Water as a Source of Hospital Acquired Infections APIC SF Bay Area Chapter, May 13, 2015

## This presentation has been modified to remove copyright protected material.



Michael Castro michael\_castro@pall.com

> Kevin Olson kolson@nalco.com

<u>Purpose</u>: Review potable water systems in the Healthcare Industry, focusing on Engineering & Plumbing Challenges (systemic vs. immunocompromised populations). Review both outbreak response and proactive risk minimization plans and best practices.

<u>Process</u>: An interactive presentation focusing on evidence based data, published best practices & common potable water system challenges.

<u>Pay-off:</u> Members of APIC SF Bay Area will have more resources & background to minimize the risk associated with waterborne pathogens (both proactively and reactively) in your healthcare facility.

## **Michael Castro**

**Disclosure:** Pall Medical Employee Western Region, Hospital Water Specialist

- B.S. in Mechanical Engineering, Bucknell University
- Enrolled in Masters in Public Health Program, GCU
- Pursuing Certification in Infection Prevention & Control (CIC®)
- Over 16 years of experience in advising and helping healthcare facilities, universities, commercial buildings & manufacturing facilities reduce their risk due to waterborne pathogens in all water systems including cooling water and domestic water.
- Specific areas of focus include microbiological filtration and secondary disinfection of domestic water systems. Previously worked with Hospital's Potable Water Systems in KY, NJ, NY & AZ.
- Member in APIC, ASHE, AWT, and ASHRAE

## **Kevin Olson**

**Disclosure:** Nalco, an Ecolab Company Environmental Hygiene Services Specialist

- BS Chemical Engineering, University of California at Davis
- 28 years with Nalco; Water treatment in Industrial, Food & Beverage, Semiconductor and Institutional markets
- Over 7 years of experience in advising and helping healthcare facilities, universities, commercial buildings & manufacturing facilities reduce their risk due to waterborne pathogens in all water systems (domestic water, industrial/HVAC water, other at-risk systems)
- Specific areas of focus include Water Safety Plans, secondary disinfection of domestic water systems, industrial water systems, and microbiological filtration
- Work with Healthcare and Institutional Facilities in Western US
- Member of APIC, ASHRAE, NSF Certified HACCP Manager

# What do Facility Managers and Infection Preventionists have in common?

Not Enough Hours In The Day!

You Each Have Your Own Perspective

You Each Have Your Own Priorities

Both Teams Can Reduce the Risk Associated with Waterborne Pathogens & HAIs

But it takes a team effort!

Water is essentia... And yet, water can cause unintended human harm if not properly engineered, managed and monitored.

#### What happens to drinking water from its origin to the tap?

In most circumstances, well-controlled, hygienic water is delivered from water plants to cities. During transport, water is cold and flows continuously through large diameter pipes. However, this situation changes dramatically at the point-of-entrance to buildings1, 2. Within buildings water stagnates and its temperature increases. It passes through complex internal distribution systems consisting of narrow pipes with possibly corroded inner surfaces and dead ends. This environment provides optimal conditions for the formation of biofilm from which bacteria and other microorganisms are continuously released into the water 3-5.

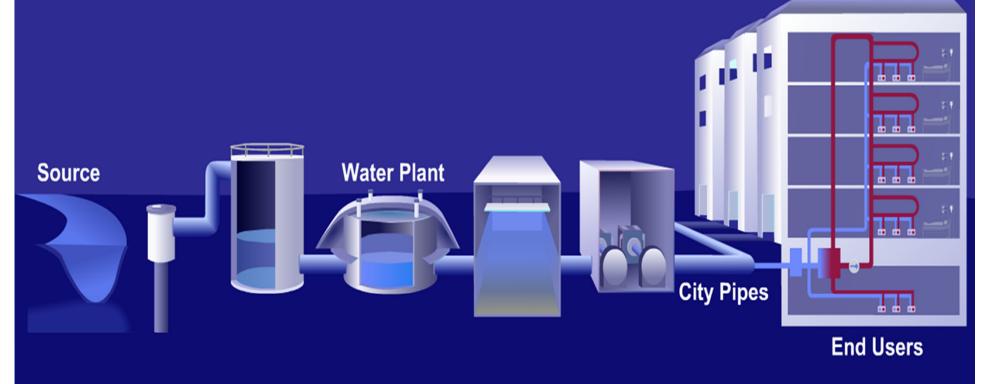


Image 1: Drinking water is derived from lakes, rivers or deep underground. It is purified in water plants and transported underground in large diameter pipes to cities and buildings, where it then runs through small diameter pipes, stagnates and warms up. These conditions are ideal for biofilm formation.

### **Complex Drinking Water Supply Chain**

Why can we not rely on the Water Treatment Plant alone?

- Domestic Water is not Sterile Upon Entry into Facility
- Building Plumbing Systems Allow Further Biofilm Growth and Proliferation
- Often Times Severely Immunocompromised Patients Contact & Use H2O



### **Complex Drinking Water Supply Chain**

#### Why can we not rely on the Water Treatment Plant alone?

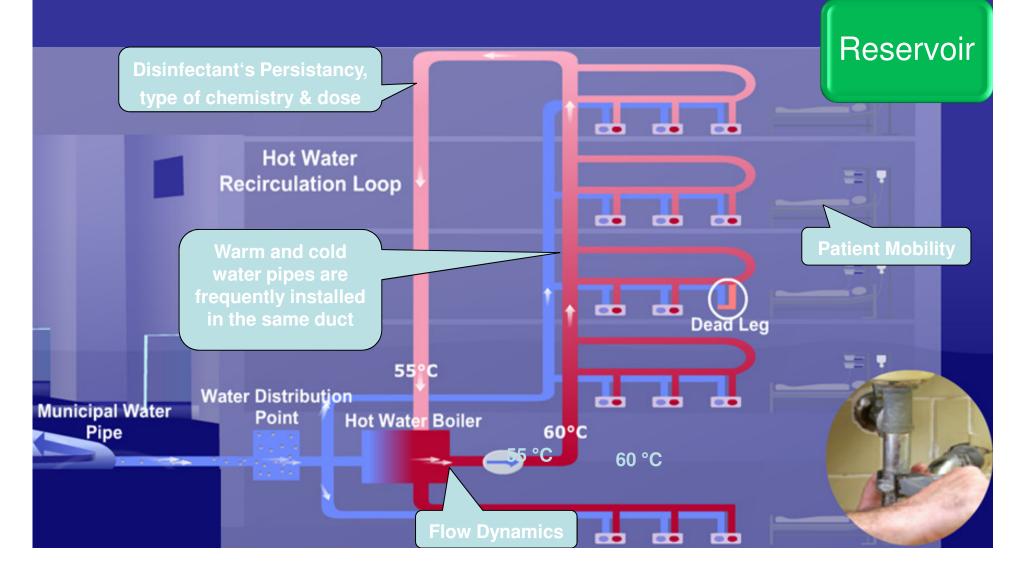
DETECTED CONTAMINANTS	UNIT	MCL	PHG OR (MCLG)	RANGE OR LEVEL FOUND	AVERAGE OR [MAX]	MAJOR SOURCES IN DRINKING WATER		
TURBIDITY								
Unfiltered Hetch Hetchy Water	NTU	5	N/A	0.2 - 0.3 <sup>(1)</sup>	[3.6] <sup>(.)</sup>	Soil runoff		
Filtered Water from Sunol Valley Water Treatment Plant (SVWTP)	NTU	1(3)	N/A		[0.98]	Soil runoff		
		min 95% of samples ≤0.3 NTU <sup>(3)</sup>	N/A	99.9 %		Soil runoff		
Filtered Water from Harry Tracy Water Treatment Plant (HTWTP)	NTU	1 <sup>(3)</sup>	N/A		[0.13]	Soil runoff		
		min 95% of samples ≤0.3 NTU <sup>(3)</sup>	N/A	100 %		Soil runoff		
City Pipes								

#### End Users

FOOTNOTES: <sup>(1)</sup> Turbidity is measured every four hours. These are monthly average turbidity values. <sup>(2)</sup> The highest turbidity of the unfiltered water in 2013 was 3.6 NTU. <sup>(3)</sup>There is no turbidity MCL for filtered water. The limits are based on the TT requirements for filtration systems.

