

Hemovigilance Module

Adverse Reaction

Delayed Serologic Transfusion Reaction

***Required for saving**

*Facility ID#: _____ NHSN Adverse Reaction #: _____

Patient Information

*Patient ID: _____

*Date of Birth: ____/____/____

*Sex: ☐ M ☐ F

Social Security #: _____ Secondary ID: _____ Medicare #: _____

Last Name: _____ First Name: _____ Middle Name: _____

Ethnicity (Specify): ☐ Hispanic or Latino ☐ Not Hispanic or Latino ☐ Unknown ☐ Declined to respond

Race (Select all that apply): ☐ American Indian or Alaska Native ☐ Asian ☐ Black or African American ☐ Middle Eastern or North African ☐ Native Hawaiian or Pacific Islander ☐ White ☐ Unknown ☐ Declined to respond

Preferred Language (Specify from the list provided): _____ Interpreter Needed: ☐ Yes ☐ No ☐ Unknown ☐ Declined to Respond

*Blood Group: ☐ A- ☐ A+ ☐ B- ☐ B+ ☐ AB- ☐ AB+ ☐ O- ☐ O+ ☐ Blood type not done

☐ Transitional ABO / Rh + ☐ Transitional ABO / Rh - ☐ Transitional ABO / Transitional Rh

☐ Group A/Transitional Rh ☐ Group B/Transitional Rh ☐ Group O/Transitional Rh ☐ Group AB/Transitional Rh

Patient Medical History

List the patient's admitting diagnosis. (Use ICD-10 Diagnostic codes/descriptions)

Code: _____ Description: _____

Code: _____ Description: _____

Code: _____ Description: _____

List the patient's underlying indication for transfusion. (Use ICD-10 Diagnostic codes/descriptions)

Code: _____ Description: _____

Code: _____ Description: _____

Code: _____ Description: _____

List the patient's comorbid conditions at the time of the transfusion related to the adverse reaction. (Use ICD-10 Diagnostic codes/descriptions)

☐ UNKNOWN

☐ NONE

Code: _____ Description: _____

Code: _____ Description: _____

Code: _____ Description: _____

Assurance of Confidentiality: The voluntarily provided information obtained in this surveillance system that would permit identification of any individual or institution is collected with a guarantee that it will be held in strict confidence, will be used only for the purposes stated, and will not otherwise be disclosed or released without the consent of the individual, or the institution in accordance with Sections 304, 306 and 308(d) of the Public Health Service Act (42 USC 242b, 242k, and 242m(d)). CDC 57.310 Rev. 3, v9.2

Public reporting burden of this collection of information is estimated to average 20 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering, and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC, Reports Clearance Officer, 1600 Clifton Rd., MS H21-8, Atlanta, GA 30333, ATTN: PRA (0920-0666).

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

*☐ Delayed serologic transfusion reaction (DSTR)

Antibody(ies): _____

***Case Definition Check all that apply:**

- ☐ Absence of clinical signs of hemolysis
- ☐ Positive direct antiglobulin test (DAT)
- ☐ Demonstration of new, clinically-significant antibodies against red blood cells
- ☐ Positive antibody screen with newly identified RBC alloantibody

Other signs and symptoms: (check all that apply)

Generalized:	☐ Chills/rigors	☐ Fever	☐ Nausea/vomiting
Cardiovascular:	☐ Blood pressure decrease	☐ Shock	
Cutaneous:	☐ Edema	☐ Flushing	☐ Jaundice
	☐ Other rash	☐ Pruritus (itching)	☐ Urticaria (hives)
Hemolysis/Hemorrhage:	☐ Disseminated intravascular coagulation	☐ Hemoglobinemia	
Pain:	☐ Abdominal pain	☐ Back pain	☐ Flank pain ☐ Infusion site pain
Renal:	☐ Hematuria	☐ Hemoglobinuria	☐ Oliguria
Respiratory:	☐ Bilateral infiltrates on chest x-ray	☐ Bronchospasm	☐ Cough
	☐ Hypoxemia	☐ Shortness of breath	

☐ Other: (specify) _____

*Severity

Since this is by definition a reaction with no clinical symptoms, severity of the reaction cannot be graded.

☒ Not determined

*Imputability

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

- ☐ Transfusion performed by your facility is the only possible cause for seroconversion.
- ☐ The patient has other exposures (e.g. transfusion by another facility or pregnancy) that could explain seroconversion, but transfusion by your facility is the most likely cause.
- ☐ The patient was transfused by your facility, but other exposures are present that most likely explain seroconversion.
- ☐ 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 was the new alloantibody identified?

- ☐ Occurred between 24 hours and 28 days after cessation of transfusion
- ☐ Occurred less than 24 hours after cessation of transfusion OR greater than 28 days after cessation of transfusion
- ☐ No new antibody was identified

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

