



Vaccine and Infectious Disease Organization
- International Vaccine Centre



Managing Aerosols from an Operations & Maintenance Perspective

Tracey Thue, Biosafety Officer

Vaccine & Infectious Disease Organization – International Vaccine Centre

(VIDO-InterVac), University of Saskatchewan

Managing Aerosols from an O&M Perspective

- ❖ Introduction to VIDO-InterVac
- ❖ Risk Assessment process
- ❖ RG3 pathogens, aerosols
- ❖ Engineering controls in CL3-Ag, HEPA filters
- ❖ Local Risk Assessment for O&M staff
- ❖ Procedure for O&M staff

International Vaccine Centre (InterVac)

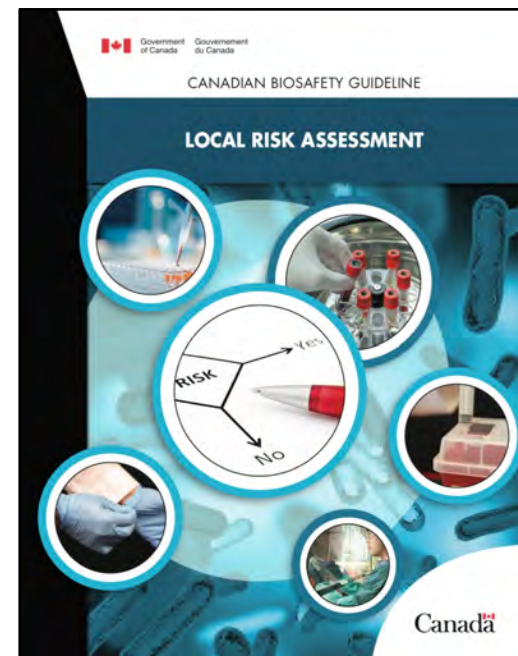
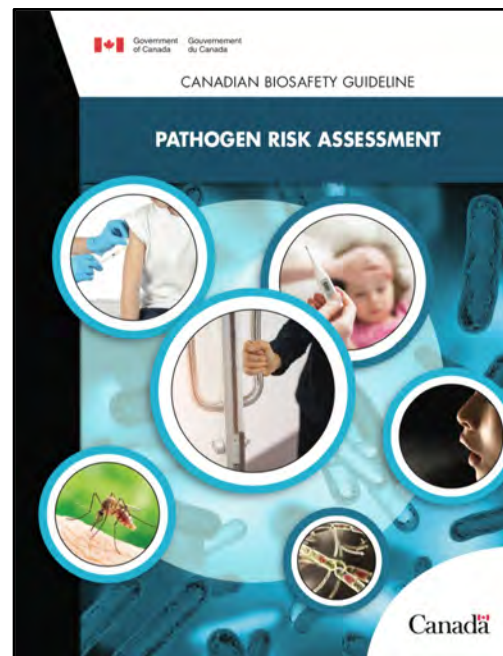
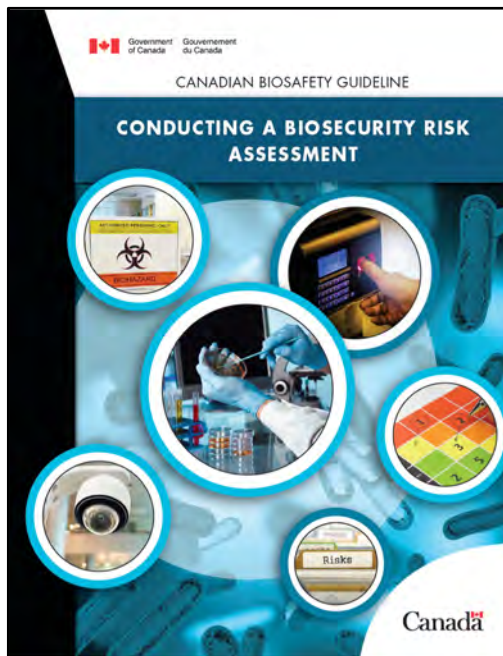


