

Infection Preventionists Role With Environmental Microbial Hazards

Greater Omaha APIC Conference

August 9, 2019

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Objectives

1. Identify opportunities for Infection Preventionists to consult with Industrial Hygienists based upon the infection control risk assessment findings.
2. Examine interventions that can be implemented into the infection control plan to mitigate environmental hazards.

No conflict of interest to disclose



What's all the concern about with construction?

Architects and engineers may not understand the workflow of the areas they are designing

Many healthcare executives may not understand the importance of Infection Preventionists in the realm of space design and construction

Many Infection Preventionists are uncomfortable with their role in construction and view their input as minor



The truth of the matter...

HEALTH

F.D.A. Details Contamination at Pharmacy

By SABRINA TAVERNISE and ANDREW POLLACK OCT. 26, 2012



RELATED COVERAGE

With Meningitis Outbreak, a Spotlight on Family Behind Compounding Pharmacy OCT. 24, 2012

Recent Spinal Shots May Pose Greatest Meningitis Risk OCT. 24, 2012

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Seattle Children's Hospital mold infections leave one dead, force closure of most operating rooms



Contaminants introduced into the healthcare environment

- During construction or renovation
 - Materials exposed to weather elements
 - Dust, debris containing mold are released if the ICRA plan is poorly executed
 - i.e. above ceiling work, wall demo, removal of millwork
- Through openings in the building envelope (doors, windows, openings) or building pressurization
- Through the HVAC system
- Through moisture intrusion (flooding, high humidity)
- Through the water distribution system
- Excavation adjacent to air intake of occupied facilities



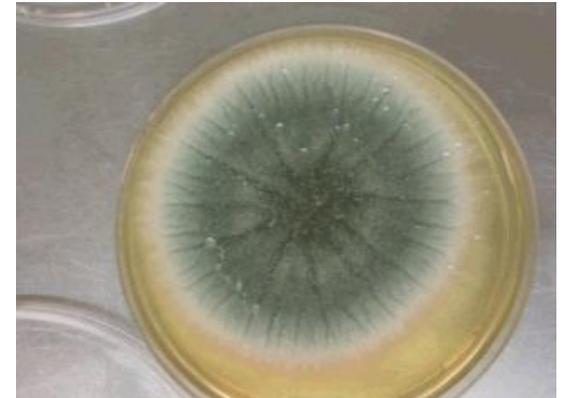
Environmental Contaminants

Mold is ubiquitous and usually found in spore form

- If conditions are favorable, spores will germinate and form reservoirs

Minimum requirements for mold growth

- Moisture (humidity, flood)
- Temperature
- nutrients



Construction Related Pathogens

- Aspergillus, Fusarium, Mold, Mucor, Penicillium, Rhizomucor, Rhizopus, Stachybotrys
- Cause superficial, locally invasive, and disseminated infections



Aspergillus species

Found in nature

Found in plaster, sheetrock and dust

Grows in damp materials/settings

Can cause illness or even death in some “at risk” individuals

- Infants
- Elderly
- Immunocompromised

Invasive disease

- Spreads rapidly to brain, heart, bone, kidney or skin
- Invasive pulmonary aspergillosis occurs only in people with diseases affecting the immune system (cancer, BMT)
- May be fatal



Roles and Responsibilities

Industrial Hygienist

Provide consultation, technical advice

Respond to inquiries about occupational and environmental health exposure issues

Provide training and technical support

Serve as subject matter experts

Provide guidance

Does not replace the IPs role

Infection Preventionist

Acquire knowledge – Learn the construction language

Educate, Raise awareness - Start with executives (buy-in), hospital staff, contractors

Collaborate – ICRA team members

Communicate – speak up, do not feel intimidated

Guide – provide expertise

Oversight – ensure all procedures and protocols are followed

Inspect and verify – prior, during, and after construction



When do Infection Preventionists consult Industrial Hygienists?

Environmental assessment & opportunities

- Construction rounds
- Moisture intrusion events
- Project in high risk settings and population

What monitoring tools are available to you?

