

## Hemovigilance Module Adverse Reaction Transfusion Associated Dyspnea

\*Required for saving NHSN Adverse Reaction #: \*Facility ID#: **Patient Information** \*Patient ID: \*Gender: M Other \*Date of Birth: / / Gender Identity (Specify): Sex at Birth: M F Unknown Male Female Male-to-female transgender Female-to-male transgender Identifies as non-conforming Other Asked but unknown\_\_\_\_\_ Social Security #: Secondary ID: Medicare #: Middle Name: First Name: Last Name: Ethnicity (Specify): Race (Specify): (Select all that apply): Hispanic or Latino American Indian or Alaska Native Not Hispanic or Latino Asian Unknown Black or African American Declined to respond Middle Eastern or North African Native Hawaiian or Pacific Islander White Unknown Declined to respond Interpreter Needed: Yes Preferred Language (Specify): Declined to Respond Unknown \*Blood Group: B-Blood type not done B+ Transitional ABO / Transitional Transitional ABO / Rh + Transitional ABO / Rh -Rh Group A/Transitional Group B/Transitional Group O/Transitional Rh Group AB/Transitional Rh Rh **Patient Medical History** List the patient's admitting diagnosis. (Use ICD-10 Diagnostic codes/descriptions) Description: \_\_\_\_\_ Description: \_\_\_\_\_ Code: Code: \_\_\_\_\_ Description: \_\_\_\_\_ List the patient's underlying indication for transfusion. (Use ICD-10 Diagnostic codes/descriptions) Description: Code: \_\_\_\_\_ 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 57.315 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).



On the	
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:	· · · · · · · · · · · · · · · · · · ·
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Code: Description:	



## **Transfusion Associated Dyspnea**

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)									
Code:            Description:									
Code:            Description:									
Code: Description:									
Additional Information									
Transfusion History									
Has the patient received a previous transfusion?									
Blood Product: WB RBC Platelet Plasma Cryoprecipitate Granulocyte									
Date of Transfusion:/ UNKNOWN									
Was the patient's adverse reaction transfusion-related?									
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									
* Transfusion associated dyspnea (TAD)									
*Case Definition									
Check all that apply:									
Acute respiratory distress occurring within 24 hours of cessation of transfusion.									
Allergic reaction, TACO, and TRALI definitions are not applicable.									
Other signs and symptoms: (check all that apply)									
Generalized: Chills/rigors Fever Nausea/vomiting									
Cardiovascular: Blood pressure decrease Shock									
Cutaneous:									
Other rash Pruritus (itching) Urticaria (hives)									
Hemolysis/Hemorrhage: Disseminated intravascular coagulation Hemoglobinemia									
Positive antibody screen									
Pain: Abdominal pain Back pain Flank pain Infusion site pair									
Renal: Hematuria Hemoglobinuria Oliguria  Rilatoral infiltratos en chest y ray Renchespasm Cough									
Respiratory: Bilateral infiltrates on chest x-ray Bronchospasm 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, inlcuding prolonged hospitalization  Disability and/or incapacitation  Other medically important conditions  Symptomatic treatment only  Life-threatening reaction  Congenital anomaly or birth defect(s) of the fetus  Unknown or not stated								
*Imputability								
Which best describes the relationship between the transfusion and the reaction?								
Patient has no other conditions that could explain symptoms.								
There are other potential causes that could explain symptoms, 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								
Module-generated Designations  NOTE: Designations for each definition, soverity, and imputability will be automatically assigned in the NUSN								
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.								
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*Do you agree with the <u>case definition</u> designation?								
^Please indicate your designation								
*Do you agree with the severity designation?								
*Do you agree with the <u>severity</u> designation? YES NO ^Please indicate your designation								
Ticase indicate your designation								
*Do you agree with the <i>imputability</i> designation?								
^Please indicate your designation								
Dationt Treatment								
Patient Treatment								
Did the patient receive treatment for the transfusion reaction?  YES 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 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?										
Transfusion Start and End Date/Time	*Component code (check system used)	Amount transfused at reaction onset	(Requ	t number uired for ion and .l)	*Unit expiration Date/Time	*Blood group of unit	Implic ated Unit?			
^IMPLICATED UNIT										
	ISBT-128 Codabar	Entire unit Partial unitmL		·———			- Y B+ /A			
	ISBT-128	Entire unit Partial unitmL		 			N B+ /A			
Custom Field	ds									
Label				Label						
Comments										