

*required for saving Facility ID#: Exposure Event#						
*HCW ID#:						
HCW Name, Last: First:	Middle:					
*Gender:	_//					
*Work Location:						
*Occupation If occupation is physician, indica	te clinical specialty_					
Section I - General Exposure Information						
1. *Did exposure occur in this facility: 1. *Did exposure occur in this facility: 1a. If No, specify name of facility in which exposure occur	red:					
2. *Date of exposure:// 3. *Time of expos	ure: 🗆 AM 🗋 I	PM				
4. Number of hours on duty: 5. Is exposed pe	erson a temp/agency	/ employee? 🗌 Y 🔲 N				
6. *Location where exposure occurred:						
7. *Type of exposure: (Check all that apply)						
7a. Percutaneous: Did exposure involve a clean, unused	needle or sharp obj	ect?				
\Box Y \Box N (If No, complete Q8, Q9, Section II and Se	ction V–XI)					
□ 7b. Mucous membrane (Complete Q8, Q9, Section III a	and Section V-XI)					
\Box 7c. Skin: Was skin intact? \Box Y \Box N \Box Unknown (If No, complete Q8, Q9, Section III & Section V-XI)						
□ 7d. Bite (Complete Q9, and Section IV-XI)						
8. * Type of fluid/tissue involved in exposure: (Check one)						
Blood/blood products	\Box Body fluids: (Check one)					
□ Solutions (IV fluid, irrigation, etc.): (Check one)	\Box Visibly bloody					
□ Visibly bloody	🗆 Not v	visibly bloody				
□ Not visibly bloody	الجلوم من حور بنام نوما	asta ana hadu fuid tuna.				
$\Box \text{ Tissue}$	If body fluid, indicate one body fluid type:					
Other (specify) Unknown		□ Sputum				
9. *Body site of exposure: (Check all that apply)	□ Pericardial					
☐ Hand/finger □ Foot	Peritoneal	🗆 Urine				
Eye 🗆 Mouth	Pleural	Feces/stool				
□ Arm □ Nose	🗆 Semen	Other (specify)				
□ Leg □ Other (specify)	□ Synovial					
	🗆 Vaginal fluid					

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Public reporting burden of this collection of information is estimated to average 1 hour 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 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		
1. *Was the needle or sharp object visibl	y contaminated with blood prior to exp	oosure? 🗆 Y 🗆 N
2. Depth of the injury: (Check one)		
Superficial, surface scratch	Deep puncture of	or wound
☐ Moderate, penetrated skin		
3. What needle or sharp object cause	d the injury? (Check one)	
Device (select one)	vice sharp object (specify)	🗌 Unknown sharp object
Hollow-bore needle		
\Box Arterial blood collection device	🗌 Biopsy needle	\Box Bone marrow needle
\Box Hypodermic needle, attached to syringe	\Box Hypodermic needle, attached to IV tubing	\Box Unattached hypodermic needle
\Box IV catheter – central line	\Box IV catheter – peripheral line	\Box Huber needle
\Box Prefilled cartridge syringe	🗌 IV stylet	\Box Spinal or epidural needle
\Box Hemodialysis needle	\Box Dental aspirating syringe w/ needle	\Box Vacuum tube holder/needle
☐ Winged-steel (Butterfly [™] type) needle	Hollow-bore needle, type unknown	□ Other hollow-bore needle
Suture needle		
Suture needle		
Other solid sharps		
🗌 Bone cutter	🗌 Bur	Electrocautery device
Elevator	Explorer	\Box Extraction forceps
□ File	🗌 Lancet	🗌 Microtome blade
🗆 Pin	🗌 Razor	Retractor
\Box Rod (orthopedic)	□ Scaler/curette	\Box Scalpel blade
□ Scissors	🗌 Tenaculum	🗆 Trocar
□ Wire		
Glass		
🗌 Capillary tube	\square Blood collection tube	\Box Medication ampule/vial/bottle
Pipette	□ Slide	\square Specimen/test/vacuum tube
Plastic		
Capillary tube	\Box Blood collection tube	□ Specimen/test/vacuum tube
Non-sharp safety device		
\Box Blood culture adapter	\Box Catheter securement device	\Box IV delivery system
\Box Other known device (specify)		
4. Manufacturer and Model:		

