



National Healthcare Safety Network Biovigilance Component Hemovigilance Module Surveillance Protocol

Division of Healthcare Quality Promotion
National Center for Emerging and Zoonotic Infectious Diseases
Centers for Disease Control and Prevention
Atlanta, GA, USA



Version History

Version	Release Date	Summary of Revisions
1.0	March 2009	First version publicly released.
1.1	June 2010	<p>Revised background and text in main body of document.</p> <p>Revised case definition criterion based on WG recommendations, pilot responses, and CDC recommendations.</p> <p>Updated FNHTR definition to allow reaction without documented fever.</p> <p>Defined hypotension for infants and small children</p> <p>Clarified TAGVHD probable and possible criteria.</p>
1.2	July 2010	Corrected definition of hypoxemia in glossary of terms.
1.3	June 2011	<p>Added version number and version history summary.</p> <p>Summarized introduction and background sections for brevity.</p> <p>Reorganized surveillance methods section for ease of use.</p> <p>Clarified reporting of “approved deviation” incidents.</p> <p>Clarified use of “other” in adverse reaction reporting.</p> <p>Clarified use of “doubtful” or “ruled out” in adverse reaction reporting.</p> <p>Added denominator summary options to list of available analysis reports.</p> <p>Replaced < and > signs with appropriate text for.</p> <p>Added “cessation of” to time frame requirements in case definitions.</p> <p>NEW probable case definition category for allergic reaction reporting.</p> <p>Updated adult hypotensive reaction case definition to align with updated ISBT definition.</p> <p>NEW possible imputability category for DHTR.</p> <p>DELETED possible case definition category for hypotensive reaction.</p> <p>NEW probable imputability category for PTP reaction.</p> <p>Updated and clarified imputability categories for TAGVHD reaction.</p> <p>DELETED possible case definition category for TRALI.</p> <p>Simplified imputability criteria for TTI.</p> <p>Clarified case definition and imputability criteria for all adverse reactions.</p>
2.0	January 2013	<p>Complete revision of organization and presentation of information</p> <p>Major change in incident reporting requirements. With this release, only incidents that relate to an adverse patient reaction are required for participation.</p> <p>Major change in adverse reaction reporting requirements. With this release, minor allergic reactions are no longer required for participation.</p> <p>Combined the signs/symptoms with laboratory/radiology columns in case definition tables for clarity. Listed criteria in alphabetical order where possible for consistency and clarity. Moved general severity requirements from the appendix to the criteria tables where they were previously missing.</p> <p>Re-ordered adverse reaction tables to put respiratory reactions first.</p> <p>Added Imputability criteria of Doubtful, Ruled Out, and Not Determined to the case definition tables as OPTIONAL reporting categories. The reporting is not a change, but including them in the table is new. They were added for clarity.</p> <p>Added specific AHTR criteria to allow for reporting of non-immune mediated reactions.</p> <p>Added a separate case definition table for Other and Unknown reactions. These categories are available for OPTIONAL use.</p> <p>Removed redundant and unnecessary appendices.</p>
2.1	August 2013	<p>Minor revisions to verbiage throughout for clarity.</p> <p>Added definitions and illustration of surveillance key terms in Section 1.</p> <p>Added clarification of surveillance vs. clinical definitions in Section 1.</p> <p>Added less-specific case definition categories for OPTIONAL reporting of cases that do not fully meet CDC case criteria for the following reactions: hypotension, febrile non-hemolytic, acute hemolytic and delayed hemolytic.</p>



Version	Release Date	Summary of Revisions
		Added a possible case definition category for TTI for OPTIONAL reporting of syndromic cases that are not laboratory confirmed.
2.1.1	September 2013	Updated diagram in Section 1 and added version history for v2.0 and v2.1.
2.1.2	January 2014	Updated the incident codes in Section 4 and included required reporting of discards and total crossmatch procedures on the Monthly Reporting Denominators form in Section 5.
2.1.3	August 2014	Added a suggested citation for the surveillance protocol in Section 1. Updated the acute hemolytic case definition in Section 3 for clarity. Updated the reporting requirements in Section 5 for clarity.
2.2	January 2016	Updated contact instructions for consistency in Section 1: User support Updated version number in Section 1: Suggested Citation Remove Root Cause Analysis Result from Section 4: Incident Glossary Updated denominator report description to include Pathogen-reduced products in Section 5: Required Reporting
2.3	June 2016	Updated denominator report description to include Table 3 description.
2.4	January 2017	Section 1: Setting – Added Non-Acute Care Facility reporting Section 2: FDA Reporting – Added new section Section 3: Annual Facility Survey – Added information about Non-Acute Care Facility Annual Facility Survey; Added links to the Annual Facility Survey – Non-Acute Care Facility form and table of instructions Section 4: Adverse Reactions – Updated Optional Reporting; Added FDA Reporting; Updated Adverse Reaction Classification



Table of Contents

Section 1. Hemovigilance Module Surveillance Overview5

Section 2. FDA Reporting7

Section 3. Hemovigilance Module Annual Facility Survey8

Section 4: Hemovigilance Module Adverse Reactions9

Adverse Reaction Case Classification Criteria Tables 11

 Transfusion-associated circulatory overload (TACO)..... 11

 Transfusion-related acute lung injury (TRALI)..... 12

 Transfusion-associated dyspnea (TAD) 13

 Allergic reaction 14

 Hypotensive transfusion reaction 15

 Febrile non-hemolytic transfusion reaction (FNHTR) 16

 Acute hemolytic transfusion reaction (AHTR) 17

 Delayed hemolytic transfusion reaction (DHTR) 18

 Delayed serologic transfusion reaction (DSTR) 19

 Transfusion-associated graft vs. host disease (TAGVHD)..... 20

 Post transfusion purpura (PTP) 21

 Transfusion-transmitted infection (TTI) 22

 Other or Unknown..... 24

Adverse Reaction Glossary..... 25

Section 5. Hemovigilance Module Incidents26

Incident Codes 27

Occupation Codes 30

Incident Glossary 311

Section 6. Hemovigilance Module Denominators.....322





Section 1. Hemovigilance Module Surveillance Overview

Purpose

The National Healthcare Safety Network (NHSN) Hemovigilance (HV) Module was created to implement national surveillance of transfusion-associated adverse events aimed at improving patient safety, minimizing morbidity and mortality of transfusion recipients, and identifying emerging complications and pathogens associated with blood transfusion.

Settings

The Hemovigilance Module may be used by any U.S. healthcare facility where blood components and manufactured blood products are transfused (e.g., adult or pediatric facilities, acute or non-acute care facilities). Surveillance must be performed facility-wide, including patient care areas for emergency, general medical, and surgical patients; obstetrics and gynecology; orthopedics, oncology, and other chronic diseases; and any other facility location where transfusions are administered.

Methods

The NHSN Hemovigilance Module requires comprehensive surveillance of patients and blood components throughout the transfusion process, from product receipt to administration to the patient. Participation in the NHSN Hemovigilance Module requires reporting of all adverse transfusion reactions and reaction-associated incidents that occur **for patients transfused at or by your facility** as well as a monthly summary of components transfused or discarded and patient samples collected for type and screen or crossmatch.

Data Collection

NHSN is a web-based application used by healthcare facilities to report surveillance data. Paper versions of all forms are used to collect data prior to data entry in the NHSN Hemovigilance Module. The paper forms are available on the [NHSN Blood Safety Surveillance website](#). A link to the appropriate form(s) and their instructions is provided in the following sections for your convenience.