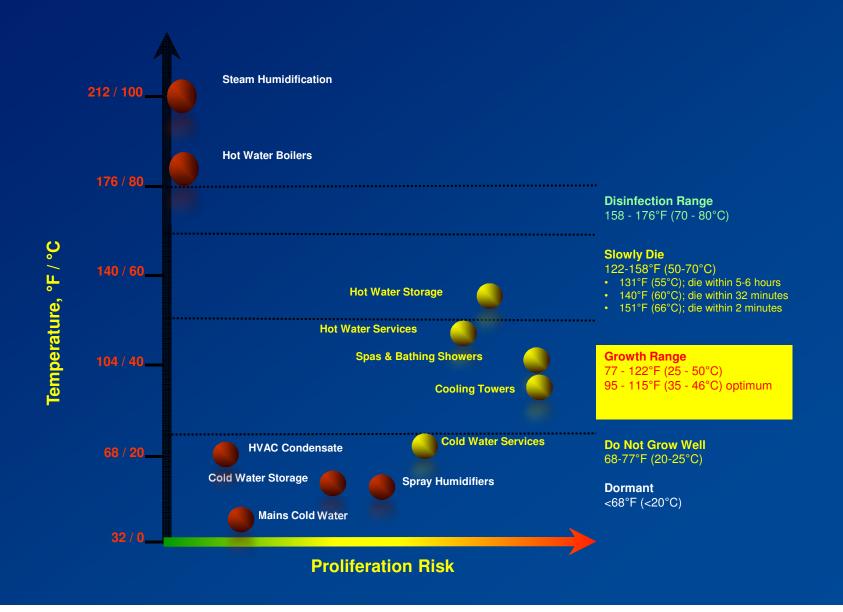
*Annual Water Quality Report 2013.* (2013). Retrieved from San Francisco Water Power Sewer: http://sfwater.org/index.aspx?page=634

**SYSTEM CONDITIONS** Complexity, age, poor temperature controls, lack of residual disinfectant, and water stagnation can provide conditions that allow formation of biofilms.



### **Utility & Domestic Services**

Temperature vs. Proliferation Risk



## **Bioffilm is our enemy** This is where most pathogens live, thrive and are often well protected from human interventions

Mycobacter

Pseudomonas

egionella

Acinetobacter

Stenotrophomonas

### Which sink option is best?

Portal of Exit



Knee/foot pedals



**Sensor faucets** 

Wrist blades

#### What do you consider when evaluating these options?

#### What does the wetted surface look like inside sink fixtures?





### Why does your healthcare facility use these?







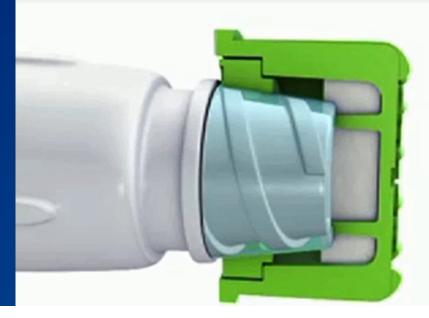
RUR CRIMING LOR (1983



## **Bioburden / Biofilm**



### Bacterial Bioburden



## Disinfects Ports

Why are these interventions appropriate for medical devices, but not effective with respect to hospital plumbing systems?



- Contact time
- High levels of disinfectant
- Disposal after a certain time

#### What is biofilm and how does it develop?

In water distribution systems biofilm can develop within a few days even if the water meets drinking water criteria 2. Biofilm can host bacteria, amoeba, algae and other microorganisms. Under low flow conditions, such as in dead legs, particularly thick biofilms can form. Under the force of water flow biofilm shears off and biofilm particles can colonize other parts of the water distribution system3. External physical stress in the pipework, such as disinfection measures, can result in an increased expression of the biofilm phenotype cell which is responsible for the strong attachment of cells to a surface 5.

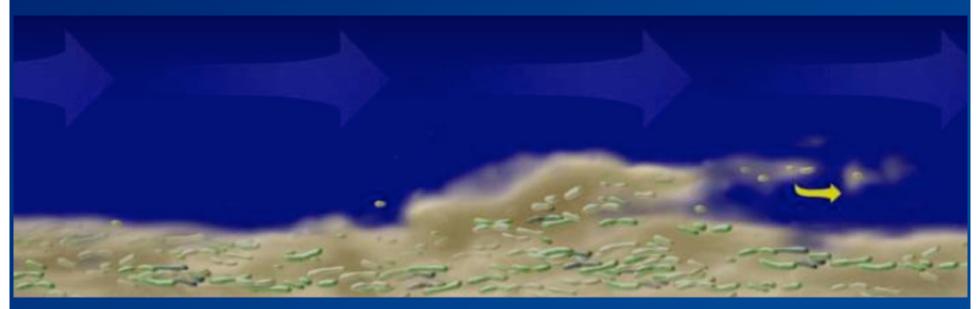


Image 2: Biofilm establishes in several phases over a few days. It contains microorganisms within its slimy matrix. With increasing thickness, biofilm particles containing large amounts of bacteria are released into the water stream.

#### Why does biofilm influence the water quality?

With increasing thickness, biofilm better protects the microorganisms within, from chemical agents and thermal disinfection procedures 2, 5. It is extremely difficult to completely eradicate the biofilm community once established. Irregular shedding from a biofilm can result in significant deviations of bacterial counts at sampling sites or points-of-use (POU) 2-4. Bacteria within biofilm communities have been shown to exhibit greater resistance against antimicrobial treatments than corresponding planktonic cells 3.



Image 3: When biofilm loaded with bacteria is released into the water stream, high microbial counts may be measured at the outlets. Annual testing provides only a snapshot of information, while regular testing is useful to monitor the bacterial risk of a pipe network.

#### Which microorganisms can be found inside biofilms?

The majority of bacteria in a water pipework live within biofilm (about 95%) and only about 5% occur in the water phase 4,6. Biofilms contain a large variety of waterborne microorganisms. These include protozoa (e.g. Acanthamoeba), fungi (e.g. *Aspergillus* spp.), viruses and a number of human pathogenic bacteria 1, 3-6,. Among those bacterial species found in biofilm that are potentially harmful for immunocompromised people are *Pseudomonas aeruginosa*, non-tuberculous Mycobacteria, *Stenotrophomonas maltophilia*, *Acinetobacter baumanii, Chrysobacterium* spp., *Sphingomonas* spp., and *Klebsiella* spp. 3-6. *Legionella pneumophil*a is perhaps the best-known bacterium colonizing biofilm, and it can be found in both central storage areas (e.g. water tanks) as well as peripheral water outlets 2,3,5. Waterborne *Pseudomonas aeruginosa* is a major cause of severe infections 7-9 including pneumonia, sepsis, wound and skin infections.