Large CL3 and
CL3-Ag facility

In-house full-time
O&M team of 7

Risk Assessment

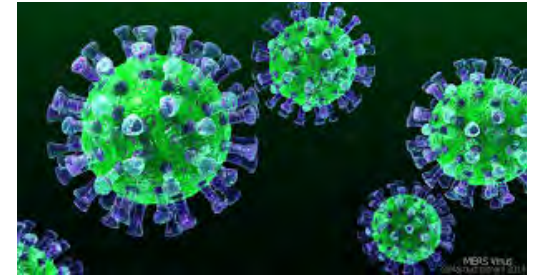
- Step 1: Identify and Characterize Hazards
- Step 2: Identify and Assess Risk
- Step 3: Develop and Implement Risk Mitigation Strategies
- Step 4: Review and Continually Improve



Risk Groups 1 - 4

RG1 *low individual, community risk*

E. coli, Baker's yeast



RG2 *moderate individual, low community risk*

Adenoviruses, PEDV, Influenza, ZV

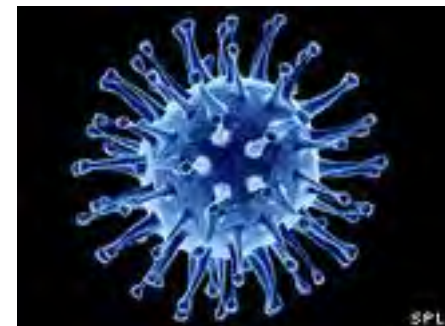


RG3 *high individual, low community risk*

HPAI, MERS-CoV, *M. tuberculosis*,
M. bovis

RG4 *high individual, community*

Ebola virus



Aerosols



- Created by any action that imparts energy into a liquid or semi-liquid
- Larger aerosol droplets (5 – 100 μm) settle quickly & contaminate surfaces: **ingestion hazard**
- Smaller droplets (< 5 μm) evaporate rapidly, particulates remain airborne for a long time: **inhalation hazard**

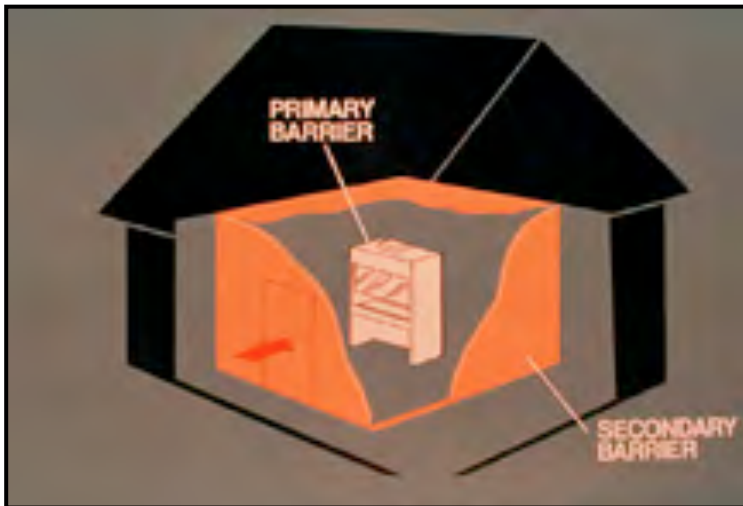
Controls of Biosafety

A photograph of a modern biosafety building. The building has a light blue or grey facade on the left side and a large glass facade on the right side. There are several trees and bushes in the foreground, and a paved walkway. The sky is clear and blue.

- **Engineering Controls**
- **Administrative Controls**
- **Procedural Controls**
- **Personal Protective Equipment**

Engineering Controls

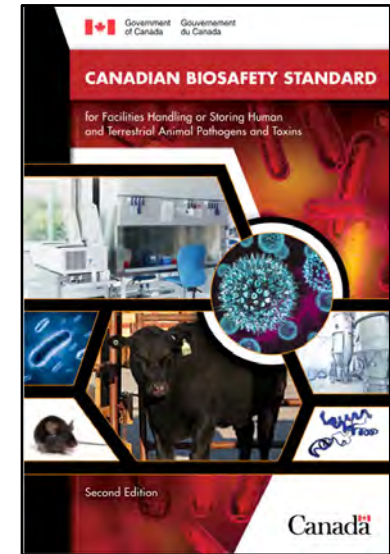
Facility Design Secondary Containment



- “Box in a box”
- 1° barrier (BSC) protects worker
- 2° barrier protects environment outside the laboratory
 - HEPA filters