Training

Training presentations are available on the NHSN Biovigilance Component website for self-paced training and must be reviewed prior to participating in the Hemovigilance Module. CDC also provides webinar and in-person training opportunities for current NHSN participants. These opportunities are communicated through the NHSN quarterly newsletter and emails from the Biovigilance Team.

User Support

CDC is available to answer your questions about the Surveillance Protocol and to help navigate the NHSN web application. Please contact us at nhsn@cdc.gov. Type **BIOVIGLANCE** in the subject line for quickest routing to the Biovigilance/Hemovigilance Team.

Suggested Citation for the Hemovigilance Module Surveillance Protocol

U.S. Centers for Disease Control and Prevention. The National Healthcare Safety Network (NHSN) Manual: Biovigilance Component v2.2. Atlanta, GA: Division of Healthcare Quality Promotion, National Center for Emerging and Zoonotic Infectious Diseases. Available at: <http://www.cdc.gov/nhsn/PDFs/Biovigilance/BV-HV-protocol-current.pdf>. Accessed [enter date].



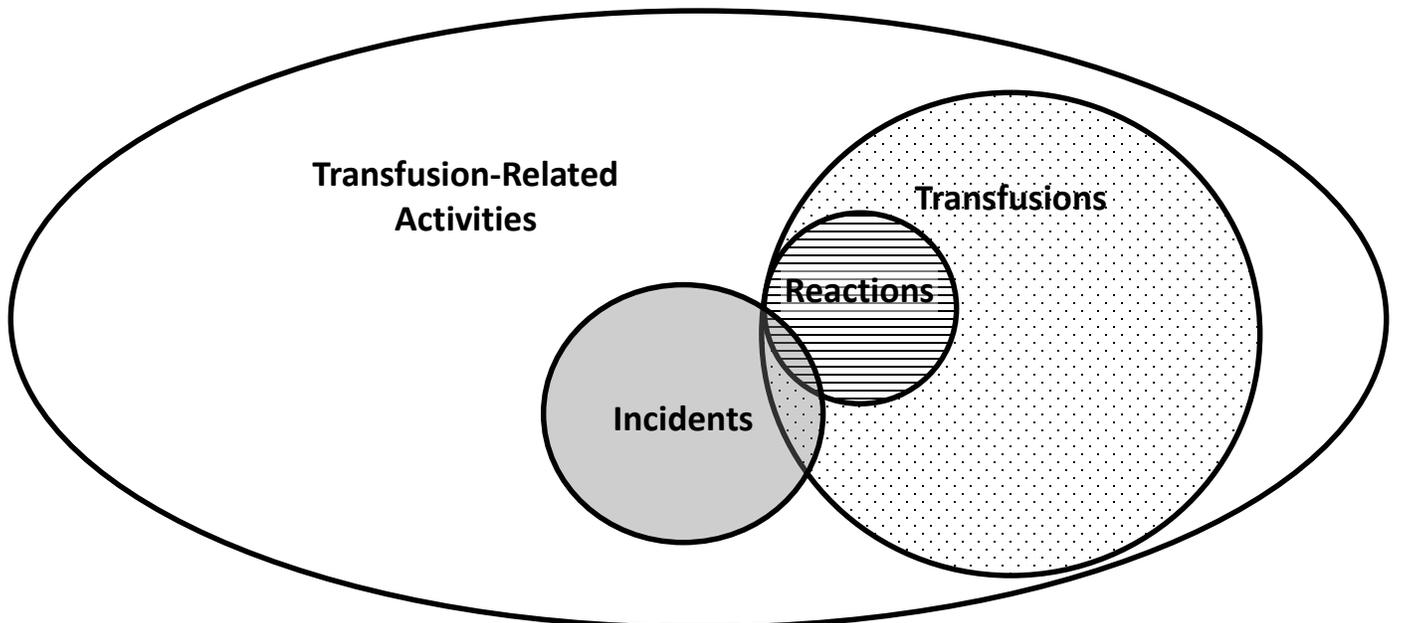
Key Terms (see Fig. 1)

- **Adverse event:** An unintended and undesirable occurrence before, during or after transfusion of blood or blood components. Adverse events include both incidents and adverse reactions.
- **Adverse reaction:** An undesirable response or effect in a patient temporally associated with the administration of blood or blood components. It may or may not be the result of an incident.
- **Incident:** Any error or accident that could affect the quality or efficacy of blood, blood components, or patient transfusions. It may or may not result in an adverse reaction in a transfusion recipient.
- **Near miss:** A subset of incidents that are discovered before the start of a transfusion that *could* have led to a wrongful transfusion or an adverse reaction in a transfusion recipient.

Data Reporting Requirements (See Fig. 1)

- An annual facility demographic and practice survey for each **calendar** year of participation
- ALL adverse reactions that follow transfusion **at or by your facility**
- ALL incidents (i.e., errors or accidents) associated with an adverse reaction
- The number of blood components transfused or discarded and patient samples collected for type and screen or crossmatch each month

Figure 1. Venn diagram of NHSN Hemovigilance Module surveillance terms.



- Transfusion-Related Activities**
 - Patient Sample Collection
 - Sample Handling and Testing
 - Inventory Management
 - Patient Monitoring

- Transfusion**
 - Number of Components
 - Number of Patients

Adverse Events

- Reactions**

Incidents

- Near Miss Incidents
- Incidents Related to Transfusion (No Adverse Reaction)
- Incidents Related to Transfusion and Adverse Reaction



Section 2. Data Sharing with Food and Drug Administration (FDA)

Submitting FDA reports via NHSN Hemovigilance Module

Starting in January 2017, users will be able to share reports that satisfy the Safety Reporting Rule requirement with FDA via the NHSN Hemovigilance module. Facility can join the 'FDA Super Group' to enable data sharing. Users may enroll in the FDA Super Group by contacting the Biovigilance team at nhsn@cdc.gov. Once enrolled, users may submit serious transfusion reactions and transfusion-related fatalities to NHSN and FDA using the Adverse Reaction form.



Section 3. Hemovigilance Module Annual Facility Survey

Required Reporting

Participating facilities must enter the Hemovigilance Module Annual Facility Survey at the time that they enroll or activate the Biovigilance Component and at the beginning of each calendar year thereafter. The survey is used by CDC to classify facilities for appropriate comparisons in aggregate data analyses and to learn more about common practices among transfusion services. The data collected in the survey covers the previous **calendar** year. For example, if the facility is enrolling in NHSN for the first time in October of 2013, report information for January 2012-December 2012 on the first Hemovigilance Module Annual Facility Survey. In January 2014, complete a new survey with data from January 2013-December 2013. CDC recommends collecting all survey information on a paper form before attempting to enter data into the web application.

As of January 2017, non-acute care facilities will be able to report hemovigilance data to NHSN. Non-acute care facilities should complete Annual Facility Survey for Non-acute care facility 57.3XX. This form contains questions tailored to non-acute care facilities. Users may refer to the Non-Acute Care Facility Table of Instructions form 57.3XX for detailed instruction about data collection.

