

PURPOSE:

To provide instructions for receiving blood components from the Montlake Transfusion Service Laboratory (TSL). Process includes inspection of shipping container and blood component, entry into Sunquest (SQ) and loading in the Haemobank or other appropriate storage device

LOCATION

Northwest Transfusion Support Service (TSS)

PRINCIPLE & CLINICAL SIGNIFICANCE:

Receipt of blood products from Montlake TSL is achieved through observation of packaging to maintain temperature, comparison of quantities shipped against quantities received, entry of the blood component into the LIS for tracking including documented visual inspection of the blood component and placement in the appropriate storage device. Included is loading of both allocated and stock red blood cell components into the Haemobank using BloodTrack software.

POLICIES:

- Any shipments with questionable storage conditions must have the temperature verified and documented prior to accepting the shipment into inventory
- Receiving of blood components must be processed in a manner such that time out of controlled storage conditions is limited.
 - It is recommended only one component type (box) is received at a time and stock components are received separate from allocated components.
 - In the event Montlake TSL needs to be contacted for resolution of a step failure, the implicated blood component should be placed in the quarantine location of the appropriate storage device to maintain appropriate temperature of the component during resolution.
- All blood components, regardless of the type, must be received from "In-Transit" status to "available" status prior to placing in appropriate storage and/or issuing
- Red blood cell components stored in the Haemobank must be scanned in BloodTrack prior to loading into the Haemobank.
- Blood components may be shipped to Northwest Lab with or without an attached Transfusion Record.

• Two different Transfusion Records are utilized at NW campus. One is generated by Sunquest and the other by Haemobank kiosk.

Transfusion Record generated by	Generated when blood component is	Refer to
Sunquest	 NOT IN the Haemobank at the time of allocation Prints for the following: Platelets Plasma Cryoprecipitate Granulocytes RBCs allocated at Montlake prior to shipping to NW TSS 	Appendix 1
Haemobank	 IN the Haemobank at the time of allocation Prints when Haemobank remotely allocated RBCs are removed from the Haemobank to issue for transfusion 	Appendix 2

REAGENTS/SUPPLIES/EQUIPMENT:

Reagents:	Supplies:	Equipment:
NA	Absorbent Material	Shipping Container
	Plastic Liners	
	Coolants depending on components:	
	 Wet ice 	
	 Frozen coolant packs 	
	 Gel packs wrapped in bubble 	
	wrap stored at 20-24°C	
	 Dry ice 	

QUALITY CONTROL:

Shipping conditions will be monitored routinely upon component receipt and shipment

INSTRUCTIONS:

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Accepting Delivery of Blood Components

STEP	ACTION				
	Open the shipp the packing slip	•	ner and ti	me stamp or write tl	he date and time of opening on
	If Packing slip (BBR9) The				
1	Is enclosed		Go to n	ext step	
•	Not enclosed		• Doc		ask for a copy to be faxed time the box was opened on
	Verify contents shipment appea			riately to maintain re	equired shipping temperature and
	lf		Pa	cking condition	Shipping Temp Range
	Red Blood Ce	lls	W	et ice is present	1-10° C
2	Platelets, Gra			oom temperature abilizing packs	20-24°C
	Fresh Frozen Cryoprecipitat	,	Dr	y Ice is present	< -18°C
*NOTE: The temperature does not need to be taken/recorded unless the condition is not met or if the tech has reason to believe that products have transported at the temperature ranges listed above If Then					
	Shipment is acceptable	Go to step 5			
3	Temperature not maintained, shipment leaking or otherwise damaged	If Shippin tempera is in que	g • iture	temperature by p in the middle of t component in a s component, the p the two blood co after 3-5 minutes component, plac component and s	prated thermometer to verify the placing the thermometer probe he component and fold the sandwich. If more than probe can be placed between mponents. Read temperature s. For a single frozen e probe between the Styrofoam protector t temperature or other shipment

STEP			ACTI	ON
	lf		Then	
4	Temperature is acceptand shipment is othe acceptable	rwise	Go to next s	step hift lead or manager and complete QI
	acceptable, shipmen leaking or otherwise damaged		Report	Montlake TSL to coordinate resolution
5		nbers mate	ch and all cor	nose listed on the packing slip and verify nponents are accounted for is noted
	If component is	Then		
	Red cell or platelet			radiated or psoralen treated
			omponent is diated	Then The blood component label
		mac	lateu	 The blood component label must state "Irradiated"
				VOLUNTEER DONOR
				E0379V00
				RED BLOOD CELLS ADENINE SALIVE (AS - 3) ADDED IRRADIATED LEUKOGYTES REDUCED From 500 mL CP2D Whole Blood Store at 1 to 6 C
6				 And the irradiator Rad Sure indicator if present appears as the following
				RAD-SURE™ OPERATOR: DATE:/_/_ 25 Gy INDICATOR IRRADIATED Lot No: XXXXXXXXXXX Exp. XXXXX
				NOTE: If the indicator displays " NOT " then the unit is not irradiated
				RAD-SUREIM OPERATOR 370 DATE ////8/2/ RAD-SUREIM OPERATOR 370 DATE ////8/2/ XR 25 Gy INDICATOR CONT IRRADIATED LOT NO 035293XR25 LOT NO 035293XR25 LOT NO 035293XR25

