

Texas Liver Institute

Employee Needlestick/Blood or Body Fluid Exposure Policy and Procedures

1. **GENERAL PROCEDURE STATEMENT:** This procedure is intended to guide the management of an employee injury from needle sticks or any contact of broken skin, subcutaneous tissue or mucous membranes with blood, body fluids or tissue. Following these procedures will minimize the risk of infection, ensure appropriate documentation, and minimize the likelihood of future exposure incidents.

2. AFFECTED DEPARTMENT(S): All staff

3. PROCEDURE:

A. The steps of the plan include:

- I. Immediate wound treatment and supervisor notification.
- II. Exposure evaluation and follow up
- III. Counseling and retraining
- B. **Immediate Wound Treatment and Notification.** When a staff member is exposed to a patient's blood, bodily fluids, tissue, or other potentially infectious material, or stuck with a contaminated sharp instrument used on a clinic patient:
 - I. As soon as it is safe to do so, the staff member will wash exposed skin thoroughly with soapy water and exposed mucous membranes (eye or mouth) will be flushed with copious amounts of water.
 - II. The staff member will immediately inform his/her supervisor or designated person.
 - III. The supervisor and affected staff member will fill out Needle Stick Injury/Body Fluid Exposure Report Form within 7 days of incident (Appendix A)

C. Exposure Action Plan

- I. The supervisor or staff member will notify the QA/Compliance Director of the exposure incident.
- II. Staff member will be offered testing for infectious diseases. Affected staff members are not required to undergo testing for infectious diseases but may forfeit a future workman's compensation claim if baseline testing was not obtained. Employee Consent Form attached as Appendix B.
- III. If the source (patient) is known, the supervisor will:
 - 1. Inform the patient of the staff members' injury
 - 2. Review the patient's health history and will ask the patient to consent to infectious disease testing
 - 3. Advise the patient that testing is voluntary and free of charge



- 4. Advise the patient that they may be informed of test results if they would like to know
- 5. Obtain informed consent and patient signature (Appendix C)
- 6. Send the patient to the appropriate lab so that a blood sample can be drawn
- 7. Action plan documented on the Needle Stick Injury/Body Fluid Exposure Report Form (**Appendix A**)
- IV. Supervisor and staff member will complete the Needlestick and Sharp Object Injury Checklist (Appendix D) or Blood and Body Fluid Exposure Checklist (Appendix E).

D. Counseling and Retraining

- I. Counseling and retraining of the staff member and/or any coworker whose actions or omissions may have contributed to the exposure will be performed.
- II. Retraining will be completed within 10 days of the incident.



ATTACHMENT A

Needle Stick Injury/ Body Fluid Exposure Report Form

I. Employee In	formation:				
Full Name			Female Date of B	irth/	/
Address: Street		City		State	_ Zip
Date Hired:/	/ /	Job Title:			
Status: DEmploye	e, 🛛 Contract Emplo	yee, D Volunteer,	■Student, Other: _		
Phone No: ()				
Hepatitis B vaccin	e: 1st//	2nd/	/3rd/	/	
Date of completion	n//				
Last Tetanus vacci	ine://///////_				
II. Exposure					
Work area of Expo	osure:				
	Tin		(am/pm)		
	Eye/mucous men			ect	
Other					
III. Patient (sour	ce) Information:				
	, 		Date of Birth		
)				
•	ce where injury/expo				
1 1	5 7 1				
	ow have or has she/h				
If yes, give Date:	Syphilis	Hepatitis A	Hepatitis B		
	HIV/AIDS	Hepati	tis C		
IV. Description of	f Incident (Complete			V Checklist, A	ttachment D or
Blood & Body Flu	id Exposure Checkli	st, Attachment E)			
V. Treatment at 7					
VI. Lab: HIV-1 PCR	&2(*run stat if source	e is positive) H H	BsAg 🛛 HBsAb 🕻	HBcAb	HCVAb HCV RNA
	to monitor for advers	a roation.			
	□CBC, Diff, Plts		N/Cr DALT/AS	T/AlkPhos/T	Bili



