

Exposure to Blood/Body Fluids

OMB No.	0920-066
Exp. Date:	02-29-200

Facility ID # :	Exposure Event #:
HCW ID:	
	First: Middle:
	Date of Birth: / /
Section I -General Exposure Information	
1. Did the exposure occur in this facility:	YN
1a. If No, specify name of facility	in which exposure occurred:
2. Date of exposure://	3. Time of exposure: AM PM
4. Number of hours on duty:	5. Is exposed person an temp/agency employee?YN
6. Location where exposure occurred:	
7. Type of exposure: (check all that apply)	
	re involve a clean, unused needle or sharp object?
•	mplete Q8, Q9, Section II, and Sections V-XI)
·	e Q8, Q9, Section III, and Sections V-XI)
7c. Skin: Was skin intact? Y	N Unknown (If No, complete Q8, Q9, Section III,
	and Sections V-XI)
7d. Bite (complete Q9 and Section	ns IV-XI)
8. Type of fluid/tissue involved in exposur Blood/blood products Solution (IV fluid, irrigation, etc.	Body Fluid: (check one)
9. Body site of exposure: (check one) Hand/Finger Eye Arm Leg Foot Mouth Nose Other (specify):	Pleural Feces/stool Semen Other Synovial (specify): Vaginal fluid
outer (opecity)	

Assurance of Confidentiality: The 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 15 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).

Section II - Percutaneous Injury					
. Was the needle or sharp object visibly contaminated with blood prior to exposure?Y N . Depth of the injury (check one): Superficial, surface scratch Moderate, penetrated skin Deep puncture or wound Unknown					
What needle or sharp object caused the injury? (check one) Deliow-bore needles:					
Suture needle Source needle Bone cutter Bovie electrocautery device Bur Elevator Explorer File Sharp object, type unknown Forceps Lancet Microtome blade Pin Razor Retractor Rod Scaler/curette Scalpel blade Scissors Tenaculum Trocar Wire					
Manufacturer and Model:					

5b. If the device had a safety feature, whe	n did the injury occur? (check one)
Before activation of the safety feat	
During activation of the safety feat	
Safety feature improperly activated	
Safety feature improperly activated Safety feature failed, after activated	
	JII
Safety feature not activated	
Other (specify):	
6. When did the injury occur: (check one)	
Before use of the item	
During use of the item	
After use of item, before disposal	
During or after disposal	
Unknown	
7. For what purpose or activity was the sharp de	evice heing used? (check one)
Obtaining a blood specimen percutaneously	<u> </u>
Performing phlebotomy	Performing a dental procedure
Performing arterial puncture	Hygiene (prophylaxis)
Performing a fingerstick/heelstick	Restoration (amalgam composite, crown)
Other blood-sampling procedure (specify)	Root canal
Giving a percutaneous injection	8- 7
Giving a percutation Giving an IM injection	Oral surgery
Giving a SC injection	Simple extraction
Placing a skin test (e.g., tuberculin, allergy, etc.)	Surgical extraction Handling device/equipment or specimen
Performing a line-related procedure	
Inserting or withdrawing a catheter	Handling equipment
Obtaining a blood sample from a central or	Transferring BBF into specimen container
peripheral I.V. line or port	Processing specimen Disassembling device/equipment
Injecting into a line or port	Disassembling device/equipment Decontaminating/processing used equipment
Connecting I.V. Line	Opening/breaking glass container (e.g., ampule)
Performing surgery/autopsy/other invasive procedure	Other
Suturing Specify procedure:	
Incising	Other (specify):
Palpating/exploring	Unknown
	Cimilowii
	the injury was different than that indicated in Q7,
what was the activity at the time of injury?	
Handling device/equipment or specimen	Housekeeping/patient-care activities, not described above
Recapping	Cleaning room
Transferring/passing/receiving device	Collecting/transporting waste
Disassembling device/equipment	Other (specify)
Disposing device	
Placing sharp in container	
9. Who was holding the device at the time the in	
Exposed person Co-worker/other person	No-one – the sharp was an uncontrolled sharp in the environment
10. What happened when the injury occurred: (check one)
Patient moved and jarred device	Overfilled/punctured sharps container
Device slipped	Improperly disposed sharp
Device rebounded	Other (specify):
Sharp was being recapped	Unknown
Collided with co-worker or other person	
. STITUCE TITULES TOTALE OF OUICE PERSON	

