

## Hemovigilance Module Adverse Reaction Transfusion Associated Circulatory Overload

*Facility ID#: NHSN Adverse Reaction #:   Patient Information      *Patient ID: *Patient ID: *Patient ID: *Sex: M F Social Security #: Secondary ID: Medicare #: Last Name: First Name: Middle Name: Last Name: First Name: Middle Name: Middle Name: Ethnicity (Specify): Hispanic or Latino Not Hispanic or Latino Ourknown Declined to respond American Indian or Alaska Native Antive Hawaiian or White Unknown Declined to respond African African Declined to respond Preferred Language (Specify from the list provided):
*Patient ID:       *Date of Birth:       /_/         *Sex:       M       F         Social Security #:       Secondary ID:       Medicare #:         Last Name:       First Name:       Middle Name:         Ethnicity (Specify):       Hispanic or Latino       Not Hispanic or         Ethnicity (Specify):       American Indian or       Asian         Black or African       Middle Eastern or North         Alaska Native       American       African         Native Hawaiian or       White       Unknown       Declined to respond         Interpreter Needed:       Yes       No
*Sex:       M       F         Social Security #:
Social Security #:       Secondary ID:       Medicare #:         Last Name:       First Name:       Middle Name:         Ethnicity (Specify):       Hispanic or Latino       Not Hispanic or         Ethnicity (Specify):       American Indian or       Asian       Black or African         Race (Select all that apply):       Alaska Native       American       African         Image: Secondary ID:       Not Hispanic or       Image: Secondary ID:       Image: Secondary ID:         Image: Secondary ID:       American Indian or       Asian       Black or African       Middle Eastern or North         Race (Select all that apply):       American Indian or       Asian       Image: Secondary ID:       Image: Secondary ID:
Last Name:
Ethnicity (Specify):       Hispanic or Latino       Not Hispanic or Latino       Unknown       Declined to respond         American Indian or Race (Select all that apply):       American Indian or Native Hawaiian or Pacific Islander       Asian       Black or African American African       Middle Eastern or North African         Interpreter Needed:       Yes       No
Ethnicity (Specify).       Hispanic of Latino       Latino       Declined to respond         Race (Select all that apply):       American Indian or Asian       Black or African       Middle Eastern or North         American Indian or Alaska Native       American       African       Declined to respond         Interpreter Needed:       Yes       No
Race (Select all that apply):       Alaska Native       American       African         Native Hawaiian or Pacific Islander       White       Unknown       Declined to respond         Interpreter Needed:       Yes       No
Preferred Language (Specify from the list provided):
*Blood Group:       A-       A+       B-       B+       AB+       O-       O+       Blood type not done         Transitional ABO / Rh +       Transitional ABO / Rh +       Transitional ABO / Rh -       Rh         Group A/Transitional       Group B/Transitional Rh       Group O/Transitional Rh       Group AB/Transitional Rh
Patient Medical History
List the patient's admitting diagnosis. (Use ICD-10 Diagnostic codes/descriptions)
Code: Description:
Code: Description:
Code: Description:
List the patient's underlying indication for transfusion. (Use ICD-10 Diagnostic codes/descriptions)
Code: Description:
Code:            Description:
Code: Description:
List the patient's comorbid conditions at the time of the transfusion related to the adverse UNKNOWN reaction. (Use ICD-10 Diagnostic codes/descriptions)
Code: Description:
Code: Description:
Code: Description:

Assurance of Confidentiality: The voluntarily provided information obtained in this surveillance system that would permit identification of any individual or institution is collected with a guarantee that it will be held in strict confidence, will be used only for the purposes stated, and will not otherwise be disclosed or released without the consent of the individual, or the institution in accordance with Sections 304, 306 and 308(d) of the Public Health Service Act (42 USC 242b, 242k, and 242m(d)). CDC 57.318 Rev. 3, v9.2

Public reporting burden of this collection of information is estimated to average 21 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering, and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC, Reports Clearance Officer, 1600 Clifton Rd., MS H21-8, Atlanta, GA 30333, ATTN: PRA (0920-0666).



