

**NEW YORK STATE DEPARTMENT OF HEALTH  
CLINICAL LABORATORY EVALUATION PROGRAM**

**Proposed Blood Services Standard  
Comment Period: September 20, 2018 to November 5, 2018**

<b><i>Blood Services</i></b>	
<b>Standard</b>	<b>Guidance</b>
<p><b>Blood Services Standard 2 (BS S2)</b></p> <p>Facilities utilizing a one-person verification process for matching recipients to the blood or blood component at the time of transfusion must:</p> <ul style="list-style-type: none"><li>a) Use an FDA-approved automated identification technology that positively identifies the recipient and matches the blood or blood component to the recipient;</li><li>b) Follow the manufacturer's instructions for the proper collection and labelling of the pre-transfusion specimen, including the placement and retention of any required secondary bar-coded wristbands;<ul style="list-style-type: none"><li>i. All required bar-coded wristbands must be placed on the patient prior to the collection of the pre-transfusion specimen;</li></ul></li><li>c) Follow the manufacturer's instructions for the automated matching of the patient to the blood or blood component prior to transfusion;<ul style="list-style-type: none"><li>i. If automated scanning mechanisms fail, including the need to perform manual data entry, or if any bar-coded identification band is removed from the patient prior to the transfusion for any reason, the facility must use a two-person patient identification process.</li></ul></li></ul>	<p>(c) The 'manufacturer' in this instance includes both the vendor of the electronic ID system and the vendor of the bar-coded wristbands, if not the same.</p>