N-HSN National Healthcare Safety Network



5. Did the needle or other sharp object involved in the in	
5a. If yes, indicate type of safety feature: (Check one)	-
Bluntable needle, sharp	Needle/sharp ejector
Hinged guard/shield	☐ Mylar wrapping/plastic
\Box Retractable needle/sharp	\Box Other safety feature (specify)
\Box Sliding/gliding guard/shield 5b. If the device had a safety feature, when did the in	\Box Unknown safety mechanism jury occur? (Check one)
\square Before activation of the safety feature was appropriate	\square Safety feature failed, after activation
\Box During activation of the safety feature	\Box Safety feature not activated
\square Safety feature improperly activated	Other (specify)
6. When did the injury occur? (check one)	
\Box Before use of the item	\Box During or after disposal
\Box During use of the item	Unknown
\Box After use of the item before disposal	
7. For what purpose or activity was the sharp device beir Obtaining a blood specimen percutaneously	ng used: (Check one)
Performing phlebotomy	\Box Performing a fingerstick/heelstick
Performing arterial puncture	\Box Other blood-sampling procedure (specify)
Giving a percutaneous injection	
\Box Giving an IM injection	\Box Placing a skin test (e.g., tuberculin, allergy, etc.)
\Box Giving a SC injection	
Performing a line related procedure	
\Box Inserting or withdrawing a catheter	\Box Injecting into a line or port
\square Obtaining a blood sample from a central or	\Box Connecting an I.V. line
peripheral I.V. line or port Performing surgery/autopsy/other invasive procedure	
□ Suturing	Palpating/exploring
	Specify procedure:
Performing a dental procedure	
Hygiene (prophylaxis)	□ Oral surgery
Restoration (amalgam composite, crown)	\Box Simple extraction
🗌 Root canal	\Box Surgical extraction
Periodontal surgery	
Handling a specimen	
Transferring BBF into a specimen container Other	Processing specimen
Other diagnostic procedure (e.g., thoracentesis)	
Other (specify)	



8. What was the activity at the time of injury? (Check one)		
Cleaning room	□ Collecting/transporting waste	
\Box Decontamination/processing used equipment	Disassembling device/equipment	
Handling equipment	\Box Opening/breaking glass container (e.g., ampule)	
Performing procedure	\Box Placing sharp in container	
	\Box Transferring/passing/receiving device	
Other (specify)		
9. Who was holding the device at the time the injury occurred?	' (Check one)	
\Box Exposed person	\Box No one, the sharp was an uncontrolled sharp in the environment	
Co-worker/other person	in the environment	
10. What happened when the injury occurred? (Check one)		
\square Patient moved and jarred device	□ Contact with overfilled/punctured sharps container	
\Box Device slipped	Improperly disposed sharp	
Device rebounded	□ Other (specify)	
\Box Sharp was being recapped	Unknown	
\Box Collided with co-worker or other person		



