

Person Completing Requisition:			
Institution:	Client #		
Dept:	Physician/Provider:		
Address:			
City:	ST:	ZIP:	
Phone (Lab):	Phone/Email (Provider)		
Special Reporting Requests:	PO#:		



MEDICARE

Is testing for outpatient Medicare enrollee or Wisconsin Medicaid recipient? **Yes** **No**
 If yes, please complete and attach our beneficiary form located at www.versiti.org/medical-professionals/products-services/requisitions

PATIENT INFORMATION

Last Name:	First Name:	MI:	DOB:
MR#:	Accession#:	Draw Date:	Draw Time:
Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other Karyotype: _____	Is patient currently pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No Due date: _____		
Has patient had an allogeneic stem cell transplant? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, send pre-transplant extracted DNA sample	Has patient had a blood transfusion in the last 2 weeks? <input type="checkbox"/> Yes <input type="checkbox"/> No Date and type of transfusion: _____		
Specimen Type: <input type="checkbox"/> ACD Blood <input type="checkbox"/> Buccal Swabs <input type="checkbox"/> EDTA Blood <input type="checkbox"/> Bone Marrow <input type="checkbox"/> DNA <input type="checkbox"/> Sodium Heparin Blood <input type="checkbox"/> Other _____			
Fetal Specimen Type: <input type="checkbox"/> Amniotic Fluid <input type="checkbox"/> Cultured Amniocytes <input type="checkbox"/> CVS <input type="checkbox"/> Cultured CVS <input type="checkbox"/> DNA <input type="checkbox"/> Other _____			

PATIENT HISTORY (Necessary for optimal interpretation of test results and recommendations)

Ethnic Background (check all that apply): <input type="checkbox"/> Caucasian <input type="checkbox"/> African American <input type="checkbox"/> Hispanic/Latino <input type="checkbox"/> Asian <input type="checkbox"/> American Indian <input type="checkbox"/> Other _____	Clinical Diagnosis:
Relevant Clinical Presentation and Laboratory Findings (attach case notes if available):	
Family history of disorder? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, please describe. Attach pedigree and additional information if available.	

TEST ORDERS (see reverse side for sample requirements)

ENGRAFTMENT / CHIMERISM	BCR-ABL TESTING
TRANSPLANT INFORMATION: <input type="checkbox"/> Bone Marrow <input type="checkbox"/> Solid Organ <input type="checkbox"/> Other _____ Transplant Date: _____	PURPOSE OF TESTING: <input type="checkbox"/> Diagnosis <input type="checkbox"/> Monitoring Therapy <input type="checkbox"/> BCR-ABL Quantitative Analysis (4502) <input type="checkbox"/> BCR-ABL Breakpoint Identification *Order with BCR-ABL Quant* (4504) <input type="checkbox"/> BCR-ABL Kinase Mutation Analysis (4507)
Pre-Transplant Testing <input type="checkbox"/> For Recipient Sample (4020) (Provide donor name) Donor Name: _____ <input type="checkbox"/> For Donor Sample (4040) (Provide recipient name) Recipient Name: _____ <input type="checkbox"/> For Twin Zygosity Analysis (4060) (Provide donor name) Donor Name: _____ <input type="checkbox"/> For Twin Zygosity Analysis (4070) (Provide recipient name) Recipient Name: _____	PRENATAL GENOTYPING LMP Date: _____ Gestational Age: _____ Sample(s) submitted from (check all that apply): (Analysis of parental samples is highly recommended) <input type="checkbox"/> Mother <input type="checkbox"/> Father <input type="checkbox"/> No Parental Sample Father's Name: _____ Father's DOB: _____
Post-Transplant Testing <input type="checkbox"/> STAT Testing (Results in 48 hours/72 hours if sorted cells) <input type="checkbox"/> Chimerism on blood or bone marrow (4199) Prepare Sorted Cells, Perform Chimerism <input type="checkbox"/> CD3 & CD33 cells (4091/4199) <input type="checkbox"/> CD19 cells (4097/4199) <input type="checkbox"/> CD56 cells (4098/4199) <input type="checkbox"/> CD3 cells (4093/4199) <input type="checkbox"/> CD3 & CD33 & CD56 cells (4107/4199) <input type="checkbox"/> CD19 & CD56 cells (4106/4199) <input type="checkbox"/> Prepare MNC, chimerism (4092/4199) <input type="checkbox"/> Prepare buffy coat, chimerism (4094/4199)	RED CELL GENOTYPING For Hemolytic Disease of Fetus and Newborn Check appropriate system and complete serological information if known <input type="checkbox"/> RhC/c (4445) <input type="checkbox"/> RhD Zygosity(4475) <input type="checkbox"/> RhD (4455) <input type="checkbox"/> Fy ^{a/b} (Duffy)(4405) <input type="checkbox"/> RhE/e (4465) <input type="checkbox"/> K1/K2 (Kell)(4415) <input type="checkbox"/> Jk ^{a/b} (Kidd)(4425) <input type="checkbox"/> M/N(4435) <input type="checkbox"/> S/s (4485)
ERYTHROID CHIMERISM <input type="checkbox"/> Erythroid Chimerism (4250) Donor Genotype _____ <input type="checkbox"/> AA <input type="checkbox"/> AS (Required) Recipient Genotype _____ <input type="checkbox"/> SS <input type="checkbox"/> AS (Required)	HEMOCHROMATOSIS <input type="checkbox"/> Hemochromatosis (4600)
	SICKLE CELL DISEASE <input type="checkbox"/> Hemoglobin SC Mutation Analysis (4624)
	MATERNAL CELL CONTAMINATION <input type="checkbox"/> Maternal sample sent for MCC only <input type="checkbox"/> Maternal sample sent for MCC and genotyping
	Versiti Use Only _____ EDTA _____ BM _____ Amnio _____ ACDA _____ ACDB _____ Clot _____ Heparin _____ CVS _____ Other _____ Opened By _____ Evaluated By _____

