TRAINING UPDATE

Lab Location:

SGAH and WAH

Date Implemented:

11.20.2012

Department:

Blood Bank

Due Date:

12.3.2012

DESCRIPTION OF PROCEDURE REVISION

Name of procedure:

Circular of Information for the Use of Human Blood and Blood Components

Description of change(s):

- The circular is approved by the FDA as an acceptable extension to the blood product container labels
- Copies of the circular are kept in the BB for anyone who requests
- Additional copies of the circular are obtained from ARC

EMPLOYEE SIGNATURES

I have read and understand the procedure described above:

| Name | Signature | Date |
|------|-----------|------|
| L | | |

Non-Technical SOP

| Title | Circular of Information for the Use of Human Blood and Blood Components | |
|-------------|--|----------------|
| Prepared by | Stephanie Codina | Date: 11.20.12 |
| Owner | Stephanie Codina | Date: 11.20.12 |

| Laboratory Approval | | | |
|---|-----------------------|------|--|
| Print Name and Title | Signature | Date | |
| Refer to the electronic signature page for approval and approval dates. | | | |
| | | | |
| | | | |
| 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 |
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1. PURPOSE

Blood and component labels contain critical but minimal information about the contents. A current Circular of Information is an extension of the container labels. It describes each component available for transfusion and gives indications and contraindications for use.

2. SCOPE

The circular prepared jointly by AABB, American Red Cross, America's Blood Centers, and the Armed Service Blood Program will be utilized by the Blood Bank.

3. RESPONSIBILITY

All blood bank staff members must be familiar with the contents and location of the Circular of Information pamphlets.

4. **DEFINITIONS**

None

5. PROCEDURE

| Step | Action | | |
|------|--|--|--|
| 1 | The Circular of Information for the Use of Human Blood and Blood Components is approved by the FDA as an acceptable extension to the blood product container labels. | | |
| 2 | Copies of the pamphlet will be maintained by blood bank and will be provided upon request. | | |
| 3 | Additional copies of the pamphlet may be obtained from the blood supplier. | | |

6. RELATED DOCUMENTS

None

7. REFERENCES

None

8. REVISION HISTORY

| Version | Date | Reason for Revision | Revised By | Approved By |
|---------|------|------------------------------|---------------|----------------|
| | | Supersedes WAH-SGAH B314.001 | | |
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9. ADDENDA AND APPENDICES

None