STEP	ACTION				
	Plasma or	Cor not Qua refe Cor Cor Go	irradiated or pa ality Improveme	state "Psora WOLUNTER FERANTING PLATELETS	Sure Irradiator indicator present ately if blood component is ed and document on a arantine the product and Final Disposition of Blood
	cryoprecipitate				
7	If component is For stock Allocated to a	Then Go to step Go to step			
	 Verify the following information matches between the component label, Transfusion Record and Unit Compatibility Label adhered to the component Donor Unit # Division (DIV) Expiration Date Verify the following information on the attached Transfusion Record matches the Unit Compatibility Label Medical Record Number Patient Full Name 				
8	Componer	nt Label	Unit Com Lat		Transfusion Record
	Donor	Unit #		or Unit #	Donor Unit #
	Divisio	n (DIV)	Divisi	on (DIV)	Division (DIV)
	Expirati	on Date		tion Date	Expiration Date
			Nu	al Record mber	Medical Record Number
			Patient	Full Name	Patient Full Name

STEP	ACTION				
	If information	Then			
	Matches Go to next step				
	Not Match Call Montlake TSL for resolution				
9	Initial the packing list and file in the appropriate location.				
10	Go to next section – <u>Receiving Blood into Sunquest Inventory</u>				

Receiving Blood into Sunquest (SQ) Inventory

STEP	ACTION				
1	Open Sunquest (SQ) function and log into location NW				
2	Click on 'Blood Status Update'				
3	Select < In-Transit to Inve	ntory> from the drop-down menu in the "Update Option" field			
	Scan the appropriate infor	mation in the appropriate fields			
	Field	Scan			
	Unit #	Donor ID Number barcode from component label			
4	Component code	Ecode barcode			
	Division #	Select or verify the correct division code, if applicable			
	NOTE : The component code should be scanned to ensure the correct component type is listed, even if it prepopulates upon scanning the unit number				
5	Tab through the date and time to enter the current date/ time, or manually enter the correct date/time if necessary				
	 Press <tab> to enter "INV ~Inventory" as the default in the "New status" field</tab> Press <tab> again and a "Temperature field" will open – do not enter temperature data</tab> 				
6	alert the user if the temper	erature data in this field. Sunquest does not have logic to rature is out of range. If there are concerns regarding product to SOP <i>Quarantine and Final Disposition of Blood</i> est <i>Campus</i>			
7	Press Tab and the "Pass v	visual inspection □Yes □No" will appear			
8		n and document the results of the inspection - refer to SOP od Components Northwest Campus			

STEP	ACTION					
	If visual inspection	n Select the following for				
	Passes	□ <u>Y</u> es				
	Fails	□ <u>N</u> o Document the reason for failure and quarantine the component - rrefer to SOP Quarantine and Final Disposition of Blood Component at Northwest Campus: Appendix A Quarantine and Discard Reason Codes				
9	 Click <<u>9</u>. Unit Location> NWBB Verify the components are listed in the correct inventory destination Click <ok></ok> 					
10	Click < <u>S</u> ave> at the	bottom of the screen to complete the transfer				
	If component is	Then				
11	 ALLOCATED with attached SQ Transfusion Record Select 'Allocated' from the New Status dropdown box wher Unit activity window opens Click <save></save> NOTE: If 'Released" status is selected in error contact Montlak TSL for resolution 					
	UNALLOCATED	Go to next step				
12	Repeat steps 4-11 for each additional unit					
	Place the blood components in the appropriate storage device refer to SOP Blood Storage and Inventory Management at Northwest Campus					
	If component is	Then				
	Frozen Plasma and Cryoprecipitate	Place in Blood Component freezer				
40	Red Blood Cells	Go to section <u>Loading Components into the</u> <u>Haemobank</u>				
13	Platelets	Place in Platelet Incubator				
	Washed Red blood Thawed Plasma	cells or Place components on allocated shelf of blood refrigerator				
	Granulocyte	Store in the shipping container it was delivered in				
	Neonate RBC (O n <7days old,≤3 day Hgb S negative)					

