Department of LABORATORY MEDICINE

University of Washington Medical Center 1959 NE Pacific Street. Seattle, WA, 98195 Transfusion Services Laboratory Policies and Procedures Manual

Original Effective Date: Nu 03-11-16 PC Revision Effective Date:

Number: PC-0006.02

FC-0000.

TITLE: Electronic Crossmatch

PURPOSE:

To provide instructions for performing electronic crossmatch (EXM) to detect ABO/Rh incompatibility between recipient and donor red blood cell and granulocyte components

PRINCIPLE & CLINICAL SIGNIFICANCE: Principle

The LIS (Sunquest) is programmed to detect ABO/Rh incompatibility between recipient and donor. This method of crossmatching can be used in place of a serologic crossmatch when the patient and donor meet the following requirements:

- Recipient requirements
 - Two consecutive concordant ABO/Rh results on file
 - One from an non-expired crossmatch eligible battery (TSCR, TSCREX, TXM) with no pending tests
 - Second ABO/Rh from an independent collection with testing performed by the UWMC TSL
 - Current antibody screen is negative
 - No history of clinically significant antibodies
 - Electronic crossmatch eligibility (EXM Elig) is NOT set to "NO"
- Donor requirements
 - o ABO/Rh serologically confirmed and the results entered into the LIS
 - No clinically significant antibodies (any antibodies will be listed on the unit label or an attached tag and these units are not accepted by the UWMC TSL)
 - Component is in an available status with no pending tests
- Patient and donor test results and interpretations used by the computer for crossmatching are reviewed and verified by a TSL technologist at the time of testing

Clinical Significance

Advantages of the electronic crossmatch include: reduced sample volume, decreased workload, efficient use of blood inventory, reduced time for providing red blood cell or granulocyte components for transfusion, and reduced exposure of personnel to blood specimens.

SPECIMEN REQUIREMENTS:

Current (in-date) specimen includes a current ABO/Rh and negative antibody screen testing (see requirements under 'Principle' above)

REAGENTS/SUPPLIES/EQUIPMENT:

Reagents:	Supplies:	Equipment:
NA	NA	Sunquest BB Module

TITI E: Electronic Crecomotoh	Number:
	PC-0006.02

QUALITY CONTROL:

Sunquest is validated to ensure the system contains logic to alert the user to ABO/Rh incompatibility between the recipient and the donor unit

INSTRUCTIONS:

STEP	ACTION						
	Log into Sunquest using the inventory location where the component is located and the Transfusion Record will print						
	If component is located at	Then log in at					
1	UW-TSL	BB					
1	UWMC 2 nd Floor	BB2					
	SCCA	SA1					
	NW	NWBB (HaemoBank) NWBB2 (non HaemoBank)					
2	Open the Blood Order Processing (BOP) module						
	Scan/enter the MR# from the product order	or delivery request					
3	NOTE : The order may also be accessed by scanning the CID on the patient sample if performing an electronic crossmatch in conjunction with other patient testing						
4	Click on the < <u>O</u> rder Selection> tab and double click on the order line with the appropriate crossmatch eligible battery						
	Perform a patient history check (HXCK)(refer to SOP Patient History Check)						
6	NOTE: Carefully review information under the 'Problems' and 'Comments' tabs because any information listed there that disqualifies the patient for electronic crossmatch will not automatically disqualify the patient for electronic crossmatch						
	Select and allocate the red cell component						
7	 Click on <allocation> tab</allocation> Click on <blood inventory="" search=""> bit</blood> 	itton					
	Enter UR (University Routine) in the 'Search <u>F</u> unction' field to view All 'available' red cell components (RBCG) in all UW TSL inventory locations						
8	To search for red cell components in	Enter					
	2 nd floor OR inventory only	BB2 in the "Unit Location" field					
	SCCA inventory only	SA1 in the "Unit Location" field					
	NW inventory only	NWBB in the "Unit Location" field					
10	 Enter any additional search criteria as n components ABO/Rh compatible with th required attributes and antigens Click <<u>S</u>earch> 	ecessary to restrict the search to he patient and matching any additional					