HEPA Filtration

		CL2	CL2 -Ag	CL3	CL3 -Ag	CL4
3.5.9	Supply and exhaust air systems to be provided with automatic mechanical/electronic interlocks that prevent sustained positive pressurization of the containment zone.			■	■	■
3.5.10	Where IDA is provided, exhaust air to be: <ul style="list-style-type: none"> passed through a filter that prevents the release of infectious material or toxins; or 100% exhausted directly to the outdoors. (Not required for SA zones.)	□	■			
3.5.11	Exhaust air to be passed through HEPA filtration.			■	■	



Institute of Environmental Sciences & Technology

Recommended practice for basic provisions for HEPA filter units as a basis for agreement b/n customers & suppliers.

Describes 11 levels of performance & 6 grades of filter construction.

		CL2	CL2 -Ag	CL3	CL3 -Ag	CL4
3.5.12	Exhaust air to be passed through two stages of HEPA filtration.					■
3.5.13	HEPA filters to conform to IEST-RP-CC001.5.			■	■	■
3.5.14	HEPA filter housings to be designed to withstand structural changes at applied pressure of 1 000 Pa (i.e., 4 inches water gauge [in. w.g.]) in accordance with ASME N511 and AG-1.			■	■	■
3.5.15	HEPA filter housings to be designed to allow <i>in situ</i> filter isolation, decontamination , and testing.			■	■	■

Engineering Controls

To researchers working in the lab



Engineering Controls

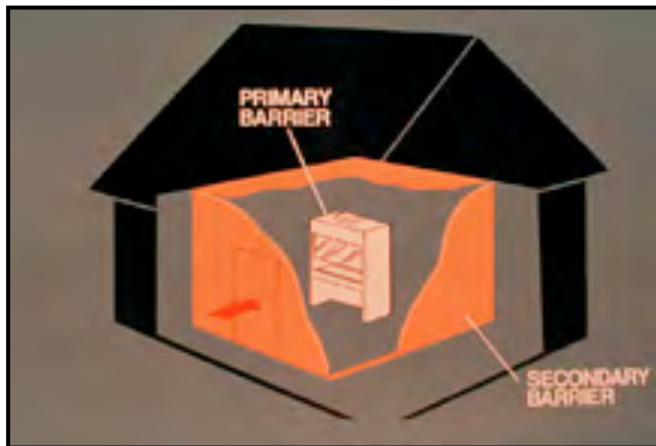
To O&M team on HEPA deck




Engineering Controls

Facility Design Secondary Containment

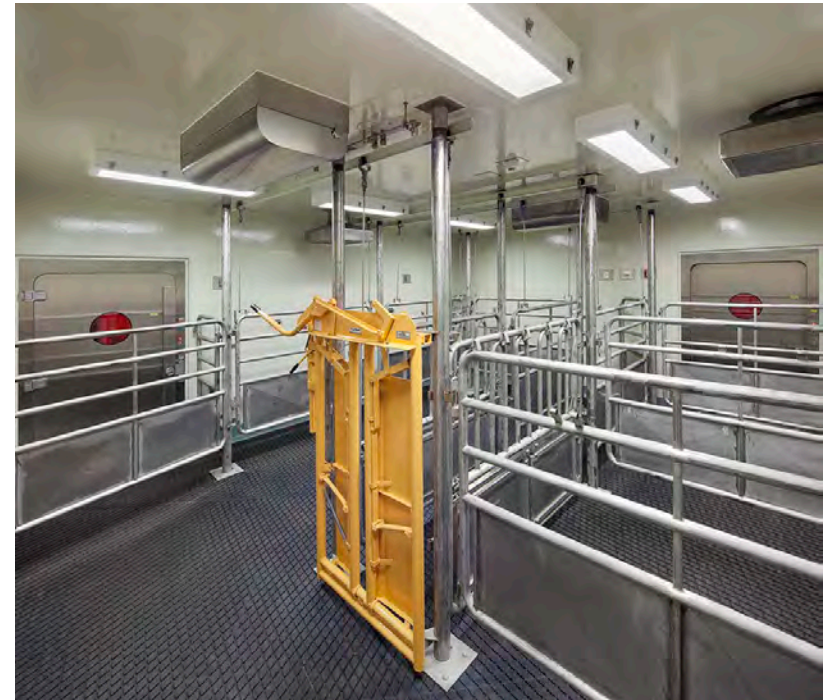
In CL3-Ag, large animals cannot be placed inside containment device, so