Scanning Blood Components into BloodTrack

STEP	ACTION				
1	Open BloodTrack software from Citrix Receiver				
2	Click on <transactions></transactions>				
3	Log in by scanning your UWMC ID Badge or entering in your EID# (Employee Identification #)				
4	Click on <activate out=""></activate>				
	Answer the question "Do you want to add patient details?				
	If component is Select				
5	Not Allocated - refer to Appendix 1• No Go to next stepAllocated with SQ Transfusion Record attached - refer to Appendix 2• Yes Go to step 7				
6	 The activate out dialog box will open to enter component information Select <cooler> from the Transport Method dropdown box</cooler> Scan the following information from the blood component ISBT label in the appropriate field (a picture on the screen will prompt which the barcode to scan in each field) Unit Number Product Code Unit Blood Group Expiration Date A green "Good" prompt will display when complete and go to step 				
	If Then Green Good • Repeat for any additional components • Go to section Loading Components in the Haemobank Red Stop Call Montlake TSL for resolution				
7	 Select <cooler> form the Transport Method dropdown box</cooler> Scan the following information from the blood component ISBT label in the appropriate field (a picture on the screen will prompt which the barcode to scan in each field) Unit Number Product Code Unit Blood Group Expiration Date 				
	\circ Expiration Date				

STEP	ACTION			
	 Medical Record Number Patient Last Name Patient First Name Review entry for accuracy and correct if necessary NOTE: Do not enter the Patient Gender, Patient Birth Date, or Patient Blood Group			
9	 Click on <execute></execute> Click <yes> when the dialog box pops up "Patient Blood Group is Empty. Do you want to continue?"</yes> A green "Good" prompt will display when complete 			
10	Repeat steps 5- 9 for any additional components			
11	Go to section Loading Components into the Haemobank			

Loading Components into the Haemobank

STEP	ACTION					
1	Log in by scanning your UWMC ID Badge or entering in your EID# (Employee Identification #)					
2	Select <putting in=""></putting>					
3	Scan the ID number of the blood product					
	window appears NOTE: This displays even	Touch < YES> when the "Temperature Indicator Check'				
4	the component. Blood components received as acceptable in SQ are then loaded in Haemobank as acceptable components. For unacceptable components- refer to SOP: <i>Quarantine and Final Disposition of Blood Components</i> <i>at Northwest Campus</i>					
5	Select <cooler></cooler>					
	If green screen	Then				
6	APPEARS prompting you to place the blood component into the storage location	Place the component into the designated location (tray will light up blue) in the storage device and close the door				
	Does NOT APPEARS	 Verify the component was entered into BloodTrack Call Montlake TSL for resolution 				
		ether another blood component will be loaded				
-	lf	Then				
7	Yes	Repeat steps 3 thru 4 to load each additional component				
	No	Go to next step				

STEP	ACTION
8	Touch <logout> when all blood components are loaded</logout>

PROCEDURE NOTES/LIMITATIONS

- For autologous or other rare or difficult to replace units, it may be necessary to preserve units that have been exposed to temperatures outside of the acceptable range. In these circumstances, the UWMC TSL Medical Director approval is required. Approval and reason for deviation to the SOP must be documented.
- The same packing processes may also be used during emergency storage events when alternative equipment storage unit is not available. Refer to SOP: Blood Storage and Inventory Management

REFERENCES:

- Technical Manual. Bethesda, MD; AABB, current edition.
- Standards for Blood Banks and Transfusion Services. Bethesda, MD; AABB, current edition.

RELATED DOCUMENTS:

SOP Visual Inspection of Blood Components Northwest Campus) SOP Blood Storage and Inventory Management at Northwest Campus SOP Quarantine and Final Disposition of Blood Component at Northwest Campus

UWMC SOP Approval:								
UWMC CLIA Medical Director								
	Andrew Bryan, MD	Date						
Transfusion Service Manager		Date						
	Nina Sen							
QA Manager		Date						
Transfusion	Tayler Reeves							
Service								
Medical Director		Date						
	Monica Pagano, MD							
UWMC Biennial Review:								
		Date						
		Date						

11/23/21- Revised to include check for irradiated or psoralen treated blood components when components are received at Northwest campus

9/23/22- Updated to reflect storage of neonate RBC in backup refrigerator

APPENDICES:

APPENDIX 1: Sunquest Printed Transfusion Record and Labeled Blood Component

Applies to platelets, thawed plasma, thawed cryoprecipitate, and red blood cell components allocated from Montlake stock prior to shipping to NW campus. **Unit Compatibility Label** will be attached to the top front of the blood component

