREGNANCY LOSS / POSTNATAL  
TEST REQUEST FORM**

Revised \ 201

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Aurora, Colorado 80045  
(303) 724-5701  
(888) 659-4932 Toll Free  
(303) 724-5795 Fax

PATIENT INFORMATION				
Patient Last Name	First Name	Middle Initial	Sex: M F	
Date of Birth	Hospital/ID #	INPATIENT	OUTPATIENT	
REFERRED BY				
PHYSICIAN	FACILITY			
Address	Address			
Phone	FAX	Phone	FAX	
Email				
BILLING				
Bill charges to:	Patient	Insurance*	Hospital	Physician/Clinic
*Attach billing information <b>Include signed waiver if microarray requested</b>				
SIGNS, SYMPTOMS, DIAGNOSIS, AND ICD10 CODES REQUIRED FOR SPECIMEN PROCESSING (PLEASE DO NOT USE "RULE OUT")				
			<p>In addition to the studies requested below, I authorize the laboratory to perform FISH (fluorescence <i>in situ</i> hybridization) if indicated by the patient's clinical history or cytogenetic results.</p>	
			<p style="text-align: center;">Physician/Authorized Signature</p>	
SPECIMEN INFORMATION		STUDIES REQUESTED		
Date Collected	Time	AM	PM	Amount Collected
<b>PRENATAL</b>		<b>CHROMOSOMES:</b>		
Gestation by U/S		Standard chromosome analysis		
LMP		Standard chromosome analysis; <b>IF NORMAL → REFLEX to SNP microarray (on cultured cells)</b>		
G P SAB TAB		<b>FISH:</b>		
Fetal Sex (if known): M F		STAT trisomy screen by FISH & standard chromosome analysis		
Amniotic Fluid		STAT trisomy screen by FISH; <b>IF ABNORMAL → REFLEX to standard chromosome analysis</b>		
<b>Alpha-fetoprotein (reflex to AChE)? Yes No</b>		<b>IF NORMAL → REFLEX to SNP microarray** &amp; 5-cell chromosome analysis</b>		
Chorionic villus		FISH for (specify): & standard chromosome analysis		
Percutaneous umbilical blood		<b>MICROARRAY:</b>		
Other (specify):		Chromosomal SNP microarray** & standard chromosome analysis		
		Chromosomal SNP microarray** & 5-cell chromosome analysis		
		<b>** If inadequate direct sample, microarray may be performed on cultured cells.</b>		
<b>PREGNANCY LOSS</b>		<b>CHROMOSOMES:</b>		
Gestation		Standard chromosome analysis		
Fetal Sex (if known): M F		Standard chromosome analysis; <b>IF NORMAL → REFLEX to SNP microarray (on cultured cells)</b>		
Products of conception		Standard chromosome analysis; <b>IF CULTURE FAILURE → REFLEX to pregnancy loss trisomy screen by FISH</b>		
Placenta		<b>FISH:</b>		
Fetal tissue (specify):		Pregnancy loss trisomy screen by FISH on paraffin embedded tissue		
Other (specify):		FISH for (specify): & standard chromosome analysis		
		<b>MICROARRAY:</b>		
		Chromosomal SNP microarray		
		Chromosomal SNP microarray & 5-cell chromosome analysis		
<b>POSTNATAL</b>		<b>CHROMOSOMES:</b>		PATIENT LABEL HERE
Peripheral blood		Standard chromosome analysis		
Cord blood		High resolution chromosome analysis		
Skin Biopsy		<b>FISH:</b>		
Saliva - Spit		STAT trisomy screen by FISH & standard chromosome analysis		
Saliva - Buccal/Mouth Swab		FISH for (specify): & standard		
Other (specify):		chromosome analysis (if not previously performed)		
<b>Is sample parental/familial follow-up for an abnormal microarray? Yes</b>		<b>MICROARRAY:</b>		
Child's name:		Chromosomal SNP microarray		
DOB:		Chromosomal SNP microarray & 5-cell chromosome analysis		
		7 u #U <sup>+</sup> - <b>Contact Laboratory if patient not initially studied by CGL</b>		
<b>ADDITIONAL SERVICES</b>		Culture for molecular or biochemical studies → <b>Include Information for Referral Specimens form, signed waiver, and payment for shipping</b>		
(Indicate sample above)		Culture and freeze		
<b>Microarray consent, specimen requirements, information for referral specimens, and waiver forms all available online.</b>				



**PATIENT INFORMATION**

Patient Last Name \_\_\_\_\_ First Name \_\_\_\_\_ Middle Initial \_\_\_\_\_ Sex: M F  
Date of Birth \_\_\_\_\_ Hospital/ID # \_\_\_\_\_ INPATIENT \_\_\_\_\_ OUTPATIENT \_\_\_\_\_