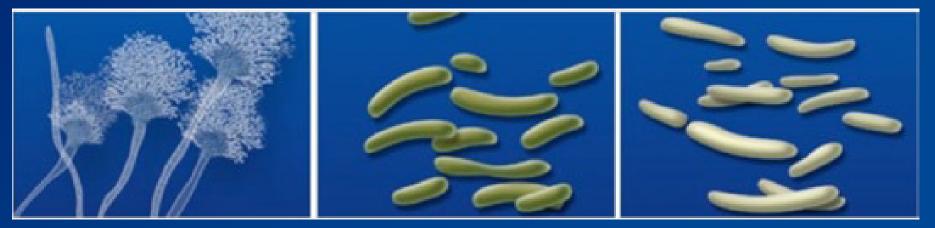


Image 4: Biofilm in water networks may contain a large variety of microorganisms such as fungi (e.g. Aspergillus spp., left), rodshaped bacteria (e.g. Legionella spp., middle) and protozoa (e.g. amoeba, right).

#### What are Viable But Non-Culturable cells?

The Viable But Non-Culturable (VBNC) cell fails to grow on routine bacteriological culture media, but is alive and capable of renewed metabolic activity - indeed it can be "resuscitated" to a culturable state with renewed virulence 6,10-12. This discovery has thrown the accuracy of quantifying culturing techniques into question. It is understood that a high proportion of biofilm dwelling cells live in the VBNC state and that the VBNC state can be induced by antibacterial material such as copper pipes 11 as well as by thermal treatments 12. As water pathogens such as *P. aeruginosa* in their VBNC state are not detectable by standard culture methods, alternative diagnostic technologies such as Polymerase Chain Reaction (PCR) or Fluorescence *In Situ* Hybridization (FISH) are required in order to confirm their presence 6.

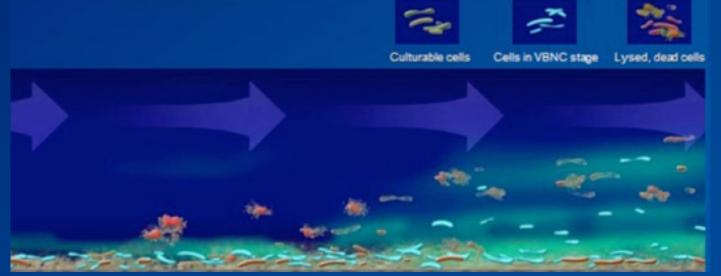


Image 5: Chemical or heat disinfection can destroy layers of biofilm and organisms within those layers. However, biofilm is rarely completely removed and bacteria can survive in a VBNC state (displayed as blue cells). Those cells can establish new colonies after chemical or heat treatment cessation.

#### What role do amoeba play in the biofilm community?

Amoeba are very important hosts for water bacteria. *L. pneumophila, Mycobacteria* spp. and other "amoeba resistant bacteria" can be carried by these protozoa 13,14. *Legionellae* are taken up into amoeba without being digested and replicate there within vacuoles. When the *Legionellae* have reached a certain density, the vacuoles release them into the water system 14.



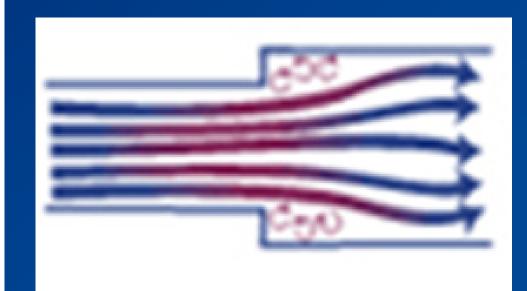
Image 6: Amoeba can incorporate Legionella which then proliferate inside vacuoles and are later released, either in the form of planktonic, free living bacteria, or packed within vacuoles.

# Are there any Mechanical Engineers in the room?

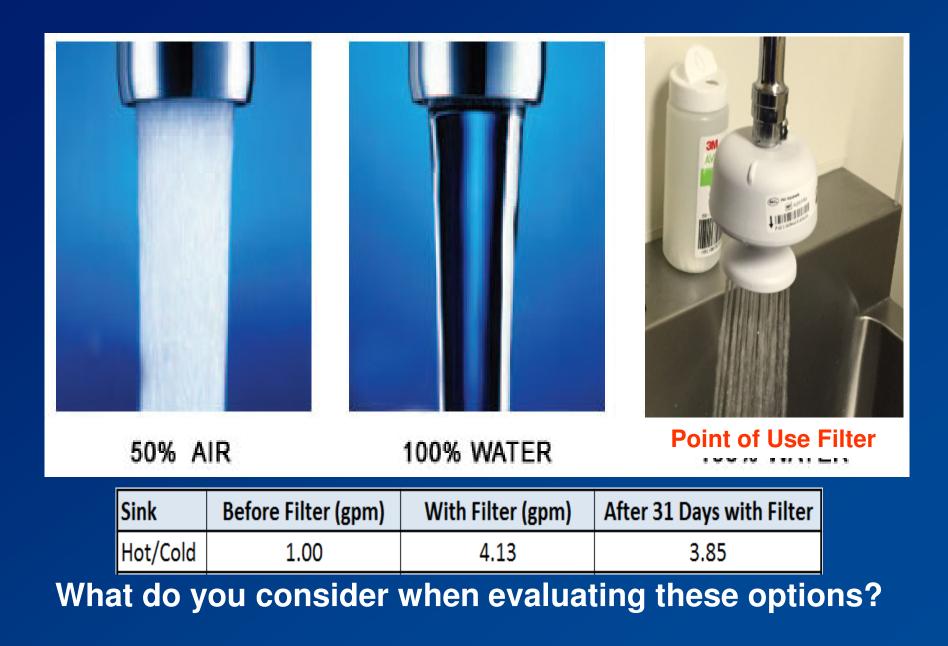
## **Civil Engineers in the room?**

## Has anyone considered Fluid Dynamics as it pertains to Infection Control?

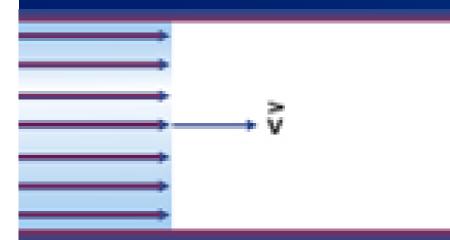




#### Which is the best option at point of use?

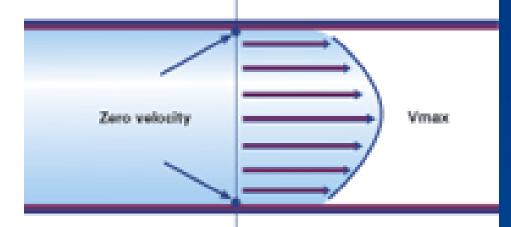


### Low Flow (use, flushing or laminar flow devices)

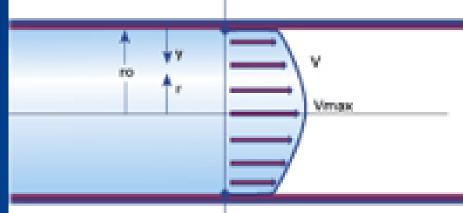




Hypothetical: Zero Liquid Viscosity & Pipe Wall Friction results in a velocity profile that is a straight line. The vertical sheet of fluid moves forward at velocity V.



Laminar Flow Profile for Newtonian Fluid The velocity is zero at the pipe wall and increases parabolically with flow, reaching its maximum at the pipe's center.



Turbulent Flow Profile for a Newtonian Fluid The velocity is zero at the pipe wall, but the face velocity is straighter and squared up.

