

Hemovigilance Module Adverse Reaction Transfusion Associated Dyspnea

*Required for saving *Facility ID#: NHSN Adverse Reaction #: **Patient Information** *Date of Birth: /___/___ *Patient ID: *Sex: M Social Security #: Secondary ID: Medicare #: Last Name: First Name: Middle Name: Not Hispanic or Ethnicity (Specify): Hispanic or Latino Unknown Declined to respond Latino American Indian or Asian Black or African Middle Eastern or North Race (Select all that Alaska Native American African apply): Native Hawaiian or White Unknown Declined to respond Pacific Islander Interpreter Needed: Yes Preferred Language (Specify from the list provided): Unknow Declined to Respond AB+ *Blood Group: $A - \overline{A+}$ Blood type not done Transitional ABO / Transitional Transitional ABO / Rh + Transitional ABO / 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: Code: _____ Description: Code: ____ Description: List the patient's underlying indication for transfusion. (Use ICD-10 Diagnostic codes/descriptions) Code: _____ Description: Code: _____ Description: Code: Description: **UNKNOWN** List the patient's comorbid conditions at the time of the transfusion related to the adverse reaction. (Use ICD-10 Diagnostic codes/descriptions) NONE 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 57.315 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).



Code: _

Form Approved
OMB No. 0920-0666
Exp. Date: 12/31/2027
www.cdc.gov/nhsn
Description:



Transfusion Associated Dyspnea

	medical procedure including past procedure ent hospital or outpatient stay. (Use ICD-10								
Code:	Description:								
Code:									
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 Spe	cify								
Reaction Details									
*Date reaction occurred:// *Time reaction occurred:: Time unknown									
*Facility location where pa	tient was transfused:								
Is this reaction associated with an incident? Yes No If Yes, Incident #:									
Investigation Res	sults								
* Transfusion associate	d dyspnea (TAD)								
*Case Definition									
Check all that apply:									
Acute respiratory distress occurring within 24 hours of cessation of transfusion.									
	ACO, and TRALI definitions are not applicate	ble.							
Other signs and symptoms									
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								
	Positive antibody screen								
Pain:	Abdominal pain Back pain	Flank pain Infusion site pain							
_Renal:	Hematuria Hemoglobinuria								
Respiratory:	Bilateral infiltrates on chest x-ray	Bronchospasm Cough							
		roath							
	Hypoxemia Shortness of br	Calli							



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 Death 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 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? *Please indicate your designation *Do you agree with the severity designation? *Please indicate your designation *Do you agree with the imputability designation? *Please indicate your designation *Please indicate your designation							
Patient Treatment							
Did the patient receive treatment for the transfusion reaction? 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	ס							
Component	Details										
*Was a partic	cular unit implicated	d in (i.e., respo	onsib	le for) the a	dverse	Yes	No [N/A			
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 grou	dr	Implic ated Unit?			
^IMPLICATED	UNIT										
	ISBT-128 Codabar 	Entire unit Partial unitmL		 		A- A+ A+ AB- O- O+	AB+	Y			
: :	ISBT-128 Codabar	Entire unit Partial unitmL		.——— .————		A- A+ A+ AB- O- O+	AB+	N			
Custom Field	ds										
Label				Label							
		'	-			/	/				
Comments											
Comments											