**REFERRED BY**

PHYSICIAN \_\_\_\_\_ FACILITY \_\_\_\_\_  
Address \_\_\_\_\_ Address \_\_\_\_\_  
Phone \_\_\_\_\_ FAX \_\_\_\_\_ Phone \_\_\_\_\_ FAX \_\_\_\_\_  
Email \_\_\_\_\_

**BILLING**

Bill charges to: Patient Insurance\* Hospital Physician/Clinic \* Attach billing information

**SIGNS, SYMPTOMS, DIAGNOSIS, AND ICD10 CODES** REQUIRED FOR SPECIMEN PROCESSING (PLEASE DO NOT USE "RULE OUT")

In addition to the studies requested, I authorize the laboratory to perform FISH (fluorescence *in situ* hybridization) if indicated by the patient's clinical history or cytogenetic results.

Physician/Authorized Signature: \_\_\_\_\_

**SPECIMEN COLLECTION**

Date Collected \_\_\_\_\_ Time \_\_\_\_\_ AM \_\_\_\_\_ PM \_\_\_\_\_ Amount Collected \_\_\_\_\_

**SPECIMEN INFORMATION**

Bone marrow aspirate \_\_\_\_\_ Peripheral blood \_\_\_\_\_  
Bone marrow core biopsy \_\_\_\_\_ Lymph node \_\_\_\_\_  
Solid tumor (source): \_\_\_\_\_ FFPE slides (4 micron on Plus Slides) \_\_\_\_\_  
(source): \_\_\_\_\_  
Other (specify): \_\_\_\_\_

CML \_\_\_\_\_ Myelodysplasia \_\_\_\_\_  
AML \_\_\_\_\_ B-cell lymphoma \_\_\_\_\_  
ALL \_\_\_\_\_ T-cell lymphoma \_\_\_\_\_  
Myeloma \_\_\_\_\_ CLL \_\_\_\_\_  
Other: \_\_\_\_\_

**Transplant Information**

Is the patient post transplant? Yes \_\_\_\_\_ If Yes, sex of donor: M F

**STUDIES REQUESTED**

**Chromosome analysis**

**FISH Studies for:**

BCR/ABL1 t(9;22) \_\_\_\_\_ ALL Panel \_\_\_\_\_  
Chimerism (for BMT patients) \_\_\_\_\_ AML Panel \_\_\_\_\_  
EGFR \_\_\_\_\_ AML Panel plus \_\_\_\_\_  
IGH (BAP) \_\_\_\_\_ PML/RARA \_\_\_\_\_  
KMT2A (MLL)(11q23) \_\_\_\_\_ Eosinophilia Panel \_\_\_\_\_  
MECOM (EVI1) \_\_\_\_\_ MDS Panel \_\_\_\_\_  
PML/RARA t(15;17) \_\_\_\_\_ MPN Panel \_\_\_\_\_  
PTEN \_\_\_\_\_ Myeloma panel \_\_\_\_\_  
TP53 (17p13.1) \_\_\_\_\_ Myeloma panel plus \_\_\_\_\_  
1p36/1q25 deletion \_\_\_\_\_ Reflex MYC; IGH/MYC \_\_\_\_\_  
19q13 / 19p13 deletion \_\_\_\_\_  
Screen for prior abnormal clone \_\_\_\_\_  
Other (specify): \_\_\_\_\_

Initial CLL Panel (ATM; 12; 13q14; IGH/CCND1;TP53)  
Follow-up CLL Panel (ATM; 12; 13q14; TP53)  
Diffuse Large B cell Lymphoma Panel (MYC; IGH/MYC; IGH/BCL2; BCL6; IGH)  
High Grade B cell Lymphoma Panel (BCL6; MYC; IGH/MYC; BCL2)  
MALT Lymphoma Panel (BCL6 for +3; MALT1; BIRC3/MALT1)  
Marginal Zone Lymphoma Panel (BCL6 for +3; 7/7q; TP53)  
Neuroblastoma Panel (MYCN/2cen; 1p36; ATM)  
Medulloblastoma Panel (MYCN; MYC; SEC63/6cen)  
Ependymoma Panel (1p36/1q25; CDKN2A; EGFR)  
IGH/CCND1 \_\_\_\_\_  
IGH/BCL2 \_\_\_\_\_  
IGH/MYC & MYC (BAP) \_\_\_\_\_

PATIENT LABEL HERE

**Additional FISH studies available.**  
**Please see our website for complete FISH Test list or call the laboratory.**