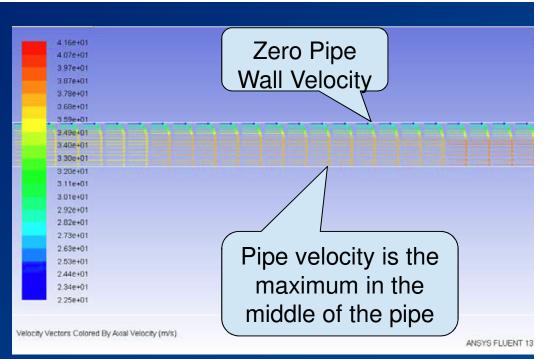
http://www.coleparmer.com/TechLibraryArticle/815

#### Improved Pipe Wall Velocity

- Larger Pipe Diameter
- Higher Flow Rates
- Smoother Surface (Coefficient of Friction)
- No Presence of Deposits
- Shorter Piping Runs

#### Poor Pipe Wall Velocity

- Smaller Pipe Diameter
- Lower Flow Rates
- Rougher Surface
   (Coefficient of Friction)
- Presence of Biofilm, Corrosion or Scaling Deposits
- Longer Piping Runs





## The importance of the viable but non-culturable state in human bacterial pathogens

#### Laam Li<sup>1</sup>, Nilmini Mendis<sup>1</sup>, Hana Trigui<sup>1</sup>, James D. Oliver<sup>2</sup> and Sebastien P. Faucher<sup>1\*</sup>

<sup>1</sup> Department of Natural Resource Sciences, Faculty of Agricultural and Environmental Sciences, McGill University, Ste-Anne-de-Bellevue, QC, Canada <sup>2</sup> Department of Biology, University of North Carolina at Charlotte, Charlotte, NC, USA

#### Table 1 | The species of human pathogens with a proven VBNC state.

Species	VBNC state inducing factor	Resuscitation condition	Resuscitation window	References
Acinetobacter calcoaceticus	Starvation			Lemke and Leff, 2006
Aeromonas hydrophila	Starvation	Temperature upshift		Rahman et al., 2001; Maalej et al., 2004
Agrobacterium tumefaciens	Starvation, chemicals (copper)			Byrd et al., 1991; Alexander et al., 1999
Arcobacter butzleri	Starvation	Rich medium, NOT temperature upshift	270 days	Fera et al., 2008

"Unlike normal cells that are culturable on suitable media and develop into colonies, VBNC cells are living cells that have lost the ability to grow on routine media, on which they normally grow" (Oliver, 2000).

#### Why is *Pseudomonas aeruginosa* of particular concern?

*Pseudomonas aeruginosa* is one of the most problematic bacteria in health care facilities and is responsible for about 10-20% of hospital-associated infections (HAIs) (pneumonia, wound infections, blood stream infections and urinary tract infections) in intensive care units (ICUs) <sup>8</sup>. Several studies have shown that up to 50% of the hospital acquired *P. aeruginosa* infections may be derived from the water distribution system <sup>15-17</sup>. Infection chains from water taps to people have been reported. *P. aeruginosa* easily colonizes all kinds of fluids (even distilled water) and rapidly forms biofilms8. *P. aeruginosa* strains have developed resistance against commonly used antibiotics, rendering effective treatment increasingly complicated and expensive <sup>18</sup>. *P. aeruginosa* is also increasingly recognized as a problematic water pathogen outside of hospital settings.



Image 7: Pseudomonas aeruginosa is an aerobic bacterial species and commonly found at the periphery of water systems such as taps, showers or sinks. A greenish color on the underside of a tap aerator may indicate colonization with Pseudomonas aeruginosa.

## What are the pathways for infection transmission from water sources to people?

Inhalation and aspiration represent transmission pathways for *Legionella* spp. whereas *Pseudomonas* spp. is mainly transmitted by contact and aspiration. During daily routines, tap water is used for personal hygiene. For example, in the healthcare environment, due to the severity of their disease states, ICU patients often have multiple access devices such as catheters, drains and tracheal tubes. These portals of entry represent potential entrance sites for bacteria. Droplets of contaminated tap water or contaminated hands of nursing staff can inadvertently come into contact with those entrance sites. Rogues *et al.* reported that 14% of ICU health care workers hands were *Pseudomonas* positive when washed with contaminated tap water and 12% were positive when the last contact was with a *Pseudomonas* positive patient 19. Contaminated bottled water or contaminated water from drinking water dispensers has also been described as a source of hospital-associated *Pseudomonas* infections in ICUs and Bone Marrow Transplantations (BMTs)

20, 21.



Image 8: Water for wound care, or the patient's personal hygiene, may contain bacteria resulting in patient colonization and infection. A patient may not necessarily have to use a water outlet to become colonized; immobilized patients, e.g. within ICUs, can come in contact with contaminated water brought by the nursing staff to the patient's bed.

## Why is complete biofilm eradication by systemic treatments so difficult?

Water distribution systems in large buildings are frequently complex networks and can be up to 50 km in length. Dead ends, corroded pipes, low throughput, insufficient temperature below 55 °C in the hot water pipes and above 20 °C in the cold pipes contribute to biofilm formation and impede eradication of biofilm. Heat & flush procedures (10-20 minutes of simultaneous flushing of all outlets with water heated to > 70 °C) may have only short term effects 22,. *Legionella* strains may even become heat resistant after thermal treatment over a long time 12,. Thermal procedures can result in warming up cold water 23 when both hot and cold water pipes are located in the same duct increasing the risk of biofilm in cold water.

Chemical treatments are bactericidal to free floating bacteria but have limited effects on biofilm and may create hazardous byproducts during use 22, 24, 25.

Therefore, areas with vulnerable users require additional interventions (e.g. point-of-use water filtration) to minimize patient's exposure to waterborne pathogens.

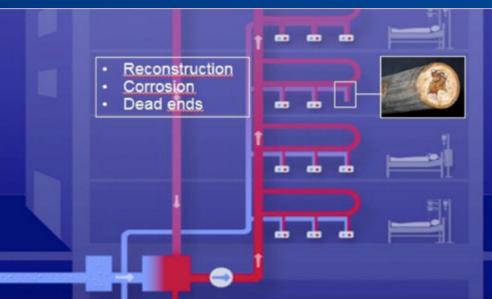


Image 9: Water systems in hospitals can contain corroded pipes and dead ends which cannot be reached by systemic disinfection. From there, bacteria can be released to recolonize the system after disinfection has been stopped. Construction work may also cause biofilm release into the network, colonizing other parts of the water system.

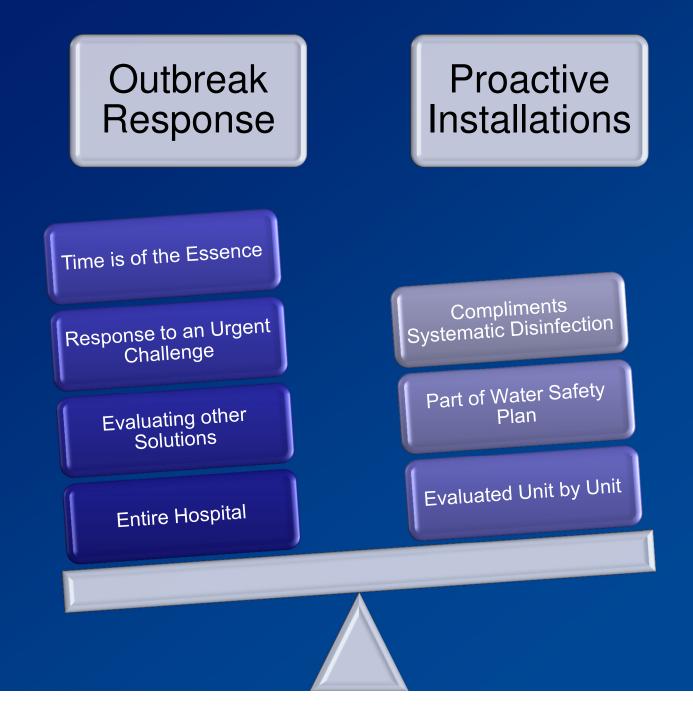
## Where are point-of-use (POU) water filters (tap filter, shower filter) typically used?

