BLOODBORNE PATHOGENS EXPOSURE CONTROL PLAN

	NAME OF FACILITY	DATE		
A.	certain other body fluids;	ol plan is to: loyee occupational exposure to blood on oodborne Pathogens standard 1910.1030		
В.	Exposure Determination OSHA requires employers to determine which employees may have work exposure to blood or other potentially infectious materials. The exposure determination is made without regard to the use of personal protective equipment (i.e. employees are considered to be exposed even if they wear personal protective equipment). This exposure determination is required to list all job classifications in which all employees may be expected to have such occupational exposure, regardless of frequency. At this facility the following job classifications are in this category: (list job classifications or job titles) (list job classifications or job titles)			
	employees may have occupational in these classifications would be e other potentially infectious materia these employees to have occupati listed in order to clearly understand are considered to have occupational	g of job classifications in which some exposure. Since not all the employees xpected to have exposure to blood or als, tasks or procedures that would cause onal exposure are also required to be al which employees in these categories all exposure. The job classifications and test are as follows (or put in appendix): TASK/PROCEDURE		
C.		ethodology an include a schedule and method of uirements of the standard. The following		

1. Compliance Methods

Universal precautions will be observed at (name of facility)
in order to prevent contact with blood or other potentially infectious materials. All blood or other potentially infectious material will be considered infectious regardless of the perceived status
of the source individual.
Engineering and work practice controls will be utilized to eliminate or minimize exposure to employees at this facility. Where occupational exposure remains after institution of these controls, personal protective equipment shall also be utilized. The following engineering controls will be utilized:(list controls, such as sharps containers, biosafety cabinets,etc) ENGINEERING PROTECTIVE
<u>DEPARTMENT</u> <u>CONTROL</u> <u>EQUIPMENT</u>
The above controls will be examined and maintained on a regular basis. The schedule for reviewing the effectiveness of the controls is as follows: DESCRIPTION REVIEW NEW TECHNOLOGY PERSON
OF CONTROL SCHEDULE CONSIDERED RESPONSIBLE
Readily accessible handwashing facilities must be made available to the employees who have exposure to blood or other potentially infectious materials. If handwashing facilities are not feasible, either an antiseptic cleanser <u>and</u> clean cloth/paper towels or antiseptic towelettes must be made available.
(name of person/position) will ensure that after
the removal of personal protective gloves, employees shall wash their hands and any other potentially contaminated skin surface as soon as feasible with soap and water.
(name of person/position) will ensure that if any
employees have exposure to their skin or mucous membranes then those areas must be washed or flushed with water as soon as feasible following contact.

2. Needles

Contaminated needles and other contaminated sharps will not be bent, recapped, removed, sheared or purposely broken. An exception is allowed if the procedure would require that the contaminated needle be recapped or removed and no alternative is feasible and the action is required by the medical procedure. If such action is required then the recapping or removal of the needle must be done by the use of a mechanical device or a one-handed technique. At this facility recapping or removal is only permitted for the following procedures:

PROCEDURE	MECHANICAL <u>DEVICE USED</u>	ONE HANDED TECHNIQUE USED

3. Containers for REUSABLE Sharps

Contaminated sharps that are reusable are to be placed immediately, or as soon as possible, after use into appropriate sharps containers. At this facility the sharps containers are puncture resistant, labeled with a biohazard label and are leak proof.

	PERSON WHO	FREQUENCY OF
LOCATION OF	REMOVES	CONTAINER
CONTAINERS	<u>SHARPS</u>	<u>CHECKS</u>
		

4. Work Area Restrictions

In work areas where there is a reasonable likelihood of exposure to blood or other potentially infectious materials, employees are not to eat, drink, apply cosmetics or lip balm, smoke, or handle contact lenses. Food and beverages are <u>not</u> to be kept in refrigerators, freezers, shelves, cabinets, or on counter tops where blood or other potentially infectious materials are present. Mouth pipetting/suctioning of blood or other potentially infectious materials is prohibited. All procedures will be conducted in a manner which will minimize splashing, spraying, splattering, and generation of droplets of blood or other potentially infectious materials. Methods which will be employed at this facility to accomplish this goal are:

WORK AREA/		METHOD
<u>DEPARTMENT</u>	<u>PROCEDURE</u>	<u>EMPLOYED</u>
5. Specimens		
Specimens of blood or	other potentially infecti	ious materials will be placed
in a container which	prevents leakage durin	ng the collection, handling,
		mens. The container used for las a biohazard. The OSHA
		he labeling/color coding of
		ecautions in the handling of
		ble as containing specimens. mens remain in the facility.
WORK AREA/		
DEPARTMENT	CODE USED	PRECAUTIONS USED
Any specimens which	could puncture a prin	nary container will be placed
within a secondary con	tainer which is punctur	re resistant. If the outside of
		he primary container must be
	torage, transport, or ship	prevents leakage during the oping of the specimen.
Contaminated Equipn	nent	
1.1		
that aggingment which h		on) is responsible for ensuring
		l with blood or other potentially ervicing or shipping and will be
	-	amination of the equipment is
not feasible.		
Personal Protective Eq	uipment	
PPE Provision		
that the fallactica access		ion) is responsible for ensuring
mat the following provi	sions are met. All PPE	used at this facility will be

6.

7.

provided without cost to employees. PPE will be chosen based on the expected exposure to blood or other potentially infectious materials. The PPE will be considered appropriate only if it does not permit blood or other potentially infectious materials to pass through or reach the employees' clothing, skin, eyes, mouth, or other mucous membranes under normal conditions of use and for the duration of time which the PPE will be used.

responsibility for distribution. You could also list which procedures		
would require the protective clothing and the recommended type of of protection required, this could also be listed as an appendix to		
PPE Use		
(name of person/position) will ensure that employees use appropriate PPE unless the supervisor shows that employee temporarily and		
briefly declined to use PPE when under rare and extraordinary circumstances, it was the employee's professional judgement that in the specific instance its use would have prevented the delivery of healthcare or posed an increased hazard to the safety of the worker or co-worker. When the employee makes this judgement, the circumstances will be investigated and documented in order to determine whether changes can be made to prevent such occurrences in the future.		
PPE Accessibility		
(name of person/position) will ensure that appropriate PPE in the appropriate sizes is readily accessible at the worksite or is issued without cost to employees. Hypoallergenic gloves, glove liners, powderless gloves, or other similar alternatives will be readily accessible to those employees who are allergic to the gloves normally provided.		
PPE Cleaning, Laundering and Disposal		
(name of person/position) will ensure that all PPE will be cleaned, laundered, repaired, replaced, and/or disposed of at no cost to the employees. All garments which are penetrated by blood must be removed immediately or as soon as feasible. All PPE will be removed before leaving the work area and placed in an appropriately designated area or container for storage, washing, decontamination or disposal.		

Gloves

Gloves will be worn where it is expected that employees will have hand contact with blood, other potentially infectious materials, non-intact skin, and mucous membranes; when performing vascular access procedures and when handling or touching contaminated items or surfaces. Hypoallergenic gloves, glove liners, powderless gloves, or other similar alternatives will be provided to employees who are allergic to the gloves normally provided.

Disposable gloves used at this facility are not to be washed or decontaminated for re-use and are to be replaced as soon as practical when they become contaminated or as soon as feasible if they are torn, punctured, or when their ability to function as a barrier is compromised. Utility gloves may be decontaminated for re-use provided that the integrity of the glove is not compromised. Utility gloves will be discarded if they are cracked, peeling, torn, punctured, or exibit other signs of deterioration or when their ability to function as a barrier is compromised.

Eye and Face Protection

Masks in combination with eye protection devices, such as goggles or glasses With solid side shields, or chin length face shields, will be worn whenever splashes, spray, splatter, or droplets of blood or other potentially infectious materials may be generated and eye, nose, or mouth contamination can reasonably be expected. Situations at this facility which would require such protection are:

SITUATION

WORK AREA/	REQUIRING	PROTECTION
<u>DEPARTMENT</u>	<u>PROTECTION</u>	<u>REQUIRED</u>
		
Additional Protection		
similar outer garments v	vill be worn in instance	gowns, aprons, clinic jackets, or es when gross contamination can ic surgery. These situations are:
DEPARTMENT	<u>PROCEDURE</u>	PPE REQUIRED

8. Housekeeping

9.

