ABSL-1 FACILITY INSPECTION FORM

Office of Safety University of North Dakota 3851, Campus Rd Stop 9031 Grand Forks, ND 58202-9031 Ph. No. 701-777-3341

Fax: 701-777-413





DATE O	F SURVEY:	COI	NDUCTED BY:		BU	ILDING:				
		,								
ROOM	ROOM NUMBER: DEPARTMENT: PRINCIPAL INVESTIGATOR:									
E-MAIL ADDRESS:										
RESPONSIBLE PERSON (OTHER THEN PI):										
PHONE NUMBER: E-MAIL ADDRESS:										
	<u> </u>									
ITEM #			ITEM		YES	NO	CTI	N/A	COMMENTS CTI=CORRECTED AT TIME OF INSPECTION	
SECT	SECTION A: ANIMAL BIOSAFETY (These questions are based on the Animal Biosafety Level 1 section of Biosafety in Microbiological and Biomedical Laboratories, 5th Edition.									
1.0 SIG										
1.1	The animal laboratory door(s) date emergency contact inform		th the current Office of Safety issued sign	age and display up-to-						
2.0 DO	CUMENTATION AND TRAIN	NING				<u> </u>				
2.1	All personnel know how to acc	cess the Office	e of Safety website.							
2.2		cess UND's <u>B</u>	loodborne Pathogens Exposure Control	Plan on the Office of						
2.2	Safety website.	LIMB! O	d III II DI d OCC		_					
2.3			ccupational Health Plan on the Office of	•	<u> </u>					
2.4	Facility specific emergency pla		astitutional Biosafety Manual on the Offi	ice of Safety website.	<u> </u>		H			
2.6			fety Training Course within the past year	and documentation is			Ш			
2.0	available.	<u> </u>	viumi me past year	and documentation is						
2.7			completed the Animal care and Use Training is independent of Laboratory Safety							

ITEM #	ITEM	YES	NO	CTI	N/A	COMMENTS CTI=CORRECTED AT TIME OF INSPECTION
3.0 STA	NDARD MICROBIOLOGICAL PRACTICES					
3.1	Biological Agent(s) used in the animal room.					
3.2	The animal facility director establishes and enforces policies, procedures, and protocols for institutional policies and emergencies. a. Each institute must assure that worker safety and health concerns are addressed as part of the animal protocol review. b. Prior to beginning a study animal protocols must also be reviewed and approved by the Institutional Animal Care and Use Committee (IACUC) and the Institutional Biosafety Committee.					
3.3	Access to the animal room is limited. Only those persons required for program or support purposes are authorized to enter the facility. All persons including facility personnel, service workers, and visitors are advised of the potential hazards (natural or research pathogens, allergens, etc.) and are instructed on the appropriate safeguards.					
3.4	A safety manual specific to the animal facility is prepared or adopted in consultation with the animal facility director and appropriate safety professionals. The safety manual must be available and accessible. Personnel are advised of potential hazards and are required to read and follow instructions on practices and procedures.					
3.5	The supervisor must ensure that animal care, laboratory and support personnel receive appropriate training regarding their duties, animal husbandry procedures, potential hazards, manipulations of infectious agents, necessary precautions to prevent exposures, and hazard/exposure evaluation procedures (physical hazards, splashes, aerosolization, etc.). Personnel must receive annual updates and additional training when procedures or policies change. Records are maintained for all hazard evaluations, employee training sessions and staff attendance.					
3.6	An appropriate medical surveillance program is in place, as determined by risk assessment. The need for an animal allergy prevention program should be considered. Facility supervisors should ensure that medical staff is informed of potential occupational hazards within the animal facility, to include those associated with research, animal husbandry duties, animal care and manipulations. Personal health status may impact an individual's susceptibility to infection, ability to receive immunizations or prophylactic interventions. Therefore, all personnel and particularly women of childbearing age should be provided information regarding immune competence and conditions that may predispose them to infection. Individuals having these conditions should be encouraged to self-identify to the institution's healthcare provider for appropriate counseling and guidance. Personnel using respirators must be enrolled in an appropriately constituted respiratory protection program.					
3.7	A sign incorporating safety information must be posted at the entrance to the areas where infectious materials and/or animals are housed or are manipulated. The sign must include the animal biosafety level, general occupational health requirements, personal protective equipment requirements, the supervisor's name (or other responsible personnel), telephone number, and required procedures for entering and exiting the animal areas. Identification of specific infectious agents is recommended when more than one agent is being used within an animal room.					
3.8	Access to the animal room is limited. Only those persons required for program or support purposes are authorized to enter the facility. All persons including facility personnel, service workers, and visitors are advised of the potential hazards (natural or research pathogens, allergens, etc.) and are instructed on the appropriate safeguards.					

