

## Hemovigilance Module Incident

*Required for saving						
*Facility ID#:	NHSN Incident #:	Local Incident # or Log #:				
Discovery						
*Date of discovery: / / /						
*Time of discovery:::(H	H:MM)	Time approximate 🛛 🗌 Time unknown				
*Where in the facility was the incide	nt discovered?					
*At what point in the process was the incident <b>first discovered</b> ? (check one)						
Product check-in Order entry Sample testing Satellite storage						
□ Product storage □ Sample collection □ Product manipulation □ Product administration						
☐ Inventory management ☐ Sample handling ☐ Request for pick-up ☐ Post-transfusion review/audit						
Product/test request Sample receipt Product issue Other						
*How was the incident first disco	overed? (check one)					
☐ Visual inventory review	Observation b	y staff of unit/reagent/sample/equipment				
☐ Routine audit or supervisory re	view 🗌 Comparison o	of product label to patient information				
Computer system alarm or war	ning 🛛 🗌 Comparison d	of product label to physician order				
Comparison of sample to pape	rwork 🛛 🗌 When checkir	ng patient ID band				
Repeat or sample re-testing	Notification or	complaint from floor (nurse, MD, etc.)				
Historical record/previous type	check 🗌 When produc	t/units returned to lab				
Communication from lab to floo	r 🗌 Patient transf	usion reaction				
🗌 Human 'lucky catch'	Other (specify	/)				
Occurrence						
*Date initial incident occurred:						
*Time initial incident occurred:	:(HH:MM)	] Time approximate 🛛 Time unknown				
Incident summary: (500 characters	s 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)).						
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 D-74, Atlanta, GA 30333 ATTN: PRA (0920-0666).						



*Incident code(s): (m	nax 20) Use NHSN inci	ident codes in the surveilland	ce protocol.				
Incident Code	Occurrence Location	n Incident Code	Occurrence Location				
1		11					
2		12					
3		13					
4		14					
5		15					
6		16					
7		17					
8		18					
9		19					
		20					
MS 99 Miscella	aneous, specify						
Job function of the	worker(s) involved in	the incident: (max 6) <i>Use I</i>	NHSN occupation codes in the protocol.				
Other Other	(OTH), specify		Worker unknown				
*Incident result: (ch	eck one)						
🗌 1 – Product tra	nsfused, reaction	🗌 3 – No product trans	fused, unplanned recovery				
$\Box$ 2 – Product transfused, no reaction $\Box$ 4 – No product transfused, planned recovery							
*Product action: (ch	eck all that apply)						
Not applicable							
Product retriev	Product retrieved and returned to inventory						
□ Product retrieved and destroyed							
^Single or mult	iple units destroyed?						
🗌 Single un	it:						
Code system used: 🗌 ISBT-128 🔲 Codabar							
Unit #:							
	nponent 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?	es 🗌 No				
^Patient ID#(s)	):						



*Record/other action: (check all that apply)						
Record corrected	Record corrected Floor/clinic notified		Attending physician notified			
Additional testing	Patient sample re-coll	ected	Other (specify)			
Investigation Results	Investigation Results					
*Did this incident receive root cause analysis?						
Custom Fields						
Label		Label				
	/ /		1 1			
			·····			
	<u> </u>					
	<u> </u>		·····			
Comments (2000 characters max)						