

Adverse Transfusion Reaction Form

PATIENT INFORMATION: <i>(Please print)</i>	
Patient's Name _____	
Patient Unique ID # <i>(i.e. SSN/MR#)</i> : _____	Patient DOB: / / _____
Physician: _____	Agency Name: _____
Agency Address: _____	Agency Phone #: _____
Phlebotomist:	
Name: (Please Print) _____	Date sample collected _____
Signature _____	Time sample collected _____

CLERICAL REVIEW:

Check box to indicate that crossmatch information tag correlates with unit label

 ABO Rh (D) Blood Unit Number

 Does patient identification match crossmatch tag? YES NO

 Was a Blood Warmer used? YES NO

TRANSFUSION SUMMARY:

Donor Unit Number (s): #1 _____ #2 _____

 What product was transfused? RBC (Leukoreduced/Washed) FFP (Fresh Frozen Plasma)

 Platelets (Pheresis) Cryoprecipitate Other _____

 Approximate **volume** of component transfused #1 _____ #2 _____

 Date and time *transfusion initiated*: #1 _____ #2 _____

 Date and time *reaction occurred*: #1 _____ #2 _____

 Date and time *transfusion discontinued*: #1 _____ #2 _____

SIGNS AND SYMPTOMS OF REACTION: (✓ those that apply)

<input type="checkbox"/> Urticaria (hives)	<input type="checkbox"/> Hemoglobinuria	<input type="checkbox"/> Nausea	<input type="checkbox"/> Hypotension	<input type="checkbox"/> Skin Flushing
<input type="checkbox"/> Chills/Rigors	<input type="checkbox"/> Fever	<input type="checkbox"/> Vomiting	<input type="checkbox"/> Headache	<input type="checkbox"/> Anxiety
<input type="checkbox"/> Coughing	<input type="checkbox"/> Back Pain	<input type="checkbox"/> Tachycardia	<input type="checkbox"/> Diarrhea	<input type="checkbox"/> Shock
<input type="checkbox"/> Dyspnea (chest tightening)	<input type="checkbox"/> Generalized Bleeding	<input type="checkbox"/> Cyanosis	<input type="checkbox"/> Orthopnea (shortness of breath)	<input type="checkbox"/> Other:

VITAL SIGNS:

At beginning of transfusion: Blood Pressure: _____ Temp: _____ Pulse: _____

At time reaction occurred: Blood Pressure: _____ Temp: _____ Pulse: _____

At 1 hour after transfusion: Blood Pressure: _____ Temp: _____ Pulse: _____

PATIENT'S GENERAL CONDITION: _____

Report prepared By: _____ Date: _____ Time: _____

Reported to QualTex-IRL By: _____ Date: _____ Time: _____

NOTE: Please submit the following: 1 lavender top tube (EDTA) and 1 red top tube, all component bags with tubing & IV solution attached (remove needles), post transfusion urine and this completed form. Label all samples with patient's name, Patient Unique ID #, date, time collected and phlebotomist's initials. Call Hospital Services (210-731-5550) for pick up of samples and call Immunohematology Reference Lab (210-731-5509) for notification of possible transfusion reaction. F(210)249-4417, Qualtexlabs.org