ITEM #	ITEM	YES	NO	CTI	N/A	COMMENTS CTI=CORRECTED AT TIME OF INSPECTION
3.9	Protective laboratory coats, gowns, or uniforms are recommended to prevent contamination of personal clothing. Gloves are worn to prevent skin contact with contaminated, infectious and hazardous materials, and when handling animals. Gloves and personal protective equipment should be removed in a manner that minimizes transfer of infectious materials outside of the areas where infectious materials and/or animals are housed or are manipulated. Persons must wash their hands after removing gloves, and before leaving the areas where infectious materials and/or animals are housed or are manipulated. Eye and face and respiratory protection should be used in rooms containing infected animals, as dictated by the risk assessment.					
3.10	Eating, drinking, smoking, handling contact lenses, applying cosmetics, and storing food for human consumption must not be permitted in laboratory areas. Food must be stored outside of the laboratory in cabinets or refrigerators designed and used for this purpose.					
3.11	All procedures are carefully performed to minimize the creation of aerosols or splatters of infectious materials and waste.					
3.12	Mouth pipetting is prohibited. Mechanical pipetting devices must be used.		П	П		
3.13	Equipment and work surfaces are routinely decontaminated with an appropriate disinfectant after work with an infectious agent, and after any spills, splashes, or other overt contamination.					
3.14	Animals and plants not associated with the work being performed must not be permitted in the areas where infectious materials and/ or animals are housed or are manipulated.					
3.15	An effective integrated pest management program is required.					
	All wastes from the animal room (including animal tissues, carcasses, and bedding) are transported from the animal room in leak-proof, covered containers for appropriate disposal in compliance with applicable institutional, local and state requirements. Decontaminate all potentially infectious materials before disposal using an effective method.					
3.16	Policies for the safe handling of sharps, such as needles, scalpels, pipettes, and broken glassware must be developed and implemented. When applicable, laboratory supervisors should adopt improved engineering and work practice controls that reduce the risk of sharps injuries. Precautions, including those listed below, must always be taken with sharp items. These include: a. Use of needles and syringes or other sharp instruments in the animal facility is limited to situations where there is no alternative for such procedures as parenteral injection, blood collection, or aspiration of fluids from laboratory animals and diaphragm bottles. b. Disposable needles must not be bent, sheared, broken, recapped, removed from disposable syringes, or otherwise manipulated by hand before disposal. Used disposable needles must be carefully placed in puncture-resistant containers used for sharps disposal. Sharps containers should be located as close to the work site as possible. c. Non-disposable sharps must be placed in a hard-walled container for transport to a processing area for decontamination, preferably by autoclaving. d. Broken glassware must not be handled directly. Instead, it must be removed using a brush and dustpan, tongs, or forceps. Plastic ware should be substituted for glassware whenever possible.					

