REQUEST FOR BLOOD AND TRANSFUSION TESTING **Transfusion Services Laboratory**

 2^{nd} Person performing Positive Patient ID

Bloodworks Labs	☐ CENTRAL PH. (206) 689-6525
	OVERLAKE PH. (425) 467-3374
LABORATORY SERVICES	☐ EVERGREEN PH. (425) 434-4949
	☐ SKL PH. (425) 656-7900

FAX (206) 343-1780

FAX (425) 688-5031

FAX (425) 899-7524

921 Terry Ave, Seattle, WA 98104

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PATIENT INFORMATION										
HOSPITAL/INSTITUTION			ORDERING PHYSICIAN			DATE				
Name on Sample LAST	ame on Sample LAST FIRST MIDDLE			LE						
						PERSON COMPLETING REQUEST				CONTACT PHONE
						TENSON COMPLETING REGOLDS				
MEDICAL RECORD NUMBER						DIVENUES				
WEDICAL RECORD NOWBER						DIAGNOSIS:				
Social Security Number		Sex (M/F)	Date of Bir	rth (mm/	/dd/yyyy)	TYPE OF IMPEN	DING PROCI	EDURE/SI	JRGERY	(if applicable):
PRIORITY										
☐ Emergency Crossr	matched / Testir	ng requested S	TAT		FOR BLOO	DWORKS USE ONI	LY			
Emergency Uncro		men required)			BW TECH	READ-BACK OF PHO	NE ORDER VER	IFIIED BY	ORDER#	ŧ
-1 1- 6					APPROVED BY	1		TIME REC	EIVED AT	BW
Planned Transfusi	on: Date:		Time:							
☐ Routine-(Release within 4 hrs after receipt of specimen at BW)										
□ Patient Waiting in clinic (2hr)										
Patient Profile To set up BW patient profile only, not for placing component order. (Check applicable boxes below, see back of from)										
	rradiated 🗆 Le	ukocyte Reduc	ed 🗆 CMV I	Negative	e 🗆 Plasm	na Reduced 🗆	Washed	□ CD3	38	
TESTING REQUEST	ED									
☐ Type and Screen – Required for Transfusion (valid for 3 days) ☐ Extended I					Postnatal Profile (for Mother) Fetal Bleed Screen to dose Rh immune globulin					
☐ HOLD specimen (valid for 3 days) ☐ Extended			Postnatal Profile (for Baby)							
☐ ABO/RhD only ☐ HSCT cross				smatch - Recipient (Enter donor information below) (RF12)						
☐ Antibody Screen ☐ HSCT cros.				smatch -Donor (Enter recipient information below) (RF13)						
□ Direct Antiglobulin Test (DAT)										
☐ Prenatal Profile				Name:		MRN:				
SPECIAL REQUIREM	MENTS / PRO	CESSING (C	neck all th		require	4)				
SPECIAL REQUIREMENTS / PROCESSING (Check all that are required				-	_					
	, , ,			ished (1st ord Ision Service	☐ HbS Neg (Sickle Cell Disease / < 4 Months)			☐ Plasma Reduced Platelets		
Reduced provided if not available) Transfusion Service MD Approval) Disease / < 4 Months) Platelets										
	(e				-		1			
Red Blood Cells, Leukocytes Reduced	RBCs	RBCs FOR HAEMOBANK CUSTOMERS: Check if RBCs needed if not electronic crossmatch (Remote Allo eligible				•	INFANT RBC DIVIDED Units held for one patient. Assigned Aliquots (~30-50 mL)			
Plasma	Adult(~25	Adult(~250mL)Pediatric Plasma (~50m			ma (~50m	L)	Pooled Plasma (Specify volume needed)L			
Platelets, Adult Dose					FOR BLOODWORKS USE ONLY					
Leukocytes Reduced ———Adult Dose					Trip#		SPECIMEN#			
Cryoprecipitate Adult dose (5 units/pool) Cryo- Single Unit		Init (~20mL)								
Other (see back of form)										
Specimen Draw for Testing				HX □ Tech ID: Date:						
Draw Date:	Draw Tir	ne:	Comments:				ABO/RhD: Hx of Ab:			
X	x					Last panel date:				
Person Collecting Specimen and performing										
Positive Patient ID										

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Explanation of items on the front of the form

Patient Profile: no current blood/component order, set up BW Patient Profile only with special attributes:

- Check the Patient Profile box to set up a patient profile in the BW computer database to establish required special attributes/modifications (e.g. CMV negative, irradiated, volume reduced, washed) for a particular patient for future blood component orders. This option is provided to allow required special attributes/modifications to be defined for a patient before components are ordered to help ensure future component orders include the correct required special attributes. All future orders will contain these special attributes/modifications even if they are not indicated on the orders. An attribute/modification may be removed from a patient's record only by a specific order from the patient's medical provider to BW to remove the attribute/modification.
- Indicate special attributes/modifications required on the Patient Profile by checking all applicable boxes listed in the Patient Profile box.
- The patient identification information entered into the BW Patient Profile will be exactly as received on this request.

Other:

Write the product name or abbreviation in the "OTHER" box on the front of form.

Indicate the number of units required.

Place a check mark in the appropriate boxes for any required attributes.

The Following "Other" components require type and crossmatch testing:

- Granulocytes (GRANS): Must receive Prior Bloodworks approval
- Red Blood Cells Resuspended, Leukocytes Reduced (~ 500 mL) (RBCR): Intended for neonatal whole blood exchange only.

The Following "Other" components do not require type and crossmatch testing:

- Platelets Apheresis, Matched [HLA or Family (MAP)]: Must receive Prior Bloodworks approval.
- Pooled Plasma for Apheresis: Order by volume (0.5 to 10 L). Specific volume is required.
- Cryo-Poor Plasma Pooled (CPPP): Order by volume (0.5 to 10 L). Specific volume is required.

Attributes/Modifications:

- When attributes/modifications are selected with blood component orders, these are added to the BW patient profile.
- · All future orders will contain these special attributes/modifications even if they are not indicated on the current orders.
- An attribute/modification may be removed from a patient's record by utilizing the "Patient Profile" section on the front of this form.

Adult Standard Dose:

- Platelets
 - Apheresis Platelets = a platelet count of > 3.0 x 10¹¹.
 - o Infant platelet dose: RN will need to draw off the dose from an apheresis platelet
- Cryoprecipitate
 - "Standard Dose" is the number of units pooled in each bag to reach a Factor VIII level of at least 400 IU, and a fibrinogen at least 750 mg per dL. Per the Circular of Information, assume 80 IU of Factor VIII and 150 mg of fibrinogen for each unit of Cryoprecipitated AHF (i.e., 400 IU Factor VIII and 750 mg per dL). The standard dose will yield an estimated rise in plasma fibrinogen of 37 mg per dL.

Tests performed in Profiles:

- Type and Screen: ABO, Rh, antibody screen. Antibody identification will be reflexively performed for a positive screen to expedite blood if needed
- Prenatal Profile: ABO, Rh, antibody screen. Antibody identification if indicated.
- Extended Postnatal Profile (for Mother) Fetal Bleed Screen to dose Rh immune globulin.
 ABO, Rh, antibody screen on mother. Antibody identification and fetal bleed screen if indicated. Sample should be drawn within a few hours post-delivery
- Extended Postnatal Profile (for Baby): ABO, RhD, direct antiglobulin test. Antibody identification performed separately if indicated/requested.