- Particle testing
- Air sampling
- Moisture sensor – drywall

Level of experience

Available expert guidance



CDC Environmental Infection Control Guidelines (Last update: April 2019)

Particulate sampling –

- Commonly used to compare indoor (clean) to outdoor (dirty) air
- Absence of air quality and action level standards

Air Sampling –

- Controversial due to unresolved technical limitations and need for substantial lab support
- Lack of standards linking fungal spore levels with infection rates (no safe level of exposure)

No recommendation is offered on routine microbiologic air sampling before, during, or after construction or before or during occupancy of areas housing immunocompromised patients. (Unresolved Issue)

Investigators have suggested the following limits for aspergillosis outbreaks:

- 15 CFU/m³ for gross colony count of fungal organisms
- <0.1 CFU/m³ for *Aspergillus fumigatus* and other opportunistic fungi for HEPA filtered areas



Assessing Indoor Air Quality

Particulate Sampling

Hand held laser particulate counter

- Instrument calibrated in accordance with ISO requirements



Air Sampling

Quantitative fungal air cultures

- Andersen single stage N6
- Sabourauds Dextrose agar used for fungal cultures



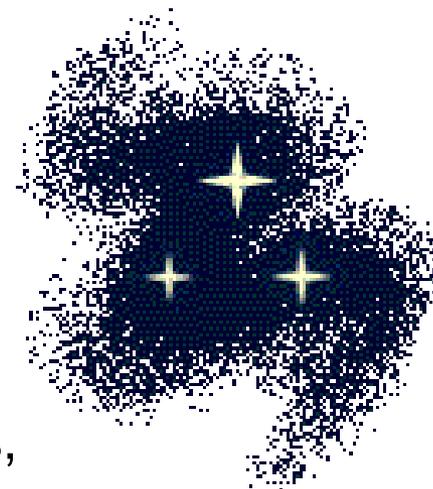
Air Particulate Sampling

Air Particles

Large dust particles travel in straight lines

Small spore particles travel on air currents

- Spores can stay suspended minutes/hour
- Settling rates vary based upon air movements, air exchange rates, and source strengths



Particles

- Collide with gas molecules & deflect
- Fibers can grab particles by electric charge

Air takes the route of least resistance

Construction dust associated with 5.0 to 10.0 micron size particles



Air Particulate Testing

Determined by high risk population near project, large project scope or the project is occurring within a patient unit.

- Measure baseline particles before project starts–
 - outside (comparative high particle count)
 - construction site (comparative high particle count)
 - occupied spaces adjacent to barriers (lower counts)
- Establish the ‘norm’ prior to construction activity
- Measure routinely during project to validate dust mitigation
 - Identifies a change in the project or integrity of barriers



ICRA Compliance



ICRA Complinnace



Air Sampling Microbial Commissioning

Fred & Pamela Buffett Cancer Center Grand Opening May, 2017

Patient Care Services –

- 36 bed Hematopoietic Stem Cell Transplant Unit
- 12 bed Intensive Care Unit
- 24 bed Progressive Care Unit
- 36 bed Medical-Surgical Unit
- Airborne isolation rooms