Form

[CDC 57.300 Hemovigilance Module Annual Facility Survey - Acute Care Facility](#)

[CDC 57.3XX Hemovigilance Module Annual Facility Survey - Non-Acute Care Facility](#)

Form Instructions

[CDC 57.300 Hemovigilance Module Annual Facility Survey - Acute Care Facility Table of Instructions](#)

[CDC 57.3XX Hemovigilance Module Annual Facility Survey - Non-Acute Care Facility Table of Instructions](#)



Section 4: Hemovigilance Module Adverse Reactions

Required Reporting

All CDC-defined transfusion-associated adverse reactions that are possibly, probably, or definitely related to a **transfusion performed by the participating facility** must be reported to NHSN. If a patient experiences more than one adverse reaction during or following the same transfusion episode, complete a separate form for each reaction. Adverse reaction reports should be entered into NHSN after an investigation of the reaction has been completed and imputability has been determined to the extent possible. Reports should be entered within 30 days of the month that the reaction occurred or when the investigation is completed.

Optional Reporting

Reporting suspected adverse reactions where imputability is determined to be doubtful or ruled out is not required. A facility may report reactions determined to be doubtful or ruled out in order to use NHSN to document transfusion reaction investigations each month. Adverse reactions that are not defined in the surveillance protocol may also be reported using the 'Other' and 'Unknown' adverse reaction categories; standard severity and imputability criteria are provided for that purpose. CDC will not aggregate or analyze these optional reports.

FDA Reporting

Users will be able to share reports with the FDA via the FDA Super Group. To mark a report for sharing, users can select the Adverse Reaction form's 'Report to FDA' checkbox. If selected, this checkbox will indicate the report should be made available to (e.g., shared) the FDA Super Group. For more information consult the Section 2. Data Sharing with Food and Drug Administration (FDA).

Adverse Reaction Classification

Each CDC-defined transfusion-associated adverse reaction **must** be classified according to the reaction-specific case definition, severity, and imputability criteria printed in this section of the protocol. It is imperative that every facility classify adverse reactions according to protocol definitions. Accurate classification will usually require a detailed review of the patient record.

Starting in January 2017, the NHSN Hemovigilance module will feature a series of question specific to each reaction that will be used to determine the case definition, severity, and imputability designations. Users will continue to identify the transfusion reaction experienced by the patient. The series of question will not determine the type of transfusion reaction. The series of questions are taken from the Adverse Reaction Case Classification Criteria Tables and follows the same designation criteria. The automated designation function will improve data quality by ensuring transfusion reactions are classified appropriately. Users will have the opportunity to accept or reject the case classification after answering the guided questions.

Surveillance definitions are distinctly different from clinical definitions. Surveillance definitions are designed to capture data consistently and reliably in order to identify trends and inform quality improvement practices. The surveillance definitions are not intended as clinical diagnostic criteria or to provide treatment guidance. The Biovigilance Team is working to provide users with access to aggregate data for bench marking purposes.

Defined Adverse Reactions

- Transfusion-associated circulatory overload (TACO)
- Transfusion-related acute lung injury (TRALI)



- Transfusion-associated dyspnea (TAD)
- Allergic reaction (where severity is severe, life threatening, or death)
- Hypotensive transfusion reaction
- Febrile non-hemolytic transfusion reaction (FNHTR)
- Acute hemolytic transfusion reaction (AHTR)
- Delayed hemolytic transfusion reaction (DHTR)
- Delayed serologic transfusion reaction (DSTR)
- Transfusion-associated graft vs. host disease (TAGVHD)
- Post-transfusion purpura (PTP)
- Transfusion-transmitted infection (TTI)

Note

Starting in January 2017, users have the option to report serious transfusion reactions to the FDA using the NHSN Hemovigilance module. Users are responsible for compliance with FDA regulations. For further information consult Section 2: Data Sharing with Food and Drug Administration (FDA).

Form

[CDC 57.304 Hemovigilance Module Adverse Reaction](#)

Form Instructions

[CDC 57.304 Hemovigilance Module Adverse Reaction Table of Instructions](#)



Adverse Reaction Case Classification Criteria Tables

Transfusion-associated circulatory overload (TACO)

Case Definition	Severity	Imputability
<p>Definitive: New onset or exacerbation of 3 or more of the following within 6 hours of cessation of transfusion:</p> <ul style="list-style-type: none"> • Acute respiratory distress (dyspnea, orthopnea, cough) • Elevated brain natriuretic peptide (BNP) • Elevated central venous pressure (CVP) • Evidence of left heart failure • Evidence of positive fluid balance • Radiographic evidence of pulmonary edema <p>Probable: N/A</p> <p>Possible: N/A</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p> <p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Definite: No other explanations for circulatory overload are possible.</p> <p>Probable: Transfusion is a likely contributor to circulatory overload AND EITHER The patient received other fluids as well OR The patient has a history of cardiac insufficiency that could explain the circulatory overload, but transfusion is just as likely to have caused the circulatory overload.</p> <p>Possible: The patient has a history of pre-existing cardiac insufficiency that most likely explains circulatory overload.</p> <p style="text-align: center;">OPTIONAL</p> <p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Transfusion-related acute lung injury (TRALI)

Case Definition	Severity	Imputability
<p>Definitive: NO evidence of acute lung injury (ALI) prior to transfusion AND ALI onset during or within 6 hours of cessation of transfusion AND Hypoxemia defined by any of these methods:</p> <ul style="list-style-type: none"> • PaO₂/FiO₂ less than or equal to 300 mm Hg • Oxygen saturation less than 90% on room air • Other clinical evidence <p>AND Radiographic evidence of bilateral infiltrates AND No evidence of left atrial hypertension (i.e., circulatory overload)</p> <p>Probable: N/A</p> <p>Possible: N/A</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p> <p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Definite: There are no alternative risk factors for ALI present.</p> <p>Probable: N/A</p> <p>Possible: There is evidence of other causes for acute lung injury such as:</p> <p>Direct Lung Injury</p> <ul style="list-style-type: none"> • Aspiration • Pneumonia • Toxic inhalation • Lung contusion • Near drowning <p>Indirect Lung Injury</p> <ul style="list-style-type: none"> • Severe sepsis • Shock • Multiple trauma • Burn injury • Acute pancreatitis • Cardiopulmonary bypass • Drug overdose
		OPTIONAL
		<p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Transfusion-associated dyspnea (TAD)

Case Definition	Severity	Imputability
<p>Definitive: Acute respiratory distress occurring within 24 hours of cessation of transfusion AND Allergic reaction, TACO, and TRALI definitions are not applicable.</p> <p>Probable: N/A</p> <p>Possible: N/A</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p> <p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Definite: Patient has no other conditions that could explain symptoms.</p> <p>Probable: There are other potential causes that could explain symptoms, but transfusion is the most likely cause.</p> <p>Possible: Other present causes are most likely, but transfusion cannot be ruled out.</p>
		<p style="text-align: center;">OPTIONAL</p> <p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Allergic reaction

Note: Minor allergic reactions (Non-severe) do not have to be reported to NHSN.

