

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

Hemovigilance Module Adverse Reaction Febrile Non-hemolytic Transfusion Reaction

*Required for saving	g							
*Facility ID#:	NHSN Adverse Reaction #:							
Patient Information	n							
*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 Latino	Unknown Declined to respond						
Race (Select all that apply):	American Indian or Asian Alaska Native	Black or African Middle Eastern or North American African						
	Native Hawaiian or White Pacific Islander	Unknown Declined to respond						
Preferred Language (Specify from the list provided): Declined to Respond n								
*Blood Group: A- A+ B- B+ AB- AB+ O- O+ Blood type not done Transitional ABO / Rh + Transitional ABO / Rh - Rh Group A/Transitional Group B/Transitional Rh Group A/Transitional Rh								
Rh	RII — ·							
Patient Medical Hi	-							
•	Imitting diagnosis. (Use ICD-10 Diagnostic o	• • •						
Code:		Description:						
Code:								
Code:	• •							
List the patient's un	derlying indication for transfusion. (Use ICE)-10 Diagnostic codes/descriptions)						
Code:	Description:							
Code:	Description:							
Code:	Description:							
	morbid conditions at the time of the transfus 10 Diagnostic codes/descriptions)	sion related to the adverse UNKNOWN NONE						
Code:	Description:							
Code:								
Code:								

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 57.311 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).





	edical procedure including past procedures and procedures to be nt hospital or outpatient stay. (Use ICD-10 Procedure NONE								
Code:									
Code:									
Code:									
Transfusion History Has the patient received a previous transfusion? YES NO UNKNOWN									
Blood Product:									
Date of Transfusion:									
Was the patient's adverse reaction transfusion-related? 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	,								
	/ / *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 reaction (FNHTR)								
	- Lanciación roaction (Francis								
*Case Definition	· · · · · · · · · · · · · · · · · · ·								
	ring or within 4 hours of cessation of transfusion:								
Ever (greater than transfusion value	or equal to 38°C/100.4°F oral and a change of at least 1°C/1.8°F) from pre-								
Chills/rigors are pre	sent								
Check all that apply:	Sch								
	d, but reported symptoms and/or available information are not sufficient.								
Other signs and symptoms: (c									
Generalized:	Nausea/vomiting								
Cardiovascular:	Blood pressure decrease Shock								
	Edema Flushing Jaundice								
Cutaneous:	Other rash Pruritus (itching) Urticaria (hives)								
l la marchine in // la marchine and	Disseminated intravascular coagulation Hemoglobinemia								
Hemolysis/Hemorrhage:	Positive antibody screen								
Pain:	Abdominal pain Back pain Flank pain Infusion site pain								
Renal:	Hematuria Hemoglobinuria Oliguria								
	Bilateral infiltrates on chest x-								
Respiratory:									
1 10 0 0 10 10 10 10 10 10 10 10 10 10 1	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 Life-threatening reaction							
Disability and/or incapacitation Congenital anomaly or birth defect(s) of the fetus							
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 signs/symptoms.							
There are other potential causes present that could explain signs/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 <u>case definition</u> designation? YES NO ^Please indicate your designation							
*Do you agree with the <u>severity</u> designation?							
^Please indicate your designation							
*Do you agree with the <i>imputability</i> 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							
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							



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=	ebotomy											
Other Specify:												
Outcome												
*Outcome: Death Major or long-term sequelae Minor or no 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												
*Was a particular unit implicated in (i.e., responsible for) the adverse reaction?												
Transfusion Start and End Date/Time	sion Amo		nt		*Unit expiration Date/Time	*Blood group of unit		Implic ated Unit?				
^IMPLICATED	UNIT											
	ISBT-128 Codabar	Entire unit Partial unitmL				B+ AB-	+ B- AB+ N/A	Y				
	ISBT-128 Codabar	Entire unit Partial unitmL				B+ AB-	+ B- AB+ D+ N/A	N				
Custom Field	ds											
Label				Label								
Comments												