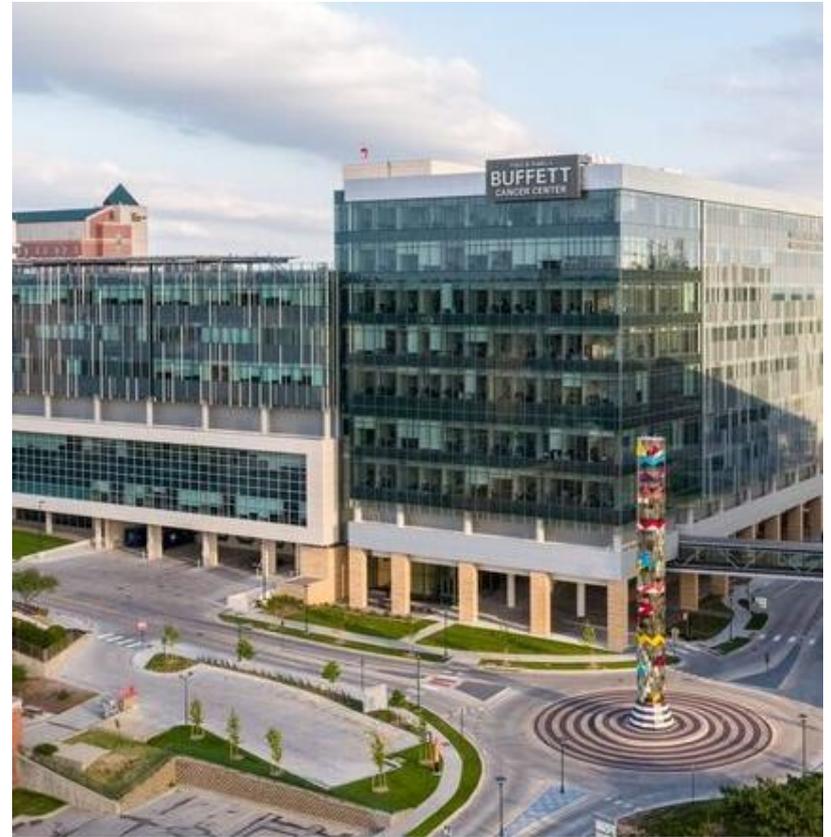
Surgical Suites

Infusion Center

Outpatient Clinic – procedural areas

Pharmacy – compounding area

All Ancillary Support Services



Clean Room Standards

ISO 14644-1: 2015 Cleanroom Standard - International Standards Organization (ISO)

- Most used standard in Pharma and Electronics controlled environment

USP 797 - Pharmaceutical cleanroom classification for compounding sterile preparations

FS209E - U.S. Federal Standard 209 E

- U.S. General Services Administration (GSA) retired the standard in 2001, superseded by ISO 14644, but it is still widely accepted



Features of a Clean Room

- Provide a secure environment with controlled air filtration lowering the contamination risks
- Maintain particulate-free air through the use of HEPA air filtration
- Classified according to the number and size of particles permitted per volume of air in a specified amount of time
- Design of air distribution system includes the downstream air return
 - Air flow is unidirectional
 - Vertical flow rooms – use of low wall air returns around the perimeter of the zone
 - Horizontal flow rooms – the use of air returns at the downstream boundary of the air flow process

