

BIOHAZARDOUS AGENT REFERENCE DOCUMENT

Alpha Toxin

The Biohazardous Agent Reference Document (BARD) is a general guidance resource that reviews and summarizes the nature of a pathogen or biotoxin, and offers safety requirements for work with the agent in the laboratory. The BARD may replace the formal SOPs used in conjunction with some IBC registrations.

The BARD is provided as an additional guidance tool, and is not a substitute for a risk assessment, biosafety training, lab-specific training, or a formal <u>IBC master protocol registration</u>. This document should be readily available in the laboratory, and it is the responsibility of the Laboratory Supervisor or Principal Investigator to ensure that all personnel have read, understood, and signed the document. The BARD is for informational purposes only, and is not intended to be a substitute for professional medical advice, diagnosis, or treatment. Please consult a health care provider for any medical questions or concerns.

INSTRUCTIONS

- 1. Review the information contained in this document.
- 2. Add any necessary information that is specific to your work in the laboratory (such as strain-specific information). Please be sure that the track changes function is turned on to indicate any changes that you make.
- **3.** Instruct all personnel to review the BARD and sign the last page, indicating that they have read and understood the information.
- 4. Submit the BARD along with your IBC master protocol registration, amendment, or continuing review.



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CHARACTERISTICS	
Morphology	Pore-forming cytolysin, belong to the aerolysin-
	like family of toxins.
Characteristics	Major virulence factor of Clostridium septicum,
	the causative agent of atraumatic "gas
	gangrene." Muscle cells exposed to the toxin
	undergo cellular oncosis, characterized by
	mitochondrial dysfunction and release of
	reactive oxygen species

HEALTH HAZARDS	
Host Range	Humans, vertebrate and invertebrate animals
Modes of	Inhalation, mucous membrane contact, sharps
Transmission	injury, ingestion, dermal contact.
Signs and	Possible swelling, necrosis, edema, blisters, and
Symptoms	restriction in blood supply at site of exposure.
Toxic Dose	LD50 = 10 µg/kg (mouse, i.p.)
Incubation	Unknown for toxin alone, 6 hours to 3 days for
Period	clostridial myonecrosis associated with bacterial
	infection

MEDICAL PRECAUTIONS / TREATMENT	
Prophylaxis	None available for toxin
Vaccines	Not recommended
Treatment	Supportive treatment
Surveillance	Monitor for symptoms. Detection of toxin may
	be achieved by ELISA
UVM IBC	Report any exposures or signs and symptoms to
Requirements	your supervisor
Additional	Immunocompromised individuals are at a higher
Medical	risk for complications associated with exposure
Precautions	

LABORATORY HAZARDS	
Laboratory	No data. At least six cases of laboratory-acquired
Exposures	infections with Clostridium spp. have been
	reported up to 1976
Sources	Clostridium septicum

CONTAINMENT REQUIREMENTS	
BSL - 2	Preparation or dilution of the agent, work with clinical specimens and cultures known or suspected to contain the agent
BSL - 3	
ABSL - 2	Administration of the agent to an animal model. Animals may be housed at ABSL-1 post-exposure
ABSL - 3	
Aerosol	Centrifugation, homogenizing, vortexing or
generating activities	stirring, pipetting, pouring liquids, filling or expelling syringes
Primary	Use a chemical fume hood, biosafety cabinet, or
containment device	glove box for preparing stocks and dilutions

EXPOSURE PROCEDURES	
Mucous	Flush eyes, mouth or nose for 15 minutes at
membranes	eyewash station.
Other	Wash area with soap and water for 15 minutes
exposures	
Medical	Contact UVMMC Infectious Disease Dept. directly at
Follow-Up	(802) 847-2700 for immediate assistance. Bring this
	document with you if seeking medical care.
Reporting	Report all exposures or near misses to:
	1. Your immediate Supervisor
	2. The UVM Biosafety Officer at (802) 777-
	9471 and Risk Management at 6-3242
	Risk Management and Safety;
	https://www.uvm.edu/riskmanagement/inc
	ident-claim-reporting-procedures

PERSONAL PROTECTIVE EQUIPMENT (PPE)	
Minimum PPE	Nitrile gloves, lab coat or gown, appropriate
Requirements	eye/face protection. Wash hands after removing
	gloves.
Additional	Store in a secure location
Precautions	
(Risk	
assessment	
dependent)	



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VIABILITY	
Disinfection	10% bleach with a contact time of 30 minutes.
Inactivation	Autoclaving at 121°C for 15 - 30 min
Stability in	Stable at normal room temperature and pressure
Environment	

SPILL CLEAN UP PROCEDURES

Small Spill	Notify others working in the lab. Allow aerosols to settle. Don appropriate PPE. Cover area of the spill with paper towels and apply approved disinfectant, working from the perimeter towards the center. Allow 30 minutes of contact time before clean up and disposal. Dispose in double biowaste bags and biobox.
Large Spill	Inside of a lab: Call UVM Service Operations at 656-2560 and press option 1 to speak to a dispatcher. Ask them to page Risk Management and Safety. Outside of a lab: Pull the nearest fire alarm and evacuate the building. Wait out front of the building for emergency responders to arrive.

REFERENCES	
Canadian PSDS	https://www.canada.ca/en/public- health/services/laboratory-biosafety- biosecurity/pathogen-safety-data-sheets-risk- assessment/clostridium.html
BMBL	https://www.cdc.gov/labs/pdf/CDC- BiosafetyMicrobiologicalBiomedicalLaboratories -2020-P.pdf
Molecular Microbiology	https://onlinelibrary.wiley.com/doi/full/10.1111 /j.1365-2958.2005.04774.x
American Society for Microbiology	https://www.ncbi.nlm.nih.gov/pmc/articles/PM C257555/
Toxins	https://www.ncbi.nlm.nih.gov/pmc/articles/PM C4344638/
Cayman Chemical	https://www.caymanchem.com/cms/caymanch em/LiteratureCMS/Detection%20of%20Clostridi um%20Septicum%20Alpha%20Toxin.pdf

STUDENT / EMPLOYEE NAME

SIGNATURE

DATE