ITEM #	ITEM	YES	NO	CTI	N/A	COMMENTS CTI=CORRECTED AT TIME OF INSPECTION
4.0 SAF	ETY EQUIPMENT (PRIMARY BARRIERS AND PERSONAL PROTECTIVE EQUIPMENT)					
4.1	A risk assessment should determine the appropriate type of personal protective equipment to be utilized.					
4.2	Special containment devices or equipment may not be required as determined by appropriate risk assessment.					
4.3	Protective laboratory coats, gowns, or uniforms may be required to prevent contamination of personal					
	clothing. Protective outer clothing is not worn outside areas where infectious materials and/or animals are housed or manipulated. Gowns and uniforms are not worn outside the facility.					
4.4	Protective eyewear is worn when conducting procedures that have the potential to create splashes of microorganisms or other hazardous materials. Persons who wear contact lenses should also wear eye protection when entering areas with potentially high concentrations or airborne particulates. Persons having contact with NHPs must assess risk of mucous membrane exposure and wear protective equipment (e.g., masks, goggles, face shields, etc.) as appropriate for the task to be performed.					
4.5	Gloves are worn to protect hands from exposure to hazardous materials. A risk assessment should be performed to identify the appropriate glove for the task and alternatives to latex gloves should be available. Change gloves when contaminated, glove integrity is compromised, or when otherwise necessary. Gloves must not be worn outside the animal rooms. Gloves and personal protective equipment should be removed in a manner that prevents transfer of infectious materials. Do not wash or reuse disposable gloves. Dispose of used gloves with other contaminated waste.					
4.6	Persons must wash their hands after handling animals and before leaving the areas where infectious materials					
	and/or animals are housed or are manipulated. Hand washing should occur after the removal of gloves.		$ \; \sqcup \;$			
	ORATORY FACILITIES (SECONDARY BARRIERS)					
5.1	The animal facility is separated from areas that are open to unrestricted personnel traffic within the building. External facility doors are self-closing and self-locking. Access to the animal facility is restricted. Doors to areas where infectious materials and/or animals are housed, open inward, are self-closing, are kept closed when experimental animals are present, and should never be propped open. Doors to cubicles inside an animal room may open outward or slide horizontally or vertically.					
5.2	The animal facility must have a sink for hand washing. Sink traps are filled with water, and/or appropriate liquid to prevent the migration of vermin and gases.					
5.3	The animal facility is designed, constructed, and maintained to facilitate cleaning and housekeeping. The interior surfaces (walls, floors and ceilings) are water resistant. Floors must be slip resistant, impervious to liquids, and resistant to chemicals. It is recommended that penetrations in floors, walls and ceiling surfaces be sealed, including openings around ducts, doors and doorframes, to facilitate pest control and proper cleaning.					
5.4	Cabinets and bench tops must be impervious to water and resistant to heat, organic solvents, acids, alkalis, and other chemicals. Spaces between benches, cabinets, and equipment should be accessible for cleaning. Chairs used in animal area must be covered with a non-porous material that can be easily cleaned and decontaminated. Furniture must be capable of supporting anticipated loads and uses. Sharp edges and corners should be avoided.					

ITEM #	ITEM	YES	NO	CTI	N/A	COMMENTS CTI=CORRECTED AT TIME OF INSPECTION
5.5	Ventilation should be provided in accordance with the Guide for Care and Use of Laboratory Animals. No					
	recirculation of exhaust air may occur.	_			l	
	It is recommended that animal rooms have inward directional airflow.					
	Ventilation system design should consider the heat and high moisture load produced during the cleaning of					
	animal rooms and the cage wash process.					
5.6	Internal facility appurtenances, such as light fixtures, air ducts, and utility pipes, are arranged to minimize					
	horizontal surface areas to facilitate cleaning and minimize the accumulation of debris or fomites.					
5.7	If floor drains are provided, the traps are filled with water, and/or appropriate disinfectant to prevent the					
7.0	migration of vermin and gases.	_	_			
5.8	Cages are washed manually or preferably in a mechanical cage washer. The mechanical cage washer should				l	
	have a final rinse temperature of at least 180°F. If manual cage washing is utilized, ensure that appropriate			ΙШ		
7.0	disinfectants are selected.					
5.9	Illumination is adequate for all activities, avoiding reflections and glare that could impede vision.			H	$\vdash \vdash \vdash$	
5.10	Emergency eyewash and shower are readily available; location is determined by risk assessment.			Ш		
	MMABLE LIQUIDS STORAGE					
6.1	Flammables stored are in an approved flammable liquids cabinet. (Contact Office of Safety with questions.)			H	$\vdash \vdash \vdash$	
6.2	Volatile liquids are stored in an explosion-proof refrigerator when required.	<u> </u>		Щ	⊢片	
6.3	Aerosol cans are kept away from heat and ignition sources.	<u> </u>				
	ME HOODS					
7.1	Inspected within last year.	<u> </u>			<u> </u>	
7.2	Undamaged.	<u> </u>		Щ.		
7.3	Used Correctly.	oxdot		$\sqcup \sqcup$	oxdot	
	LOGICAL SAFETY CABINETS					
8.1	All active BSCs have been certified within the last 12 months by a vendor approved by UND.					
8.2	The certification label is attached and initialed by a vendor approved by UND.		Щ			
8.3	Intake and rear grilles are clear of obstructions.		Ш			
8.4	Bunsen burners and/or open flames are not used in biological safety cabinets. (Open flames are not permitted					
	inside BSCs; consider an alternative, such as an electrical Bacti-Cinerator).					
8.5	Work surfaces are clean and free of visible biological residue.		Щ	Щ		
8.6	The sash alarm is not muted.					
	CTRICAL					
9.1	Extension cord use is temporary.		Ш			
9.2	Proper grounding is used.					
9.3	Cord and equipment in good condition.					
9.4	No outlet overloading.					
9.5	Outlets near water GFCI protected.					
9.6	Electrical Panels Accessible.					
9.7	Shock hazards have proper signage.					
1			l L	\sqcup		

