

Immunochemistry Reference Laboratory Requisition



Versiti Illinois: Phone 630-264-7832 | Fax 630-892-8648

Versiti Indiana: Phone 317-916-5188 | Fax 317-916-5189

Versiti Michigan: Phone 616-233-8583 | Fax 616-233-8687

Versiti Wisconsin: Phone 414-937-6205 | Fax 414-937-6461

NOTE: Versiti does NOT bill patients or insurance. Test orders must be placed through a medical facility that has an account with Versiti. Client # required.

Ordering Institution Information					
Person Completing Requisition:			Provider Name:		
Dept:			Provider Contact (Phone/Email):		
Institution:				Client #:	
Address:			City:	State:	Zip Code:
Phone (Lab):		Special Reporting Requests (Fax Number/Email):			
Patient Information					
Last Name:			First Name:		MI:
DOB:	MRN:		Accession #:		
Sample Collection Date:		Time:	Sex Assigned at Birth: <input type="checkbox"/> Male <input type="checkbox"/> Female		Ethnicity:
Patient Clinical History – Fill Out Below or Attach Patient Clinical History					
Patient Status: <input type="checkbox"/> Inpatient <input type="checkbox"/> Outpatient		Hgb/HCT:	Diagnosis:		
ABO/RH:		Known Antibodies:			
Number of Pregnancies:		Currently Pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No If Yes, Weeks Pregnant: _____			
Antibody Therapies: <input type="checkbox"/> RhIG <input type="checkbox"/> IVIG <input type="checkbox"/> Anti-CD38 <input type="checkbox"/> Anti-CD47 <input type="checkbox"/> Other: _____ Date Last Given: _____					
History of Stem Cell Transplant? <input type="checkbox"/> Yes <input type="checkbox"/> No		Date:	Patient Prior ABO/Rh:		Donor ABO/Rh:
Prior Transfusions: <input type="checkbox"/> Yes <input type="checkbox"/> No		# of Units:	ABO/Rh of Units	Date(s) Transfused:	
Specimen Type – See Page 2 for Specimen Requirements, DO NOT USE TUBES THAT CONTAIN SILICONE SEPARATOR GEL					
<input type="checkbox"/> EDTA/Whole Blood (Lavender/Pink Top) <input type="checkbox"/> Clot/Serum (Red Top) <input type="checkbox"/> Other: _____					
Test Orders – Additional Testing May Be Performed As Required. Attach Patient Results and List Medications on Page 2.					
<input type="checkbox"/> Antibody Identification (3060) <input type="checkbox"/> Antibody Titration (3080) <input type="checkbox"/> Positive DAT/Elution (3020) <input type="checkbox"/> ABO Discrepancy <input type="checkbox"/> Crossmatch Problem (3050) <input type="checkbox"/> Transfusion Reaction Investigation <input type="checkbox"/> HDFN Investigation (3100) <input type="checkbox"/> Other: _____					
Units Requested – *Compatibility Screen is NOT Intended as Crossmatch of Record, Refer to Your Facility's Policy Prior to Unit Issue*					
# Units Needed: _____ <input type="checkbox"/> Compatibility Screened <input type="checkbox"/> Irradiated <input type="checkbox"/> CMV Neg <input type="checkbox"/> Other: _____					
Serology Testing Performed at the Wisconsin Location					
<input type="checkbox"/> DAT Negative Workup (3111) <input type="checkbox"/> Thermal Amplitude (3021) <input type="checkbox"/> Donath Landsteiner (3011) <input type="checkbox"/> Drug-Dependent RBC Antibody Study (3110) – Drug(s): _____					
Complete MEDICATION Section on Page 2. For Each Drug, Send Dry Powder Weight of 500mg or More (Not Reconstituted).					
Molecular Testing Performed at the Wisconsin Location					
<input type="checkbox"/> Red Cell Genotyping Panel (3530) <input type="checkbox"/> Weak RhD Analysis (3040) <input type="checkbox"/> Weak RhD Analysis (3040) – Reflex to <input type="checkbox"/> Fya/Fyb (3860) – For Duffy Null Associated Neutropenia <input type="checkbox"/> Partial RhD Analysis (3240) Partial RhD Analysis (3240), If Indicated					
VERSITI USE ONLY: _____ EDTA/Whole Blood _____ Clot/Serum _____ Other: _____ Evaluated By: _____					

