

TRAINING UPDATE

Lab Location: SGAH and WAH **Date Implemented:** 11.8.2012
Department: Blood Bank **Due Date:** 11.30.2012

DESCRIPTION OF PROCEDURE REVISION

Name of procedure:
HLA Matched / Crossmatched Platelet Pheresis Products
Description of change(s):
<ul style="list-style-type: none">• Added definitions for HLA-matched and crossmatched platelets• Changed HLA testing lab from Quest to ARC; added instructions for sending testing to ARC• Removed appendices instructing techs how to determine HLA match, because ARC no longer labels platelets with the HLA type.• Added appendix for matched support pathway (ARC flowchart)

EMPLOYEE SIGNATURES

I have read and understand the procedure described above:

Name	Signature	Date
------	-----------	------

Non-Technical SOP

Title	HLA Matched/Crossmatched Platelet Pheresis Products	
Prepared by	Leslie Barrett	Date: 1/21/2009
Owner	Stephanie Codina	Date: 5/11/2010

Laboratory Approval		
Print Name and Title	Signature	Date
<i>Refer to the electronic signature page for approval and approval dates.</i>		
Local Issue Date:		Local Effective Date:

12 month (or new) management review and approval: Signature acknowledges SOP version remains in effect with NO revisions.		
Print Name	Signature	Date

Form revised 3/31/00

TABLE OF CONTENTS

1. PURPOSE.....	2
2. SCOPE	2
3. RESPONSIBILITY.....	2
4. DEFINITIONS.....	2
5. PROCEDURE.....	3
6. RELATED DOCUMENTS	5
7. REFERENCES	5
8. REVISION HISTORY.....	5
9. ADDENDA AND APPENDICES.....	5

1. PURPOSE

HLA matched or crossmatched apheresis platelet products are used to treat patients that have become refractory to platelet transfusions due to the presence of HLA or platelet specific antibodies.

2. SCOPE

Platelets bear a variety of antigens, including HLA and platelet-specific antigens. Patients who have been transfused may develop antibodies towards platelet antigens. When platelets are transfused to a patient with an antibody directed towards an antigen expressed on the platelets, the survival time of the transfused platelets may be markedly shortened. Matching the HLA antigens on the platelets to the HLA antibodies that the recipient possesses or performing a platelet crossmatch may help to provide platelets with longer periods of survival. Both crossmatched and HLA-matched platelets require additional time for preparation and orders should be coordinated with the blood supplier.

3. RESPONSIBILITY

All Blood Bank staff must understand and follow this procedure when HLA-matched and/or crossmatched platelets are requested for a patient.

4. DEFINITIONS

Platelet Transfusion Refractoriness - When the recipient experiences a “less-than-expected” increase in platelets following a platelet transfusion. Platelets can be destroyed by immune mechanisms (HLA and platelet-specific antibodies) or non-immune mechanisms (splenomegaly, sepsis, fever, intravascular devices, and DIC). A 1-hour post-transfusion platelet count should be used to differentiate between immune-mediated and non-immune-mediated destruction. Immune refractory states will demonstrate poor recovery in the early post-infusion interval.

Form revised 3/31/00

Corrected Count Increment (CCI) - A calculation used to determine the recipient's response to platelet transfusion. A 1-hour CCI >7500 indicates an adequate response to platelet transfusion, while a CCI <5000 on two separate occasions indicates platelet refractoriness. CCI values that are adequate 1 hour post transfusion and continue to decrease are more suggestive of non-immune causes (splenomegaly, fever, infection, DIC, amphotericin B use, bleeding, etc.). Neither crossmatched nor HLA-matched platelets are needed for non-immune platelet destruction.

$$CCI = \frac{\text{Body Surface Area (m}^2\text{)} \times (\text{Post Transfusion Platelet Count} - \text{Pre Transfusion Platelet Count}) \times 10^{11}}{\text{Number of Platelets Transfused}}$$

Crossmatched Platelets – This test is performed similarly to crossmatching for red cells. Donor platelets and recipient plasma are tested against each other. Donor platelets that react with the recipient plasma are considered incompatible. Donor platelets that do not react with recipient plasma are considered compatible. Crossmatched platelets may be used for brief support of the patient. However, results may not be accurate and testing relies upon a large supply of donor platelets which is not always available. All platelet crossmatch orders should be followed with HLA typing of the recipient for long-term platelet support.