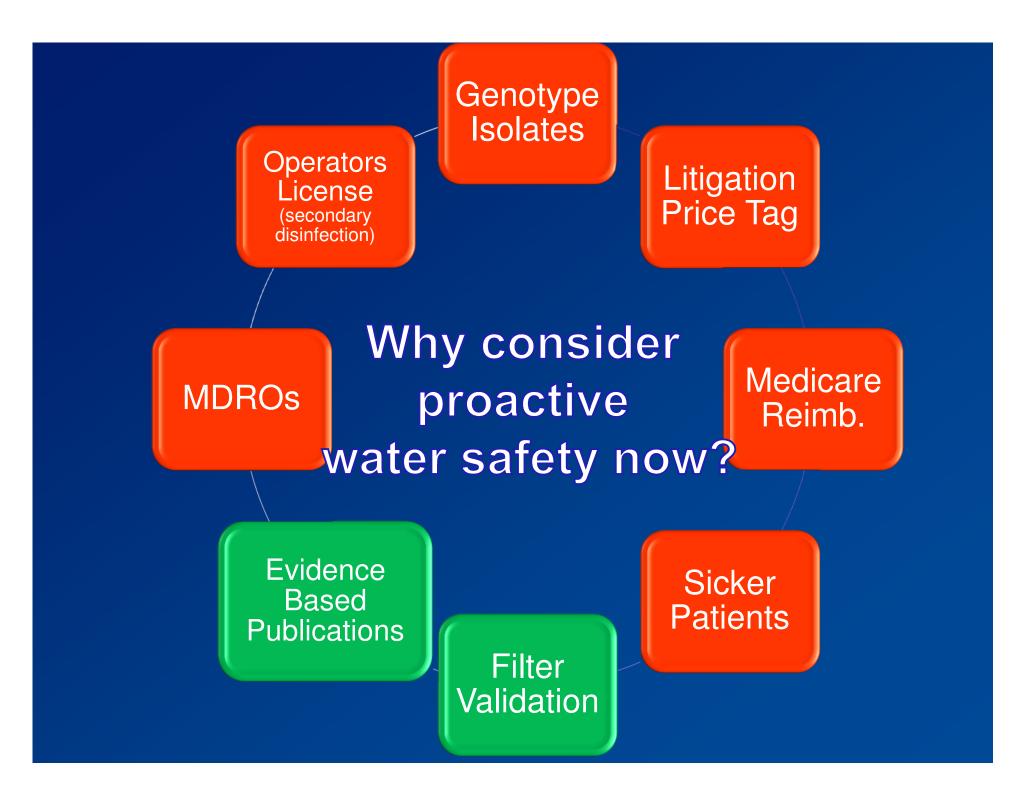
Point-of-use water filters are used as an additional physical barrier in those areas where immunocompromised people come into contact with water <sup>26-47</sup> and in outbreak or critical contamination situations. They can be flexibly installed at faucets (tap filters) or connected to shower hoses (shower filters). In medical facilities the most common areas for POU water filter installation are bone marrow transplant units, hematology/oncology units, ICUs, transplantation units, burn units, neonatology, manual endoscopic reprocessing, birth tubs, kitchen (for food preparation and drinking water provision), and geriatric departments. Based on the clinical experiences POU water filtration is also increasingly used in other areas with immunocompromised patients such as nursing homes or home care settings.

POU filters are quickly installed which makes them an effective management tool in acute situations such as outbreaks e.g. in public buildings, apartment houses, swimming pools, sports centers and hotels.



Image 10: Point-of-use water filters can be installed at taps, showers or in-line applications in various health care settings, such as hospitals, day surgery units, dental practices, dialysis units, or rehabilitation centers. They may also be used for immunocompromised homecare patients.





## 721 Hospitals Penalized For Patient Safety

By Jordan Rau | December 19, 2014

Medicare Reimb.



Medicare is penalizing 721 hospitals with high rates of potentially avoidable mistakes that can harm patients, known as "hospital-acquired conditions." Penalized hospitals will have their Medicare payments reduced by 1 percent over the fiscal year that runs from October 2014 through September 2015. To determine penalties, Medicare evaluated three types of HACs. One is central-line associated bloodstream infections, or CLABSIS. The second is catheter-associated urinary tract infections, or CAUTIS. The final one, Serious Complications, is based on eight types of injuries, including blood clots, bed sores and falls. Here are the hospitals that are being penalized:

### Why Consider Point of Use (POU) Filtration Today?

## Reimbursement

## Risk

## What is Point of Use Filtration?



## There are various types of POU filters & manufactures



## The Definition of Point of Use Filtration may differ





## Point-of-use filters may be

- Immediate and efficient barrier
- Documented validation
- Suited for cold and hot water
- Compatible with chemical & thermal disinfection
- Quick & easy to install
- Comfortable handling

## Can infection/colonisation rates be reduced?

## Ice Machines



## TRIB LIVE | News

**Opinion/The Review** News Investigative Neighborhoods State Politics U.S./World Sports Home Columnists Allegheny Butler Fayette Indiana Somerset Washington Armstrong Beaver Pittsburgh Eagle Cam Pennsylvania Contact Us Education Blog

## Legionella bacteria found in hospital ice machines at UPMC Presbyterian



Pittsburgh Tribune-Review

#### By Luis Fábregas and Adam Smeltz

Published: Friday, May 2, 2014, 12:01 a.m.

Legionella bacteria in ice machines at UPMC Presbyterian contributed to one patient's death and sickened two others, hospital officials disclosed on Thursday, calling it an unusual episode uncovered because a patient aspirated ice chips.

## Ice Machines Maintenance









## Ice Machine Statement from CDC

"Guidelines for Environmental Infection Control in Health-Care Facilities Recommendations of CDC and the Healthcare Infection Control Practices Advisory Committee (HICPAC)"

Ice Machines are a possible source of infection due to microorganism contamination

"Microorganisms may be present in ice, ice storage chests, and icemaking machines. The two main sources of microorganisms in ice are the potable water from which it is made and a transferral of organisms from hands. Ice from contaminated ice machines has been associated with patient colonization, blood stream infections, pulmonary and gastrointestinal illnesses, and pseudoinfections. Microorganisms in ice can secondarily contaminate clinical specimens and medical solutions that require cold temperatures for either transport or holding."

## Ice Machine Statement from CDC

"Guidelines for Environmental Infection Control in Health-Care Facilities Recommendations of CDC and the Healthcare Infection Control Practices Advisory Committee (HICPAC)"

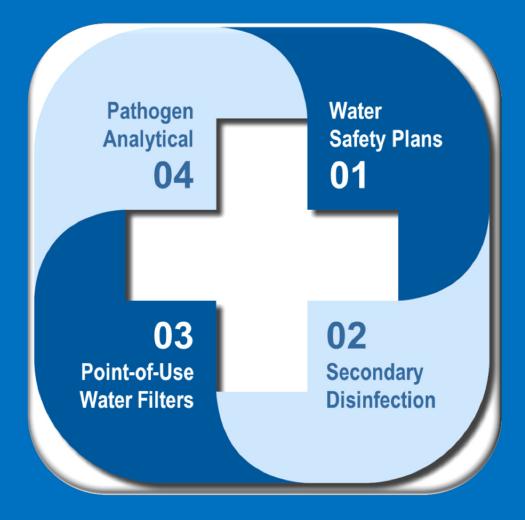
Microorganisms associated with ice machine contamination from potable water:

- Legionella spp.
- Nontuberculous mycobacteria (NTM)
- Pseudomonas aeruginosa
- Burkholderia cepacia
- Stenotrophomonas maltophilia
- Flavobacterium spp.