Case Definition	Severity	Imputability
<p>Definitive: 2 or more of the following occurring during or within 4 hours of cessation of transfusion:</p> <ul style="list-style-type: none"> • Conjunctival edema • Edema of lips, tongue and uvula • Erythema and edema of the periorbital area • Generalized flushing • Hypotension • Localized angioedema • Maculopapular rash • Pruritus (itching) • Respiratory distress; bronchospasm • Urticaria (hives) <p>Probable: ANY 1 of the following occurring during or within 4 hours of cessation of transfusion:</p> <ul style="list-style-type: none"> • Conjunctival edema • Edema of lips, tongue and uvula • Erythema and edema of the periorbital area • Localized angioedema • Maculopapular rash • Pruritus (itching) • Urticaria (hives) 	<p>Severe, Life-threatening, Death: Involves respiratory and/or cardiovascular systems and presents like an anaphylactic reaction. There is anaphylaxis when, in addition to mucocutaneous symptoms, there are airway symptoms, hypotension, or associated symptoms like hypotonia and syncope. The respiratory signs and symptoms may be laryngeal (tightness in the throat, dysphagia, dysphonia, hoarseness, stridor) or pulmonary (dyspnea, cough, wheezing, bronchospasm, hypoxemia). Such a reaction usually occurs during or shortly after cessation of transfusion.</p> <p>Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Definite: Occurs during or within 2 hours of cessation of transfusion AND No other evidence of environmental, drug or dietary risks.</p> <p>Probable: Occurs during or within 2 hours of cessation of transfusion AND There are other potential causes present that could explain symptoms, but transfusion is the most likely cause.</p> <p>Possible: Occurs 2 - 4 hours after cessation of transfusion OR Other present causes are most likely, but transfusion cannot be ruled out.</p>
OPTIONAL	OPTIONAL	OPTIONAL
<p>Possible: N/A</p>	<p>Non-severe: There is no immediate risk to the life of the patient, and the patient responds quickly to symptomatic treatment.</p>	<p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Hypotensive transfusion reaction

Case Definition	Severity	Imputability
<p>Definitive: All other adverse reactions presenting with hypotension are excluded AND Hypotension occurs during or within 1 hour after cessation of transfusion.</p> <ul style="list-style-type: none"> • Adults (18 years and older): Drop in systolic BP of greater than or equal to 30 mmHg and systolic BP less than or equal to 80 mmHg. • Infants, children and adolescents (1 year to less than 18 years old): Greater than 25% drop in systolic BP from baseline (e.g., drop in systolic BP of 120mmHg to below 90mmHg). • Neonates and small infants (less than 1 year old OR any age and less than 12 kg body weight): Greater than 25% drop in baseline value using whichever measurement is being recorded (e.g., mean BP). <p>Probable: N/A</p>	<p>Non-severe: The recipient required no more than discontinuation of transfusion and symptom management and no long-term morbidity resulted from the reaction.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to hypotension, or hypotension led directly to long-term morbidity (e.g., brain damage) AND Vasopressors were not required.</p> <p>Life-threatening: The recipient required vasopressors.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p>	<p>Definite: Occurs less than 15 minutes after the start of the transfusion AND Responds rapidly (i.e., within 10 minutes) to cessation of transfusion and supportive treatment AND The patient has no other conditions that could explain hypotension.</p> <p>Probable: Onset is between 15 minutes after start and 1 hour after cessation of transfusion OR The patient does not respond rapidly to cessation of transfusion and supportive treatment OR There are other potential causes present that could explain hypotension, but transfusion is the most likely cause.</p> <p>Possible: Other conditions that could readily explain hypotension are present.</p>
OPTIONAL		OPTIONAL
<p>Possible: Hypotension occurs, does not meet the criteria above. Other, more specific reaction definitions do not apply.</p>	<p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Febrile non-hemolytic transfusion reaction (FNHTR)

Note: Reactions may be classified as FNHTRs in the absence of fever if chills or rigors occur.

Case Definition	Severity	Imputability
<p>Definitive: Occurs during or within 4 hours of cessation of transfusion AND EITHER Fever (greater than or equal to 38°C/100.4°F oral and a change of at least 1°C/1.8°F) from pre-transfusion value OR Chills/rigors are present.</p> <p>Probable: N/A</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p>	<p>Definite: Patient has no other conditions that could explain signs/symptoms.</p> <p>Probable: There are other potential causes present that could explain signs/symptoms, but transfusion is the most likely cause.</p> <p>Possible: Other present causes are most likely, but transfusion cannot be ruled out.</p>
OPTIONAL		OPTIONAL
<p>Possible: FNHTR is suspected, but reported symptoms and/or available information are not sufficient to meet the criteria defined above. Other, more specific adverse reaction definitions do not apply.</p>	<p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Acute hemolytic transfusion reaction (AHTR)

Note: Report hemolytic reactions resulting from immune or non-immune causes, including when the recipient is **intentionally** transfused with incompatible blood components.

Case Definition	Severity	Imputability
<p>Definitive: Occurs during, or within 24 hours of cessation of transfusion with new onset of ANY of the following signs/symptoms:</p> <ul style="list-style-type: none"> • Back/flank pain • Chills/rigors • Disseminated intravascular coagulation (DIC) • Epistaxis • Fever • Hematuria (gross visual hemolysis) • Hypotension • Oliguria/anuria • Pain and/or oozing at IV site • Renal failure <p>AND 2 or more of the following:</p> <ul style="list-style-type: none"> • Decreased fibrinogen • Decreased haptoglobin • Elevated bilirubin • Elevated LDH • Hemoglobinemia • Hemoglobinuria • Plasma discoloration c/w hemolysis • Spherocytes on blood film <p>AND EITHER (IMMUNE-MEDIATED) Positive direct antiglobulin test (DAT) for anti-IgG or anti-C3</p> <p>AND Positive elution test with alloantibody present on the transfused red blood cells</p> <p>OR (NON-IMMUNE MEDIATED) Serologic testing is negative, and physical cause (e.g., thermal, osmotic, mechanical, chemical) is confirmed.</p> <p>Probable: Meets signs and symptoms criteria for acute hemolysis</p> <p>AND EITHER (IMMUNE MEDIATED) Physical cause is excluded but serologic evidence is not sufficient to meet definitive criteria</p> <p>OR (NON-IMMUNE MEDIATED) Physical cause is suspected and serologic testing is negative.</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p> <p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Definite: ABO or other allotypic RBC antigen incompatibility is known</p> <p>OR Only transfusion-related (i.e., immune or non-immune) cause of acute hemolysis is present.</p> <p>Probable: There are other potential causes present that could explain acute hemolysis, but transfusion is the most likely cause.</p> <p>Possible: Other causes of acute hemolysis are more likely, but transfusion cannot be ruled out.</p> <hr/> <p style="text-align: center;">OPTIONAL</p> <p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>
<p style="text-align: center;">OPTIONAL</p> <p>Possible: AHTR is suspected within 24 hours of cessation of transfusion, but symptoms, test results, and/or information are not sufficient to meet the criteria defined above. Other, more specific adverse definitions do not apply.</p>		



Delayed hemolytic transfusion reaction (DHTR)

Note: Report all hemolytic reactions, including when the recipient is **intentionally** transfused with incompatible blood components.

Case Definition	Severity	Imputability
<p>Definitive: Positive direct antiglobulin test (DAT) for antibodies developed between 24 hours and 28 days after cessation of transfusion AND EITHER Positive elution test with alloantibody present on the transfused red blood cells OR Newly-identified red blood cell alloantibody in recipient serum AND EITHER Inadequate rise of post-transfusion hemoglobin level or rapid fall in hemoglobin back to pre-transfusion levels OR Otherwise unexplained appearance of spherocytes.</p> <p>Probable: Newly-identified red blood cell alloantibody demonstrated between 24 hours and 28 days after cessation of transfusion BUT Incomplete laboratory evidence to meet definitive case definition criteria.</p> <p>NOTE: Patient may be asymptomatic or have symptoms that are similar to but milder than AHTR; symptoms are not required to meet case definition criteria.</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p> <p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Definite: No other explanation for symptoms or newly-identified antibody is present.</p> <p>Probable: An alternate explanation for symptoms or newly-identified antibody is present, but transfusion is the most likely cause.</p> <p>Possible: Other explanations for symptoms or newly-identified antibody are more likely, but transfusion cannot be ruled out.</p>
OPTIONAL		OPTIONAL
<p>Possible: DHTR is suspected, but reported symptoms, test results, and/or available information are not sufficient to meet the criteria defined above. Other, more specific adverse reaction definitions do not apply.</p>		<p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Delayed serologic transfusion reaction (DSTR)

Note: Delayed serologic reactions should only be reported for patients **transfused by your facility**.

