

OF	R Needlestick & Sharp Object Injury Report SAFETY CENTER
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Ema	name: First name: First name: First name:
	y ID: (for office use only) S Facility ID: (for office use only) Completed by: FOR MICROSOFT®ACCESS
1.	Date of injury: ☐ ☐ ☐ 2. Time of injury: ☐ ☐ EXPOSURE PREVENTION ►
3.	Surgical service: Information Network ►
	□ 1 General □ 6 ENT □ 11 Transplants □ 2 Cardiovascular □ 7 Neurosurgery □ 12 Ophthalmology □ 3 OB/C-section □ 8 Plastic □ 13 Thoracic □ 4 Gynecology □ 9 Urology
	□ 5 Orthopedic □ 10 Oral/Dental □ 99 Other service, describe:
3a	Surgical procedure being performed:
3b.	Was it an endoscopic/laparoscopic/robotic/minimally invasive procedure?
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4. 	2 Surgeon (resident) specify specialty 10 Scrub nurse at time of incident → □ 1 RN □ 2 ORT □ 3 UAP 16 Surgeon (fellow) specify specialty 11 Other Nurse 3 Ob/Gyn (attending) 12 Nursing student 4 Ob/Gyn (resident) 13 OR assistant/attendant 5 Anesthesiologist (attending) 14 Housekeeper 6 Anesthesiologist (resident) 15 Physician assistant
4a.	If the injury was sustained by an anesthesia team member, what anesthesia task was being performed at the time of exposure? describe:
5.	Where did the injury occur? (check one box only) 1 Pre-operative area
6.	Was the source patient's identity known? (check one box only) 1 Yes □ 2 No □ 3 Unknown □ 4 Not applicable
7.	At the time of the injury, was the sharp instrument/item? (check one box only) 1 Held by another person
8.	The sharp item was: (check one box only) 1 Contaminated (known exposure to patient or contaminated equipment) 2 Uncontaminated (no known exposure to patient or contaminated equipment) 3 Unknown was there blood on the device? □ 1 Yes □ 2 No □ 3 Unknown
9.	For what purpose was the sharp item originally used? (check one box only) 1 Unknown/not applicable
	2 Injection, intra-muscular/subcutaneous, or other injection 13 Suturing muscle/fascia tech/
	through the skin (syringe) 14 Suturing skin To start IV or set up heparin lock (IV catheter or winged set-
	type needle) □ 15 Electrosurgery 4 To connect IV line (intermittent IV/piggyback/IV infusion/other □ 16 Drilling/sawing
	 IV line connection) Injection into (or aspiration from) IV injection site or IV port 17 Retracting tissue/bone 18 Wiring/fixing
	6 To place an arterial line/catheter □ 19 Using as a tool, not on patient
	7 To place a central line/catheter 20 To contain a specimen or pharmaceutical (glass items)
	8 To place other non-vascular line/catheter 9 To draw venous blood sample 99 Other; describe
	10 To draw arterial blood sample → if used to draw blood was it? □ 1 Direct stick? □ 2 Drawn from a line?