## Ice Machine Best Practices

- Manufacturers guidelines need to be adhered to. Cleaning and disinfection procedures and frequency.
- Filter incoming water with sediment filters (1-10 micron) and change-outs as per manufacturer recommendation. Be aware of bacteria bleed by of sediment filters and change accordingly.
- Do not use carbon filters on ice machines unless only required if disinfectant (chlorine, chlorine dioxide) odor/taste is objectionable
- Consider 0.2 micron absolute point-of-use filtration on high risk ice machines placed downstream of sediment/carbon filters.
- Use sterile or sterile grade filtered water with 0.2 micron for all water used to clean and disinfect machine components.
- Use sterile or sterile disposable basin for soaking components. Do not use a sink due to drain contamination.
- Avoid contamination of ice machine drains trays. This increases risk retrograde contamination on ice chute and water dispenser.

## Systematic Risk Management



## Water Safety Plans Expert Risk Management

» Pathogen Water Safety Plans
• Hazard Analysis
• Process Flow Diagrams
• Plan Design & Implementation
• Awareness Training
• Verification & Validation

ANALYZE ACT Cycle of Safety

ADIUS

» Web-based Data Monitoring & Management

» Consulting & Support

## Best Practices: Domestic Water Services

Strategy	Purpose	Reference
New Construction/ Renovation	Goal is to be aware of design features (cross connections, need for piping insulation, dead-legs, low flow zones, water hammer arrestors, etc.) or stagnant conditions that can increase risk if not properly managed.	ASHRAE 12, 188P OSHA
New Systems, Startup/Shutdown	Goal is to define practice to manage the water system to limit stagnation, implement practices to flush systems after lengthy shutdown or interruption of water service, and requirements for clean and disinfection before commissioning new systems.	ASHRAE 12, 188P
System Maintenance	Goal is to define practice (Clean and disinfect, flushing, repair, etc.) for system maintenance of hot and cold water tanks, ice machines, water filters, shower heads and hoses, faucets, etc.	ASHRAE 12, 188P
Water Temperature	<ul> <li>Water temperature recommendations for legionellae control are:</li> <li>Maintain water heater outlet temperatures at or above 140°F (60°C);</li> <li>Maintain the hot water temperature at coldest point in the water heater, the storage tank, or the distribution system at or above 124°F (51°C);</li> <li>Maintain the cold water temperature in any part of system at or below 77°F (25°C).</li> </ul>	ASHRAE 12, 188P OSHA
Water Disinfection	Where water disinfection or treatment is performed, a defined program must be followed to assure it meets EPA requirements for potable water applications.	ASHRAE 12, 188P OSHA
Emergency Disinfection	Goal is to define practice to be followed if there are suspected legionellosis health problems associated with the use of potable water in a building system.	ASHRAE 12, 188P OSHA
Legionella Monitoring	Recommended to verify control of the hazard. * <i>Typically recommended for investigative or post remedial verification purposes.</i>	ASHRAE 12 OSHA*

## **Best Practices:** Cooling Tower

Strategy	Purpose	Reference
System Operation	Goal is to operate in a manner that keeps the system treated and limits stagnant conditions. Startup/Shutdown; Intermittent operation; New system startup	ASHRAE 12, 188P CTI
Inspection & Maintenance	Goal is to maintain mechanical design intent to limit aerosol release, to maintain balanced water flows and to eliminate dead zones.	ASHRAE 12, 188P CTI OSHA
Design & Siting	Be aware of design features (sumps, drift eliminators, location of tower) that can increase risk if not properly managed.	ASHRAE 12, 188P CTI OSHA
Scale & Corrosion Control	A comprehensive scale and corrosion program is necessary to limit scale and corrosion formation to within specified critical limits.	ASHRAE 12, 188P CTI
Biocide Control	A comprehensive biocide program applied to within critical limits is necessary to maintain microbial control. Biocides must be applied in a manner that demonstrates control.	ASHRAE 12, 188P CTI OSHA
Clean & Disinfect (C&D)	Goal is to prevent accumulation of slimes and sludge which can allow microbial proliferation and increase <i>Legionella</i> risk. Twice annual C&D Off-line and On-line	ASHRAE 12, 188P CTI OSHA
Legionella Monitoring	Recommended to verify control of the hazard. * <i>Typically recommended for investigative or post remedial verification purposes.</i>	ASHRAE 12 CTI OSHA*
Aerobic Bacteria Monitoring	Monitoring is essential to verify biocide program is sufficient to control microbial growth.	СТІ

## ASHRAE 188p

#### 4.3. Health Care Facility Requirements

**4.3.1** Health care facilities that do not meet all of the qualifications of 4.3.2 shall comply with the requirements in Section 4.2, Section 6, and Section 7.

**4.3.2** Health care facilities that meet all of the following qualifications shall comply with either the requirements in Sections 4.2, 6 and 7 or the requirements in normative Appendix A "Health Care Facilities":

- a. The health care facility is accredited by a regional, national or international accrediting agency or by the Authority Having Jurisdiction (AHJ) over the health care facility Infection Prevention and Control (IC) activities; and
- b. The health care facility Infection Control program has an Infection Preventionist that is Certified in Infection Control (CIC) by the Certification Board of Infection Control and Epidemiology (CBIC) or other regional, national or international certifying body or the health care facility has an Epidemiologist with a minimum of a Master's degree or equivalent.

PROGRAM TEAM – Identify persons responsible for Program development and implementation.

DESCRIBE WATER SYSTEMS/FLOW DIAGRAMS – Describe the potable and non-potable water systems within the building and on the building site and develop water system schematics.

ANALYSIS OF BUILDING WATER SYSTEMS – Evaluate where hazardous conditions may occur in the water systems and determine where control measures can be applied.

CONTROL MEASURES – Determine locations where control measures must be applied and maintained in order to stay within established control limits.

MONITORING – Establish procedures for monitoring whether control measures are operating within established limits and if not, take corrective actions

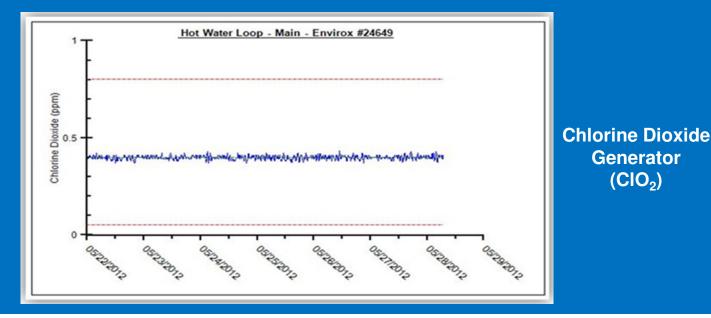
CORRECTIVE ACTIONS/CONFIRMATION – Establish procedures to confirm that:

- The Program is being implemented as designed. (verification)
- The Program effectively controls the hazardous conditions throughout the building water systems (validation)

DOCUMENTATION – Establish documentation and communication procedures for all activities of the Program.

