



Patient's sticker

REPORT OF SUSPECTED TRANSFUSION REACTION

STOP THE TRANSFUSION AND VERIFY INFORMATION BELOW

INFUSION RECORD (Please tick mark the appropriate option)

- Yes No The identity of the patient agrees with the Name and CR no. of the patient.
- Yes No The Name and CR no. on the Blood Centre Compatibility label agrees with those of the patient.
- Yes No The ABO and Rh type on the Blood Centre Compatibility label agrees with those on the component label.
- Yes No The Unit Number on the attached Compatibility label agrees with the Unit Number on the component label.

Clinical diagnosis: _____ Indication for Transfusion _____

- Previous H/O Transfusion(s) YES NO If YES, Give details _____ and Last transfusion date _____
- H/O Previous Transfusion Reactions: YES NO. If YES, give details _____
- Equipment used: Blood infusion set / Blood warmer Reinfusion device Pressure device Pump / Bed side filter
- Was Patient under anesthesia at time of transfusion? YES NO
- Was patient pretreated for transfusion? YES NO If YES, Pretreatment Medications _____ Time: _____

SYMPTOMS OF THE SUSPECTED REACTION:

Patient Data				Check all that apply:
Vital Signs	Pre-Transfusion	During Reaction	Post-Transfusion	
Time				<input type="checkbox"/> Chills <input type="checkbox"/> Rigors <input type="checkbox"/> Fever (1°C or 2°F rise in temperature) <input type="checkbox"/> Flushing <input type="checkbox"/> Urticaria <input type="checkbox"/> Wheeze <input type="checkbox"/> Stridor <input type="checkbox"/> Cough <input type="checkbox"/> Dyspnea <input type="checkbox"/> Chest Pain <input type="checkbox"/> Raised JVP <input type="checkbox"/> Shock <input type="checkbox"/> Back Pain <input type="checkbox"/> Heat / Pain at the IV site <input type="checkbox"/> Bleeding <input type="checkbox"/> Hemoglobinuria <input type="checkbox"/> Oliguria <input type="checkbox"/> Jaundice <input type="checkbox"/> Anxiety <input type="checkbox"/> Nausea <input type="checkbox"/> Vomiting <input type="checkbox"/> Other: _____
Temperature				
Blood Pressure				
Pulse				
Respiratory Rate				
O ₂ Sat.	sPo ₂ : <input type="checkbox"/> Room Air <input type="checkbox"/> O ₂ Therapy _____ L/min	sPo ₂ : <input type="checkbox"/> Room Air <input type="checkbox"/> O ₂ Therapy _____ L/min	sPo ₂ : <input type="checkbox"/> Room Air <input type="checkbox"/> O ₂ Therapy _____ L/min	

GIVE A BRIEF DESCRIPTION OF THE REACTION:

DONOR BLOOD SUSPECTED OF CAUSING REACTIONS

UNIT NO. _____ COMPONENT: _____ VOLUME TRANSFUSED: _____

Date and time of issue from blood centre	
Date and time of start of transfusion	
Date and time of reaction	
Date and time of recovery	

MEASURES TAKEN:

- Transfusion temporarily stopped Analgesics Diuretics Antipyretics Steroids Antibiotics Supplementary O₂
- Antihistamines Blood culture sent Mechanical ventilation ICU required Chest X-ray Vasopressors ABG/VBG Urine R/M
- Others (Please specify) _____

SEND THE FOLLOWING TO THE BLOOD CENTRE FOR ALL SUSPECTED TRANSFUSION REACTIONS (Please tick mark):

- 5 mL blood specimen each in an EDTA (DARK LAVENDER TOP) tube and in CLOT ACTIVATOR (YELLOW TOP) tube labelled with:
 - Patient Sticker, Phlebotomist Signature, Date & time of sampling and the note "POST REACTION"
- The blood component under investigation with attached IV set.
- Urine sample labelled with patient sticker, Date & time of sampling and the note "POST REACTION".
- A COPY OF THIS COMPLETED FORM. PLACE THE ORIGINAL IN THE PATIENT'S MEDICAL RECORD.

Date:

**Name of Nursing officer:
Sign:**

**Name of Resident Doctor/ Faculty
Sign:
Contact No:**



Department of Transfusion Medicine AIIMS Jodhpur

SUSPECTED TRANSFUSION REACTION-BLOOD CENTRE REPORT *(To be filled by blood centre only)*

Patient's Name: _____ Age/Gender: _____ CR no.: _____

Ward: _____ Consultant: _____ Resident: _____

Date of receiving transfusion reaction form _____

Patient's Pretransfusion Records:

ABO & Rh: _____ Crossmatch: _____ Crossmatch Date: _____ Crossmatch done by: _____

Date Issued: _____ Time Issued: _____

Returned Component Record: BLOOD BAG

Component type _____ Amount Returned _____ Expiration Date _____ Compatibility Label on Product: Yes No

ABO & Rh: _____ Donor unit no.: _____ Appearance: _____

Manufacturer of bag: _____ Lot no.: _____ Expiry of bag: _____ Segment no.: _____

Check blood product for bacterial contamination (i.e., peculiar odor, brownish or purple color, clots, or abnormal masses in bag) Yes / No _____

Blood product bag sent for culture if contamination is suspected or if any of the following clinical indicators are present:

- shock
- hypertensive (systolic rises \geq 30mm Hg)
- hypotensive (systolic falls \geq 30mm Hg)
- fever (2°C or 3.5°F rise in temperature)
- rigors (shaking chills)
- tachycardia (heart rate is \geq 120/min, or rises \geq 40/min above pre-transfusion rate).

COMPATIBILITY LABEL

Patient ABO & Rh _____ Unit ABO & Rh _____ Unit no. _____

Patient Name: _____ CR no.: _____

Patient's Specimen

	Visual Hemolysis Check (Pos/Neg)	anti-A	anti-B	anti-AB	anti-D	A ₁ c	Bc	Oc	Auto-ctrl	Blood group	DAT	IAT	Ab screen	Crossmatch		
														Saline	37°C	AHG
PRE																
POST																

Other Results/Comments: _____

Laboratory investigations

	Hb/Hct	Total WBC	Platelet count	Peripheral smear report (Any fragmented cells/ spherocytes/ RBC agglutination)	LDH	LFT			Urine microscopy/ Hb	Any other lab. finding/ Comments
						TB/IB	Total protein/ albumin	ALT/AST		
PRE										
POST										

Unit Culture results: _____ Unit Gram stain results if indicated: _____ Patient culture results _____

Impression:

Imputability:

Advice:

TRANSFUSION MEDICINE RESIDENT NAME & SIGNATURE: _____

Date: _____

TRANSFUSION MEDICINE FACULTY NAME & SIGNATURE: _____

Date: _____