	Other:				
VII.	 Follow-up Service Chosen (circle one): a. Evaluation by TLI Medical Staff b. See own private physician. c. Go to Emergency Room. d. Obtain appropriate testing at County e. Decline Further Evaluation f. Name of health care professional: g. If treated away from worksite, where 	was it given	-		
	Facility:				
	Street: City:		Zin		
Emp	loyee Signature:				
Prov	ider Signature:			Date:	
Follo	ow-up dates: 1 Month: 2 Month: 3 Month: 6 Month:		-		

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ATTACHMENT B

HIV, HBV, and HCV Testing Employee Consent Form

As an employee/volunteer of Texas Liver Institute (TLI), I have been exposed to blood or other potentially infectious blood/body fluid. I agree to a blood draw for detection of antibodies to the Human Immunodeficiency Virus (HIV), Hepatitis B antigens and antibodies, and Hepatitis C performed by an outside laboratory. I understand that these tests may not be conclusive because a positive result means additional tests may be needed and a negative result does not necessarily eliminate consideration of AIDS. I have also been informed that the results of this blood test will only be released to those health care personnel and insurance companies providing medical care and coverage to me as allowed by federal and state law. I understand that these test results will be part of my medical record and will not be released unless I have signed an authorization for release of medical information.

I consent to the release of all medical records, information, results for evaluation to the following persons or organizations:

- a. TLI is required by law to keep the above information for the length of my employment plus thirty years.
- b. Supervisors and managers who may also need to be informed of any work or duty restrictions.
- c. Any hospital, clinic, physician, nurse, or other health care professional to whom the results of any medical treatment may be needed to provide care or treatment to me, including any physician or health care provider to whom I may be referred.

I voluntarily agree to these tests and understand that I will not be charged for any of the costs incurred by TLI for these lab tests.

I agree not to hold TLI authorized personnel or referral physicians and their authorized representatives responsible for any action that may be taken because of this release of information.

This consent may be revoked by me when received in writing to TLI authorized representative.

1. I hereby <u>give my consent</u> for the performance of the HIV, HEP B, and HEP C blood tests and to the release of results as outlined above.

(Print Name of TLI Employee/Volunteer)	
(Signature)	
(Date)	
(Print Name of Witness)	
(Witness Signature)	
(Date)	

2. I <u>decline</u> the opportunity for the HIV, HEP B, and HEP C blood tests at this time.

(Print Name of TLI Employee/Volunte	er)
(Signature)	
(Date)	
(Print Name of Witness)	
(Witness Signature)	
(Date)	



ATTACHMENT C

SOURCE CONSENT

Initials: ______1. I understand that an employee of this facility has been contaminated with my blood or other potentially infectious blood product or body fluid. I authorize a representative for this facility to draw blood for the following tests: Hepatitis B surface antigen, Hepatitis B surface antibody, Hepatitis C antibody, HIV.

Initials: _____ 2. I consent to the release of all medical records, information, results for evaluation to the following persons or organizations:

- a. This facility's officials that are required by law to keep the above information for the length of the exposed employee's employment plus thirty years.
- b. Supervisors and managers of this facility who may also need to be informed of any work or duty restrictions for the employee.
- c. Any hospital, clinic, physician, nurse, or other health care professional to whom the results of any medical treatment may be needed in order to provide care or treatment to me, including any physician or health care provider to whom the employee or myself may be referred

Initials:______ 3. I voluntarily agree to these tests and understand that I will not be charged for any of the costs incurred by this facility for these lab tests.

Initials:______ 4. I agree not to hold this facility's authorized personnel or referral physicians and their authorized representatives responsible for any action that may be taken as a result of this release of information.

Initials:_____ 5. This consent may be revoked by me when received in writing by the TLI authorized representative.

