



#### Hemovigilance Module Adverse Reaction Hypotensive Transfusion Reaction

*Required for saving		
*Facility ID#: NHS	N Adverse Reaction #:	
Patient Information		
*Patient ID:	*Gender: M F	Other *Date of Birth://
Social Security #:		Medicare #:
Last Name:	First Name:	Middle Name:
Ethnicity Hispanic or Latin	o Not Hispanic or Not Latino	0
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 (Us	e worksheet on page 4 for additiona	al codes and descriptions.)
(part 1) List the patient's adr	nitting diagnosis. (Use ICD-10 Diagn	ostic codes/descriptions)
Code:	Description:	
Code:		
Code:	Description:	
	erlying indication for transfusion. (Us	e ICD-10 Diagnostic codes/descriptions)
Code:	Description:	
Code:		
Code:	Description:	
	norbid conditions at the time of the tra O Diagnostic codes/descriptions)	ansfusion related to the UNKNOWN NONE
Code:	Description:	
Code:		
Code:	Description:	
		Continued >>
of any individual or institution is collections and will not otherwise be discleded by the sections 304, 306 and 308(d) of the Fublic reporting burden of this collection reviewing instructions, searching exist collection of information. An agency runless it displays a currently valid OM	ted with a guarantee that it will be held in so sed or released without the consent of the tublic Health Service Act (42 USC 242b, 24 on of information is estimated to average 2 ing data sources, gathering and maintaining not conduct or sponsor, and a person in B control number. Send comments regard to gestions for reducing this burden to CDC,	is surveillance system that would permit identification strict confidence, will be used only for the purposes individual, or the institution in accordance with 42k, and 242m(d)).  To minutes per response, including the time for any the data needed, and completing and reviewing the is not required to respond to a collection of information ling this burden estimate or any other aspect of this Reports Clearance Officer, 1600 Clifton Rd., MS D-74,



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?YESNOUNKNOWN
**If yes, provide information about the transfusion event. If not, skip to Reaction Details section.
Blood Product:
Date of Transfusion:// UNKNOWN
Did the patient experience a transfusion adverse reaction?
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
* Hypotensive transfusion reaction
*Case Definition
Check all that occurred during or within 1 hour of cessation of transfusion:
All other adverse reactions presenting with hypotension are excluded.
Hypotension
Check all that apply:
Hypotension occurs, does not meet the criteria above. Other, more specific reaction definitions do not
apply.
None of the above
Continued >>



Investigation Results (d	ontinued)							
Other signs and symptoms:	(check all that apply)							
Generalized:	Chills/rigors	Fever	Nausea/vomiting					
Cardiovascular:	Shock							
Cutaneous:	Edema	Flushing	Jaundice					
Odta 100doi	Other rash	Pruritus (itching)	Urticaria (hives)					
Hemolysis/Hemorrhage:	Disseminated intra	vascular coagulation	Hemoglobinemia					
, , , , , , , , , , , , , , , , , , ,		Positive antibody screen						
Pain:	Abdominal pain	Abdominal pain Back pain Flank pain Infusion site pain						
Renal:	Hematuria	Hemoglobinuria	Oliguria Oliguria					
Respiratory:	Bilateral infiltrates		nchospasm Cough					
	Hypoxemia	Hypoxemia Shortness of breath						
Other: (specify)								
<u>*Severity</u>								
		• ,	definitions listed in protocol)					
Symptomatic trea			prolonged hospitalization					
Life-threatening r			and/or incapacitation					
Congenital anom	aly or birth defect(s) of	the fetus	Death					
Other medically i	mportant conditions		Unknown or not stated					
<u>*Imputability</u>								
Which best describes the	e relationship between	the transfusion and the	reaction?					
The patient has no	other conditions that c	ould explain hypotensio	n.					
	tential causes present	that could explain hypot	ension, but transfusion is the most likely					
cause.								
		n hypotension are prese						
			but transfusion cannot be excluded.					
	•		e other than the transfusion.					
The relationship be	tween the adverse rea	action and the transfusio	n is unknown or not stated.					
How did the patient resp	ond the cessation of tra	ansfusion and supportive	e treatment?					
Responds rapidly (i.e., within 10 minutes) to cessation of transfusion and supportive treatment.								
The patient does not respond rapidly to cessation of transfusion and supportive treatment.								
Did the transfusion occu	at your facility?	YES NO						
When did the reaction od	cur in relation to the tra	ansfusion?						
Occurs less than 2	L5 minutes after the sta	art of the transfusion.						
Onset is between	Occurs less than 15 minutes after the start of the transfusion.  Onset is between 15 minutes after start and 1 hour after cessation of transfusion.							
			Continued >>					



Investigation Results (continued)
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
Additional Information
Patient Treatment
*Did the patient receive treatment for the transfusion reaction? YES NO UNKNOWN
If yes, select treatment(s):  Medication (Select the type of medication)
Medication (Scient 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
Phlebotomy
Other Specify:
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



Manual Galety Network			www.cuc.gov/iiis	311
Cause of death:				
Was an autopsy performed?	Yes	No		
			Continu	ıed >>



Component Details (Use worksheet on page 4 for additional units.)									
*Was a partic	ular unit implicated ir		ble for) the adver	se reaction?	Yes	;	No	N/A	
Transfusion <b>Start</b> and <b>End</b> Date/Time	*Component code (check system used)	Amount transfused at reaction onset	Unit number	*Unit expiration Date/Time	*Blood	l group	of unit	Implic ated Unit?	
^IMPLICATED UNIT									
	ISBT-128 Codabar	Entire unit Partial unitmL	 		A- B+	A+ AB- O+	B- AB+ N/A	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
(part 1) List the pa	ient's admitting diagnosis. (Use ICD-10 Diagnostic codes/descriptions)
Code:	Description:
Code:	
(part 2) List the pat	ent's underlying indication for transfusion. (Use ICD-10 Diagnostic codes/descriptions)
Code:	Description:
Code:	Description:
Code:	
Code:	
Code:	
Code:	
	ient's comorbid conditions at the time of the transfusion related to the
adverse reaction. (	Jse ICD-10 Diagnostic codes/descriptions)
adverse reaction. (	Jse ICD-10 Diagnostic codes/descriptions)
	Jse ICD-10 Diagnostic codes/descriptions) NONE  Description:
Code:	Jse ICD-10 Diagnostic codes/descriptions) NONE  Description:  Description:
Code: Code:	Jse ICD-10 Diagnostic codes/descriptions)  Description: Description: Description:
Code: Code: Code:	Description: Description: Description: Description: Description:
Code: Code:	Description: Description: Description: Description: Description: Description: Description:
Code: Code: Code: Code: Code: Code: (part 4) List the pa	Description:
Code: Code: Code: Code: Code: Code: (part 4) List the partocedures to be possible for the partocedures to be partocedures.	Description:   Description:
Code: Code: Code: Code: Code: Code: Code: Procedures to be portional procedure codes/displayed.	Description:
Code: Code: Code: Code: Code: (part 4) List the partocedures to be perfocedure codes/de Code:	Description:
Code: Code: Code: Code: Code: Code: (part 4) List the partocedures to be performed by the performance by the performa	Description:
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?
**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?
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?
**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?
**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?
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



### 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- AB+ O- O+ N/A	N	
	SBT-128	Entire			A+ B-	IN	