HLA-Matched Platelets – Both the donor and recipient are tested for HLA antigens. The antigens are matched for HLA-A and-B locus as best as possible. If the patient has HLA antibodies, donor platelets that lack the corresponding antigen are also selected. HLA-matched platelets provide more successful transfusion response in up to 75% of patients with immune-mediated refractoriness.

5. PROCEDURE

Step	Action
1	All initial requests for HLA-matched and crossmatched platelets should be approved by the Blood Bank Medical Director or clinical pathologist on-call. <ul style="list-style-type: none"> A. Approval or rejection should be documented in the Blood Bank Communication Log. Include the date, time, and pathologist's name. B. Once approved, all subsequent platelet transfusion will meet the HLA-matched or crossmatched platelet transfusion criteria. C. Crossmatched and HLA-matched platelet products are not normally stored in inventory and will take extra time to prepare. <ul style="list-style-type: none"> a. Crossmatched platelet products take approximately 24 hours from the time the specimen is received by the reference laboratory. b. HLA-matched platelet products take approximately 5 days from the time the specimen is received by the reference laboratory.
2	Document the need for special platelet transfusion by placing a comment in the patient's blood bank historical data. Refer to procedure, "Entering Special Transfusion Attributes into the LIS." <ul style="list-style-type: none"> A. Enter ";HLA" for the comment "Irradiated and HLA matched platelets required." B. Enter ";CIRPP" for the comment "Patient requires crossmatched, irradiated platelet products."

From revised 3/31/00

Step	Action	
3	<p>An HLA type and antibody screen should be referred to the American Red Cross (ARC) the first time HLA-matched platelets are requested.</p> <ul style="list-style-type: none"> A. Complete a "Hisotcompatibility Testing Requisition" form. <ul style="list-style-type: none"> a. For HLA typing, order test HLA 072, HLA-A and B Typing For Platelet Transfusion b. For HLA antibody screen, order test HLA 025, HLA Antibody Screen for Transfusion, Class I c. Order both tests for refractoriness. B. Have the appropriate specimen collected from the patient (1 tube per test). <ul style="list-style-type: none"> a. HLA typing uses EDTA or ACD whole blood b. HLA antibody screen uses serum without clot activator or gel separator C. Notify the ARC reference lab that the testing is coming D. Send the specimens to the ARC reference laboratory 	
4	<p>Complete the "American Red Cross Special Products Request Form." Submit the order form and specimen (if applicable) to the ARC Reference Laboratory.</p> <ul style="list-style-type: none"> A. Complete the Customer Information section of the form. <ul style="list-style-type: none"> a. Facility name b. Name of person completing form c. Phone number d. Fax number B. Complete the Patient Information section of the form. <ul style="list-style-type: none"> a. Patient name b. HLA type (if known) c. ABO/Rh d. Medical record number e. Date of birth f. Physician's name g. Diagnosis C. Complete the Product Criteria section of the form with the special requirements being requested. Request that ARC irradiates crossmatched and HLA-matched platelets prior to shipment. For crossmatched platelets, send one 7mL EDTA tube stored at room temperature. 	
5	<ul style="list-style-type: none"> A. When HLA results are returned, enter the patient's HLA type into the blood bank historical data file. B. Fax a copy of the result to the ARC order management department to obtain HLA-matched platelet products. 	

Step	Action
6	The blood supplier will send the platelets when available. A. All HLA-matched and crossmatched platelet products MUST be irradiated prior to transfusion. Irradiate upon arrival if ARC did not irradiate prior to shipment. B. HLA-matched platelets are entered into the computer using product code "PPHL." C. Crossmatched platelet products must be billed at the time of issue using code "PLAXM."

6. RELATED DOCUMENTS

Form: American Red Cross Histocompatibility Testing Requisition

7. REFERENCES

1. Roback, J.D., Combs, M.R., Grossman, B.J., Hillyer, C.D. 2008. Technical Manual of the AABB, 16th ed. AABB Publishing, Bethesda, Maryland.
2. Standards for Blood Banks and Transfusion Services, 2009. AABB, 26th ed. AABB Publishing, Bethesda, Maryland.
3. AABB, ABC, ARC, and ASBP. 2009. Circular of information for the use of human blood and blood components.
4. Vassallo, R. R. 2008. Changing Paradigms in Matched Platelet Support. American Red Cross.

