

Reference Laboratory Request Form NOTE: Incomplete forms may delay testing

Rootenai Health Blood Bank Phone Number: 208-625-5820 2003 Kootenai Health Way	For Reference Lab Only Specimen ID/Order No Date/Time Received:				
Coeur d'Alene, ID 83814	Butto, Fillio Modelved.				
Submitting Facility Information					
Facility Name					
Address					
Account Number					
	Fax				
Urgency of Request <u>Complete Clinical Status Information and Transfusion History</u>					
□Routine □ASAP □STAT Transf	sfusion or Surgery Date				
Patient Name	Patient ID (MRN)				
Birthdate Ethnicity Sex □M □F □Unknown					
ABO/Rh					
Sample Collection: Date Time _	Encounter/Visit/Admission #				
Clinical Status Diagnosis Pregnancy History Number of Pregnancies: Gravida/Para					
Medication	Due Dete				
Provide list if available	—— Due Date				
□IVIG □Anti-CD47 □Anti-CD38	Rhlg Given? □Y □N				
DAT Positive? □Y □N					
Transfusion History Within the last 3 months? Dates and Products					
Prior to last 3 months? ☐Y ☐N Date	tes				
History of transfusion reactions? N Dates Reaction Type					
History of HPC transplant? N Dates Patient's Prior ABO/Rh					
Donor's ABO/Rh Previous antibodies detected, check below. Other non-listed					
Anti- D C E c e f K k Fy	Fyb Jka Jkb M N S S CW WAA* CAA*				
	Autoantibody				
Red Cell Testing Request ☐ ABO discrepancy ☐ Antib	body Titer				
, ,	Cuanastad Transficaion				
☐ Anti-A₁ ☐ DAT ☐ Suspected Transfusion Reaction					
□ Elutio					
☐ Extended phenotype (serological) ☐ dTT	Red Cell Treatment □Class II				
(Da	Daracellex) □Class III				



Instructions:

- 1. Contact Reference Laboratory before sending samples.
- 2. Fill out this request form completely. Attach copies of any work performed at your facility. Incomplete forms may delay testing and require further communication. See Pages 3 and 4 for detailed instructions.
- 3. Label all samples with: Full patient name, second unique patient identifier number, date collected. Incorrectly or unlabeled specimens may be rejected and cannot be tested.
- 4. Update the Reference Laboratory with any changes in the status of the request.
- 5. Contact your local blood center to request antigen negative units.

Sample Preferences:

Test Request	Sample Preferences
ABO Discrepancy	
D (Rh) Discrepancy Resolution	1 6ml EDTA tube
DAT	I offili EDTA tube
Extended Phenotype	
Antibody ID	
Antibody Titer	3 6ml EDTA tubes
Elution	3 OIIII EDTA lubes
dTT Red Cell Treatment	

Turnaround Time:
Approximate Turnaround Time for Preliminary Results
Routine: Within 1-2 days
ASAP: Within 24 hours
STAT: Within 8 hours

Notes:

- All TATs are measured from the time the sample is received by the testing laboratory
- Complex workups may require additional time to resolve. A preliminary report will be provided



Form Instructions

Field Title	How the information you supply is used to focus testing efforts
Requesting Physician	Significance in testing: The request cannot proceed without a physician's order
	How to complete: Enter the physician's first and last name
Ethnicity	Significance in testing: The patient's race/ethnicity may help guide the workup and selection of rare red cells to test when the presence of an antibody to a high prevalence antigen is suspected Example:
	African American may indicate anti-Jsb, Hy, Ata and others Caucasian may indicate anti-Kpb, k, Yta and others Hispanic may indicate anti-Dib, Ge and others Asian may indicate anti-Dib, Jra and others
	How to complete: enter race/ethnicity (e.g., African American, Caucasian, Hispanic/Mexican, Hispanic/Puerto Rican, Asian, Native American, Pacific Islander, etc.)
Encounter/Visit/Admission#	Significance in testing: The encounter/visit/admission# is entered in our LIS and may aid in tracking patients with multiple workups How to complete: Enter the encounter, visit, or admission number for the
	patient's hospital stay
Diagnosis	Significance in testing: Knowing the patient's diagnosis can save time by eliminating repeat testing when the initial results are unusual
	Example: Patients with Multiple Myeloma may be receiving Daratumumab/Dazalex (DARA), which binds to antigen CD38. This antigen is present in, in smaller quantities, on the surface of red blood cells including blood bank reagent red cells. This causes results for antibody screens and antibody IDs to be panreactive and requires Dithiothreitol (DTT) treatment to determine if the patient has any underlying allo-antibodies. Knowing the patient's diagnosis along with their treatment saves the blood bank tech time and resources during testing
	How to complete: Indicate the major underlying diagnosis. Please, do not use "anemia." Examples include Multiple Myeloma, AML, etc.
Medications	Significance in testing: Information about medications and pregnancy status can help to focus the investigation whenever the results are unusual.
	Example: WinRhoD in the medication list, together with a diagnosis of thrombocytopenia, ITP, can be a strong predictor of anti-D in a D+
	How to complete: List all current and recent medications, especially Rh Immune Globulin, IVIG, and other monoclonal antibody therapies. Provide pregnancy information, if applicable
Transfusion History	Significance in testing: Information about previous transfusions determine the type of procedure that can or cannot be performed Example: Autologous vs allogeneic (differential) absorptions. Autologous adsorptions and routine phenotype cannot be performed if the patient has been transfused within the past 3 months
	How to complete: Indicate "Y" if the patient has ever received a prior blood transfusion. Of all prior transfusions, enter the number of transfusions received in the last 90 days. Indicate the date (MM/DD/YYYY) of the last transfusion

Continued on next page



Field Title	How the information you supply is used to focus testing efforts
Transfusion Reactions	Significance in testing: Transfusion reactions can help to focus the investigation whenever the results are unusual Example: The presence of anti-E was detected by Gel and PEG-tube methods. The hospital reported transfusing E- blood, but the patient still had a hemolytic transfusion reaction. The sample was tested again by extended incubation and enzyme methods, which then detected anti-c. Transfusion with E- c- units resulted in no further transfusion reactions. How to complete: Determine if patient has experienced transfusion reactions and classify the type of reaction. Enter post-transfusion bilirubin, if
	available. Significance in testing: Information about previous antibodies may determine the type of testing that should be performed and may influence the transfusion recommendations.
Previous Antibodies	Example: Patient history indicates previous anti-Jk ^a and anti-E. Testing would proceed for other antibodies, and a transfusion recommendation would be made for the known and newly detected antibodies. How to complete: Select antibodies previously identified for that patient,
	e.g., anti-K, -E. Use <i>Other non-listed</i> to indicate other specifies not listed. Example anti-V