## **Secondary Disinfection A Continuous Treatment Strategy**

- » Evaluation of alternatives: Pros & Cons
- » Hot and cold potable water treatment
- » NSF-61 certified equipment
- » NSF-60 certified chemistry
- » Water treated per EPA regulations
- » 360 24/7 monitoring of disinfectant residuals
- » 360 automation of system alarms





 $(CIO_2)$ 

## **Summary of Secondary Disinfection Choices**

	Chlorine Dioxide	Chloramine	Chlorine	Copper- Silver	Ozone	UV-Light	Thermal Disinfect
Effective against legionellae	YES	YES	YES	YES	YES	YES	YES
Effective against most bacteria	YES	YES	YES	YES	YES	YES	NO
Effective against biofilm	YES	YES	YES	NO	YES	NO	NO
No Legionella resistance	YES	NO	NO	NO	YES	YES	YES
Protects whole system	YES	YES	YES	YES	YES	NO	NO
Not affected by pH	YES	NO	NO	NO	YES	YES	YES
Not affected by water hardness	YES	YES	YES	NO	YES	YES	YES
Easy to monitor	YES	YES	YES	NO	NO	NO	YES
Low corrosion rates	YES	YES / NO	NO	NO	NO	YES	YES
No Trihalometanes (THM's)	YES	NO	NO	YES	NO	YES	YES
Low disinfection by-products (DBP)	YES	YES	NO	YES	NO	YES	YES

## Point of Use (POU) Filters A Point Control Strategy

- » An absolute barrier for waterborne pathogens
- » "Sterilizing Grade Filtered Water"
- » For high risk patient areas (BMT, ICU, NICU, BURN, ONCOLOGY, ETC.)
- » For immediate response to an outbreak or incident





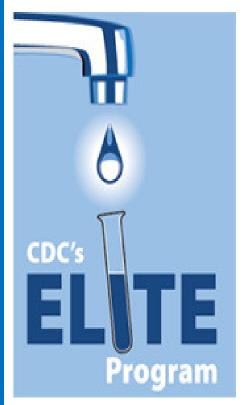




## Pathogen Analytical Validation of the Control Strategy

# » A Certified CDC-ELITE Proficient Lab » Legionella Culture Test per ISO 11731 » Interpretation & Consulting » Testing Plans





## PLAN REQUIREMENT: EMERGENCY REMEDIATION DOMESTIC WATER SYSTEMS

If a possible HCA infection case is identified and if *waterborne pathogens* are detected in building's water distribution system, an Action Plan must be in place.

Strategy	Purpose
Thermal Eradication	Increasing temperature in the hot water distribution system of 160°F – 170°F while continuously flushing each outlet in the system for at least 30 minutes
Shock Chlorination	Increasing chlorine level of the hot and cold water distribution systems to 2 mg/L and maintaining that level throughout the system at least 2 hours
High Level Disinfection	Increasing chlorine level of the hot and cold water distribution systems to 50 mg/L and maintaining at least 10 mg/L throughout the system and at outlets for 24 hours

## **Remediation Strategies**

### » Cleaning & Disinfection Services

- Cooling Water Systems
- Potable Hot & Cold Water Systems
- Therapy Pools
- Decorative Fountains
- Ice Machines





BEFORE



DURING



AFTER

## Water as a Source of Hospital Acquired Infections: Summary

- Potable water is not sterile
- Healthcare facilities inherently amplify waterborne pathogens
- A Water Safety Plan should be implemented
- Systemic solutions can be added to provide overall/general risk reduction
- Critical care units need additional interventions to improve point of use water quality to the immunocompromised population
- Publications are available by critical care unit & by pathogen to show how sterilizing grade filtered water can help reduce risk of HAI.

# Questions??

#### Michael Castro

- 480-636-0405
- michael\_castro@pall.com

#### Kevin Olson

- 510-928-9097
- kolson@nalco.com

#### References

- Cunliffe D et al., "Water Safety in Buildings", WHO Press, World Health Organization, Geneva/Switzerland, March, 2011
- 2. Exner M et al., "Prevention and control of health care-associated waterborne infections in health care facilities", AJIC, 33:S26-S40, 2005
- 3. Lindsay D & von Holy A, "Bacterial biofilm within the clinical setting: What healthcare professionals should know", J Hosp Infect, 64:313-325, 2006
- 4. Kreysig D, "Der Biofilm Bildung, Eigenschaften und Wirkungen", published in Bioforum, GIT Verlag, Darmstadt/Germany 24:40-43, 2001
- 5. Flemming HC & Wingender J. "The biofilm matrix", Nat Rev Microbiol, 8:623-633, 2010
- 6. Moritz M *et al.*, "Integration of *Pseudomonas aeruginosa* and *Legionella* pneumophila in drinking water biofilms grown on domestic plumbing materials", Int J Hyg Env Health, 213:190-197, 2010
- 7. Anaissie EJ, et al., "The hospital water supply as a source of nosocomial infections: a plea for action", Arch Intern Med, 162:1483-1492, 2002
- 8. Wunderink RG & Mendoza DL, " Epidemiology of *Pseudomonas aeruginosa* in the intensive care unit", published in "Infectious diseases in critical care", Springer Verlag, 218-225, 2007
- 9. Trautmann M *et al.*, "Common RAPD pattern of *Pseudomonas aeruginosa* from patients and tap water in medical intensive care unit", Int J Hyg Env Health, 209:325-331, 2006 10. Oliver JD "Recent findings on the viable but non-culturable state in pathogenic bacteria", FEMS Microbiol Rev, 34:415-425, 2010
- 11. Dwidjosiswojo Z *et al.*, "Influence of copper ions on the viability and cytotoxicity of *Pseudomonas aeruginosa* under conditions relevant to drinking water environments", Int J Hyg Env Health, in press, 2011
- 12. Allegra S *et al.*, "Longitudinal evaluation of the efficacy of heat treatment procedures against *Legionella* spp. In hospital water systems by using a flow cytometric assay", Appl Env Microbiol, 77:1268-1275, 2011.
- 13. Drancourt M, Adékambi T & Raoult D, "Interactions between Mycobacterium xenopi, amoeba and human cells", J Hosp Infect, 65:138-142, 2007
- 14. Winiecka-Krusnell J & Linder E, "Free living amoeba protecting Legionella in water: The tip of an iceberg?", Scand J Infect Dis, 31:383-385, 1999
- 15. Reuter S et al., "Analysis of transmission pathways of Pseudomonas aeruginosa between patients and tap water outlets", Crit Care Med, 10:2222-2228, 2002
- 16. Blanc DS et al., "Faucets as a reservoir of endemic Pseudomonas aeruginosa colonization/infections in intensive care units", Intensive Care Med, 30: 1964-1968, 2004
- 17. Vallés J *et al.*, "Patterns of colonization by *Pseudomonas aeruginosa* in intubated patients: a 3-year prospective study of 1.607 isolates using pulsed-field gel electrophoresis with implications for prevention of ventilator-associated pneumonia", Intensive Care Med, 30:1768-1775, 2004
- 18. Jung R et al., "Surveillance of multi-drug resistant Pseudomonas aeruginosa in an urban tertiary-care teaching hospital", J Hosp Infect, 57:105-111, 2004
- 19. Rogues AM et al., "Contribution of tap water to patient colonisation with Pseudomonas aeruginosa in a medical intensive care unit", J Hosp Infect, 67:72-78, 2007
- 20. Eckmanns T *et al.*, "An outbreak of hospital-acquired *Pseudomonas aeruginosa* infections caused by contaminated bottled water in intensive care units", Clin Microbiol Infect, 14:454-458, 2008
- 21. Wong V et al., "Spread of Pseudomonas fluorescens due to contaminated drinking water in a bone marrow transplant unit", J Clin Microbiol, 49:2093-2096, 2011
- 22. Blanc D et al., "Water disinfection with ozone, copper and silver ions, and temperature increase to control Legionella: seven years of experience in a university teaching hospital", J Hosp Infect 60:69-72, 2005
- 23. Patterson WJ et al., "Colonization of transplant unit water supplies with Legionella and protozoa: precautions required to reduce the risk of legionellosis", J Hosp Infect 37:7-17, 1997
- 24. Eckmanns T et al., "Prevention of nosocomial Legionnaire's disease", Dtsch Ärztebl, 103:1294-1300, 2006
- 25. Lin E, Stout J, Yu V., "Controlling *Legionella* in hospital drinking water: an evidence-based review of disinfection methods", Infect Control Hosp Epidemiol, 32:166-173, 2011 26. Vianelli N *et al.*, "Resolution of a *Pseudomonas aeruginosa* outbreak in a haematology unit with the use of disposable sterile water filters", Haematologica, 91:983-985, 2006 27. Van der Mee-Marquet N *et al.*, "Water Microfiltration: A procedure to prevent *Pseudomonas aeruginosa* infection", XVIe Congrès National de la Société Francaise d'Hygiène Hospitalière, Reims, Livre des Résumés, S137, June 4th 2005
- 28. Hall J et al., "Provision of safe potable water for immunocompromised patients in hospital", J Hosp Infect, 58:155-158, 2004
- 29. Trautmann M *et al.*, "Point-of-use water filtration reduces endemic *Pseudomonas aeruginosa* infections on a surgical intensive care unit", Am J Infect Control, 36:421-429, 2008
- 30. Legrand ICS *et al.*, "Impact of terminal water microfiltration on the reduction of nosocomial infections due to *Pseudomonas aeruginosa* in burnt patients", presented at 11ème Journées Internationales de la Qualité Hospitalière et en Santé (JIQHS), Paris, poster n°204, 2009
- 31. Wright CL *et al.*, "A two year double cross-over study investigating point-of-use filters for reducing Gram-negative nosocomial pathogens from hospital water", J Hosp Infect 76(Suppl. 1):S38, 2010
- 32. Hell M *et al.*, "Water microfiltration at the point of use a procedure to prevent infection/colonization with water borne pathogens in ICU patients?", J Hosp Infect, 76(Suppl. 1):S38, 2010