8. REVISION HISTORY

Version	Date	Reason for Revision	Revised By	Approved By
		Supersedes SOP WAH-SGAH B321.002		
000	5/11/2010	Updated owner Section 4: Definitions added Section 5: Computer process revised to reflect computer upgrade Section 9: added Addenda A	S. Codina	N. Cacciabeve
001	10.12.12	Section 4: Added definitions for HLA-matched and crossmatched platelet products. Section 5: Changed HLA testing lab from Quest to ARC. Section 9: Removed appendices instructing techs how to HLA-match platelet products. Added Appendix for Matched Platelet Support Pathway.	SCodina	NCacciabeve

9. ADDENDA AND APPENDICES

A: Matched Platelet Support Pathway

Form revised 3/21/09



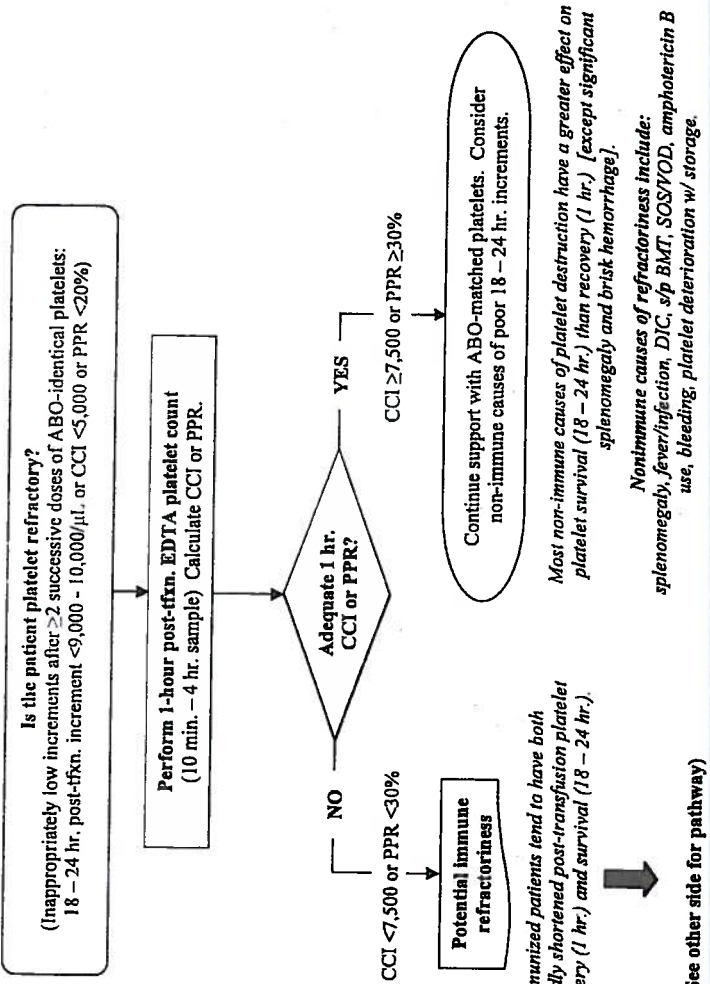
**Matched Platelet
 Support Pathway**
 v. 03132008

$$\text{PPR (\%)} = \frac{\text{Percent Predicted Recovery (PPR)}}{\text{observed increment } (x10^9/L)} = \frac{\# \text{ pls. transf. } (x10^9)}{\text{blood volume (L)}}$$

Gilcher's Rule of 5s for blood volume (mL/kg):

	Obese	Thin	Normal	Muscular
Male:	60	65	70	75
Female:	55	60	65	70

$$\text{CCI (m}^3\text{/}\mu\text{L/10}^4\text{)} = \frac{\text{Corrected Count Increment (CCI)}}{\text{increment } (\mu\text{L}) \times \text{BSA (m}^2\text{)}} = \frac{\# \text{ pls. transf. } (x10^4)}{\text{BSA (m}^2\text{)}}$$



Most alloimmunization occurs against Class I Human Leukocyte Antigens (HLA-A & HLA-B loci), with occasional Human Platelet Antigen (HPA) co-immunization, or rarely, HPA-only alloimmunization (e.g., HPA-1b, HPA-5b, etc.).

