

Incident #:	[s	ystem generated]
Local Incide	nt # or Log) #:
known		he facility was the incident ?
Comparis ork Compari ck Human ' eagent/sample/e utine audit or su	son of produ lucky catch' quipment pervisory rev	Patient transfusion reaction Visual inventory review
iest 🗌 Sample co Product stora bick-up 🗌 Product	bllection 🤤 🤄 ge 🗌 Availat t issue 🗌 Pro	Sample handling ole for issue Product selection Oduct administration
: (HH:MM ne approximate		*Where in the facility did the incident occur?
incident (Use Cl	DC Occupatio	on Type codes on page 5)
e be disclosed or released with n(d)). 10 minutes per response, inclu prmation. An agency may not ments regarding this burden es	ding the time for review conduct or sponsor, and timate or any other aspo	or institution is collected with a guarantee that it will be held in individual, or the institution in accordance with Sections 304, ing instructions, searching existing data sources, gathering and d a person is not required to respond to a collection of ect of this collection of information, including suggestions for
	Local Incide known Check one) Check one) Comparis ork Comparis ork Comparis ork Human ' eagent/sample/e utine audit or su dent first discove uest Sample composition Dick-up Product stora Dick-up Incident occomposition the incident occomposition Incident (Use CI when approximate Incident (Use CI vstem that would permit identified State of the incident occomposition. An agency may not ments regarding this burden estimation	discovered? known Check one) G Comparison of product ork Comparison of product ork Human 'lucky catch' eagent/sample/equipment description utine audit or supervisory reveated dent first discovered? (Check uest Sample collection Sa

xx-xx-20xx

*Where in the process did the incident first occur? (Check one)
Product check-in Product/test request Sample collection Sample handling
Sample receipt Sample testing Product storage Available for issue
Product selection Product manipulation Request for pick-up Product issue
Product administration Other (specify)
*Enter Incident Code (See Incident Codes on Page 4 of Form):
OR 🗌 Incident detail not specified
Incident summary:
*Incident result: (Check one)
\Box 1 = No recovery, harm \Box 2 = No recovery, no harm
\Box 3 = Near miss, unplanned recovery \Box 4 = Near miss, planned recovery
*Product action: (Check all that apply)
Product retrieved
Product destroyed
Code system used: (Check one) 🗌 ISBT-128 📄 Codabar
Indicate whether single or multiple units were destroyed:
*Single unit: a. Unit #:
OR b. Component Code:
*Multiple units: Component Code(s) # of Units
Code # of Units Code # of Units (Add add'l)
Product issued but not transfused
Product transfused
If the unit was transfused was a patient reaction associated with this incident?
YES NO
If YES, Patient ID#: Patient ID#:
*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? [Future]
YES NO Severity Code: (check one)
If YES, indicate result of analysis: (Check all that apply)
Technical Organizational Human Patient-related
Other (specify)
Custom Fields