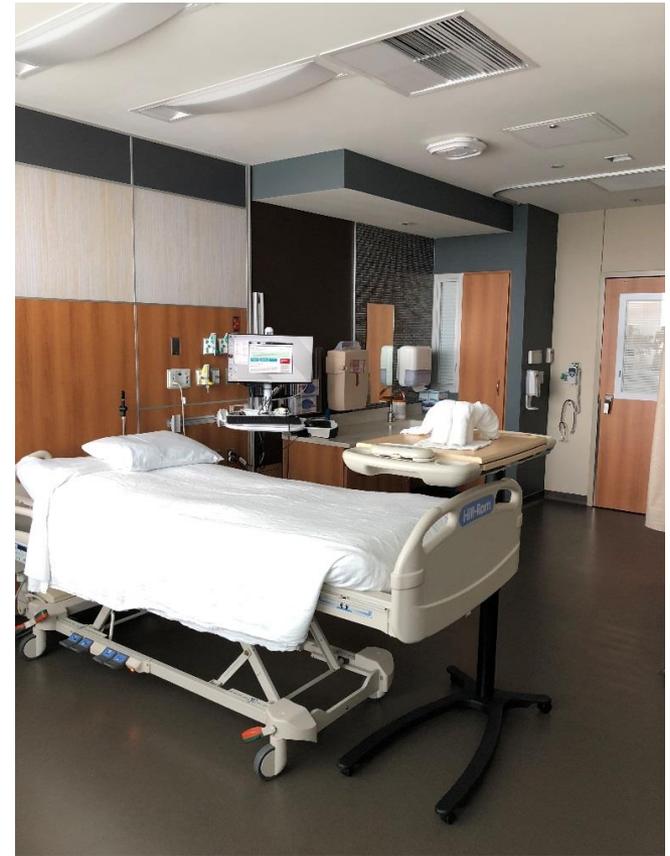


Air Distribution and Directional Flow

Vertical Flow: Surgery Suite



Patient Room



Clean Room Classifications

- Cleanrooms are classified according to the number and size of non-viable particles permitted per volume of air
- Classifications use **non-viable** particle measurements
 - Rationale: The fewer particulates present in a clean room the less likely it is that airborne microorganisms will be present
- ISO 14644-1 requires specific particle counts to classify the cleanroom
 - Sampling plans use several randomized locations allowing for different particle concentration levels in different parts of the cleanroom
 - 5.0 micron size particles are allowable for ISO class 6-9 only



Class	Maximum particles/m ³ ^a						FED STD 209E equivalent
	≥0.1 μm	≥0.2 μm	≥0.3 μm	≥0.5 μm	≥1 μm	≥5 μm	
ISO 1	10 ^b	d	d	d	d	e	
ISO 2	100	24 ^b	10 ^b	d	d	e	
ISO 3	1,000	237	102	35 ^b	d	e	Class 1
ISO 4	10,000	2,370	1,020	352	83 ^b	e	Class 10
ISO 5	100,000	23,700	10,200	3,520	832	d,e,f	Class 100
ISO 6	1,000,000	237,000	102,000	35,200	8,320	293	Class 1,000
ISO 7	c	c	c	352,000	83,200	2,930	Class 10,000
ISO 8	c	c	c	3,520,000	832,000	29,300	Class 100,000
ISO 9	c	c	c	35,200,000	8,320,000	293,000	Room air

^a All concentrations in the table are cumulative, e.g. for ISO Class 5, the 10 200 particles shown at 0,3 μm include all particles equal to and greater than this size.

^b These concentrations will lead to large air sample volumes for classification. Sequential sampling procedure may be applied; see Annex D.

^c Concentration limits are not applicable in this region of the table due to very high particle concentration.

^d Sampling and statistical limitations for particles in low concentrations make classification inappropriate.

^e Sample collection limitations for both particles in low concentrations and sizes greater than 1 μm make classification at this particle size inappropriate, due to potential particle losses in the sampling system.

^f In order to specify this particle size in association with ISO Class 5, the macroparticle descriptor M may be adapted and used in conjunction with at least one other particle size. (See C.7.)



ISO Classifications & settings

ISO 1 = Cleanest room possible
(nanotechnology)

ISO 5 = Critical area in immediate proximity of exposed sterilized containers (USP 797)

ISO 6-8 considered cleanrooms

ISO 7 = Area immediately adjacent to the aseptic processing line under dynamic conditions (USP 797)

ISO 8 = Supporting area or clean room area used as an ante room for PPE donning (USP 797)

ISO 9 = Ordinary room air



Viability Particulates – Microbial Air Sampling

USP 797 requirements

- All viable air sampling must be volumetric by impaction on a media plate
- Settle plates are not permitted
- Multiple locations are required for air sampling
- Recommend sampling around contamination-prone areas

Viable colony counts that are considered elevated or actionable are as follows:

- ISO Class 5 = Active airborne $>1 \text{ CFU/m}^3$
- ISO Class 7 = Active airborne $>10 \text{ CFU/m}^3$
- ISO Class 8 = Active airborne $>100 \text{ CFU/m}^3$



ISO Classifications applied to non-traditional clean rooms in hospitals

Gormley, T., et al. (2017). Methodology for analyzing environmental quality indicators in a dynamic operating room environment. *AJIC*. 45, 354-359.