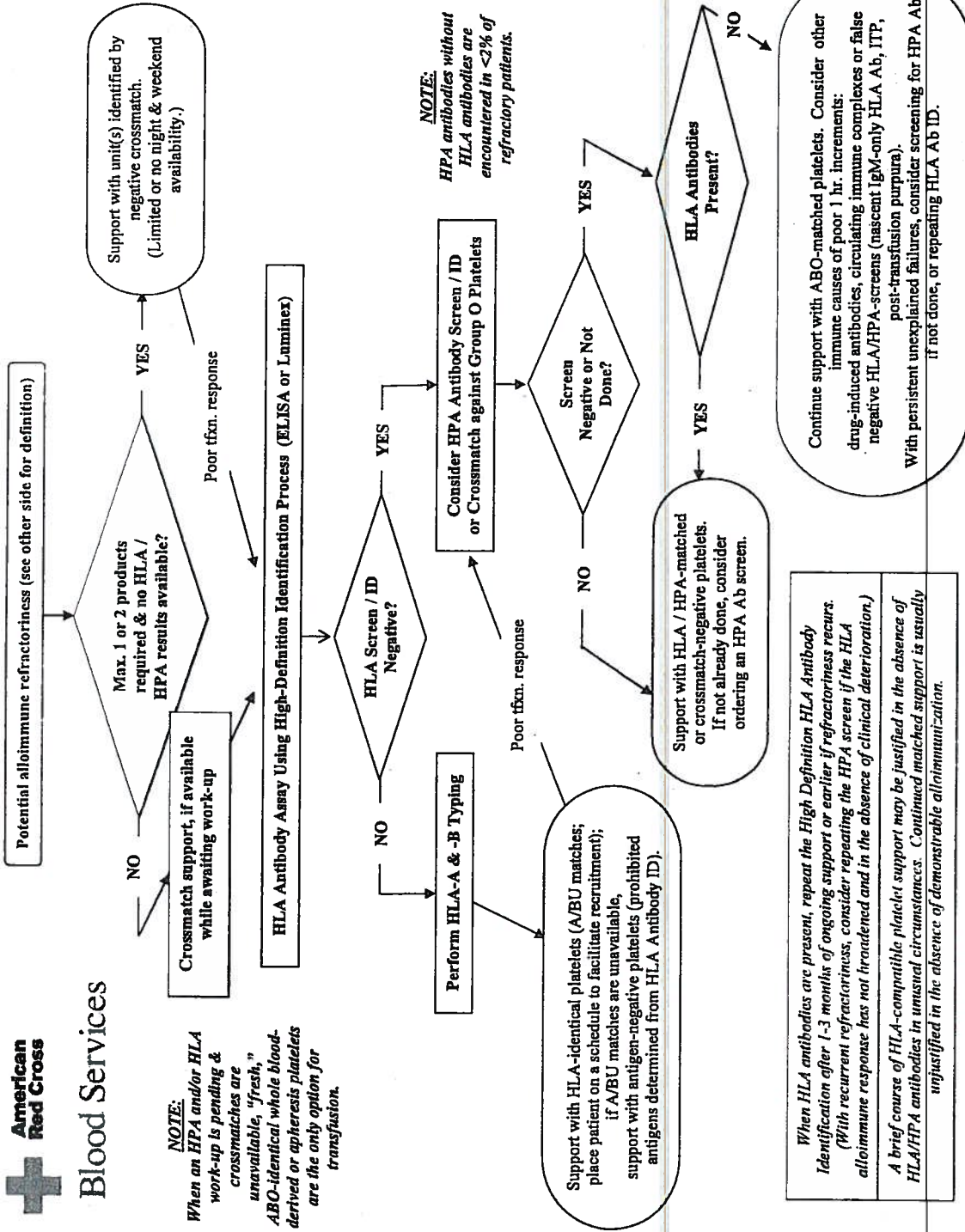
Likelihood of successful transfusion for HLA-alloimmunized patients (in descending order):

1. HLA-identical platelet selection (A/BU matches when available; selective recruitment for pts. w/ broad alloimmunization)
2. HLA antigen-negative units or platelets compatible by crossmatching
3. HLA type-selected non-identical matches (well-chosen BX, BUX and C matches)
4. ABO-identical whole blood-derived or apheresis platelets when HPA and/or HLA work-up is pending & crossmatches are unavailable



Blood Services

NOTE:
 When an HPA and/or HLA work-up is pending & crossmatches are unavailable, "fresh," ABO-identical whole blood-derived or apheresis platelets are the only option for transfusion.



NOTE:
 HPA antibodies without HLA antibodies are encountered in <2% of refractory patients.

When HLA antibodies are present, repeat the High Definition HLA Antibody Identification after 1-3 months of ongoing support or earlier if refractoriness recurs. (With recurrent refractoriness, consider repeating the HPA screen if the HLA alloimmune response has not broadened and in the absence of clinical deterioration.)
 A brief course of HLA-compatible platelet support may be justified in the absence of HLA/HPA antibodies in unusual circumstances. Continued matched support is usually unjustified in the absence of demonstrable alloimmunization.