- 2° barriers become 1° barriers
- This is why risk  when working in animal cubicles

Engineering Controls

To researchers working in a large animal cubicle



Engineering Controls

To O&M team on HEPA deck above
large animal cubicle



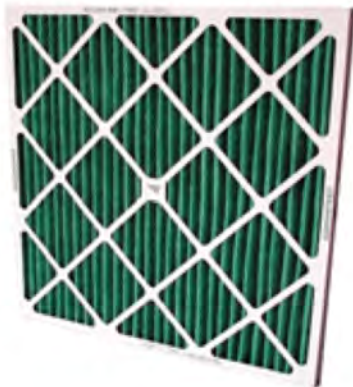
HEPA Filters

Camfil HEPA/ULPA Filter

P/N 5210192 XH Absolute

99.97 – 99.9995% filtering efficiency @ 0.3 μ m

Micro-fine glass media formed into pleats separated by a corrugated aluminum separator.



Camfil 30/30 Pre-filter

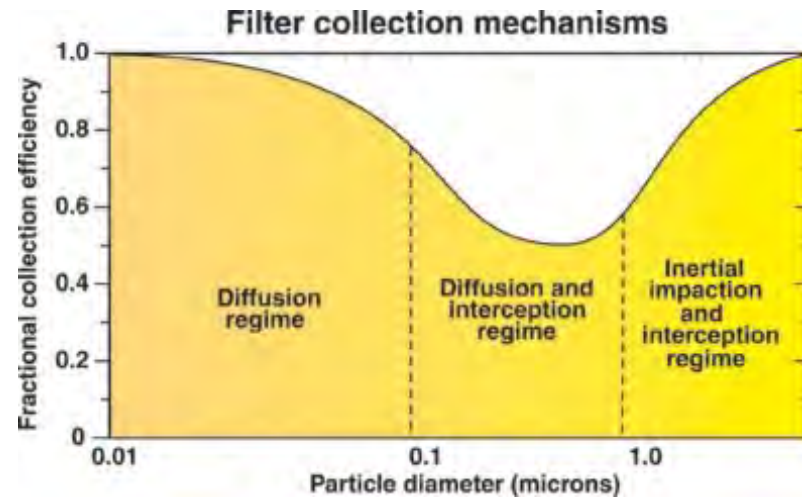
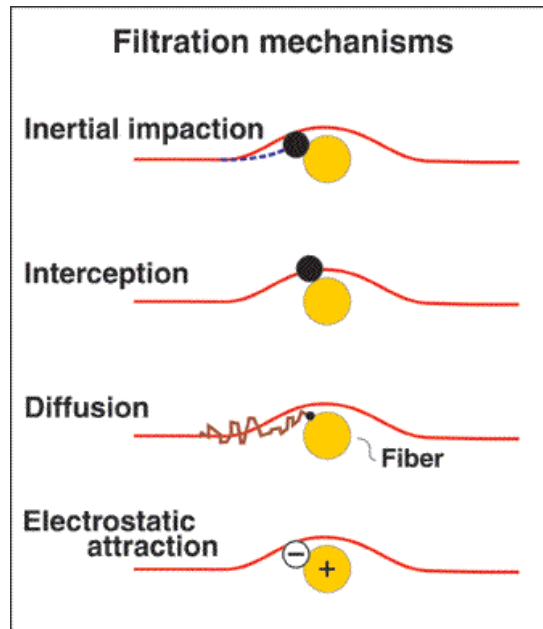
MERV 8 ASHRAE Std 52.2-2007, App J.

Filters particles of sizes 3.0 – 10 μ m (and larger)
i.e. Mold, spores, cement dust, **hair, dander**

Cotton and synthetic media with welded wire support grid, beverage board enclosing frame.

