

Exposure to Blood/Body Fluids

OMB No. 0920-0666 Exp. Date: xx-xx-20xx

*Facility ID # :	*Exposu	re Event # :	
*HCW ID:			
HCW Name, Last:	First:	Middle:	
*Gender:		h://	
*Occupation: If occ			
		,	
Section I -General Exposure Inform	mation		
1			
1. *Did the exposure occur in this fac	ility: V I	NĪ	
_	-		
1a. If No, specify name of fac	-		
2. *Date of exposure://_		1	PM
4. Number of hours on duty:	5. Is exposed p	oerson an temp/agency e	employee?YN
6. *Location where exposure occurred	d:		
7. *Type of exposure: (check all that	apply)		
7a. Percutaneous: Did the ex		an, unused needle or sh	arn object?
		Section II, and Sections	
7b. Mucous membrane (Com	_	\ \	
	- / \	\ /	
7c. Skin: Was skin intact?	$-^{Y}$ $-^{N}$	7 _ /	
		and Sec	tions V-XI)
7d. Bite (complete Q9 and Se	ections IV-XI		
	`)		
8. *Type of fluid/tissue involved in ex	(posure: (check one)		
Blood/blood products		dy Fluid: (check one)	
Solution (IV fluid, prigation		Visibly bloody	
(check one)	, , _		visibly bloody
Visibly blood	dy		y y
Not visibly bloody	•	dy fluid, indicate one bo	ody fluid type:
Tissue		Amniotic	Saliva
Other (specify):		CSF	Survu
Unknown		Pericardial	Tears
		Peritoneal	Urine
0 *Dody site of avnosures (sheels one		Pleural	Feces/stool
9. *Body site of exposure: (check one Hand/Finger	•)	Semen	Other
Fland/Finger Eye		Synovial Vaginal fluid	(specify):
Eye Arm		v agiliai ilulu	
Aiii Leg			
Leg Foot			
Mouth Nose			
Other (specify):			

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 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 to CDC, Reports Clearance Officer, 1600 Clifton Rd., MS D-79, Atlanta, GA 30333, ATTN: PRA (0920-0666).

Section II - Percutaneous Injury
1. *Was the needle or sharp object visibly contaminated with blood prior to exposure?Y
N
2. Depth of the injury (check one):
Superficial, surface scratch
Moderate, penetrated skin
Deep puncture or wound
Unknown 3. What needle or sharp object caused the injury? (check one)
Hollow-bore needles:
Hypodermic needle attached to a syringe
Unattached hypodermic needle
Prefilled cartridge syringe needle
I.V. stylet Vacuum tube collection holder with needle (includes Vacutainer® type devices)
Spinal or epidural needle
Bone marrow needle
Biopsy needle
Other type of hollow-bore needle (specify):
— Hollow-bore needle, type unknown — Huber needle
—— Huber needle —— Winged-steel (Butterfly ™ type) needle
Hemodialysis needle
Solid sharp/Object: Other sharp object/device:
Suture reedle Capillary tube
Bone cutter Medication ampule/vial/I.V. bottle Bovie electrocautery device Pipette (glass)
Bur Slide
Elevator Specimen/test/vacuum tube
Bone chip/chipped tooth
Sharp object, type unknown
Other device (specify):
Lancet Microtome blade
Pin
Razor
Retractor
Rod
Scalar/curette
Scalpel blade Scissors
Tenaculum
Trocar
Wire
4. Manufacturer and Model:
5. Did the needle or other sharp object involved in the injury have a safety feature?Y N
5a. If Yes, indicate type of safety feature: (check one); If No, skip to Q6.
Sliding/gliding guard/shield Needle/sharp ejector
Stiding/gliding guard/shield Needle/sharp ejector Hinged guard/shield Mylar wrapping/plastic
Bluntable needle/sharp — Other safety feature (specify):
Retractable needle/sharp Unknown safety mechanism

5b. If the device had a safety feature, who Before activation of the safety feature During activation of the safety feature improperly activate Safety feature failed, after activate Safety feature not activated Other (specify):	ature was appropriate ature ed
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 Obtaining a blood specimen percutaneously Performing phlebotomy Performing arterial puncture Performing a fingerstick/heelstick Other blood-sampling procedure (specify) Giving a percutaneous injection Giving an IM injection Giving a SC injection Placing a skin test (e.g., tuberculin, allergy, etc.) Performing a line-related procedure Inserting or withdrawing a catheter Obtaining a blood sample from a central or peripheral I.V. line or port Injecting into a line or port Connecting I.V. Line Performing surgery/autopsy/other invasive procedure Suturing Specify procedure: Incising Palpating/exploring	Performing a dental procedure Hygiene (problylaxis) Restoration (amalgan composite, crown) Root canal Peridontal surgery Oral surgery Simple extraction Surgical extraction Handling specimen Transferring BBF into specimen container Processing specimen Other Other diagnostic procedure (e.g., thoracentesis) Other (specify): Unknown
8. What was the activity at the time of injury? Handling device/equipment or specimen Handling equipment Recapping Transferring/passing/receiving device Disassembling device/equipment Decontamination/processing used equipment Opening/breaking glass container (e.g., ampule) Performing procedure	Disposing device Placing sharp in container Housekeeping/patient-care activities, not described above Cleaning room Collecting/transporting waste Other (specify)
9. Who was holding the device at the time the ir Exposed person Co-worker/other person	ujury occurred? (check one) No-one – the sharp was an uncontrolled sharp in the environment
10. What happened when the injury occurred: (comparison of the injury occurred) (comparison	check one) — Overfilled/punctured sharps container — Improperly disposed sharp — Other (specify): Unknown

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, endotsacheal, NG, sectal, urine catheter) Vaginal delivery Other (specify): Unknown 3. Barriers used by the worker at the time of exposure: (check all that apply) Face shield Gloves 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

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

Section V - Source Information								
Vas the source patient known? Y	_ N							
Vas HIV status known at the time of exposur	e?	Y	N					
Check the test results for the source patient (Pa	-	_		ninate, U=	unknown, l	R=refused,		
Hopotitic D		= not tested)		T.T.		NITT		
Hepatitis B HBsAg	P	N	I	U	R	NT	\wedge	
HBeAg								
Total anti-HBc								
ant <u>i-HBs</u>								
Hepatitis C							V	
anti-HCV EIA								
ant <u>i-HCV supplemental</u>								
PCR-HCV RNA				\supset				
HIV								
EIA, EL <u>ISA</u>	$\overline{)}$							
Rapid HIV								
Confirmatory test								
Section VI-For HIV Infected Sou	ırce							
togo of disease table and	. AIF	NG.	O.I			AIDC		
tage of disease (check one): End-s		OS _			natic HIV, no sympt			
		illness _			no sympe	01110		
s the source patient taking anti-retroviral dru	gs?	Y _	N		_U			
2a. If Yes, indicate drug(s):								
Most recent CD4 count:mm³ Date: _	/_							
				mo / y	r			
Viral load: copies/mlUndetectabl	e	D	ate:	_/				
				1110 / y				

Section VII - Initial Care Given to Healthcare Worker				
1. HIV postexposure prophylaxis: Offered? Y N U Taken? Y N U				
2. HBIG given? Y N U				
3. Hepatitis B vaccine given? Y U				
4. Is the HCW pregnant? Y N U				
4a. If Yes, which trimester? 1 2 3 U				
Section VIII - Baseline Lab Testing				
Was baseline testing performed? Y N U				
Section IX - Follow-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				
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				