TITLE: Electronic Crossmatch

STEP	ACTION						
	Select appropriate unit(s) by checking the box at the left of the unit row						
	Flag HID/Ar	ea		95	Comp	Dv	AB
			W1416 01 0005 W1416 01 0005	05 20	E0101 E0332	A0	0/1
11			W1416 01 0005	22	E0336	AO	0/1
		1	W1416 01 0005	23	F0332	00	0/1
	NOTE : Units are normally selected based on shortest expiration which sort to the top of the screen						
	Click <ok> to cor</ok>	nplete t	he allocation				
12	2 WARNING: If any QA warnings appear, the electronic crossmatch cannot be completed						
13	Verify the components selected appear in the Compatibility Testing field and "Pend" appears in the XM And TS fields						
	Click < <u>S</u> ave>						
14	WARNING: If any QA warnings appear, the electronic crossmatch cannot be completed						
	Review the "Electronic Crossmatch Eligibility Report						
	If Patient is Then						
	Eligible for	The u	nit number wi	II appear unde	er the "l	Jnit(s)) to be resulted" on
	electronic the Electronic Crossmatch Eligibility Report						
	crossmatch	CA	x				×
			Ele	ctronic Crossmatch	n Eligibility	Report	
				W1479	99		
		Unit(s) to be resulted W1416 01 000365 E4527:00					
			*** END OF REPORT ***				
15					UK		
		• CI	ick <ok>. Tł</ok>	e crossmatch	grid wil	l be a	utomatically
		populated (see table below)and the Transfusion Record will print at the location selected in Step 1					
			Field	Result			
			XM	ECMP			
			TS	OK			
		NOTE prints	Refer to 'Pr at the wrong	ocedure Notes location	s' below	if the	Transfusion Report

TITLE: Electronic Crossmatch	Number:
	PC-0006.02

STEP	ACTION					
	Not eligible for electronic crossmatch	 Ine Electronic Crossmatch Eligibility Report will list the reason(s) for the electronic crossmatch failure Example: Filectronic Crossmatch Eligibility Report will list the reason(s) for the electronic crossmatch Eligibility Report will list the reason(s) for the electronic crossmatch Eligibility Report will list the reason(s) for the electronic crossmatch Eligibility Report will list the reason(s) for the electronic crossmatch Eligibility Report will list the reason(s) for the electronic crossmatch Eligibility Report will list the reason(s) for the electronic crossmatch Eligibility Report will list the reason(s) for the electronic crossmatch Eligibility Report will exit save? Click <cancel>. The Blood Order Processing box will appear with the message "CAX was canceled, will exit save"</cancel> Blood Order Processing (CAX was canceled, will exit save) Click <ok></ok> Investigate and resolve the cause of the failure Proceed to serologic crossmatch testing if the patient does not 				
		NOTE: Any QA failu electronic crossmate	res on the component or the patient will fail the			
16	Click on the appropriate button when the "Call BPI" (Blood Product Issue) pop up box appears:					
	If the compone	omponent will be issued Then click on				
	Immediately		< <u>I</u> ssue> (refer to SOP <i>Issuing Blood</i> Components)			
	Later or at a diffe	rent location	< <u>N</u> o>			

CALCULATIONS/INTERPRETATIONS/RESULTS REPORTING/NORMAL VALUES/CRITICAL VALUES

CALIBRATION: NA

PROCEDURE NOTES AND LIMITATIONS:

Procedure Notes

- The presence of rouleaux in not associated with antibody formation and does not disqualify a patient from electronic crossmatch
- Electronic Crossmatch may not be used on patients when the Electronic crossmatch eligibility (EXM Elig) is set to NO in Blood Bank Administrative Data Entry. This safeguard is used when there is some question regarding unexplained reactions with the patient's plasma thought to be due to developing antibodies that may be clinically significant and prevents electronic crossmatching
- To reprint a Transfusion Record printed at the wrong location, log into 'Blood Order Processing' (BOP) using the correct inventory/print location and use code ;UR in the 'Add Unit Test (<u>x</u>)' field for the selected blood component
- If the patient does not qualify for electronic crossmatch, refer to the appropriate SOP for serologic testing or issuing uncrossmatched blood as appropriate
- Electronic crossmatch is also referred to as computer-assisted crossmatch (CAX) in Sunquest

Limitations

- Patients lacking two consecutive ABO/Rh types in agreement are not eligible for electronic crossmatch until a second matching ABO/Rh is resulted. This will occur following any ABO/Rh tests resulted as 'BBCAN'. Rh negative patients receiving Rh positive units will not qualify for the electronic crossmatch
- Patients in SQ BAD file listed as NTD Pos/Neg and NTD interpretation in the specimen ABO/Rh test result are not eligible for electronic crossmatch. A serologic crossmatch must be performed
- Only RBCs in "Available Status" without pending testing are eligible for electronic crossmatch
- Serologic crossmatch must be performed on units issued during computer downtime

REFERENCES:

- Technical Manual. Bethesda, MD: AABB Press, current edition
- Standards for Blood Banks and Transfusion Services. Bethesda, MD: AABB Press, current edition
- Guidance for Industry: "Computer Crossmatch" (Computerized Analysis of the Compatibility between the Donor's Cell Type and the Recipient's Serum or Plasma Type) U.S. Department of Health and Human Services, Food and Drug Administration Center for Biologics Evaluation and Research, April 2011
- Guidelines for Implementing an Electronic Crossmatch. Bethesda, MD; AABB, 2003

RELATED DOCUMENTS:

SOP Sample Acceptability& Order Receipt SOP Issuing Blood Components

APPPENDIX:

NA

TITI E: Electronic Crocomotoh	Number:
TTLE. Electronic Crossination	PC-0006.02

UWMC Biennial Review:					
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REVISION HISTORY:

01/15/21: Add NW Sunquest login and inventory locations