Shipping Addresses			
Versiti-IL 1200 N. Highland Ave Aurora, IL 60506		Versiti-IN 3450 N. Meridian Street Indianapolis, IN 46208	
Versiti-MI 1036 Fuller Ave NE Grand Rapids MI 49503		Versiti-WI 638 N. 18 th Street Milwaukee, WI 53233	
Sample Identification			
All samples must include sample identification clearly marked on each specimen container. Proper identification includes full name of individual, date obtained, hospital and/or patient identification number and the identification of the individual obtaining the specimen. Federal regulations mandate that a completed laboratory requisition form accompany each sample. Blood samples must be packaged to comply with requirements of mail or overnight courier service, if used.			
Specimen Requirements – Ship Refrigerated or Room Temperature, DO NOT SEND FROZEN			
SUSPECTED SEROLOGIC PROBLEM		REQUESTED AMOUNT	
Warm Autoimmune Hemolytic Anemia – IAT Positive with all panel cells tested and a positive DAT (1+ - 4+) *For patients under 20 kg body weight, sample requirements will be adjusted depending on communication with patient's provider.		No transfusion within the past 3 months: 24mL EDTA whole blood (lavender or pink top) AND 21mL clotted whole blood (red top) Transfused within the past 3 months: 5mL EDTA whole blood (lavender or pink top) AND 30mL clotted whole blood (red top)	
Antibody Identification	ABO/Rh Discrepancy	5mL EDTA whole blood (lavender or pink top) AND	
Antibody Titration	Suspected Transfusion Reaction	21mL clotted whole blood (red top)	
Crossmatch Problem	Antibody Confirmation		
Positive DAT/Elution		10mL EDTA whole blood (lavender or pink top) AND 10mL clotted whole blood (red top)	
DAT Negative Autoimmune Hemolytic Anemia Study		10mL EDTA whole blood (lavender or pink top) AND 21mL clotted whole blood (red top)	
Thermal Amplitude or Donath-Landsteiner Test		5mL EDTA whole blood AND 21mL clotted whole blood prewarmed and maintained at 37°C during clotting and serum separated immediately	
Drug-Dependent RBC Antibody Study (Complete the medication history listed below)		5mL EDTA whole blood AND 21mL clotted whole blood (red top) Consult with IRL M-F 8am - 4pm at 414-937-6205. <u>For each suspected drug, dry powder weight of 500mg or more (not reconstituted) must be sent with sample(s).</u>	
Hemolytic Disease of the Fetus and Newborn (HDFN)		Child – Cord blood sample (if available) Mother – 5mL EDTA whole blood (lavender or pink top) AND 21mL clotted whole blood (red top)	
MOLECULAR TESTS		REQUESTED AMOUNT	REASON/INDICATION FOR SUBMISSION
Weak RhD Analysis		5mL EDTA whole blood (lavender or pink top)	Used for investigation of RhD discrepancies and determination of RhIG candidacy
Partial RhD Analysis		5mL EDTA whole blood (lavender or pink top)	Used for investigation of anti-D or risk of anti-D alloimmunization in Rh Positive patients
Red Cell Genotyping Panel		5mL EDTA whole blood (lavender or pink top)	
*Prenatal Molecular Tests - Use the Prenatal Molecular Requisition Form			
Medication – List All Medications, Prescription and Non-Prescription, Taken in the Past 30 Days (Include: Aspirin, Antibiotics, etc.). Attach Full List of Medication if Needed.			
Medication	Dose	Date Begun	Last Taken