10.	Did the injury occur? (check one box only) 1 Before use of item (item broke/slipped, assembling device, etc.) 2 During use of item (item slipped, patient/colleague jarred item, etc) 3 While manually retracting tissue in operative site 4 While retracting tissue using retractor or other instrument 5 Passing instruments, hand-to-hand 6 Passing instruments, hand-free transfer 7 Between incremental injections 8 In between uses of devices 9 Disassembling device or equipment 10 Sorting, disinfecting, cleaning and/or sterilizing instruments 11 While recapping a used needle		13 14 15 16 17 18	Withdrawing needle from rubber or resistant material Other after use-before disposal (in transit to trash, cleaning, left on bed, table, floor, or other inappropriate place, etc.) From item left on or near disposal container While putting item into disposal container After disposal, stuck by item protruding from opening of disposal container Item pierced side of disposal container After disposal, item protruded from trash bag or inappropriate waste container Other, describe:
11.	What type of device caused the injury? (check one box only)		Sι	ollow-bore Needle urgical and solid needle lass
Whic	h device caused the injury? (check one box from one of the three sec	ction	ns c	only)
	□ 2. Tuberculin □ 6. 21-gauge needle		9 10 11 12 13 14 15 16 17 28 29 41 42 43 44 46 47 48 49 50 51 52 53 58	Spinal or epidural Needle Unattached hypodermic needle Arterial catheter introducer needle Central line catheter needle (cardiac, etc.) Drum catheter needle Other vascular catheter needle (cardiac, etc.) Other non-vascular catheter needle Huber-type needle Pen needle Needle, not sure what kind Other needle: describe: Trocar /trocar obdurator Vacuum tube (plastic) Specimen/Test tube (plastic) Fingernails/Teeth Retractors, skin/bone hooks Staples/Steel sutures Wire (suture/fixation/guide wire) Pin (fixation, guide pin) Drill bit/bur Pickups/Forceps/Hemostats/Clamps Surgical implant/explant Sharp item, not sure what kind Other sharp item: describe:
	40 Microtome blade		55	Other sharp term. describe.
Glass	60 Medication ampule 61 Medication vial (small volume with rubber stopper) 64 Vacuum tube (glass) 65 Specimen/Test tube (glass) 66 Capillary tube		68 78	7 Glass slide 8 Automobile glass/windshield 9 Glass item, not sure what kind 9 Other glass item: describe:
11a.	Brand/Manufacturer of product: (e.g. ABC Medical Company)			
11b.	Model/serial/lot number:			
12. 13.	Was this a re-usable device? 1 Yes If the item causing the injury was a needle or sharp medical device.			No
				No Site a safety design with a sileded, recessed, retractable, or No Unknown
13a.	Was the protective mechanism activated? 1 Yes, fully □ 2 Yes, partially		3	No 🗆 4 Unknown
13b.	Did the injury incident happen? 1 Before activation □ 2 During activation		3	After activation
13c.	Safety mechanism type: 1 Sliding sheath (hinged) 2 Sliding sheath (single barrel) 3 Retracting		4 5 6	Blunting/Blunted Hinged arm Other

14.	Did the device have needles on two ends (e.g. phlebotomy, pen needle)? 1 Yes 1 No 1 Unknown
14a.	If yes, which end caused the injury? Patient end Non-patient or 'back' end Both patient and 'back' ends Unknown or N/A
14b.	If yes and it was a safety engineered device, was the protective mechanism activated on both ends? Yes, both patient end and 'back' end No, only patient end No, only 'back' end Neither end had the protective mechanism activated Was not a device with needles at both ends
15.	Did the incident result in an exposure to a hazardous drug (e.g. chemotherapy, antineoplastic)? 1 Yes 2 No 3 Unknown
16. 	What was the location of the injury? (check one box only) Right hand Left hand Other, describe:
17. 	Was the injury? Superficial (little or no bleeding) Moderate (skin punctured, some bleeding) Severe (deep stick/cut, or profuse bleeding)
18.	If injury was to a hand, did the sharp item penetrate? Single pair of gloves Double pair of gloves No gloves
19.	Dominant hand of the injured worker: 1 Right-handed 2 Left-handed
20.	Employment status of injured worker: 1 Employee
21.	Describe the circumstances leading to this injury (please note if a device malfunction was involved):
office	s incident OSHA recordable on the sharps injury log? All injuries from contaminated needlesticks are required to be recorded on the OSHA log. (for use only)
1	□ 1 Yes □ 2 No □ 3 Unknown If yes: Days away from work: Days of restricted work activity:
Was	prophylaxis provided? (for office use only) 1 Yes 2 No 3 Unknown
	this incident meet the FDA medical device reporting criteria? (Yes if a device defect caused serious injury necessitating medical or surgical rention, or death occurred within 10 works days of incident.) (for office use only) □ 1 Yes (If yes, follow FDA reporting protocol.) □ 2 No □ 3 Unknown
Cost:	(optional, for office use only) Lab charges (HBV HCV, HIV, other) Healthcare worker Source Treatment/prophylaxis (HBIG, HBV vaccine, tetanus, other) Healthcare worker Source Service charges (Emergency Dept, Employee Health, other) Other costs (Worker's Comp, surgery, other) Paid Time Off
	TOTAL