- Adapted ISO standards to compare 3 ORs in 2 states as no building code standards exist related to air particles and air sampling
- Clean space assessment using air particle, air sampling and ISO standards
- ORs were equivalent to ISO Class 7 and 8

Holy & Matouskova (2012). The importance of clean rooms for the treatment of haemato-oncological patients. *Contemporary Oncology*. 16(3), 266-272.

- Discusses the application of FS209E (ISO class) standards to clean rooms for the care of haemato-oncological patients

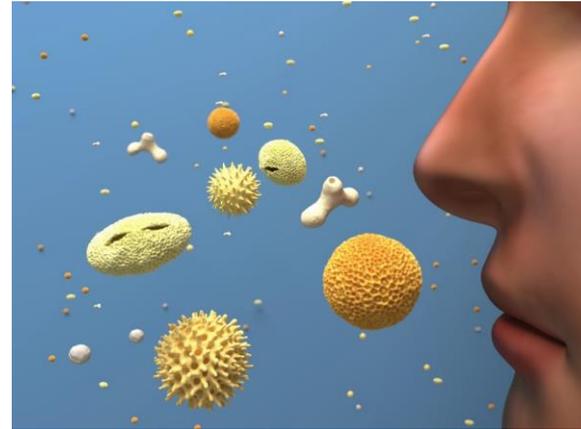


Environmental Readiness for Air Sampling

- All surfaces in patient care areas terminally clean by Environmental Services (3 separate times)
 - All areas thoroughly clean and dust-free
- Minimum number of boxes
- No outside shipping boxes in patient care areas
- Activity restricted for 24 hours in all areas planned for air cultures
- HVAC operating at normal capacity (not on setback mode) continuously for at least 24 hours after final clean and prior to collecting air cultures



Initial Particle Counts & Cultures



Location	Particle count (5 micron)	Culture (cfu/m ³)
Outdoor	918	ND
East Elevator (3 floors)	240- 2120	23-28
Center Elevator (3 floors)	252- 2360	51-65
6 th floor room (5 rms)	30- 1175	0-2
7 th floor (5 rms)	40-866	0- 17
7 th floor hallway	1670	17



Remediation

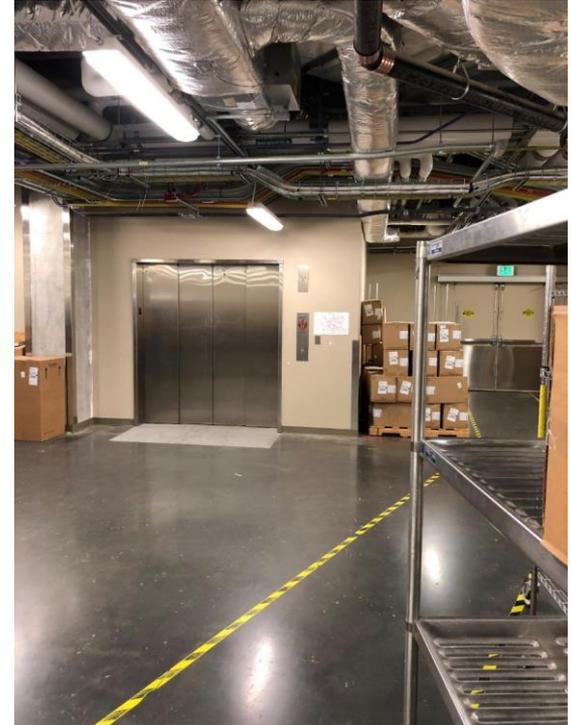
Problem:
Helicopter port
and particle entry
into building

Solution: vestibule
doors



Remediation

Problem:
Loading dock
and particle entry
Solution: Confirm
loading dock
procedures and
air curtain check