DRAWING INSTRUCTIONS: Tubes must be individually labeled with **FULL NAME OF INDIVIDUAL, DATE AND TIME OF DRAW. Samples will be accepted from 8:00 A.M. to 5:00 P.M. Monday through Friday and Saturday morning.** Emergency testing **MUST** be arranged through the laboratory by calling 1-800-245-3117, ext. 6218.

Test	Sample Type	Store and Ship
BCR-ABL Quantitative Analysis BCR-ABL Kinase Mutation Analysis BCR-ABL Breakpoint Identification Erythroid Chimerism	3-5 mL EDTA Bone Marrow (lavender top) OR 10 mL EDTA Whole Blood (lavender top)	Room temperature via an overnight courier. Samples must be received within 48 hours of being drawn.
Hemoglobin SC Mutation Analysis Prenatal Testing Red Cell Antigen RhD Zygosity Maternal Cell Contamination	FETAL: 7-15 mL Amniotic Fluid or 5-10 mg CVS, backup culture of Amniocytes or CVS is highly recommended; Two T25 flasks Cultured Amniocytes or CVS (2x10 ⁶ minimum) PARENTAL & PATIENTS: 3-5 mL EDTA whole blood (lavender top). Maternal sample for maternal cell contamination 1µg DNA (25ng/µl and 25µl)	Room temperature.
Engraftment/Chimerism	PRE-TRANSPLANT: 3-5 mL EDTA (lavender top) whole blood or bone marrow OR 4-8 Buccal Swabs POST-TRANSPLANT: 3-5mL EDTA (lavender top) whole blood or bone marrow	Room temperature.
Cell Sort Enrichment CD3, CD19, CD33, CD56	4-7 mL Na Heparin (green top) whole blood per cell type sort. (Preferred) OR 4-7mL EDTA (lavender top) OR 4-7 mL ACDA (yellow top) whole blood per cell type sort. OR 3-5 mL bone marrow per cell type sort.	Room temperature. Samples must be received within 24 hours of draw and may be drawn Monday through Thursday for delivery Tuesday through Friday.
Hemochromatosis	3-5 mL EDTA (lavender top) whole blood	Room temperature.
Tissue	50-150 mg tissue (Call lab prior to shipping.)	Freeze and ship on dry ice or place in transport media and ship on ice or cold pack.

Blood samples should be shipped by overnight carrier. The package must be shipped in compliance with carrier's guidelines. Please contact your carrier for current biohazardous shipping regulations.

Shipping address: Versiti Wisconsin – Molecular Diagnostic Laboratory
638 N. 18th Street
Milwaukee, WI 53233
Phone: (414) 937-6250

Label Box: Refrigerate, Room Temperature, or Frozen (whichever is appropriate)

Verification of Informed Consent.

It is recommended that healthcare providers obtain a signed informed consent from the patient when genetic testing is ordered. By signing the informed consent, the patient agrees that they have received and understand the indications and implications of the genetic test and are voluntarily agreeing to have the test performed. In some states, informed consent is **required** by existing laws and regulations. Versiti recommends that ordering healthcare providers verify their state laws and regulations regarding informed consent for genetic testing. An informed consent form may be available from your institution or one can be found at <http://www.versiti.org/hg> under *forms*. Information regarding a general description of the test, purpose, sensitivity, analytical limitations, and the features and genetics of the condition(s) is also available [in the Versiti test catalog](#).

New York State patients: New York state healthcare providers are required to provide verification that informed consent (complying with New York State Department of Health Genetic Testing Standard 5 [GT S5] and New York State Civil Rights Law, Section 79-l) has been obtained from their patient. In order for genetic testing to be performed in our laboratory, please sign the verification below or submit a signed informed consent form. The sample will be destroyed not more than 60 days after the sample was obtained, unless a longer period of retention is expressly authorized in the consent

Verification of Informed Consent: I am a healthcare provider for the patient named on this requisition. I have obtained the required informed consent from the patient or the patient's legal guardian for each genetic test(s) ordered above and I authorize the testing of the enclosed specimen(s). I understand that no tests other than those authorized will be performed on genetic samples.

Signature of healthcare provider

Date