Case Definition	Severity	Imputability
<p>Definitive: Absence of clinical signs of hemolysis AND Demonstration of new, clinically-significant antibodies against red blood cells BY EITHER Positive direct antiglobulin test (DAT) OR Positive antibody screen with newly identified RBC alloantibody.</p> <p>Probable: N/A</p> <p>Possible: N/A</p>	<p>Not Determined: Since this is by definition a reaction with no clinical symptoms, severity of the reaction cannot be graded.</p>	<p>Definite: New alloantibody is identified between 24 hours and 28 days after cessation of transfusion AND Transfusion performed by your facility is the only possible cause for seroconversion.</p> <p>Probable: New alloantibody is identified between 24 hours and 28 days after cessation of transfusion AND 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.</p> <p>Possible: New alloantibody is identified between 24 hours and 28 days after cessation of transfusion AND The patient was transfused by your facility, but other exposures are present that most likely explain seroconversion.</p>
		OPTIONAL
		<p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Transfusion-associated graft vs. host disease (TAGVHD)

Case Definition	Severity	Imputability
<p>Definitive: A clinical syndrome occurring from 2 days to 6 weeks after cessation of transfusion characterized by:</p> <ul style="list-style-type: none"> • Characteristic rash: erythematous, maculopapular eruption centrally that spreads to extremities and may, in severe cases, progress to generalized erythroderma and hemorrhagic bullous formation. • Diarrhea • Fever • Hepatomegaly • Liver dysfunction (i.e., elevated ALT, AST, Alkaline phosphatase, and bilirubin) • Marrow aplasia • Pancytopenia <p>AND Characteristic histological appearance of skin or liver biopsy.</p> <p>Probable: Meets definitive criteria EXCEPT Biopsy negative or not done.</p> <p>Possible: N/A</p>	<p>Non-severe: N/A</p> <p>Severe: Patient had marked symptoms and responded to treatment.</p> <p>Life-threatening: Patient had severe symptoms and required life-saving treatment (e.g., immunosuppression).</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Definite: WBC chimerism present in the absence of alternative diagnoses.</p> <p>Probable: WBC chimerism present BUT Other potential causes are present (e.g., stem cell transplantation).</p> <p>Possible: WBC chimerism not present or not done OR Alternative explanations are more likely (e.g., solid organ transplantation).</p> <hr/> <p style="text-align: center;">OPTIONAL</p> <p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Post transfusion purpura (PTP)

Case Definition	Severity	Imputability
<p>Definitive: Alloantibodies in the patient directed against HPA or other platelet specific antigen detected at or after development of thrombocytopenia AND Thrombocytopenia (i.e., decrease in platelets to less than 20% of pre-transfusion count).</p> <p>Probable: Alloantibodies in the patient directed against HPA or other platelet specific antigen detected at or after development of thrombocytopenia. AND Decrease in platelets to levels between 20% and 80% of pre-transfusion count.</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p> <p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p>	<p>Definite: Occurs 5-12 days post-transfusion AND Patient has no other conditions to explain thrombocytopenia.</p> <p>Probable: Occurs less than 5 or more than 12 days post-transfusion OR There are other potential causes present that could explain thrombocytopenia, but transfusion is the most likely cause.</p> <p>Possible: Alternate explanations for thrombocytopenia are more likely, but transfusion cannot be ruled out.</p>
OPTIONAL		OPTIONAL
<p>Possible: PTP is suspected, but laboratory findings and/or information are not sufficient to meet defined criteria above. For example, the patient has a drop in platelet count to less than 80% of pre-transfusion count but HPA antibodies were not tested or were negative. Other, more specific adverse reaction definitions do not apply.</p>	<p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Transfusion-transmitted infection (TTI)

Case Definition	Severity	Imputability
<p>Definitive: Laboratory evidence of a pathogen in the transfusion recipient.</p> <p>Probable: N/A</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p>	<p>Definite: ONE or more of the following:</p> <ul style="list-style-type: none"> Evidence of the pathogen in the transfused component Evidence of the pathogen in the donor at the time of donation Evidence of the pathogen in an additional component from the same donation Evidence of the pathogen in an additional recipient of a component from the same donation <p>AND No other potential exposures to the pathogen could be identified in the recipient.</p> <p>AND EITHER Evidence that the recipient was not infected with the pathogen prior to transfusion OR Evidence that the identified pathogen strains are related by molecular or extended phenotypic comparison testing with statistical confidence (p<0.05).</p> <p>Probable: ONE or more of the following:</p> <ul style="list-style-type: none"> Evidence of the pathogen in the transfused component Evidence of the pathogen in the donor at the time of donation Evidence of the pathogen in an additional component from the same donation Evidence of the pathogen in an additional recipient of a component from the same donation. <p>AND EITHER: Evidence that the recipient was not infected with this pathogen prior to transfusion OR No other potential exposures to the pathogen could be identified in the recipient.</p> <p>Possible: Case fails to meet definite, probable, doubtful, or ruled out imputability criteria.</p>
OPTIONAL		OPTIONAL
<p>Possible: Temporally associated unexplained clinical illness consistent with infection, but no pathogen is detected in the recipient. Other, more specific adverse reactions are ruled out.</p> <p>Note: Possible cases cannot meet the definite or probable imputability criteria.</p>	<p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Doubtful: Laboratory evidence that the recipient was infected with this pathogen prior to transfusion OR Evidence is clearly in favor of a cause other than transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: ALL of the following (where applicable):</p> <ul style="list-style-type: none"> Evidence that the transfused component was negative for this pathogen at the time of transfusion Evidence that the donor was negative for this pathogen at the time of donation Evidence that additional components from the same donation were negative for this pathogen <p>OR There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Transfusion-transmitted infection (TTI)

(continued)

Pathogens of well-documented importance in blood safety.

These pathogens have public health significance for hemovigilance, are well-documented blood stream pathogens, and/or are routinely screened for in blood donors. A full list of potentially infectious organisms is available in the drop-down pathogen list in NHSN.			
Bacterial	Viral	Parasitic	Other
<i>Enterobacter cloacae</i> <i>Escherichia coli</i> <i>Klebsiella oxytoca</i> <i>Klebsiella pneumoniae</i> <i>Pseudomonas aeruginosa</i> <i>Serratia marcescens</i> <i>Staphylococcus aureus</i> <i>Staphylococcus epidermidis</i> <i>Staphylococcus lugdunensis</i> Syphilis (<i>Treponema pallidum</i>) <i>Yersinia enterocolitica</i>	Cytomegalovirus (CMV) <i>Enterovirus spp.</i> Epstein Barr (EBV) Hepatitis A Hepatitis B Hepatitis C Human Immunodeficiency Virus 1 (HIV-1) Human Immunodeficiency Virus 2 (HIV-2) Human Parvovirus B-19 Human T-Cell Lymphotropic Virus-1 (HTLV-1) Human T-Cell Lymphotropic Virus-2 (HTLV-2) West Nile Virus (WNV)	Babesiosis (<i>Babesia spp.</i>) Chagas disease (<i>Trypanosoma cruzi</i>) Malaria (<i>Plasmodium spp.</i>)	Creutzfeldt-Jakob Disease, Variant (vCJD)