#### Continued from previous page

33. Rolling AM, "Handling of antimicrobial water filter and disposable shower rose used for wound cleansing", 19th Conf European Wound Management Assoc, Helsinki, Finland, (P27), May 2009

34. Barna Z et al., "Tap water as a potential source of nosocomial *Pseudomonas aeruginosa* infections in an intensive care unit", 19th European Congress Clin Microbiol & Infect Dis, Helsinki, Finland, Poster No<sup>o</sup> 859, May 2009

35. Warris A et al., "Point-of-use filtration method for the prevention of fungal contamination of hospital water", J Hosp Infect, 76:56-59, 2010.

36. Junker L *et al.*, "Nosocomial infections with *Legionella* spp. and other water borne bacteria can be reduced by control of shower water", Int J Infect Contr, 5:S60 (P17), 2009 37. Williams MM *et al.*, "Point-of-use membrane filtration and hyperchlorination to prevent patient exposure to rapidly growing mycobacteria in the potable water supply of a skilled nursing facility", Infect Contr Hosp Epidemiol, 32:837-844, 2011

38. Wang SH *et al.*, "Pseudo-outbreak of "Mycobacterium paraffinicum" infection and/or colonization in a tertiary care medical center", Infect Contr Hosp Epidemiol, 30:848-853, 2009

39. Harpel S et al., "Performance of a new 14 day water filter during daily use in clinical routine at two university medical centers", J Hosp Infect, 64 (Suppl.1):S47, 2006

40. Sheffer PJ *et al.*, "Efficacy of new point-of-use water filter for preventing exposure to *Legionella* and waterborne bacteria", Am J Infect Control, 33:S20-S25, 2005

41. Warris A et al., "Point-of-use filtration method for the prevention of fungal contamination of hospital water", J Hosp Infect, 76:56-59, 2010.

42. Cervia J et al., "Point-of-use water filtration reduces healthcare-associated infections in bone marrow transplant recipients", Transpl Infect Dis, 12:238-241, 2010

43. Holmes C et al., "Preventive efficacy and cost-effectiveness of point-of-use water filtration in a subacute care unit", Am J Infect Control, 38:69-71, 2010

44. Williams MM *et al.*, "Point-of-use membrane filtration and hyperchlorination to prevent patient exposure to rapidly growing *mycobacteria* in the potable water supply of a skilled nursing facility", Infect Contr Hosp Epidemiol, 32:837-844, 2011

45. Hankwitz PE *et al.*, "Non-tuberculous mycobacterial hypersensitivity pneumonitis related to a home shower: treatment and secondary prevention", BMJ Case Reports 2011 46. Zhou ZJ *et al.*, "Removal of waterborne pathogens from liver transplant unit water taps in prevention of healthcare-associated infections: a proposal for a cost-effective, proactive infection control strategy", Clin Microbial Infect, June 20, 2013

47. Teare L & Millership S, "Legionella pneumophila serogroup 1 in a birthing pool", J Hosp Infect, 82:58-60, 2012

48. American Standard Test Method (ASTM) F838-05 "Determining Bacterial Retention of Membrane Filters Utilised for Liquid Filtration", 2005

49. Circulaire DGS/SD7A/SD5C-DHOS/E4 n° 2002/243 du 22/04/2002 relative à la prévention du risque ilé aux legionelles dans les établissements de santé, April 22nd, 2002 50. Empfehlung der Kommission für Krankenhaushygiene und Infektionsprävention beim Robert Koch-Institut (RKI) "Anforderung an die Hygiene bei der Aufbereitung flexibler Endoskope und endoskopischen Zusatzinstrumentariums", Bundesgesundheitsbl-Gesundheitsforsch-Gesundheitsschutz, 45:395-411, 2002

51. Empfehlung der Kommission für Krankenhaushygiene und Infektionsprävention beim Robert Koch-Institut (RKI) "Anforderungen an die Hygiene bei der medizinischen Versorgung von immunsupprimierten Patienten", Bundesgesundheitsbl-Gesundheitsforsch-Gesundheitsschutz, 53:357-388, 2010

52. Yorkshire Cancer Network, "Provision of safe drinking water for cancer patients with immunocompromised", www.yorkshire-cancer-net.org.uk, 2005

53. UK Department of Health, "Water sources and potential for *Pseudomonas aeruginosa* infection from taps and water systems; advice for augmented care units. Department of Health Gateway Ref 17334, 2012

54. Legionella and the prevention of Legionellosis", WHO Press, World Health Organization, Geneva/Switzerland, Editors: J. Bartram, Y. Chartier, JV Lee, K. Pond & S. Surman-Lee, 2007

55. Public Health Agency Canada, "Infection Prevention and Control Guideline for Flexible Gastrointestinal Endoscopy and Flexible Bronchoscopy", www.publichealth.gc.ca, 2011

56. Gastroenterological Society in Australia (GESA), "Infection Control in Endoscopy", Edited by A Taylor et al., 3rd Edition, Digestive Health Foundation, 2010

57. Eagye KJ et al., "Impact of superinfection on hospital length of stay and costs in patients with ventilator-associated pneumonia", Semin Resp Crit Care Med, 30:116-123, 2009

58. Murphy D et al., "Dispelling the myths: The true cost of health care associated infections", APIC Briefing, Febr. 2007, www.apic.org

59. Warren KD *et al.*, "Attributable cost of catheter-associated bloodstream infections among intensive care patients in a nonteaching hospital", Crit Care Med, 34:2084-2089, 2006

60. Kilgore ML et al., "The costs of nosocomial infections", Med Care, 4:101-104, 2008

61. Tu HZ et al., "Use of disposable water filter for prevention of false-positive results due to Nontuberculosis Mycobacteria in a clinical laboratory performing routine acid-fast staining for tuberculosis", Appl Env Microbiol, 73:6296-6298, 2007