HEPA Filters

99.97% Filtering efficiency lowest at
0.3 μ m particle diameter





Building Design & Engineering Approach to Airborne Infection Control

AIR FILTRATION

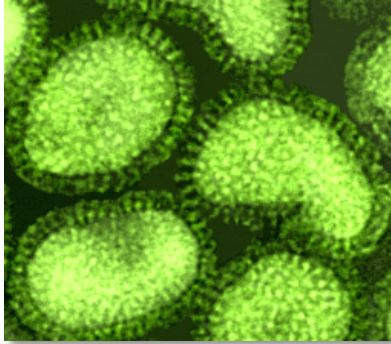
Steve Rudnick*

2010

Book chapter describing construction and function of HEPA filters.

*Exposure, Epidemiology and Risk Program, Dept. of Environmental Health, Harvard School of Public Health, Boston, Massachusetts

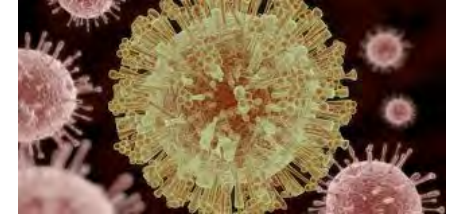
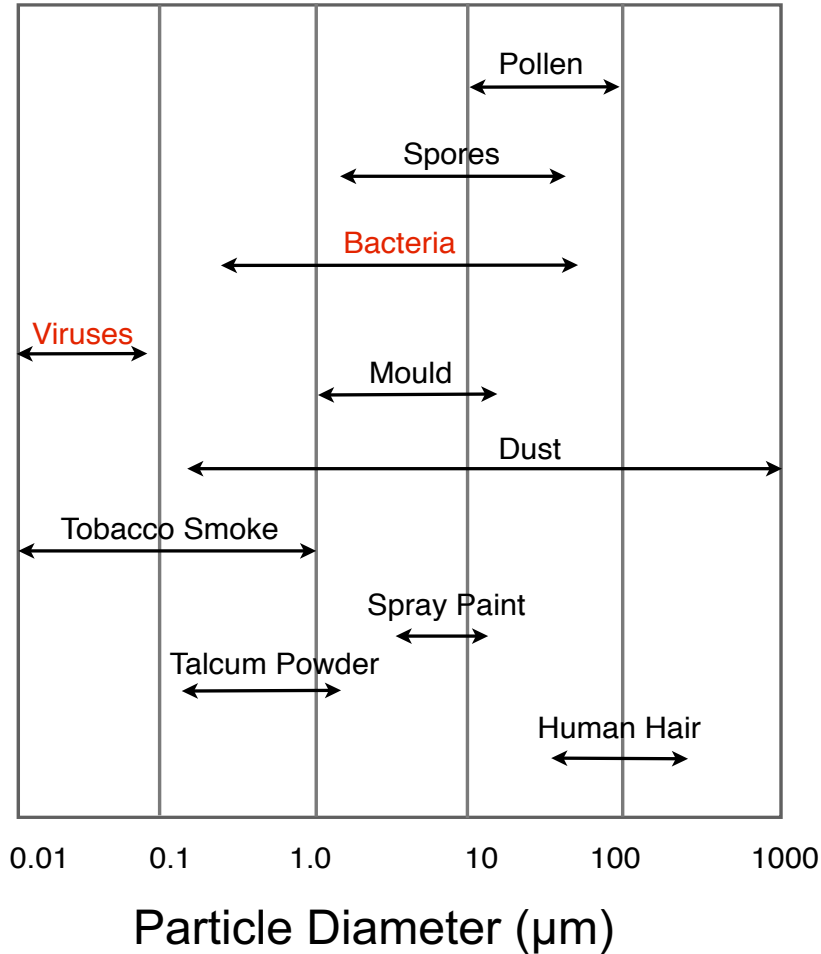
Particle Sizes



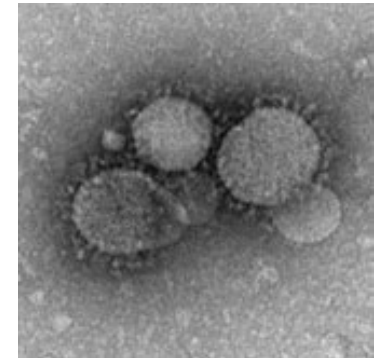
Influenza virus
0.1 - 0.2 μm



M. tuberculosis
1 - 4 μm x 0.25 μm



Zika virus
0.04 μm



MERS Co-V
0.1 - 0.3 μm



CL3 Lab single HEPA filter housing on exhaust air



CL3-Ag Animal Cubicle single HEPA filter housing on supply air



CL3-Ag Animal Cubicle double HEPA filter housing on exhaust air

Local Risk Assessment

1. Hazard Identification

Activity: Scan the HEPA filters



➤ Open HEPA housing if scan fails

Bag in/bag out design

Local Risk Assessment


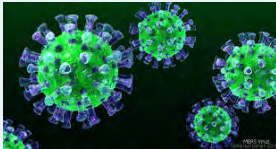
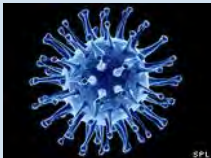
2. Identify and Assess Risk