Investigation triggers for potential transfusion-transmitted infections:

1. Identification by testing (e.g., gram stain, other smear/staining, culture, or other method) of a bacterial, mycobacterial, or fungal pathogen in a recipient within the time period from exposure (i.e., transfusion) to onset of infection appropriate for the suspected pathogen.
2. Identification of an unexpected virus in the transfusion recipient by testing (e.g., culture, direct fluorescent antibody, or polymerase chain reaction) within the time period from exposure (i.e., transfusion) to onset of infection appropriate for the suspected virus.
3. Identification of an unexpected parasite in the recipient by testing (e.g., blood smear, histopathology, serologic testing, or polymerase chain reaction) within the time period from exposure (i.e., transfusion) to onset of infection appropriate for the suspected parasite.
4. Any of the above laboratory findings in the recipient unit upon residual testing.
5. Unexplained clinical events occurring after transfusion that are consistent with transfusion-transmitted infection, such as:
 - a. Encephalitis, meningitis, or other unexplained central nervous system abnormalities.
 - b. Sepsis with or without multi-organ system dysfunction.
 - c. Hemolytic anemia and/or fever (e.g., in cases of transfusion-associated babesiosis or malaria).
 - d. Recipient death.
6. For pathogens routinely screened in the blood donor, any infection in the recipient occurring within 6 months after transfusion if:
 - a. The index donation testing was negative but
 - b. The donor was subsequently found to be infected, and
 - c. The recipient had no pre-transfusion history of the same infection.



Other or Unknown

Other: Use this option if the recipient experienced an adverse reaction that is not defined in the Hemovigilance Module surveillance protocol (e.g., transfusion-associated acute gut injury (TRAGI), transfusion-associated immunomodulation (TRIM), iron overload, microchimerism, hyperkalemia, thrombosis).

Unknown: Use this category if the patient experienced transfusion-related symptoms, but the medical event that caused those symptoms could not be classified.

Note: Reporting 'Other' and 'Unknown' reactions is not required by CDC.

REPORTING OPTIONAL		
Case Definition	Severity	Imputability
<p>Not Applicable: CDC does not specifically define the 'Other' or 'Unknown' adverse reaction categories, therefore the case definition criteria may only be reported as N/A.</p>	<p>Non-severe: Medical intervention (e.g. symptomatic treatment) is required but lack of such would not result in permanent damage or impairment of a bodily function.</p> <p>Severe: Inpatient hospitalization or prolongation of hospitalization is directly attributable to the adverse reaction, persistent or significant disability or incapacity of the patient occurs as a result of the reaction, or a medical or surgical intervention is necessary to preclude permanent damage or impairment of a body function.</p> <p>Life-threatening: Major intervention required following the transfusion (e.g. vasopressors, intubation, transfer to intensive care) to prevent death.</p> <p>Death: The recipient died as a result of the adverse transfusion reaction. Death should be used if death is possibly, probably or definitely related to transfusion. If the patient died of a cause other than the transfusion, the severity of the reaction should be graded as appropriate given the clinical circumstances related to the reaction.</p> <p>Not Determined: The severity of the adverse reaction is unknown or not stated.</p>	<p>Definite: Conclusive evidence exists that the adverse reaction can be attributed to the transfusion.</p> <p>Probable: Evidence is clearly in favor of attributing the adverse reaction to the transfusion.</p> <p>Possible: Evidence is indeterminate for attributing the adverse reaction to the transfusion or an alternate cause.</p> <p>Doubtful: Evidence is clearly in favor of a cause other than the transfusion, but transfusion cannot be excluded.</p> <p>Ruled Out: There is conclusive evidence beyond reasonable doubt of a cause other than the transfusion.</p> <p>Not Determined: The relationship between the adverse reaction and the transfusion is unknown or not stated.</p>



Adverse Reaction Glossary

Antibodies often associated with AHTR, DHTR, DSTR:

Anti-A	Anti-B	Anti-A,B	Anti-C	Anti-c	Anti-D	Anti-E	Anti-e	Anti-Fy ^a
Anti-Fy ^b	Anti-Jk ^a	Anti-Jk ^b	Anti-K	Anti-k	Anti-M	Anti-S	Other	

Bronchospasm (wheezing): A contraction of smooth muscle in the walls of the bronchi and bronchioles, causing acute narrowing and obstruction of the respiratory airway. This constriction can result in a rasp or whistling sound while breathing.

Chills/rigors: A feeling of cold with shivering or shaking and pallor.

Disseminated intravascular coagulation (DIC): Bleeding disorder characterized by reduction in the factors involved in blood clotting due to their use in widespread clotting within the vessels. The intravascular clotting ultimately produces hemorrhage because of rapid consumption of clotting factors.

Edema: Swelling of soft tissues as a result of excessive fluid accumulation.

Epistaxis: Bleeding from the nose.

Fever: For the purposes of hemovigilance, an increase of at least 1°C in temperature over the pre-transfusion value.

Hematuria: Presence of blood or red blood cells in the urine.

Hemoglobinemia: The presence of free hemoglobin in the blood plasma.

Hemoglobinuria: Presence of free hemoglobin in the urine.

Hypoxemia: Abnormal deficiency in the concentration of oxygen in arterial blood. PaO₂ / FiO₂ less than or equal to 300 mm Hg OR oxygen saturation is less than 90% on room air.

Jaundice: New onset or worsening of yellow discoloration (icterus) of the skin or sclera (scleral icterus) secondary to an increased level of bilirubin.

Oliguria: New onset of decreased urinary output (less than 500cc output per 24 hours).

Other rash: Non-urticarial skin rash.

Pruritus: Itching.

Shock: A drop in blood pressure accompanied by a drop in cardiac output including rapid heart rate (increase to 100 beats per minute or more), rapid breathing, cutaneous vasoconstriction, pallor, sweating, decreased or scanty urine production, agitation and/or loss of consciousness that required fluid resuscitation, with or without inotropic support.

Shortness of breath (dyspnea): New onset or significant worsening of shortness of breath; or a significant increase in respiratory rate (with or without hypoxemia).

Urticaria (hives): Raised wheals on the skin.



Section 5. Hemovigilance Module Incidents

Required Reporting

All incidents (i.e., accidents or errors) that are **associated with a reported adverse reaction** must be reported to NHSN using a detailed Incident form (CDC 57.305). If multiple incidents occur in association with an adverse reaction then report all. Incidents may occur before (e.g., wrong product released) or after (e.g., failure to report adverse reaction to blood bank) an adverse reaction. Each reaction must be reported using the detailed incident form; the incident result must be coded as 'Product transfused, reaction' to enter the associated patient identifier on the form. After the incident record is entered, the adverse reaction record must be linked to the incident record in the NHSN web application.

Incident Classification

Use the incident codes provided at the end of this section to classify incidents. If there is uncertainty then please contact NHSN User Support

Optional Reporting

Any incident may be optionally reported to NHSN using the detailed Incident form (57.305) or the Monthly Incident Summary form (57.302). Approved deviations from standard operating procedure are not considered incidents because they did not occur by accident or in error. However, approved deviations may be optionally reported for a facility's use. Incidents that are optionally reported will not be aggregated or analyzed by CDC.

