


<u>Name of Policy: Blood and Component Label Verification</u>  <u>Policy Number: 3364-108-204</u>  <u>Approving Officer: Senior Hospital Administrator Director, Blood Transfusion Service</u>  <u>Responsible Agent: Blood Transfusion Service Supervisor Administrative Director, Lab</u>  <u>Scope: University of Toledo Medical Center Pathology/Laboratory – Blood Bank</u>		  <u>Effective date: 03/01/2025</u>  <u>Original effective date: 06/1996</u>	
<u>Key words: Blood, Component, Label Verification, Blood Label, FFP, Cryo</u>			
<input type="checkbox"/>	<u>New policy proposal</u>	<input checked="" type="checkbox"/>	<u>Minor/technical revision of existing policy</u>
<input type="checkbox"/>	<u>Major revision of existing policy</u>	<input type="checkbox"/>	<u>Reaffirmation of existing policy</u>

(A) ~~(A)~~ — Policy Statement

The Blood Transfusion Service labels modified and “crossmatched” blood and components accurately, by a uniform procedure and with a second verification of accuracy of product labels.

(B) ~~(B)~~ — Purpose of Policy

To provide accurate, uniformly labeled blood components for distribution to patients.

(C) ~~(C)~~ — Procedure

- ~~1.~~ (1) Attach labels used as secondary bag labels to the primary bag label. Avoid obscuring FDA-mandated portions of the primary label (Name of component, Instructions to the transfusionist, “Volunteer Donor,” etc.)
- ~~2.~~ (2) Label all blood and blood components according to the appropriate format in Procedure Manual Section 400.
- ~~3.~~ (3) When blood or components are modified requiring a new ABO/Rh type label, as for ~~thawed/pooled~~ components, a second check shall be done to compare and verify that the following information is correct on the blood unit label and the compatibility label: unit source number ~~or UTMIC assigned number~~, ABO/Rh type assigned, product code, expiration date and time, UTMIC component preparation product label.
- (4) The unit bag labels shall be compared to the information on the Transfusion Record and the compatibility label at the time of product preparation/ crossmatch by scanning bar-codes for confirmation. The information shall be compared and verified a second time at product issue by scanning bar-codes for confirmation.

(D) References

- 4.———Current Edition, Standards for Blood Banks and Transfusion Services, AABB.



<del>—Danielle Weilmann, MT(ASCP)</del>	
Policies Superseded by This Policy:	

Reference: ~~—— Current Edition, Standards for Blood Banks and Transfusion Services, AABB.~~