This facility will be schedule: WORK AREA/ DEPARTMENT	SCHEDULE	PERSON DEC	ONTAMINATION MATERIALS
All contaminated wo bleach (solutions of with water) or EPA and immediately or a infectious materials, become contaminate wrap may be used to	5.25% sodium hypo registered germic s soon as feasible af as well as at the end d since the last clean	ochlorite diluted between cides after completer any spill of blood of the work shift if the coverage. Protective coverage of the work shift if the coverage of the cover	veen 1:10 and 1:100 tion of procedures I or other potentially the surface may have erings such as plastic
DEPT RECEPTACE Any broken glasswar	to the following sche INSPECTION CLE SCHEDULE The which may be con	edule: I DECON FREQUENCY	PERSON RESPONSIBLE
Reusable sharps that materials will not be reach by hand into reusable sharps will	stored or processed the containers wh	l in a manner that renere these sharps	quires employees to have been placed. All
Regulated Waste Di	sposal		
Disposable Sharps			
Contaminated sharps containers that are cand labeled or color cand	losable, puncture rea		
The following items a	are considered reusal	ble sharps at this faci	llity:

During use, containers for contaminated sharps will be easily accessible to personnel and as close as feasible to the immediate area where sharps are used or can be reasonably expected to be found, such as laundry and exam rooms.

The containers will be maintained upright throughout use and replaced routinely and not be allowed to overfill.

WORK AREA/ DEPARTMENT	TYPE OF WASTE	TYPE CONTAINER	PERSON WHO MAINTAINS CONTAINERS

When moving containers of contaminated sharps from the area of use, the containers will be closed immediately prior to removal or replacement to prevent spillage or protrusion of contents during handling, storage, transport, or shipping.

The container will be placed in a secondary container if leakage of the primary container is possible. The second container will be closeable, constructed to contain all contents and prevent leakage during handling, storage and transport, or shipping. The second container will be labeled or color coded to identify its contents.

Reusable containers will not be opened, emptied, or cleaned manually or in any other manner which would expose employees to the risk of percutaneous injury.

Other Regulated Waste

All other regulated wastes will be placed in containers which are closeable, constructed to contain contents and prevent leakage of fluids during handling, storage, transportation or shipping.

The waste will be labeled or color coded and closed prior to removal to prevent spillage or protrusion of contents during handling, storage, transport or shipping.

NOTE: Disposal of all regulated wastes will be in accordance with applicable federal, state and local regulations. DHEC is the controlling agency in South Carolina.

10.Laundry Procedures (name of person/position) will ensure that laundry
contaminated with blood or other potentially infectious materials will be handled as little as possible and will be placed in biohazard -labeled or color coded red bags at the point of use and will not be sorted or rinsed there.
NOTE: If your facility uses <u>Body Substance Isolation</u> or <u>Universal Precautions</u> in the handling of <u>all</u> soiled laundry (i.e. all laundry is assumed to be contaminated) no labeling or color coding is necessary if all employees recognize the hazards associated with the handling of the laundry.
Laundry at this facility will be cleaned at (name of laundry).

NOTE: If your facility ships contaminated laundry off-site to a second facility which does not observe <u>Universal Precautions</u> in the handling of all laundry, contaminated laundry must be placed in bags or containers which are labeled or color coded. One possible solution would be to include a requirement in the contract laundry <u>scope of work</u> requiring the laundry to observe <u>Universal Precautions</u>.

11. Hepatitis B Vaccine and Post-Exposure Evaluation and Follow Up

General
(name of company) will make available the Hepatitis B vaccine and vaccination series to all employees who have occupational exposure, and post exposure follow-up to employees who have had an exposure incident.
evaluations and procedures including the Hepatitis B vaccine and the vaccination series and post exposure follow-up, including prophylaxis are

- a. made available at no cost to the employee;
- b. made available to the employee at a reasonable time and place;
- c. performed by or under the supervision of a licensed physician or by / under the supervision of another licensed healthcare professional;
- d. provided according to the recommendations of the U.S. Public Health Service.

All laboratory tests will be conducted by an accredited laboratory at no cost to the employee.