Form

[CDC 57.305 Hemovigilance Module Incident](#)

Form Instructions

[CDC 57.305 Hemovigilance Module Incident Table of Instructions](#)

Summary Form (Optional)

[CDC 57.302 Hemovigilance Module Monthly Incident Summary](#)

Summary Form Instructions (Optional)

[CDC 57.302 Hemovigilance Module Monthly Incident Summary Table of Instructions](#)



Incident Codes

Note: Incident codes are based on MERS TM (US) and TESS (Canada) incident classification schemes.

<p>Product Check-In <i>(Transfusion Service)</i> Events that occur during the shipment and receipt of products into the transfusion service from the supplier, another hospital site, satellite storage, or clinical area.</p> <ul style="list-style-type: none"> PC 00 Detail not specified PC 01 Data entry incomplete/incorrect/not performed PC 02 Shipment incomplete/incorrect PC 03 Products and paperwork do not match PC 04 Shipped/transported under inappropriate conditions PC 05 Inappropriate return to inventory PC 06 Product confirmation incorrect/not performed PC 07 Administrative check not incorrect/not performed (record review/audit) PC 08 Product label incorrect/missing <p>Product Storage <i>(Transfusion Service)</i> Events that occur during product storage by the transfusion service.</p> <ul style="list-style-type: none"> US 00 Detail not specified US 01 Incorrect storage conditions US 03 Inappropriate monitoring of storage device US 04 Unit stored on incorrect shelf (e.g., ABO/autologous s/directed) US 05 Incorrect storage location <p>Inventory Management <i>(Transfusion Service)</i> Events that involve quality management of the blood product inventory.</p> <ul style="list-style-type: none"> IM 00 Detail not specified IM 01 Inventory audit incorrect/not performed IM 02 Product status incorrectly/not updated online (e.g., available/discarded) IM 03 Supplier recall/traceback not appropriately addressed/not performed IM 04 Product order incorrectly/not submitted to supplier IM 05 Outdated product in available inventory IM 06 Recalled/quarantined product in available inventory 	<p>Product/Test Request <i>(Clinical Service)</i> Events that occur when the clinical service orders patient tests or blood products for transfusion.</p> <ul style="list-style-type: none"> PR 00 Detail not specified PR 01 Order for wrong patient PR 02 Order incompletely/incorrectly ordered (online order entry) PR 03 Special processing needs not indicated (e.g., CMV negative, autologous) PR 04 Order not done PR 05 Inappropriate/unnecessary (intended) test ordered PR 06 Inappropriate/unnecessary (intended) blood product ordered PR 07 Incorrect (unintended) test ordered PR 08 Incorrect (unintended) blood product ordered <p>Product/Test Order Entry <i>(Transfusion Service)</i> Events that occur when the transfusion service receives a patient order. This process may be excluded if clinical service uses online ordering.</p> <ul style="list-style-type: none"> OE 00 Detail not specified OE 01 Order entered for wrong patient OE 02 Order incompletely/incorrectly entered online OE 03 Special processing needs not entered (e.g., CMV-, autologous) OE 04 Order entry not done OE 05 Inappropriate/unnecessary (intended) test order entered OE 06 Inappropriate/unnecessary (intended) blood product order entered OE 07 Incorrect (unintended) test ordered OE 08 Incorrect (unintended) blood product ordered <p>Sample Collection <i>(Service collecting the samples)</i> Events that occur during patient sample collection.</p> <ul style="list-style-type: none"> SC 00 Detail not specified SC 01 Sample labeled with incorrect patient name SC 02 Not labeled SC 03 Wrong patient collected SC 04 Collected in wrong tube type SC 05 Sample QNS SC 06 Sample hemolyzed SC 07 Label incomplete/illegible/incorrect (other than patient name) SC 08 Sample collected in error SC 09 Requisition arrived without samples SC 10 Wristband incorrect/not available SC 11 Sample contaminated
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Incident Codes

(continued)

Note: Incident codes are based on MERS TM (US) and TESS (Canada) incident classification schemes.

<p>Sample Handling <i>(Service collecting the samples)</i> <i>Events that occur when a patient sample is sent for testing.</i></p> <ul style="list-style-type: none">SH 00 Detail not specifiedSH 01 Sample sent without requisitionSH 02 Requisition and sample label don't matchSH 03 Patient ID incomplete/illegible on requisitionSH 04 No Patient ID on requisitionSH 05 No phlebotomist/witness identificationSH 06 Sample sent with incorrect requisition typeSH 07 Patient information (other than ID) missing/incorrect on requisitionSH 08 Requisition sent without sampleSH 09 Data entry incorrect/incomplete/not performedSH 10 Sample transport issue (e.g., sample broken/inappropriate conditions)SH 11 Duplicate sample sent in error <p>Sample Receipt <i>(Transfusion Service)</i> <i>Events that occur when a sample is received by the transfusion service.</i></p> <ul style="list-style-type: none">SR 00 Detail not specifiedSR 01 Sample accepted in errorSR 02 Historical review incorrect/not performedSR 03 Demographic review/ data entry incorrect/not performedSR 04 Sample incorrectly accessioned <p>Sample Testing <i>(Transfusion Service)</i> <i>Events that occur during patient sample testing by the transfusion service.</i></p> <ul style="list-style-type: none">ST 00 Detail not specifiedST 01 Data entry incomplete/incorrect/not performedST 02 Appropriate sample checks incomplete/incorrect/not performedST 03 Computer warning overridden in error or outside SOPST 05 Sample test tube incorrectly accessionedST 07 Sample test tubes mixed upST 09 Sample test tube mislabeled (wrong patient identifiers)ST 10 Equipment problem/failure/not properly QC'dST 12 Sample testing not performedST 13 Incorrect sample testing method chosenST 14 Sample testing performed incorrectlyST 15 Sample test result misinterpreted	<p>Sample Testing (continued)</p> <ul style="list-style-type: none">ST 16 Reagents used were incorrect/inappropriate/expired/not properly QC'dST 17 ABO/Rh error caught on final checkST 18 Current/historical ABO/Rh mismatchST 19 Additional testing not performedST 20 Confirmatory check incorrect/not performed (at time work performed)ST 21 Administrative check incorrect/not performed (record review/audit)ST 22 Sample storage incorrect/inappropriate <p>Product Manipulation/Processing/Testing <i>(Transfusion Service)</i> <i>Events that occur while testing, manipulating (e.g., pooling, washing, aliquoting, irradiating), processing, or labeling blood products.</i></p> <ul style="list-style-type: none">UM 00 Detail not specifiedUM 01 Data entry incomplete/incorrect/not performedUM 02 Record review incomplete/incorrect/not performedUM 03 Incorrect product (type) selectedUM 04 Incorrect product (patient) selectedUM 05 Product labeled incorrectly (new/updated)UM 06 Computer warning overridden in error or outside SOPUM 07 Special processing needs not checkedUM 08 Special processing needs misunderstood or misinterpretedUM 09 Special processing needs performed incorrectlyUM 10 Special processing needs not performedUM 11 Equipment problem/failure/not properly QC'dUM 12 Reagents used were incorrect/inappropriate/expired/not properly QC'dUM 13 Confirmatory check incorrect/not performed (at time work performed)UM 14 Administrative check incorrect/not performed (record review/audit)
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Incident Codes

(continued)

Note: Incident codes are based on MERS TM (US) and TESS (Canada) incident classification schemes.