Opening contaminated HEPA housing



Risk	Likelihood	Consequence
Exposure to worker	Medium - High	Low – High <i>Depends on pathogen survival in housing, contact</i>
Release to environment	Low - Medium	Low – High <i>Depends on pathogen survival, mobility</i>

Pathogen Characteristics

Pathogen	Survival @ RT on dry surface	Heat inactivation	Chemical inactivation
<p>M tuberculosis</p> 	Months	<p>> 65C 30 min 121C 15 min</p>	Cavicide 3 min
<p>MERS-CoV</p> 	24hr – 6 days	60C 30 min	5% MicroChem Plus, 10 min
<p>Influenza A</p> 	<p>Hard surface: 24-48hrs Porous surface: 8-12hrs SST+OM: 7days</p>	<p>70C 5 min 80C 2.5 min 90C 1 min</p>	Cavicide 3 min

Local Risk Assessment

3. Develop Risk Mitigation Strategies

❖ Administrative Controls:

- Training
- Documentation of room use
- Communication

❖ Procedural Controls:

- Post signage to communicate status of housing
- VHP decontaminate housing prior to opening
- PPE
- Disposal of HEPA filter



HEPA Filters & Housings

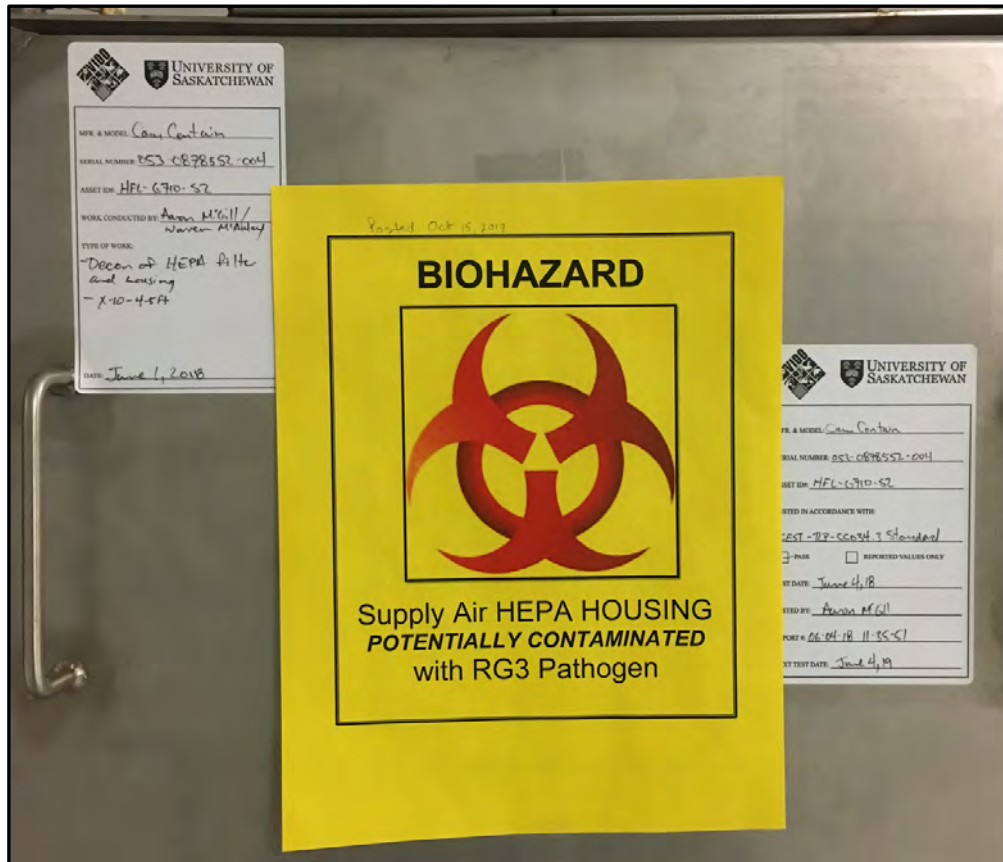


Exhaust Air HEPA Housing

Biohazard sign dated & posted by BSO on day of animal challenge, or shortly before.