Label		Label	
	//		//
Comments			



xx-xx-20xx

Hemovigilance Incident

INCIDENT CODES Based on MERS-TM and TESS	Sample Receipt (Transfusion Service)	Product Manipulation (Transfusion Service)
Product Check-In (Products Received	SR 01 Sample processed in error	UM 01 Data entry incomplete/incorrect
from Outside Source) PC 01 Data entry incomplete/not	SR 02 Historical review incorrect/not done	UM 02 Record review incomplete/incorrect
performed/incorrect	SR 03 Demographic review/data entry incorrect/	UM 03 Wrong component selected
PC 02 Shipment incomplete/incorrect	not done	UM 04 Administrative check (at time of manipulation
	SR 04 Sample incorrectly accessioned	UM 05 Labeling incorrect
PC 03 Product & paperwork do not match	(test/product) SR 05 Duplicate sample sent	UM 07 Special processing needs not checked
PC 04 Shipped under inappropriate conditions	Sample Testing (Transfusion Service)	UM 08 Special processing misunderstood or
PC 05 Inappropriate return to inventory	ST 01 Data entry incorrect/not performed	misinterpreted
PC 06 Product confirmation	ST 02 Appropriate sample checks not done	UM 09 Special processing not done/incorrectly done
PC 07 Administrative check (2 nd check)	ST 03 Computer warning overridden	Request for Pick-up (Clinical Service)
Product/Test Request (Clinical Service)	ST 05 Sample tube w/ incorrect accession label	RP 01 Request for pick-up on wrong patient
PR 01 Order for wrong patient	ST 07 Sample tubes mixed up	RP 02 Incorrect product requested for pick-up
PR 02 Order incorrectly entered on-line	ST 09 Test tubes mislabeled (wrong patient	RP 03 Product requested prior to obtaining consent
PR 03 Special needs not indicated on order	name/number)	RP 04 Product requested for pick-up pt not availabl
(e.g., CMV negative, auto) PR 04 Order not done/incomplete/incorrect	ST 10 Equipment problem	RP 05 Product requested for pick-up IV not ready
PR 05 Inappropriate/incorrect test ordered	ST 12 Patient testing not performed	RP 06 Request for pick-up incomplete
PR 06 Inappropriate/incorrect blood product	ST 13 Incorrect testing method chosen	RP 10 Product transport issues
ordered	ST 14 Testing performed incorrectly	Product Issue (Transfusion Service)
Sample Collection	ST 15 Test result misinterpreted	UI 01 Data entry incomplete/incorrect
	ST 16 Inappropriate/expired reagents used	UI 02 Record review incomplete/incorrect
SC 01 Sample labeled with incorrect patient	ST 17 ABO/Rh error caught on final check	UI 03 Pick-up slip did not match patient information
name	ST 18 Current & historical ABO/Rh don't match	UI 04 Incorrect unit selected (wrong person or right
SC 02 Not labeled	ST 19 Additional testing not performed	person wrong order)
SC 03 Wrong patient collected	ST 20 Administrative check at time work performed	UI 05 Issue delayed
SC 04 Collected in wrong tube type	ST 22 Sample storage incorrect/inappropriate	UI 06 LIS warning overridden
SC 05 Sample QNS	Product Storage (Transfusion Service)	UI 07 Computer issue not completed
SC 06 Sample hemolyzed	US 01 Incorrect storage of unit in transfusion	UI 09 Not checking/incorrect checking of unit and/o
SC 07 Label incomplete/illegible/incorrect (other than patient name)	service	patient information
SC 08 Sample collected in error	US 02 Expired product in stock	UI 11 Unit delivered to incorrect location
SC 09 Requisition arrives without samples	US 03 Inappropriate monitoring of storage device	UI 19 Wrong product issued
SC 10 Wristband incorrect/not available	US 04 Unit stored on incorrect ABO shelf	UI 20 Administrative review (self, 2 nd check at issue
SC 11 Sample contaminated	Available for Issue (Transfusion Service)	UI 22 Issue approval not obtained/documented
Sample Handling (Service Collecting	AV 01 Inventory audits	Product Administration (Clinical Service)
Samples)	AV 02 Product status not/incorrectly updated in	UT 01 Administered product to wrong patient
SH 01 Sample arrives without requisition	computer	UT 02 Administered wrong product to patient
SH 02 Requisition & sample label don't	AV 03 Supplier recall	UT 03 Product not administered
match	AV 04 Product ordered incorrectly/not submitted	UT 04 Incorrect storage of product on floor
SH 03 Patient ID incorrect/illegible on	Product Selection (Transfusion Service)	UT 05 Administrative review (unit/patient at bedside
requisition	SE 01 Incorrect product/component selected	UT 06 Administered product w/ incompatible IV fluid
SH 05 No phlebotomist/witness	SE 02 Data entry incomplete/incorrect	UT 07 Administration delayed
identification	SE 03 Not checking/incorrect checking of product	UT 08 Wrong unit chosen from satellite refrigerator
SH 06 Sample arrives with incorrect	and/or patient information	UT 10 Administered components in inappropriate o
requisition	SE 05 Historical file misinterpreted/not checked	UT 11 Appropriate monitoring of patient not done
SH 07 Patient information (other than ID)		UT 12 Floor/clinic did not check for existing product
missing/incorrect on requisition	SE 07 Special processing needs not checked	their area
CI 110 Comple transport issues	SE 09 Special processing needs not understood or	UT 13 Labeling problem on unit
SH10 Sample transport issues	misinterpreted	51

NHSN National Healthcare Safety Network OMB No. 0920-0666

	SE 11 Special processing n	SE 11 Special processing not done				
	NHSN Occupation 1	ype (Job Funct	tion) Codes			
Lab						
CLT	Clinical lab technician					
IVT	IVT Team Staff					
PHL	Phlebotomist/IV Team					
Nursing Staf	F					
CNA	Nurse Anesthetist					
LPN	Licensed Practical Nurse					
NMW	Nurse Midwife					
NUA	Nursing Assistant					
NUP	Nurse Practitioner					
RNU	Registered Nurse					
Physician						
FEL	Fellow					
MST	Medical Student					
PHY	Physician					
RES	Intern/Resident					
Technicians						
EMT	EMT/Paramedic					
HEM	Hemodialysis Technician					
ORS	OR/Surgery Technician					
PCT	Patient Care Technician					
Other Personn						
CLA	Clerical/administrative					
TRA	Transport/Messenger/Porter					
	cupation Types					
	Attendant/orderly	PHA	Pharmacist			
CSS	Central Supply	PHW	Public Health Worker			
CSW	Counselor/Social Worker	PLT	Physical Therapist			
DIT	Dietician	PSY	Psychiatric Technician			
DNA	Dental Assistant/Tech	RCH	Researcher			
DNH	Dental Hygienist	RDT	Radiologic Technologist			
DNO	Other Dental Worker	RTT	Respiratory Therapist/Tech			
FOS	Food Service	STU	Other Student			
HSK	Housekeeper	VOL	Volunteer			
ICP	Infection Control Professional					
LAU	Laundry Staff					
MNT	Maintenance/Engineering					
MOR	Morgue Technician	OTH	Other (Specify)			
OAS	Other Ancillary Staff					
OFR	Other First Responder					
OH	Occupational Health Professional					
OMS	Other Medical Staff					



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Other Technician/Therapist