

To: Transfusion Services Managers **From:** Hospital Relations Department

Date: March 13, 2020

Re: 32nd Edition AABB Standards Update

Update:

Due to unexpected changes to inspection timeframes, the effective date has been revised to June 1, 2020.

Please accept our sincerest apologies for the short notice.

Background:

The 32nd edition of AABB standards requires the following to be met for patient testing. *Standard 5.14.5 Pretransfusion Testing for Allogeneic Transfusion*There shall be two determinations of the recipient's ABO group as specified in Standard 5.14.1. The first determination shall be performed on a current sample, the second

- determination by one of the following methods:

 1) Comparison with previous records
 - 2) Testing a second sample collected at a time different from the first sample, including a new verification of patient identification.
 - 3) Retesting the same sample if patient identification was verified using a *validated electronic identification system.*

The previous edition of standards allowed for retesting of the same sample if patient identification was verified using an electronic identification system *or another process validated to reduce the risk of misidentification.* The blood bank armband system-unless electronic and validated - will no longer qualify and consequently submission of a secondary patient sample is required.

Impact to Client:

In order to comply with the updated AABB standard, your response to the statement

Samples were collected using an electronic ID system □Yes □ No

will be required on *RTF101.01A*, *Reference and Transfusion Services Request Form*. If samples were **not** collected using an electronic system, then a 2nd sample collected at a separate phlebotomy must be submitted for testing in order to provide crossmatched products.

The second sample must be properly labeled and can be an EDTA, heparin, ACD, CPD or red-top without serum separator sample.

This change is effective July 15, 2020.