Section III - Mucous Membrane and/or Skin Exposure
1. Estimate the amount of blood/body fluid exposure: (check one) Small (< 1 tsp or 5cc) Moderate (> 1 tsp and up to 1/4 cup, or 6-50 cc) Large (> 1/4 cup or 50 cc) Unknown 2. Activity/event when exposure occurred: (check one) Airway manipulation (e.g., suctioning airway, inducing sputum) Bleeding vessel Changing dressing/wound care Cleaning/transporting contaminated equipment Endoscopic procedures IV or arterial line insertion/removal/manipulation Irrigation procedure Manipulating blood tube/bottle/specimen container Patient spit/coughed/vomited Phlebotomy Surgical procedure (e.g., all surgical procedures including C-section) Tube placement/removal/manipulation (e.g., chest, endotracheal, NG, rectal, urine catheter) Vaginal delivery Other (specify): Unknown 3. Barriers used by the worker at the time of exposure: (check all that apply) Face shield Other (specify): Gloves None of the above Goggles Gown Mask
Section IV - Bite
1. Wound description: (check one) No spontaneous bleeding Spontaneous bleeding Tissue avulsed Unknown 2. Activity/event when exposure occurred: (check one) During dental procedure During oral examination Providing oral hygiene Providing non-oral care to patient Assault by patient Other (specify): Unknown

Section V - Source Information Was the source patient known? Y N Was HIV status known at the time of exposure? Y N Check the test results for the source patient (P=positive, N=negative, I=Indeterminate, U=unknown, R=refuse NT= not tested): Hepatitis B								
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Hepatitis B P N I U R NT	Was HIV	status known at the time of ex	posure?	Y _	N			
Hepatitis B P N I U R NT HBsAg HBeAg Total anti-HBc anti-HBs Hepatitis C anti-HCV EIA anti-HCV supplemental PCR-HCV RNA HIV EIA, ELISA Rapid HIV Confirmatory test Section VI -For HIV Infected Source Stage of disease (check one): End-stage AIDS Other symptomatic HIV, not AII AIDS HIV infection, no symptoms actual HIV illness Unknown s the source patient taking anti-retroviral drugs? Y N U 2a. If Yes, indicate drug(s): Most recent CD4 count: mm³ Date:/	Check th	e test results for the source pati		_		minate, U=	unknown,	R=refused,
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Rapid HIV Confirmatory test Section VI -For HIV Infected Source Stage of disease (check one): End-stage AIDS Other symptomatic HIV, not AII AIDS HIV infection, no symptoms Acute HIV illness Unknown s the source patient taking anti-retroviral drugs? Y N U 2a. If Yes, indicate drug(s):	EL	A. ELISA						
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AIDS HIV infection, no symptoms Acute HIV illness Unknown s the source patient taking anti-retroviral drugs? Y N U 2a. If Yes, indicate drug(s):	Stage of	disease (check one):	End-stage AII	OS _	Othe	r sympton	natic HIV	, not AIDS
s the source patient taking anti-retroviral drugs? Y N U 2a. If Yes, indicate drug(s):	J	, , ,						
2a. If Yes, indicate drug(s):		-	Acute HIV	/ illness _	Unkr	nown		
Most recent CD4 count:mm³ Date:/	s the so	rce patient taking anti-retrovira	al drugs?	Y _	N		_U	
	2a.	If Yes, indicate drug(s):						
	Most re	cent CD4 count:mm³ I	Oate:/_			,		
mo / yr						mo / yı	r	
Viral load: copies/ml Date:/	Viral lo	ad: copies/ml I	Oate:/_					
mo / yr						mo / w		

Sec	ction VII - Initi	al Care Given to H	lealthcare W	ork	er		
1. F	IIV postexposui	re prophylaxis: Offe			U N	U	
2. H	IBIG given?	YN	U				
3. H	Iepatitis B vacc	ine given? Y _	N U	J			
4. Is	s the HCW preg	nant? Y	N U				
4	a. If Yes, which	ı trimester? 1	23	_ U			
		aseline Lab Resul					
		ng performed?			U (If Yes. i	ndicate below)	
	Serologic Test	Date	Result		Other Test	Date	Value
Н	HIV EIA	/ /	PNIR		ALT	/ /	IU/L
I V	Confirmatory	/ /	PNIR		Amylase	/ /	IU/L
H e p	anti-HCV EIA	/	PNIR	O t	Blood glucose	//	mg/dl
e patitis	anti-HCV supp	//	PNIR	h e	Hematocrit	/	%
s C	PCR HCV RNA	//	PNR	r	Hemoglobin	//	mg/dl
H e	HBs Ag	/	P N R NT	L a	Platelet	/	10 ⁴ /L
p a t	IgM anti-HBc	/	P N R	b s	# Blood cells in urine	//	#/mm³
t i	Total anti-HBc	//	PNR		WBC	/	10 ⁴ /L
s B	NT	, ,			Creatinine	/	mg/ml
Res	anti-HBs//						
P = Positive $N = Negative$ $I = Indeterminate$ $R = Refused$ $NT = Not Tested$							
Se	ction IX - Foll	low-up					
1. Is it recommended that the HCW return for follow-up of this exposure? Y N							
1.a. If Yes, will follow-up be performed at this facility? Y N							
	,	F			,	-	
Sec	ction X - Narra	ative					
In	the worker's w	yords how did the in	niury occur?				
In the worker's words, how did the injury occur?							

Section XI- Prevention					
In the worker's words, what could have prevented the injury?					
Custom Fields					
Label	<u></u>	Label	/		
Comments					
Comments					