Hepatitis B Vaccination
(name of person/position) is in charge of the Hepatitis B vaccination program (name of physician or other healthcare professional) will provide the vaccinations.
Hepatitis B vaccinations will be made available after the employee has been
provided the training in occupational exposure (see information and training) and within 10 working days of initial assignment to all employees who have occupational exposure unless the employee has previously received the complete Hepatitis B vaccination series, antibody testing has revealed that the employee is immune, or the vaccine is contraindicated for medical reasons.
Participation in a pre-screening program will not be a prerequisite for receiving Hepatitis B vaccination.
If the employee initially declines Hepatitis B vaccination but at a later date while still covered under the Bloodborne Pathogens standard decides to accept the vaccination, the vaccination will then be made available.
All employees who decline the Hepatitis B vaccination must sign the OSHA required waiver indicating their refusal. (See page 16)
If a routine booster dose of Hepatitis B vaccine is recommended by the U.S. Public Health Service at a future date, such booster doses will be made available.
Post Exposure Evaluation and Follow Up
All exposure incidents must be reported, investigated, and documented. When an employee has an exposure incident, it must be reported to (name of person/position) immediately.
Following a report of an exposure incident, the exposed employee will receive a confidential medical evaluation and follow-up immediately performed by (name of licensed physician or other healthcare professional)
that includes at least:
a.) documentation of the route of exposure, and the circumstances under which the exposure incident occurred.b.) identification and documentation of the source individual, unless it can be established that identification is not feasible or prohibited by state or local law. (this provision may need to be modified to meet requirement of applicable state or local law and if so, modifications may be added

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c.) the source individual's blood will be tested as soon as feasible and after

consent is obtained in order to determine	HBV and HIV infectivity. If
consent is not obtained,	(name of person/position)
will establish that legally required conse	nt cannot be obtained. When
the source individual's consent is not	required by law, the source
individual's blood, if available, will be t	tested and results documented.

- d.) when the source individual is already known to be infected with HBV or HIV, testing for the source individual's known HBV or HIV status need not be repeated.
- e.) results of the source individual's testing will be made available to the exposed employee, and the employee will be informed of applicable laws and regulations concerning disclosure of the identity and infectious status of the source individual.

Collection and testing of blood for HBV and HIV serological status will comply with the following:

- a.) The exposed employee's blood will be collected as soon as feasible and tested after consent is obtained;
- b.) The employee will be offered the option of having their blood collected for testing of the employees' HBV/HIV serological status. The blood sample will be preserved for up to 90 days to allow the employee to decide if the blood should be tested for HIV serological status.

All employees who may have had an exposure incident will be offered post-
exposure evaluation and follow-up in accordance with the OSHA standard
All post exposure follow-up will be performed by
(name of clinic or health professional).
Information Provided To the Healthcare Professional

______ (name of person/position) will ensure that _____ (name of clinic or health professional) who is responsible for the employee's Hepatitis B vaccination is provided with :

- a.) a copy of the OSHA Bloodborne Pathogens Standard 29 CFR 1910.1030. (It might helpful to remind the health professional of confidentiality requirements of the standard);
- b.) a written description of the exposed employee's duties as they relate to the exposure incident;
- c.) written documentation of the route of exposure and circumstances under which exposure occurred;
- d.) results of the source individual's blood testing, if available; and
- e.) all medical records relevant to the appropriate treatment of the employee including vaccination status.

Healthcare Professional's Written Opinion
(name of person/position) will obtain and provide the employee with a copy of the healthcare professional's written opinion within 15 days of the completion of the evaluation. The written opinion for HBV vaccination and post exposure follow-up must be limited to:
 a.) whether HBV vaccination is indicated for an employee, and if the employee has received such vaccination; b.) a statement that the employee has been informed of the results of the evaluation of the post exposure follow-up; and c.) a statement that the employee has been told about any medical conditions resulting from exposure to blood or other potentially infectious materials which require further evaluation or treatment. NOTE: All other findings or diagnoses must remain confidential and must
not be included in the written report. 12. Labels and Signs
The universal biohazard symbol that is fluorescent orange or orange-red will be used.
Red bags or containers may be substituted for labels. However, regulated wastes must be handled according to DHEC rules and regulations.
Blood products that have been released for transfusion or other clinical use are exempted from these labeling requirements.
13. Information and Training
(name of person/position) will ensure that training is given at the time of initial assignment to tasks where occupational exposure may occur, and that it will be repeated within 12 months of the previous training.

a.) a copy of the OSHA standard and an explanation of its contents;b.) a discussion of the epidemiology and symptoms of bloodborne diseases;

and cover:

c.) an explanation of the modes of transmission of bloodborne pathogens;

training will be tailored to the education and language level of the employee, and given during during the normal work shift. The training will abe interactive

d.) an explanation of the _____ (name of company) Bloodborne Pathogens Exposure Control Plan, and a method for obtaining a copy;