Patient Signature

Date

Witness Signature

Date



ATTACHMENT D

NEEDLESTICK AND SHARP OBJECT INJURY CHECKLIST

Employee Name:_____

	NT 11	
	Needle	Other Sharp
	□Insulin syringe with needle	
	Tuberculin syringe with needle	□Suture Needle □Scalpel
	Other syringe with needle	\Box Other surgical instrument (nonglass)
	□Needle connected to IV line	GLASS
	Uvinged steel needle (butterfly)	□Blood Tube
	□IV catheter, loose	□Other tube
	□Vacuum tube collection	□Slide
	Other:	□Ampule, □Other glass:
2.	Brand name (or □unknown):	
3.	Intended use of sharp? (\checkmark as appropriate)	
	□Injection, IM □Inc	ision
	□Obtain body fluid/tissue sample	
	□Injection, SC/ID	□Start IV/hep lock □Suturing, skin
	□Other Injection	Aspiration IV Contain specimen/pharmaceutical
	Draw venous sample	□Heparin/saline flush □Unknown/NA □Other
4.	Injury occurred: Before intended use During inte	
	5.5	
5.	If exposure occurred "During" or "After" intended use	, was it (some)
	Because patient moved during procedure	□While dissembling
	□While recapping	
	□While putting into sharp container	
	□Found in inappropriate place	
	Other:	
6.	Did device used have engineered sharps injury protect	on? 🛛 Yes 🖾 No 🖾 Don't Know
7.	Was the protective mechanism activated?	s \Box Yes, Partially \Box No \Box Don't Know \Box N/A
8.	When, during activation of the protective mechanism of	lid exposure occur?
	\square Before \square During \square After \square N/A	
9.	Was the injured person wearing gloves? \Box Yes	s 🗆 No 🖾 Don't Know
10.	Had the injured person completed a Hep B vaccination	n series?
11.	Was there a sharps container readily available for shar	ps disposal?□Yes □No □Don't Know



 Was the injury? (✓ one) □Superficial (little or no bleeding) □Moderate(skin punctured, some bleeding) □Severe (deep stick/cut or profuse bleeding) 				
13. What body part was involved? □Fin	nger D Hand	□Arm □Leg	/foot	
14. Where was the work area of the exp	osure incident?	□Patient Rm	□Procedure Rm □Lab □Other:	
15. What is the job classification of the	injured person?	DMD DPA/F	NP DMA DLab Tech DOther:	
16. What is the employment status of the injured person? Employee Volunteer Other:				
Employee Signature:		Date: _		
Employer Signature: D				



ATTACHMENT E

BLOOD AND BODY FLUID EXPOSURE CHECKLIST

1.	Where was the work area of the exposure incident?			
2.	What is the job classification of the exposed person? \Box MD \Box PA/FNP \Box MA \Box Lab Tech \Box Other:			
3.	Employment status of the injured person?			
4.	Was the identity of the fluid source (patient) known? (\checkmark one) \Box Yes \Box No \Box Unknown \Box N/A			
5.	Which body fluids were involved in the exposure? (\checkmark all that apply)			
6.	□Blood □Gastric contents □Sputum □Mucous □Saliva □Urine □Other: Was the body fluid visibly contaminated with blood? (✓ one) □Yes □No □Unknown What body part was involved? □Finger □Hand □Arm □Leg/foot □Face/head/neck □Torso			
7.	Was the body part? (\checkmark all that apply) \Box Intact skin, \Box Non-intact skin, \Box Eyes (conjunctiva),			
8.	□Nose (mucosa), □Mouth (mucosa), □Other: Did the blood or body fluid? (✓ all that apply) □Touch unprotected skin, □Touch skin in gap between barrier garments □Touch skin through tear in glove, □Soak through barrier garment, □Soak through clothing			
9.	Which barrier garments were worn at time of exposure? (\checkmark all that apply)			
	□Single pair latex/vinyl gloves □Double pair latex/vinyl gloves □Goggles □Eyeglasses (not protective equipment) □Eyeglasses with side shields □Face Shield □Surgical Mask □Surgical Gown Was the exposure the result of □Direct patient contact □Needle stick □Touched contaminated equipment/surface □Touched contaminated drape/sheet/gown, etc □Specimen container leaked/spilled/broke □Tubing leaked/disconnected/broke □Bag/pump leaked/spilled/broke □Equipment/operator failure □Other: If equipment failure: Type/Manufacturer:			
-				
Emp	ployee Signature: Date:			
Emp	ployer Signature: Date:			