List the patient's relevant medical procedure including past procedures and procedures to be performed during the current hospital or outpatient stay. (Use ICD-10 Procedure ONONE NONE	WN
Code: Description:	
Code: Description:	
Code: Description:	
Additional Information	
Transfusion History	
Has the patient received a previous transfusion?	
Blood Product: WB RBC Platelet Plasma Cryoprecipitate Granule	ocyte
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:	
HTR TTI PTP TACO TAD TA-GVHD TRALI UNKNO	WN
Reaction Details	
*Date reaction occurred: / / *Time reaction occurred: : Time unknown	
*Facility location where patient was transfused:	_
	_
*Facility location where patient was transfused:	_
*Facility location where patient was transfused:         Is this reaction associated with an incident?         Yes         No         If Yes, Incident #:	
*Facility location where patient was transfused: Is this reaction associated with an incident? Yes No If Yes, Incident #: Investigation Results	-
*Facility location where patient was transfused: Is this reaction associated with an incident? Yes No If Yes, Incident #: Investigation Results * Transfusion associated circulatory overload (TACO)	
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Hemolysis/Hemorrhage:	Disseminated intravascular coagulation Hemoglobinemia
Pain:	Abdominal pain Back pain Flank pain pain
Renal:	Hematuria Hemoglobinuria Oliguria
	Bilateral infiltrates on chest x-ray Bronchospasm Cough
Respiratory:	Hypoxemia Shortness of breath
Other: (specify)	
*Severity	
Did the patient receive or	experience any of the following?
No treatment requ	
	cuding prolonged hospitalization
Disability and/or in	
	portant conditions Death Unknown or not stated
<b>_</b>	
*Imputability	
	relationship between the transfusion and the reaction?
	ns for circulatory overload are possible.
	ly contributor to circulatory overload
	story of a pre-existing condition(s) that most likely explains circulatory overload.
	n favor of a cause other than the transfusion, but transfusion cannot be excluded.
	evidence beyond reasonable doubt of a cause other than the transfusion.
I he relationship bei	ween the adverse reaction and the transfusion is unknown or not stated.
Did the transfusion occur	at your facility? YES NO
Does the patient have a h	istory of cardiac insufficiency?
Yes, the patient ha	s a history of cardiac insufficiency that could explain the circulatory overload, but
	is likely to have caused the circulatory overload.
	s a history of pre-existing cardiac insufficiency that most likely explains circulatory
overload.	
No, the patient doe	s not have a history of cardiac insufficiency.
Did the patient received o	ther fluids in addition to the transfusion?
Module-generated Desig	nations
NOTE: Designations for case	definition, severity, and imputability will be automatically assigned in the NHSN
application based on response	es in the corresponding investigation results section above.
*Do you gargo with the	case definition designation?
	case definition designation?
T lease mulcale your des	
*Do you agree with the	severity designation?
	signation
	mputability designation?
^Please indicate your des	signation



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Patient He	atment					_		
Did the pati	ent receive treatment	for the transfusi	ion reaction?	YES	NO	U	INKNO\	ΝN
If yes, sele	ect treatment(s):							
Me	dication (Select the typ	be of medication	1)					
	Antipyretics Ar	ntihistamines	Inotropes/Vasor	oressors Bro	onchodil	ator	Di	uretics
	Intravenous							
l In	nmunoglobulin		ntravenous steroids		teroids	A	ntibiotic	s
	Antithymocyte globu		•	Other				
	ume resuscitation (Intr	ravenous colloid	s or crystalloids)					
Res	piratory support (Sele	ect the type of su	ipport)					
	Mechanical ventilati	ion Nonii	nvasive ventilation	Oxygen				
Rer	al replacement therap	by (Select the ty	pe of therapy)					
	Hemodialysis	Peritoneal	Continuous Ven	io-Venous Hemo	filtration	Ì		
Phle	ebotomy							
Oth	-							
Outcome								
*Outcome:	Death M	lajor or long-tern	n sequelae	] Minor or no sec	uelae	No	t detern	nined
Date of	Death: /	1						
^lf	recipient died, relation	ship of transfusi	ion to death:					
	Definite Probabl	·		Ruled Out		Not de	etermine	ed
Cause	of death:							
Was an autopsy performed? Yes No								
Component	Details							
*Was a partic reaction?	cular unit implicated	d in (i.e., respo	onsible for) the a	dverse	Yes		No	
			^Unit number					N/A
Transfusion Start and <b>End</b>	*Component code	Amount	(Required for	*Unit				
Start and Enu		trancfuend at			*Plood	arour		Implic
Date/Time	(check system used)	transfused at reaction onset	Infection and	expiration Date/Time	*Blood of unit		)	
Date/Time ^IMPLICATED	(check system used)			expiration			)	Implic ated
	(check system used)		Infection and	expiration			)	Implic ated
	(check system used) UNIT		Infection and	expiration Date/Time			) В-	Implic ated
	(check system used)	reaction onset	Infection and	expiration	of unit	_		Implic ated
	(check system used) UNIT	reaction onset	Infection and	expiration Date/Time	of unit	_		Implic ated Unit?
	(check system used) UNIT	Entire unit	Infection and	expiration Date/Time	of unit	A+ [	B	Implic ated Unit?
	(check system used) UNIT	Entire unit	Infection and	expiration Date/Time	of unit	A+	B-	Implic ated Unit?
	(check system used) UNIT ISBT-128 Codabar	Entire unit	Infection and	expiration Date/Time	of unit	A+	B-	Implic ated Unit?
	(check system used)         UNIT         ISBT-128         Codabar         ISBT-128         ISBT-128	reaction onset	Infection and	expiration Date/Time	of unit       A-       B+       O-       A-	A+	B- AB+ N/A B-	Implic ated Unit?
	(check system used)         UNIT         ISBT-128         Codabar         ISBT-128         ISBT-128	reaction onset         Entire unit         Partial unit        mL	Infection and	expiration Date/Time	of unit       A-       B+       O-       A-	A+ 	B-	Implic ated Unit?
	(check system used)         UNIT         ISBT-128         Codabar         ISBT-128         ISBT-128	reaction onset	Infection and	expiration Date/Time	of unit       A-       B+       O-       A-	A+	B- AB+ N/A B-	Implic ated Unit?



Label		Label	
	11		//
Comments			