

*Required for saving

Form Approved OMB No. 0920-0666 Exp. Date: 12/31/2026 www.cdc.gov/nhsn

Hemovigilance Module Incident

required for saving				
*Facility ID#:	NHSN Incident #:	Local Incident # or Log #:		
Discovery				
*Date of discovery: / / /				
*Time of discovery: :(H	H:MM)	e approximate 🔲 Time unknown		
*Where in the facility was the incide	nt discovered?			
*At what point in the process was the incident first discovered ? (check one)				
Product check-in Ord	ler entry Sample testing Product	Satellite storage		
Product storage Sar	nple collection manipulation	Product administration		
☐ Inventory management ☐ Sar	nple handling 🔲 Request for pic	k-up Post-transfusion review/audit		
Product/test request Sar	nple receipt Product issue	Other		
*How was the incident first disc	overed? (check one)			
Visual inventory review	Observation by sta	uff of unit/reagent/sample/equipment		
Routine audit or supervisory re	view Comparison of pro	duct label to patient information		
Computer system alarm or war	rning Comparison of pro	duct label to physician order		
Comparison of sample to paperwork When checking patient ID band				
Repeat or sample re-testing	☐ Notification or com	plaint from floor (nurse, MD, etc.)		
Historical record/previous type	check When product/unit	s returned to lab		
Communication from lab to floo	or Patient transfusion	reaction		
Human 'lucky catch'	Other (specify)			
Occurrence				
*Date initial incident occurred:	1 1			
*Time initial incident occurred:	:(HH:MM)	ne approximate 🔲 Time unknown		
Incident summary: (500 characters max)				

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 57.305, Rev. 7, v9.2

Public reporting burden of this collection of information is estimated to average 10 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).



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*Incident code(s): (r	max 20) Use NHSN incident	codes in the surveillance	protocol.
Incident Code	Occurrence Location	Incident Code	Occurrence Location
1		11	
2		12	
3		13	
4		14	
5		15	
6		16	
		17	
8		18	
9		19	
10		_ 20	
MS 99 Miscell	laneous, specify		
Job function of the	worker(s) involved in the i	ncident: (max 6) Use NHS	SN occupation codes in the protocol.
Other Othe	r (OTH), specify		Worker unknown
*Incident result: (ch	heck one)		
1 – Product tra	ansfused, reaction	3 – No product transfus	sed, unplanned recovery
2 – Product tra	ansfused, no reaction	4 – No product transfus	sed, planned recovery
*Product action: (c	heck all that apply)		



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☐ Not applicable					
Product retrieved and returned to inventory					
Product retrieved and destroyed					
^Single or multiple units destroyed?					
Single unit:					
Code system used: SBT-128 Codabar					
Unit #:					
OR Component code:					
Multiple units: (select code system used)					
☐ ISBT-128 ☐ Codabar Component code: Number of units:					
ISBT-128 Codabar Component code: Number of units:					
☐ ISBT-128 ☐ Codabar Component code: ☐ ☐ ☐ ☐ Number of units: ☐					
Product issued but not transfused					
Product transfused					
^Was a patient reaction associated with this incident? UYes UNO					
^Patient ID#(s):					
*Record/other action: (check all that apply)					
Record corrected Floor/clinic notified Attending physician notified					
Additional testing Patient sample re-collected Other (specify)					
Investigation Results					
*Did this incident receive root cause analysis? Yes No					
·					
Custom Fields					
Label					
	_				
	-				
	_				
	-				
Comments (2000 characters max)					