From this date until VHP decon, all internal parts of housing considered **contaminated**.

HEPA Filters & Housings



Supply Air HEPA Housing

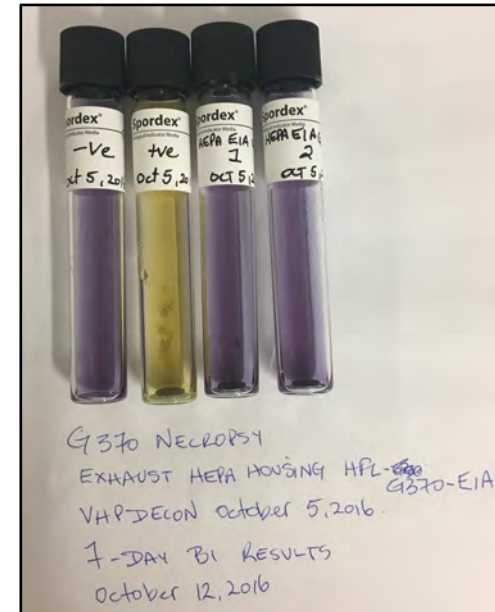
Biohazard sign posted:

“Potentially contaminated” unless the room pressure goes positive for any reason.

- Signage is changed to “contaminated”, date and reason.

HEPA Filters & Housings

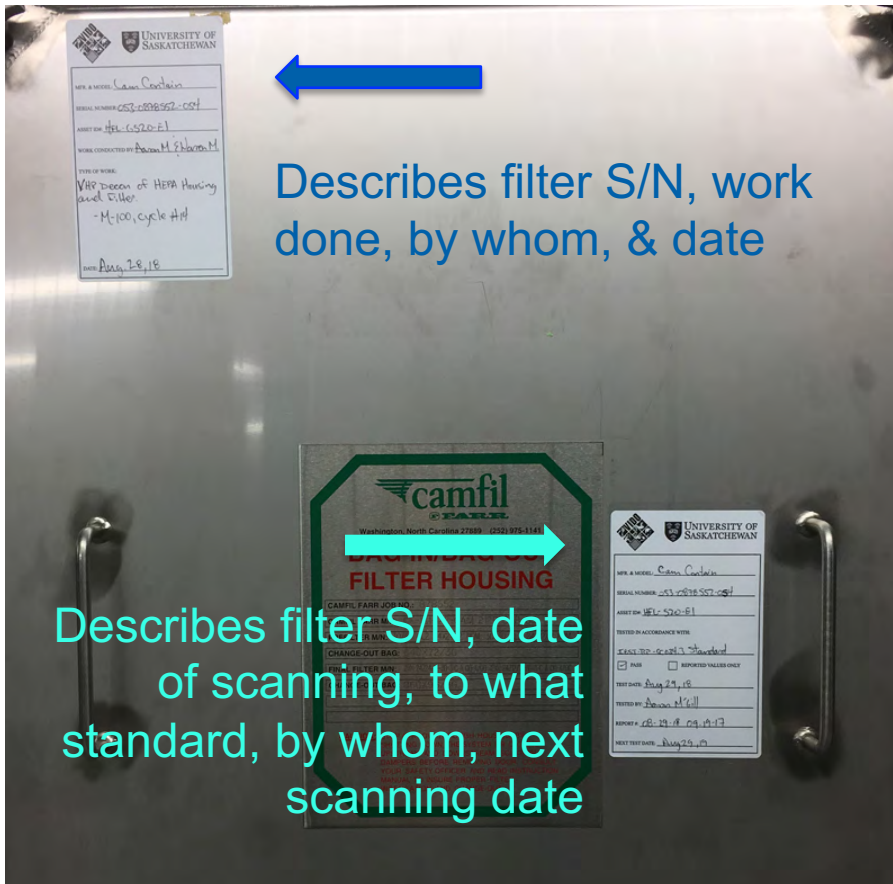
VHP Decontaminate filters and housing



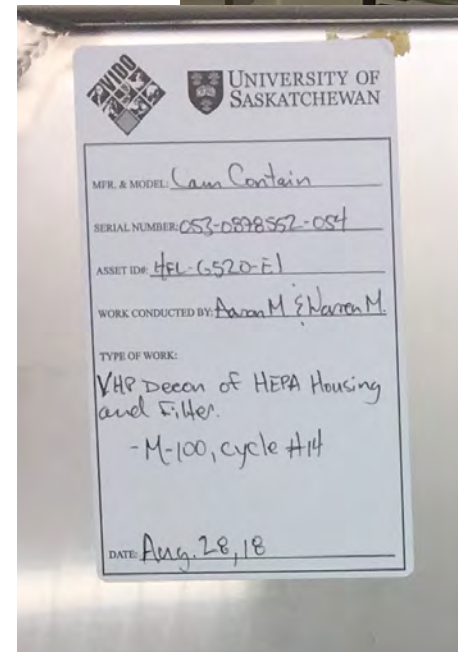
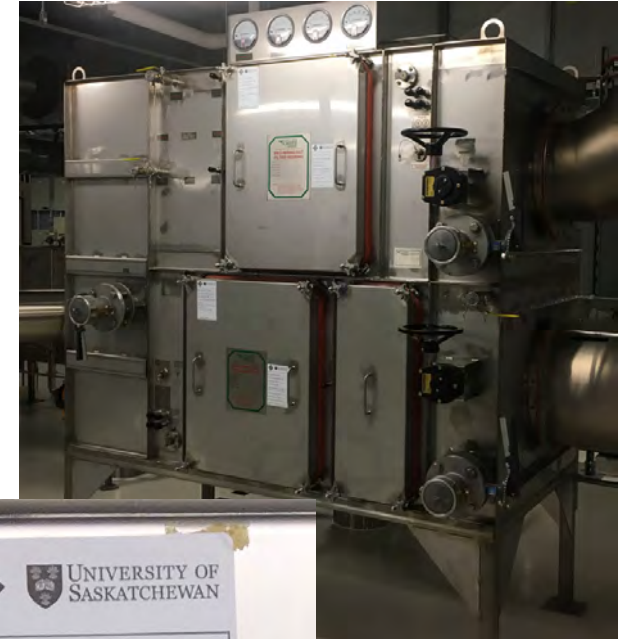
Autoclave filters
& dispose

HEPA Filters & Housings

Signage:
Testing and maintenance work



Describes filter S/N, date of scanning, to what standard, by whom, next scanning date





HEPA Filters & Housings



MMWR™

Morbidity and Mortality Weekly Report

Recommendations and Reports

December 30, 2005 / Vol. 54 / No. RR-17

Guidelines for Preventing the Transmission of *Mycobacterium tuberculosis* in Health-Care Settings, 2005

88

MMW

or other condition that interferes with the seal of the respirator to the face.

- **How long can I use my respirator for TB exposures before I discard it?** Disposable respirators can be functional for weeks to months and reused by the same HCW. Reuse is limited by hygiene, damage, and breathing resistance, and manufacturer instructions should be considered.

- **Should persons who perform maintenance on and replace filters on any ventilation system that is likely to be contaminated with *M. tuberculosis* wear a respirator?**

Laboratory studies indicate that re-aerosolization of viable mycobacteria from HEPA filters and N95 disposable respirator filter media is unlikely under normal conditions; however, the risks associated with handling loaded HEPA filters in ventilation systems under field-use conditions have not been evaluated. Therefore, persons performing maintenance and replacing filters on any ventilation system that is likely to be contaminated with *M. tuberculosis* should wear a respirator (see Respiratory Protection) and adhere to local recommendations for eye protection and gloves.

Local Risk Assessment

4. Review and Improve

❖ Administrative Controls:

- Training
- Documentation of room use
- Communication

❖ Procedural Controls:

- Post signage to communicate status of housing
- VHP decontaminate housing prior to opening
- PPE – nitrile gloves, safety glasses
- Disposal of HEPA filter



CL3 SA →

Thank you!