<p>Request for Pick-up <i>(Clinical Service)</i> <i>Events that occur when the clinical service requests pick-up of a blood product from the transfusion service.</i></p> <ul style="list-style-type: none">RP 00 Detail not specifiedRP 01 Request for pick-up on wrong patientRP 02 Incorrect product requested for pick-upRP 03 Product requested prior to obtaining consentRP 04 Product requested for pick-up, but patient not availableRP 05 Product requested for pick-up, but IV not readyRP 06 Request for pick-up incomplete (e.g., patient ID/product type missing)RP 07 Pick-up slip did not match patient information on product <p>Product Issue <i>(Transfusion Service)</i> <i>Events that occur when the transfusion service issues blood product to the clinical service.</i></p> <ul style="list-style-type: none">UI 00 Detail not specifiedUI 01 Data entry incomplete/incorrect/not performedUI 02 Record review incomplete/incorrect/not performedUI 03 Product issued for wrong patientUI 04 Product issued out of orderUI 05 Product issue delayedUI 06 LIS warning overridden in error or outside SOPUI 07 Computer issue not completedUI 08 Issued visibly defective product (e.g., clots/aggregates/particulate matter)UI 09 Not/incorrect checking of unit and/or patient informationUI 10 Product transport issues (e.g., delayed) by transfusion serviceUI 11 Unit delivered to incorrect location by transfusion serviceUI 12 Product transport issue (from transfusion service to clinical area)UI 18 Wrong product issued for intended patient (e.g., incompatible)UI 19 Inappropriate product issued for patient (e.g., not irradiated, CMV+)UI 20 Confirmatory check incorrect/not performed (at time work performed)UI 21 Administrative check incorrect/not performed (record review/audit)UI 22 Issue approval not obtained/documentedUI 23 Receipt verification not performed (pneumatic tube issue)	<p>Satellite Storage <i>(Clinical Service)</i> <i>Events that occur while product is stored and handled by the clinical service.</i></p> <ul style="list-style-type: none">CS 00 Detail not specifiedCS 01 Incorrect storage conditions of product in clinical areaCS 02 Incorrect storage location in the clinical areaCS 03 Labeling issue (by clinical staff)CS 04 Floor/clinic did not check for existing products in their areaCS 05 Product transport issues (to or between clinical areas)CS 06 Monitoring of satellite storage incorrect/incomplete/not performedCS 07 Storage tracking/documentation incorrect/incomplete/not performed <p>Product Administration <i>(Clinical Service)</i> <i>Events that occur during the administration of blood products.</i></p> <ul style="list-style-type: none">UT 00 Detail not specifiedUT 01 Administered intended product to wrong patientUT 02 Administered wrong product to intended patientUT 03 Transfusion not performed in errorUT 05 Bedside check (patient ID confirmation) incomplete/not performedUT 06 Transfused product with incompatible IV fluidUT 07 Transfusion delayed beyond pre-approved timeframeUT 09 Transfused unsuitable product (e.g., outdated/inappropriately stored)UT 10 Administered components in wrong orderUT 11 Appropriate monitoring of patient not performedUT 14 Transfusion volume too low (per order or SOP)UT 15 Transfusion volume too high (per order or SOP)UT 16 Transfusion rate too slow (per order or SOP)UT 17 Transfusion rate too fast (per order or SOP)UT 18 Inappropriate preparation of productUT 19 Transfusion protocol not followed (not otherwise specified)UT 22 Order/consent check incorrect/not performedUT 23 Transfusion documentation incorrect/incomplete/not performedUT 24 Transfusion documentation not returned to transfusion serviceUT 26 Transfusion reaction protocol not followed <p>Other</p> <ul style="list-style-type: none">MS 99 Other
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Occupation Codes

Laboratory		Additional Occupation Types	
IVT	IVT Team Staff	ATT	Attendant/Orderly
MLT	Medical Laboratory Technician	CSS	Central Supply
MTE	Medical Technologist	CSW	Counselor/Social Worker
PHL	Phlebotomist/IV Team	DIT	Dietician
Nursing		DNA	Dental Assistant/Technician
LPN	Licensed Practical Nurse	DNH	Dental Hygienist
CNA	Nurse Anesthetist	DNO	Other Dental Worker
CNM	Certified Nurse Midwife	DNT	Dentist
NUA	Nursing Assistant	DST	Dental Student
NUP	Nurse Practitioner	FOS	Food Service
RNU	Registered Nurse	HSK	Housekeeper
Physician		ICP	Infection Control Professional
FEL	Fellow	LAU	Laundry Staff
MST	Medical Student	MNT	Maintenance/Engineering
PHY	Attending/Staff Physician	MOR	Morgue Technician
RES	Intern/Resident	OAS	Other Ancillary Staff
Technicians		OFR	Other First Responder
EMT	EMT/Paramedic	OH	Occupational Health Professional
HEM	Hemodialysis Technician	OMS	Other Medical Staff
ORS	OR/Surgery Technician	OTH	Other
PCT	Patient Care Technician	OTT	Other Technician/Therapist
Other Personnel		PAS	Physician Assistant
CLA	Clerical/Administrative	PHA	Pharmacist
TRA	Transport/Messenger/Porter	PHW	Public Health Worker
		PLT	Physical Therapist
		PSY	Psychiatric Technician
		RCH	Researcher
		RDT	Radiologic Technologist
		RTT	Respiratory Therapist/Technician
		STU	Other Student
		VOL	Volunteer



Incident Glossary

Incident Result

Product transfused; reaction (No recovery; harm):

A product related to this incident was transfused; the patient experienced an adverse reaction.

Product transfused; no reaction (No recovery; no harm):

A product related to this incident was transfused; the patient did not experience an adverse reaction.

No product transfused; unplanned recovery (Near miss; unplanned recovery):

No product related to this incident was transfused; the incident was discovered ad hoc, by accident, by human lucky catch, etc.

No product transfused; planned recovery (Near miss; planned recovery):

No product related to this incident was transfused; the incident was discovered through a standardized process or barrier designed to prevent errors.



Section 6. Hemovigilance Module Denominators

Required Reporting

Facilities must report the total number of units and aliquots of specified blood components transfused and total number of discards each month. When reporting aliquots, the units from which they are made should **NOT** be counted as a transfused unit. The components transfused count should include autologous units. The total number of patient samples collected and total crossmatch procedures must also be reported. This form must be completed each month that surveillance is conducted and data can only be entered once the calendar month is over. For instance, February data must be entered after March 1st. Additionally, data cannot be entered for upcoming months.

Pathogen Reduced Blood Products

The total number of transfused units of blood components which are produced with pathogen-reduction technology (PRT) should be reported each month, if applicable. These PRT units are reported in Table 2 and are a subset of total number of units and aliquots transfused that are reported in Table 1. Table 3 relates to pathogen reduced apheresis platelets, if reported in table 2. For more guidance please refer to the Denominator QuickLearn on the [NHSN Blood Safety Surveillance website](#).

Electronic Reporting

In January 2017, the NHSN Hemovigilance module can accept electronically reported denominator data via clinical documentation architecture (CDA). Compared to manual reporting, electronic reporting will decrease the time required for data collection and reporting, reduce data entry errors, and increase data granularity. In order to electronically report data, facilities' software system must have CDA functionality. For more information about electronic reporting and CDA, review CDA guidance on the [NHSN Blood Safety Surveillance website](#).

Form

[CDC 57.303 Hemovigilance Module Monthly Reporting Denominators](#)

Form Instructions

[CDC 57.303 Hemovigilance Module Monthly Reporting Denominators Tables of Instructions](#)