Remediation

Air pressure
determination

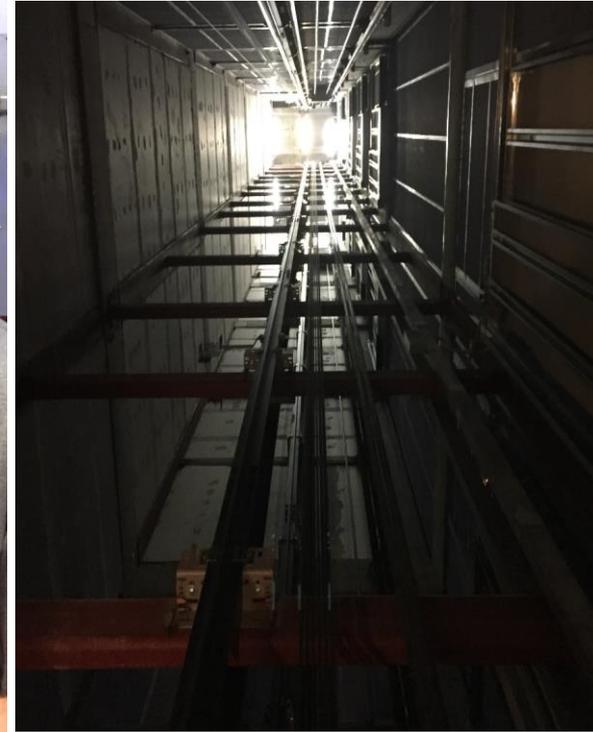
Positive flow out of
patient care areas



Remediation

Problem: Elevator shafts

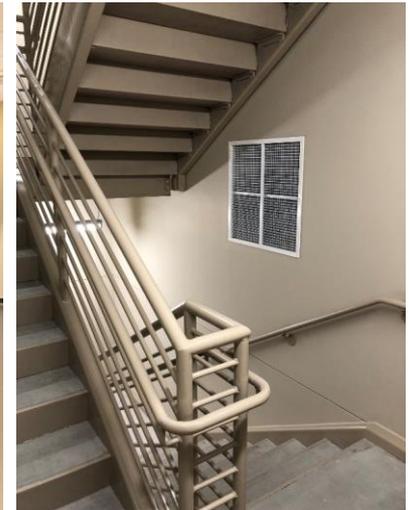
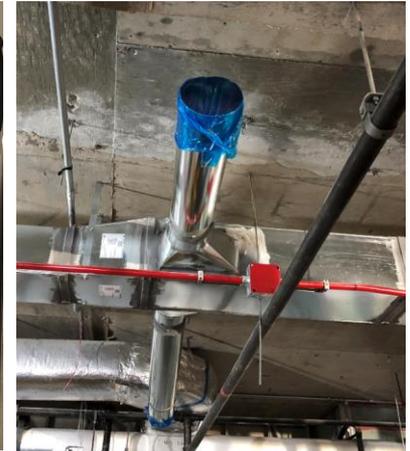
- Thoroughly cleaned with vacuum and moist towels x 2
- Roof hatch checked and weather tight
- Desperate times call for desperate measures – irradiated the problematic shaft



Remediation

Other actions:

- Thorough “terminal” cleaning
- Checked all air filters, duct work, and air handling equipment
- Sweeps added to stairwell doors, patient doors, nurse server gasket
- Stairwell fan coils inspected and cleaned
- Ceiling sealed, plenum pressurized



Location	CFU/m ³	Final 5.0 µm Particle Count (outdoor 2270)	Final ISO Classification
8 th Floor – Med-Surg			
Patient Room	---	Not Applicable	
Corridor	---	120-380	ISO Class 7
Elevator Vestibule	---	820-1290	ISO Class 7
7 th Floor - OHSCU			
Patient Room (5)	0-1.7	70-273	ISO Class 6
Corridor (Elevator)	0-5.3	70-130	ISO Class 6
Elevator Vestibule	12.3-26.5	1158-1170	ISO Class 7
6 th Floor – ICU/Prog. CU			
Patient Room (2)	0	130-808	ISO Class 7
Corridor (Elevator)	35.3	123-380	ISO Class 7
Elevator Vestibule	19.4-35.3	1180-1596	ISO Class 7
2 nd Floor			
Surgery	0-1.7	10-60	ISO Class 6
Pharmacy Comp.	0	10	ISO Class 6
Infusion Ctr – Procedure	8.8	10	ISO Class 6

Can air particles replace microbial air sampling?

