

***Required for saving**

List the patient's relevant medical procedure including past procedures and procedures to be performed during the current hospital or outpatient stay. (Use ICD-10 Procedure codes/descriptions) ☐ UNKNOWN ☐ NONE

Code: _____ Description: _____

Code: _____ Description: _____

Code: _____ Description: _____

Additional Information _____

Transfusion History

Has the patient received a previous transfusion? ☐ YES ☐ NO ☐ UNKNOWN

Blood Product: ☐ WB ☐ RBC ☐ Platelet ☐ Plasma ☐ Cryoprecipitate ☐ Granulocyte

Date of Transfusion: ____/____/____ ☐ UNKNOWN

Was the patient's adverse reaction transfusion-related? ☐ YES ☐ NO

If yes, provide information about the transfusion adverse reaction.

Type of transfusion adverse reaction: ☐ Allergic ☐ AHTR ☐ DHTR ☐ DSTR ☐ FNHTR

☐ HTR ☐ TTI ☐ PTP ☐ TACO ☐ TAD ☐ TA-GVHD ☐ TRALI ☐ UNKNOWN

☐ OTHER Specify _____

Reaction Details

*Date reaction occurred: ____/____/____ *Time reaction occurred: ____:____ ☐ Time unknown

*Facility location where patient was transfused: _____

Is this reaction associated with an incident? ☐ Yes ☐ No If Yes, Incident #: _____

Investigation Results

*☐ Allergic reaction, including anaphylaxis

*Case Definition

Check the following that occurred during or within **4 hours** of cessation of transfusion:

☐ Conjunctival edema ☐ Edema of lips, tongue and uvula ☐ Localized angioedema ☐ Hypotension

☐ Erythema and edema of the periorbital area ☐ Respiratory distress; bronchospasm ☐ Urticaria

☐ Generalized flushing ☐ Maculopapular rash ☐ Pruritus

Other signs and symptoms: (check all that apply)

Generalized: ☐ Chills/rigors ☐ Fever ☐ Nausea/vomiting

Cardiovascular: ☐ Shock

Cutaneous: ☐ Jaundice

Hemolysis/Hemorrhage: ☐ Disseminated intravascular coagulation ☐ Hemoglobinemia
☐ Positive antibody screen

Pain: ☐ Abdominal pain ☐ Back pain ☐ Flank pain ☐ Infusion site pain

Renal: ☐ Hematuria ☐ Hemoglobinuria ☐ Oliguria

Respiratory: ☐ Bilateral infiltrates on chest x-ray ☐ Cough
☐ Hypoxemia ☐ Shortness of breath

☐ Other: (specify) _____

*Severity

Did the patient receive or experience any of the following?

- | | |
|---|---|
| ☐ No treatment required | ☐ Symptomatic treatment only |
| ☐ Hospitalization, including prolonged hospitalization | ☐ Life-threatening reaction |
| ☐ Disability and/or incapacitation | ☐ Congenital anomaly or birth defect(s) of the fetus |
| ☐ Other medically important conditions | ☐ Death |
| | ☐ Unknown or not stated |

*Imputability

Which best describes the relationship between the transfusion and the reaction?

- ☐ No other evidence of environmental, drug or dietary risks.
- ☐ There are other potential causes present that could explain acute hemolysis, but transfusion is the most likely cause.
- ☐ Other present causes are most likely, but transfusion cannot be ruled out.
- ☐ Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.
- ☐ There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.
- ☐ The relationship between the adverse reaction and the transfusion is unknown or not stated.

Did the transfusion occur at your facility? ☐ YES ☐ NO

When did the reaction occur in relation to the transfusion?

- ☐ Occurred during or within 2 hours of cessation of transfusion.
- ☐ Occurred 2 - 4 hours after cessation of transfusion.

Did the same reaction occur after the transfusion was restarted (rechallenge)? ☐ YES ☐ NO

Module-generated Designations

NOTE: Designations for case definition, severity, and imputability will be automatically assigned in the NHSN application based on responses in the corresponding investigation results section above.

***Do you agree with the case definition designation?** ☐ YES ☐ NO

^Please indicate your designation _____

***Do you agree with the severity designation?** ☐ YES ☐ NO

^Please indicate your designation _____

***Do you agree with the imputability designation?** ☐ YES ☐ NO

^Please indicate your designation _____

Patient Treatment

Did the patient receive treatment for the transfusion reaction? ☐ YES ☐ NO ☐ UNKNOWN

If yes, select treatment(s):

- ☐ Medication (*Select the type of medication*)
- | | | | | |
|---|---|---|---|------------------------------------|
| ☐ Antipyretics | ☐ Antihistamines | ☐ Inotropes/Vasopressors | ☐ Bronchodilator | ☐ Diuretics |
| ☐ Intravenous Immunoglobulin | ☐ Intravenous steroids | ☐ Corticosteroids | ☐ Antibiotics | |
| ☐ Antithymocyte globulin | ☐ Cyclosporin | ☐ Other | | |
- ☐ Volume resuscitation (Intravenous colloids or crystalloids)
- ☐ Respiratory support (*Select the type of support*)
- | | | |
|---|--|---------------------------------|
| ☐ Mechanical ventilation | ☐ Noninvasive ventilation | ☐ Oxygen |
|---|--|---------------------------------|

- ☐ Renal replacement therapy (*Select the type of therapy*)
- ☐ Hemodialysis ☐ Peritoneal ☐ Continuous Veno-Venous Hemofiltration
- ☐ Phlebotomy
- ☐ Other Specify: _____

Outcome

- *Outcome:** ☐ Death ☐ Major or long-term sequelae ☐ Minor or no sequelae ☐ Not determined
- Date of Death: ____/____/____
- ^If recipient died, relationship of transfusion to death:
- ☐ Definite ☐ Probable ☐ Possible ☐ Doubtful ☐ Ruled Out ☐ Not determined
- Cause of death: _____
- Was an autopsy performed? ☐ Yes ☐ No

Component Details

***Was a particular unit implicated in (i.e., responsible for) the adverse reaction?**

☐ Yes ☐ No ☐ N/A

Transfusion Start and End Date/Time	*Component code (check system used)	Amount transfused at reaction onset	^Unit number (Required for Infection and TRALI)	*Unit expiration Date/Time	*Blood group of unit	Implicated Unit?
^IMPLICATED UNIT						
____/____/____ ____:____ ____/____/____ ____:	☐ ISBT-128 ☐ Codabar _____	☐ Entire unit ☐ Partial unit _____ mL	_____ _____ _____	____/____/____ ____:	☐ A- ☐ A+ ☐ B- ☐ B+ ☐ AB- ☐ AB+ ☐ O- ☐ O+ ☐ N/A	Y
____/____/____ ____:____ ____/____/____ ____:	☐ ISBT-128 ☐ Codabar _____	☐ Entire unit ☐ Partial unit _____ mL	_____ _____ _____	____/____/____ ____:	☐ A- ☐ A+ ☐ B- ☐ B+ ☐ AB- ☐ AB+ ☐ O- ☐ O+ ☐ N/A	N

Custom Fields

Label	Label
_____ _____ _____	_____ _____ _____

Comments