- e.) the recognition of tasks that may involve exposure;
- f.) an explanation of the use and limitations of methods to reduce exposure, for example engineering controls, work practices, and personal protective equipment (PPE);
- g.) information on the types, use, location, removal, handling, decontamination, and disposal of PPE;
- h.) an explanation of the basis of selection of PPE;
- i.) information on the Hepatitis B vaccination, including efficacy, safety, method of administration, benefits, and that it will be given free of charge;
- j.) information on the appropriate actions to take and persons to contact in an emergency involving blood or other potentially infectious materials;
- k.) an explanation of the procedures to follow if an exposure incident occurs, including the method of reporting and medical follow-up;
- l.) information on the evaluation and follow-up required after an employee exposure incident; and
- m.) an explanation of the signs, labels and color coding system used .

The person conducting the training must be knowledgeable in the subject matter.

Employees who have received training in bloodborne pathogens in the 12 months before the effective date of this policy must only receive training in provisions that were not covered in the previous training.

Additional training will be provided to employees when there are changes of tasks or procedures affecting the employees' occupational exposure.

14. Recordkeeping

Medical Record	
(name of person/position) is	responsible for keeping medical
records . These records will be kept	(insert location)
(If you contract for post exposure follow-up and He	patitis B vaccination evaluation
make sure that your contract language includes pro	visions for recordkeeping which
are consistent with OSHA standard 1910,1020.)	

Medical records must be maintained according to OSHA standard 1910.1020. These records must be kept confidential, and must be maintained for at least the duration of employment plus 30 years. The records must include:

- a.) the name and social security number of the employee.
- b.) a copy of the employee's HBV vaccination status, including the dates of vaccination.
- c.) a copy of all results of examinations, medical testing, and follow-up procedures, and
- d.) a copy of the information provided to the healthcare professional, including

a description of the employee's duties as they relate to the exposure incident, and documentation of the routes of exposure and circumstances of the exposure.

Sharps Injury Log (Needle sticks Log)

Employers which fall under the scope of OSHA's Injury and Illness record keeping System (OSHA 300 Log) must maintain a sharps injury log in away that ensures employee privacy and contains at least the following information.

- a. Type and brand of device involved in the incident, if known;
- b. Location of the incident (e.g., department or work area); and
- c. A description of the incident

The log may contain additional information as long as an employee's privacy is protected. The format of the log can be determined by the employer.

Training 1	Records
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	(name of person/position)	is responsible for	r keeping	the
training records.	These records will be kept	(ir	sert location	on)

Training records must be kept for 3 years from the date of training and the following must be documented:

- a.) The dates of the training;
- b.) An outline describing the material presented;
- c.) The names and qualifications of persons conducting the training; and
- d.) The names and job titles of all persons attending the training sessions.

Availability

All employee records will be made available to the employee or his representative in accordance with OSHA standard 1910.1020.

All employee records will be made available to OSHA and the National Institute for Occupational Safety and Health under 1910.1020.

Transfer of Records

If this facility is closed or there is no successor employer to receive and retain the records for the prescribed period, the Director of NIOSH will be contacted for final disposition.

15. Evalu	ation and Re	view		
		s program, and	l its effectivene	n) is responsible for ss, and for updating on(date)
		SAFER MED DEVICE		
<u>DATE</u>	PROCEDU	RE CON	<u>SIDERED</u>	IMPLEMENTED
engineeri engineere solicited receiving	ing controls, inged sharps injut from non-magnification input from engine EMPLOYEES INVOLVED	including self- iry protections anagement hea inployees must PROCESS BY WHICH INPUT	sheathing need, and needlele lth care work be documented.	REASONS
16. Dates				
All prov	isions require (insert	•	standard will	be implemented by
17. Outsi	de Contracto	rs		
obtained establish	from and pr	ovided to outs erating proced	ide contractor.	address information s, you may wish to ituations and attach

HEPATITIS B VACCINE DECLINATION

(MANDATORY)

I understand that due to my occupational exposure to blood or other potentially infectious materials I may be at risk of acquiring Hepatitis B virus (HBV) infection. I have been given the opportunity to be vaccinated with Hepatitis B vaccine, at no charge to myself. However, I decline Hepatitis B vaccination at this time. I understand that by declining this vaccine, I continue to be at risk of acquiring Hepatitis B, a serious disease. If in the future I continue to have occupational exposure to blood or other potentially infectious materials and I want to be vaccinated with Hepatitis B vaccine, I can receive the vaccination series at no charge to me.

 Signature	
Date	