

Exposure to Blood/Body Fluids

OMB No.	0920-066
Exp. Date:	xx-xx-20x

*Facility ID # : *Exposu	ıre Event # :
*HCW ID:	
HCW Name, Last: First:	Middle:
*Gender: *Date of Bir	rth: / /
*Occupation: If occupation is a physici	an, indicate clinical specialty:
Section I -General Exposure Information	
1. *Did the exposure occur in this facility: Y 1a. If No, specify name of facility in which expose 2. *Date of exposure: / *3. Time of exposure of hours on duty: 5. Is exposed 4. Number of hours on duty: 5. Is exposed 6. *Location where exposure occurred: 7. *Type of exposure: (check all that apply) 7a. Percutaneous: Did the exposure involve a classification of the product of the	exposure: AN PM person an temp/agency employee? Y N ean, unused needle or sharp object? , Section II, and Sections V-XI) on III, and Sections V-XI)
Solution (IV fluid, hrigation, etc.): (check one) Visibly bloody	ody Fluid: (check one) Visibly bloody Not visibly bloody ody fluid, indicate one body fluid type: Amniotic Saliva CSF Sputum Pericardial Tears Peritoneal Urine
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

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®xtype devices)
Spinal or epidural needle
Bone marrow needle
Biopsy needle
Other type of hollow-bore needle (specify): Hollow-bore needle, type unknown
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
Forceps — Other device (specify):
Lancet
Microtome blade Pin
Razor
Retractor
Rod
Scaler/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
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, whe Before activation of the safety feature During activation of the safety feature improperly activate Safety feature improperly activate Safety feature failed, after activated Safety feature not activated Other (specify):	ture 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 Palpating/exploring	Performing a dental procedure Hygiene (prophylaxis) Restoration (amalgam composite, crown) Root canal Peridonal 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 in Exposed person Co-worker/other person	jury occurred? (check one) _ No-one – the sharp was an uncontrolled sharp in the environment
10. What happened when the injury occurred: (c Patient moved and jarred device Device slipped Device rebounded Sharp was being recapped Collided with co-worker or other person	heck one) Overfilled/punctured sharps container Improperly disposed sharp Other (specify): Unknown

Section III - Mucous Membrane and/or Skin Exposure
P
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, 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 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

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 (P	-	N=negative, = not tested)		ninate, U=	unknown, I	R=refused,	
Hepatitis B	P	N	I	U	R	NT	
HBsAg		14	1		10	111	
HBeAg							
Total anti-HBc							
an <u>ti-HBs</u>							
Hepatitis C							
anti-HCV EIA							
an <u>ti-HCV supplemental</u>							
PCR-HCV RNA			\rightarrow	\supset			
HIV							
EIA, EL <u>ISA</u>	$\frac{1}{2}$						
Rapid HIV							
Confirmatory test							
Section VI-For HIV Infected Sou	ırce						
tage of disease tabelly one.	A ID	NC.	O.L.			A IDC	
tage of disease (check one): End-s		os _			no sympt		
		illness _			cy p c		
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 / yı	r		
Viral load: copies/mlUndetectabl	e	D	ate:	_/	_		
				mo / yı	r		

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 N 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