Landrin, A., Bissery, A., & Kac, G. (2005). Monitoring air sampling in operating theatres: Can particle counting replace microbiological sampling? *Journal of Hospital Infection* 61, 27-29.

- Methods of microbiological and particle counting did not correlate ($p=0.06$)
- No particle count value could be predictive of microbiological count higher than 5 CFU/m³

Cristina, M.L., et al. (2012). Can particulate air sampling predict microbial load in operating theatres for arthroplasty? *PLOS One*, 7(12), e52809.

- No statistical significant correlation between microbial loads and particle counts for $\geq 0.5\mu\text{m}$ and $>5.0\mu\text{m}$ particle size.
- Microbiological monitoring remains the most suitable method of evaluating air quality in operating rooms



Conclusions

- The correlation between air particle counts and microbial air sampling has yet to be established, therefore particle monitoring can not replace microbial monitoring
- The method recommended to measure hygienic air quality is the combination of air particle counts and microbial air sampling
- Applying ISO clean room standards to non-traditional clean room settings in a hospital is emerging in the literature



Water, Moisture, External Environment

Disease Transmission Associated with Potable Water Systems



- Disruption of water utility systems during construction or renovation can disrupt biofilm present in water delivery pipes
- Poses a threat to patients, including those far away from an active construction zone.
- Dead end pipes allow water to stagnate where microbes can grow to very high concentrations.



Potable Waterborne Pathogens

ORGANISM	SITE OF INFECTION	SOURCES OF INFECTION
<i>Pseudomonas aeruginosa</i>	Blood, catheter insertion site, lungs, urinary tract	Potable water, contaminated liquid solutions and disinfectants, endoscopes
<i>Stenotrophomonas maltophilia</i>	Blood (septicemia), pneumonia, urinary tract, wound infections skin, stools, throat, trachea, urine	Potable water, distilled water, contaminated liquid solutions and disinfectants
<i>Acinetobacter baumannii</i>	Skin, wound	Room humidifiers, distilled water, moisture in mechanical ventilators
<i>Chryseobacterium</i> spp.	Blood	Potable water (burn unit), ice machine
Non-tuberculous <i>Mycobacteria</i> (NTM) species (<i>avium</i> and <i>fortuitum</i>)	Abscesses and wound infections, disseminated	Hospital hot water system, shower
<i>Legionella pneumophila</i> (other species rarely cause disease)	Lung (pneumonia), sternal wound infection	Hospital hot water system
<i>Aspergillus</i> species and <i>Fusarium</i> species	Wound infection, disseminated disease	Hospital water system

Moisture Intrusion & Remediation

Investigate and evaluate moisture & mold problems

Assess size of moldy area, consider hidden mold, repair small mold problems before they become large problems

Communicate with building occupants at all stages of process; Plan Remediation

Dry wet, non-moldy materials within 48 hours to prevent mold growth; remove if drying delayed

Select clean-up methods for moldy items; discard moldy porous items that can't be cleaned

Use PPE, containment, negative air

Implement repair plan



External Excavation Precautions

Excavation produces an enormous amount of dust

Ideally conducted off-hours

- Reduced traffic, opening of doors, restrict entrances

- Reduces the amount of non-filtered air entering the building

Protect air handler intake

Moisten excavated soil

Protect materials delivered for later use (shrink wrap)

Address health issues of patients & employees

- Face masks, tissues, hand sanitizer at entrances



Summary

- Environmental microorganisms can cause human disease
- Construction ICRA plan is critical in mitigating contaminants and microbial transmission risks
- Infection Preventionists and Industrial Hygienists play a role in protecting patients and employees from environmental hazards
- Infection Preventionists have variable experiences and expertise with environmental hygiene
- Partner with an Industrial Hygienist



Questions





**NebraskaSM
Medicine**