Section III - Mucous Membrane and/or Skin Exposure					
1. Estimate the amount of blood/body fluid exposure: (Check one)					
\Box Small (<1 tsp or 5cc)	\Box Large (>¼ cup or 50cc)				
\Box Moderate (>1 tsp and up to ¼ cup, or 6-50cc)					
2. Activity/event when exposure occurred: (Check one)					
\Box Airway manipulation (e.g., suctioning airway, inducing sputum)	Patient spit/coughed/vomited				
□ Bleeding vessel	Phlebotomy				
\Box Changing dressing/wound care	\Box Surgical procedure (e.g., all surgical procedures including C-section)				
\square Cleaning/transporting contaminated equipment	Tube placement/removal/manipulation (e.g., chest, endotracheal, NG, rectal, urine catheter)				
Endoscopic procedures	\Box Vaginal delivery				
\Box IV or arterial line insertion/removal/manipulation	Other (specify)				
□ Irrigation procedures	Unknown				
\Box Manipulating blood tube/bottle/specimen container					
3. Barriers used by the worker at the time of exposure: (C	heck all that apply)				
□ Face shield	□ Mask/respirator				
□ Gloves	Other (specify)				
□ Goggles	No Barriers				
Gown					
Section IV - Bite					
1. Wound description: (Check one)					
\Box No spontaneous bleeding	Tissue avulsed				
□ Spontaneous bleeding	Unknown				
2. Activity/event when exposure occurred: (Check one)					
During dental procedure	□ Assault by patient				
During oral examination	Other (specify)				
Providing oral hygiene	Unknown				
\Box Providing non-oral care to patient					



OMB No. 0920-0666 Exp. Date: xx-xx-xxxx

Note: Section V-IX are required when following the protocols for Exposure Management.

Section V - Source Information							
1. Was the source patient known? \Box Y \Box N							
2. Was HIV status known at the time of a	exposure?						
3. Check the test results for the source	•		e, I=indeterminat	e, U=unknown, P	erefused, NT=n،	ot tested)	
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	2						
1. Stage of disease: (Check one)							
End-stage AIDS Other symptomatic HIV, not AIDS							
			HIV infection, no symptoms				
Acute	HIV illness	🗌 Unk	nown				
2. Is the source patient taking anti-retro	viral drugs?	□y □n	U				
2a. If yes, indicate drug(s):							
3. Most recent CD4 count:mm	3	Date:	/ mo/yr				
4. Viral load: copies/ml undet	ectable	Date:	/ 				
Section VII - Initial Care Given to Healthcare Worker							
1. HIV postexposure prophylaxis:							
Offered? I Y I N I U Taken: Y N I U (If Yes, complete PEP form)							
2. HBIG given?							
3. Hepatitis B vaccine given: \Box Y \Box N \Box U Date 1 st dose administered://							
4. Is the HCW pregnant? $\Box Y \Box N \Box U$							
4a. If yes, which trimester? \Box 1 \Box 2 \Box 3 \Box U							



Section VIII - Baseline Lab Testing

Was baseline testing performed on the HCW? \Box Y \Box N \Box U If Yes, indicate results						
Test Date	Re	sult			Test Date	Result
HIV EIA//	Р	Ν	Ι	R	ALT//	IU/L
HIV Confirmatory//	Р	Ν	I	R	Amylase//	IU/L
Hepatitis C anti-HCV-EIA/	Р	Ν	I	R	Blood glucose//	mmol/L
Hepatitis C anti-HCV-supp//	Р	Ν	Ι	R	Hematocrit//	%
Hepatitis C PCR HCV RNA//	Р	Ν	R		Hemoglobin//	gm/L
Hepatitis B HBs Ag//	Р	Ν	R		Platelets//	x10º/L
Hepatitis B lgM anti-HBc//	Р	Ν	R		Blood cells in Urine_//	#/mm³
Hepatitis B Total anti-HBc//	Р	N	R		WBC//	x10º/L
Hepatitis B Anti-HBs//	mIU/mL		-	Creatinine//	µmol/L	
Result Codes: P=Positive, N=Negative, I=Indeterminate, R=Refused				Other://		

Section IX - Follow-up

1.	Is it recommended	that the HCW	return for	follow-up c	of this exposure?	ΠY	🗆 N
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1a. If yes, will follow-up be performed at this facility? $\ \ \Box$ Y $\ \ \Box$ N

Section X - Narrative

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	Label						
	//						
Comments							
Comments							