

Unknown Transfusion Reaction

*Required for saving
*Facility ID#: NHSN Adverse Reaction #:
Patient Information
*Patient ID: *Gender: M F Other *Date of Birth://
Social Security #: Secondary ID: Medicare #:
Last Name:
Ethnicity Hispanic or Latino Not Hispanic or Not Latino
Race American Indian/Alaska Native Asian Black or African American
Native Hawaiian/Other Pacific Islander White
*Blood Group: A- A+ B- B+ AB- AB+ O- O+ Blood type not done
Patient Medical History (Use worksheet on page 4 for additional codes and descriptions.)
(part 1) List the patient's admitting diagnosis. (Use ICD-10 Diagnostic codes/descriptions)
Code: Description:
Code: Description:
Code: Description:
(part 2) List the patient's underlying indication for transfusion. (Use ICD-10 Diagnostic codes/descriptions)
Code: Description:
Code: Description:
Code: Description:
(part 3) 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:
Continued >>
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)). 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 D-74, Atlanta, GA 30333 ATTN: PRA (0920-0666).

CDC 57.319 (Front) Rev 1, v8.8



Patient Medical History (Use worksheet on page 4 for additional codes and descriptions.)
(part 4) 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:
(part 5) Additional Information
Transfusion History (Use worksheet on page 4 for additional transfusion history.)
*Has the patient received a previous transfusion? YES NO UNKNOWN
**If yes, provide information about the transfusion event. If not, skip to Reaction Details section. Blood Product: WB RBC Platelet Plasma Cryoprecipitate Granulocyte Date of Transfusion:// UNKNOWN
Did the patient experience a transfusion adverse reaction? 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?
After recognition of the transfusion reaction, was the current transfusion:
Continued Stopped and restarted Stopped indefinitely
Investigation Results
* Unknown
Diagnosis of case:
List tests relevant to reaction investigation:
Test name: Testing date: Test result:
Test name: Testing date: Test result:
Continued >>



Investigation Results (co	entinued)					
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 (ite	ching) Urticaria (hives)				
Hemolysis/Hemorrhage:	Disseminated intravascular coag	ulation Hemoglobinemia				
	Positive antibody screen					
Pain:	Abdominal pain Back pain	Flank pain Infusion site pain				
Renal:	Hematuria Hemoglobi					
Respiratory:	Bilateral infiltrates on chest x-ray	Bronchospasm Cough				
	Hypoxemia Shortness	of breath				
Other: (specify)						
*Severity	anna sia na a ann af tha fallan ia no (Daa	nana definitiona lista dia mosta an				
	experience any of the following? (Res	•				
Symptomatic treat	,	cuding prolonged hospitalization				
Life-threatening re		and/or incapacitation				
	ly or birth defect(s) of the fetus	Death				
Other medically in	portant conditions	Unknown or not stated				
<u>*Imputability</u>						
	relationship between the transfusion a					
	e exists that the adverse reaction can					
	n favor of attributing the adverse react					
	•	ion to the transfusion or an alternate cause.				
		fusion, but transfusion cannot be excluded.				
	evidence beyond reasonable doubt of					
	ween the adverse reaction and the tra					
Did the transfusion occur	at your facility? YES NO)				
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 de		YES NO				
Please indicate your designati	on					
Do you agree with the severity	designation?	YES NO				
Please indicate your designati	on					
Do you agree with the imputat	oility designation?	YES NO				
Please indicate your designati	on					
Additional Information						
		Continued >>				



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 H1 receptor blockers 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
Hemodialysis Peritoneal Continuous Veno-Venous Hemofiltration
Phlebotomy
Other Specify:
Outcome
Minor or no
*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
Continued >>



Component Details (Use worksheet on page 4 for additional units.)								
*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 expiration			Blood group of unit		
^IMPLICATED UNIT								
	ISBT-128 Codabar	Entire unit Partial unitmL	 		A- B+	A+ AB- O+	B- AB+	Y
	ISBT-128 Codabar	Entire unit Partial unitmL	 		A- B+	A+ AB- O+	B- AB+ N/A	N
	☐ ISBT-128 ☐ Codabar — — — — —	Entire unit Partial unitmL			A- B+	A+ AB- O+	B- AB+ N/A	N
Custom Field	ds							
Label			Label					
		<u> </u>						
Comments								



Hemovigilance Module Additional Worksheet

	story
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Code: Code: Code: Code: Code: Code: (part 4) List the partocedures to be porcedure codes/dictional code: Code: Code: Code: Code: Code: Code:	Description:



Hemovigilance Module Additional Worksheet

Transfusion History
Has the patient received a previous transfusion? YES NO
**If yes, provide information about the transfusion event. If not, skip to Reaction Details section.
Blood Product: WB RBC Platelet Plasma Cryoprecipitate Granulocyte
Date of Transfusion:// UNKNOWN
Did the patient experience a transfusion adverse reaction? 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
Has the patient received a previous transfusion? YES NO
**If yes, provide information about the transfusion event. If not, skip to Reaction Details section.
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Date of Transfusion://UNKNOWN
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If yes, provide information about the transfusion adverse reaction.
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Hemovigilance Module Additional Worksheet

Component Details							
*Was a particul	ar unit implicated in ((i.e., responsi	ible for) the adverse read	ction? Y	es No N	I/A	
Transfusion Start and End Date/Time	*Component code (check system used)	Amount transfused at reaction onset	Unit number	*Unit expiration Date/Time	*Blood group of unit	Implic ated Unit?	
	ISBT-128 Codabar	Entire unit Partial unit mL		<i>l 1</i>	A- A+ B- B AB- AB+ O- O+ N/A	N	
	ISBT-128 Codabar	Entire unit Partial unit mL			A- A+ B- B- AB- AB+ O- O+ N/A	N	
	ISBT-128 Codabar	Entire unit Partial unit mL			A- A+ B- B- AB- AB+ O- O+ N/A	N	
	ISBT-128 Codabar	Entire unit Partial unitmL			A- A+ B- B- AB- AB+ O- O+ N/A	N	
	ISBT-128 Codabar	Entire unit Partial unit mL	 	<i>I. I.</i>	A- A+ B- B- AB- AB+ O- O+ N/A	N	
	ISBT-128 Codabar	Entire unit Partial unitmL		<u> </u>	A- A+ B- B AB- AB+ O- O+ N/A	N	
	SBT-128	Entire			A+ B-	IN	

National Healt Safety Net	hcare work			I	OMB No Exp. Date	m Approv o. 0920-06 e: xx/xx/20 :dc.gov/nh	666 Oxx
:	Codabar	unit Partial unitmL		A- B +	AB-	AB+	