

# BLOOD PRODUCTS: TRANSFUSION FORM (PAPER)

Specimen Drawn and Labeled By: _____		(1)	Armband ID _____	
Date:    /    /      Time:    :    am /pm			Patient Name: _____	
Patient ABO/Rh Type:	Antibody Screen:	Acct.#: _____		
Unit #:	<input type="checkbox"/> Negative <input type="checkbox"/> Positive	DOB: _____		
	<input type="checkbox"/> See Comments Below	Physician: _____		
Expiration Date:	Crossmatch:	Date: _____		(2) Product Needed:
Unit ABO/Rh Type:	<input type="checkbox"/> Compatible	Time: _____		
	<input type="checkbox"/> Incompatible	Tech: _____		
	<input type="checkbox"/> Not complete			
Comments:(Other Tests)	<input type="checkbox"/> Not Required			(3)
		<input type="checkbox"/> Red Cells, Packed <input type="checkbox"/> Platelet Pheresis (6 units) <input type="checkbox"/> Fresh Frozen Plasma <input type="checkbox"/> Other		

## Emergency Release Request

**The Following Test Cannot Be Completed Prior To The Transfusion:**

Group & Type    Crossmatch    Antibody screen or Identification

I am fully aware of the possible complications from using this blood, but because of urgency, request its immediate release.

Ordering Physician \_\_\_\_\_ am / pm      Date: \_\_\_\_ / \_\_\_\_ / \_\_\_\_

## See back of form for signs and symptoms of transfusion reaction

### Transfusion Record

Mark Box Below As Completed By Transfusionist Prior To Starting The Transfusion!

- Transfusion instruction sheet has been explained and given to patient/family member.
- Physician order has been verified for correct component and amount to be given.
- Only 0.9% Sodium Chloride, injection (USP) may be added to or infused through the same tubing as blood or components.
- I have established the identity of the patient.
- ABO/Rh type and unit number on this form are identical with the unit label.
- Patient Name and Blood Bank ID number on this form and unit label are identical to those on the Blood Bank ID band on the patient.
- Transfusion has been started within 30 minutes of release of unit from Blood Bank.

- Blood Warmer Used (Do Not Exceed 42° C) Temperature\_
- Blood Product Administered in O.R.

I certify the above statements to be true:

\_\_\_\_\_  
Transfusionist (RN, Physician/Anesthesia provider) signature      RN Co-signature      Date and Time

**Refer to back of the form for any vital sign deviation exceeding the limits indicated in red**

VS Recorded	Time	Temp	Pulse	Resp	B/P	Patient Assessment	Signature
		≥2° F rise or 1° C	< > 25	>10	< > 35	Notify RN/MD When Vital Signs Exceed Limit	
<b>START</b>							
<b>15 min</b>							
<b>30 min</b>							
<b>1 hr</b>							
<b>1.5 hr</b>							
<b>2 hr</b>							
<b>2.5 hr</b>							
<b>3 hr</b>							
<b>3.5 hr</b>							
<b>4 hr</b>							
<b>Complete</b>							
<b>1 hr Post</b>						<input type="checkbox"/> No signs or symptoms of transfusion reaction	

Return completed Yellow form and empty blood bag to Blood Bank.    Entire Unit Given    Partial Unit Given \_\_\_\_\_ cc

If...	Then...
<ul style="list-style-type: none"> <li>• Temperature rise of 2° F or 1° C</li> <li>• Shaking chills</li> <li>• Back pain (lumbar or flank pain) - new or worsening</li> <li>• Dyspnea (shortness of breath), wheezing or coughing - new or worsening</li> <li>• Chest pain - new or worsening</li> <li>• Strider or difficulty breathing - new or worsening</li> <li>• Pain or heat at the transfusion site or along the vein</li> <li>• Nausea, vomiting, diarrhea or abdominal cramps - new or worsening</li> <li>• Unexplained loss of consciousness</li> <li>• Unexplained signs of shock</li> <li>• Red urine or hematuria - new or worsening</li> <li>• Headache - new or worsening</li> <li>• Joint pain - new or worsening</li> <li>• Oozing or increased bleeding from wound</li> <li>• Flushing of skin - new or worsening</li> <li>• Death</li> </ul>	<ol style="list-style-type: none"> <li>1. The transfusion <b>MUST BE STOPPED IMMEDIATELY!</b></li> <li>2. Transfusion reaction investigation initiated               <ol style="list-style-type: none"> <li>a. Verify the patient's identity. If an error noted - notify Blood Bank IMMEDIATELY</li> <li>b. Notify the Blood Bank and the Attending Physician.</li> <li>c. Fill out Transfusion Reaction Investigation Form (LAB009)</li> <li>d. Obtain an immediate post-transfusion urine specimen and retrieve the blood transfusion unit with infusion set attached and sealed off. Do not return set if only IV access.</li> </ol> </li> <li>3. Lab will notify Medical Director</li> <li>4. Lab will contact the Nursing Staff as to whether or not another unit is to be transfused based on the results of the transfusion reaction workup and the Medical Director.</li> </ol>
If...	Then...
<ul style="list-style-type: none"> <li>• Change in pulse from initial reading (either decrease or increase) <b>of greater than 25 beats per minute</b></li> <li>• Change in Blood Pressure from initial reading (either increase or decrease in systolic or diastolic) <b>of greater than 35 mmHg</b></li> <li>• Increased respirations from initial reading, <b>greater than 10 breaths per minute</b></li> </ul>	<p>The transfusion <b>MUST BE TEMPORARILY DISCONTINUED</b> and the attending physician notified of the following symptoms. The attending physician must assess the hemodynamic status of the recipient to determine if symptoms are consistent with the patient's condition. If symptoms are unrelated to the patient's hemodynamic status, a transfusion reaction investigation is recommended.</p>
<ul style="list-style-type: none"> <li>• Urticarial (hives) reaction</li> </ul>	<p><b>NOT AN INDICATION FOR AUTOMATICALLY DISCONTINUING</b> the transfusion; although it must be reported to the Blood Bank on Transfusion Reaction Investigation Form (LAB009). The patient should be treated for the urticarial reaction.</p>