NAME: TEST	L'ARN		MRN: 010141981			
	Patient Informatio	and the second se	Donor Information	-12	UNIT COMP/	ATIBILITY LABEL
Patient ABO/Rh	R-POSITIVE	Donor ABO/Rh	0-POSITIVE	NAME:	TEST, SEM	
Antibody Screen	NEGATIVE	Donor Unit#	W1415 20 012358	MED RECH ABO/RH:	U10141981	DONOR ABOIRTE
Location	NW2E	Component	RNC IL DIV 00	CROSSMATC	E Compatible	DONOR UNITS: 0-POSITIVE W1416 20 012358
Physician	UNKNOWN	Crossmatch	Compatible Exp 09/19/2020	DATE:	09/17/2020	Unt EXP: 10/15/2020 2359
Date	09/17/2020	Unit Expiration	10/15/2020 2359			
Accession #	W1000874	# of Units in Pool		24		
		Volume	350	IDE	V1415 20 012250 9 2	
Comments		Unit Antigens		A DECEMBER OF THE OWNER	11410 20 012000 0.2	5100
		Onit Anagens			Blood works Sentis, WA 98104	
 Patient's nar are identical 	resence: ne & medical record number on the unit compatibility	 physician and the Refer to the Nursin 	SEUSION IMMEDIATELY and call the Transfusion Service Laboratory g Blood Administration Policy us of Superiod Temperaturian Densitian Form	in p	roperly identify intended Recipient Cristics of Information for Indications, Salarry, candidors and periods of Infonian, radact may transmit intention agents. Rr Only	Rh POSITIVE
 Patient's nar are identical label, wrist b record. Donor ABO// on the transic compatibility label are ide Patient ABO compatibility special required. Unit is norm 	me & medical record number on the unit compatibility band(s), and transfusion Rh & the donor unit number fusion record, unit y label and donor unit face	physician and the Refer to the Nursin Complete the Report Draw a 6mL Pink to Send the complete sample, blood bag (remove needle), a Service as soon as	Transfusion Service Laboratory g Blood Administration Policy ort of Suspected Transfusion Reaction Form op EDTA blood sample from the patient d Suspected Transfusion Reaction form, blood with attached tubing and remaining contents nd the Transfusion Record to the Transfusion	RED BLL ADENINE- IRRADIATE LEUKOET	A CON A CON COUNTEER DONOR 3757000 000 CELLS SALINE (AS - 3) ADDED 58 REDUCED 10 CEDD IN CON	D202892359 15 OCT 2020
 Patient's nar are identical label, wrist b record. Donor ABO/l on the transi compatibility label are ide Patient ABO/ compatibility special requ verified. 	me & medical record number on the unit compatibility and(s), and transfusion Rh & the donor unit number fusion record, unit y label and donor unit face mical. /Rh, interpretation of t testing (if performed), and irements (if applicable) are	physician and the • Refer to the Nursin • Complete the Repo • Draw a 6mL Pink to • Send the complete sample, bipolod (remove needle), a Service as soon as	Transfusion Service Laboratory g Blood Administration Policy ort of Suspected Transfusion Reaction Form op EDTA blood sample from the patient d Suspected Transfusion Reaction form, blood with attached tubing and remaining contents and the Transfusion Record to the Transfusion possible.	ED RED BL ADENINE- IRADIANE LEUKOCYT From 1900	A CON A CON COUNTEER DONOR 3757000 000 CELLS SALINE (AS - 3) ADDED 58 REDUCED 10 CEDD IN CON	D202892359 15 OCT 2020
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APPENDIX 2: Haemobank *Transfusion Recorded* for stock blood components

Applies to red blood cell components remotely allocated from Haemobank inventory. Blood Component will not be labeled with a **compatibility label**.

UW MEDICINE T	RANSFUSION RECORD	
Bedside Verification Before administering the unit, verify in the patient's presence that: • Patient's name & medical record number are identical on the unit compatibility label, wrist band(s), and transfusion record. • Donor ABO/Rh & the donor unit number on the transfusion record, unit compatibility label and donor unit face label are identical. • Patient ABO/Rh, hiterpretation of compatibility testing (if performed) & special requirements (if applicable) are verified • Unit is normal in appearance & not expired. Date Time Transfusionist Witness	IF A TRANSFUSION REACTION IS SUSPECTED • STOP THE TRANSFUSION IMMEDIATELY and call the physician and the Transfusion Service Laboratory • Refer to the Nursing Blood Administration Policy • Complete the Report of Suspected Transfusion Reaction Form • Draw a 6mL Pink top EDTA blood sample from the patient • Send the completed Suspected Transfusion Reaction form, blood sample, blood bag with attached tubing and remaining contents (remove needle), and copy of the Transfusion Record to the Transfusion Service as soon as possible.	
Attach patient label here ONLY if there is no patient name or MRN above	UW Medicine Harborview Medical Center – University of Washington Medical Center UW Neighbordod Clinics – Valley Medical Center University of Washington Physicians Seattle Washington UW MEDICINE TRANSFUSION RECORD UH3919 REV AUG 20	B 1000 L 200 S