ITEM #	ITEM	YES	NO	CTI	N/A	COMMENTS CTI=CORRECTED AT TIME OF INSPECTION
	ERGENCY EQUIPMENT				1	
10.1	FIRE EXTINGUISHER					
	Correct type Fire Extinguisher present.					
	Fire Extinguisher easily accessible.					
	Fire Extinguisher tagged within the last year by Office of Safety.					
10.2	SAFETY SHOWERS					
	Safety showers are unobstructed.					
	Safety showers are tested monthly.					
	Safety showers are functional and installed properly.					
10.3	EYEWASHES					
	Eyewashes are unobstructed.					
	Eyewashes are tested monthly.					
	Eyewashes are functional and installed properly.					
10.4	SPILL KITS AND FIRST AID					
	Spill kits and first aid are stocked appropriately.					
	Spill kits and first aid are readily accessible.					
	Disinfectant available.					
	Broom, dustpan, forceps available.					
	Calcium gluconate available for HF.					
11.0 CH	EMICAL WASTE					
11.1	Office of Safety picks up all chemical waste from the facility.					
11.2	Chemicals are not put down the drain, in the regular trash, or in biomedical waste.					
11.3	All chemical / chemical waste containers are closed except when in use.					
11.4	Chemical wastes are compatible with their containers and are stored by compatibility (i.e. acid waste is not					
	stored with alkaline waste).		Ш		Ш	
11.5	Office of Safety picks up all empty P-listed chemical containers from the facility.					
11.6	Office of Safety picks up expired pharmaceutical wastes (excluding DEA controlled substances) from the	П				
	facility.					
	DLOGICAL WASTE					
12.1	Biomedical waste containers are labeled with the Biohazard symbol and the word "Biohazard".	<u> </u>	\Box		\sqcup	
12.2	An orange / red Biohazard bag is used to dispose of biohazardous waste.					
12.3	Biohazard waste containers are closed except when adding waste.	$\perp \perp$				
12.4	Biohazards are not put down the drain or in regular trash.	$\perp \perp$		$\perp \sqcup$	$\sqcup \sqcup$	
12.5	Biohazard waste is not mixed with chemical waste.			$\sqcup \sqcup$		
12.6	Facility-specific SOPs for the treatment and removal of biohazard waste from the facility are available and adhered to.					
12 A CTT	ARPS HANDLING AND WASTE					
13.0 SH 13.1	Sharps are disposed of in a sharps disposal container and the containers are no greater than ³ / ₄ full.					
13.1		+ $+$	$\vdash \vdash \vdash$	╁╫	┝╫╴	
	Sharps containers are tightly lidded to prevent the contents from spilling.	+	$\vdash \vdash \vdash$	├ ├	⊢	
13.3	Office of Safety picks up sharps waste for disposal.	\sqcup		\sqcup		

ITEM #	ITEM	YES	NO	CTI	N/A	COMMENTS CTI=CORRECTED AT TIME OF INSPECTION		
14.0 AU	14.0 AUTOCLAVE USE							
14.1	A facility specific SOP for autoclave validation is available and adhered to.							
14.2	Documentation of autoclave validation is maintained and made available upon request.							
14.3	Autoclaves are validated at least monthly.							