

# Microbe of the month

Breaking The Chain of Infection



MAY 2023 NEWSLETTER

Compiled by  
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Featured  
this  
month:

## PSEUDOMONAS AERUGINOSA

World Hand Hygiene Day 5<sup>th</sup> May 2023

15-minute read + QUIZ

Hello readers!

**Microbe of the Month** aims to provide a concise clinical resource, to help you keep up to date about pathogens of importance, in an easy-to-read and understand format. Each issue covers the aetiology (sources) and epidemiology of topical bacteria, viruses, or fungi - their mode/s of transmission and the infections they cause; alerts on any antimicrobial resistance (AMR) capability they may have, and the relevant Infection Prevention and Control measures which should be routinely implemented for the safety of patients and healthcare personnel.

*There is a quick quiz at the end of the newsletter to test your grasp of the content – please use this newsletter as a teaching tool in your workplace and start an ‘infectious dialogue’ about topical issues in infection control!*

With **World Hand Hygiene Awareness Day** on 5<sup>th</sup> May, this month’s focus is on **contact transmission** and the well-known Gram-negative bacterium ***Pseudomonas aeruginosa***.

This microorganism is found everywhere in the environment, and is considered to be the kingpin of ‘**opportunistic pathogens**’, which means that it exploits breaks in immune defences to cause infection. **It is among the ‘top four’ most commonly-isolated nosocomial bacteria** implicated in a variety of healthcare-associated infections (HAIs), such as septicaemia, bronchopneumonia, necrotising enterocolitis, and bone, joint, urinary and wound infections; especially in critically ill and immunocompromised patients; e.g., premature neonates, diabetics, ICU, cystic fibrosis and burn patients, and those receiving corticosteroids or cytotoxic chemotherapy.<sup>1,2</sup>

****Pseudomonas aeruginosa* (*P. aeruginosa*) also readily develops resistance mechanisms to antiseptics and multiple classes of antibiotics.***<sup>1-3</sup>

**Key words:** reservoir, mode of transmission, biofilm, immune-compromised, opportunistic pathogen, healthcare associated infections (HAIs), pandrug-resistant (PDR), infection prevention and control (IPC).



## RESERVOIRS AND MODES OF TRANSMISSION <sup>2-5</sup>

Within the healthcare setting, there are numerous reservoirs which provide ideal conditions for the growth and transmission of *Pseudomonas aeruginosa* (*P. aeruginosa*). It is also constantly reintroduced into the hospital environment on plants, fruit and vegetables, as well as by visitors and patients transferred from other facilities.

**This is why fresh flowers and pot plants are prohibited in critical care units, as well as salads, non-peelable fruit, vegetables and ice machines in haematology and transplant units.**

Cross-infection occurs from patient to patient via the unwashed hands of hospital personnel ('direct contact'), via 'indirect contact' with contaminated surfaces and reservoirs, or by the ingestion of contaminated foods and water.

### Common reservoirs where you will find *P. aeruginosa*

- ✓ Water sources – taps, shower heads, hand basins, hydrotherapy baths, mops and cleaning equipment.
- ✓ Plastic basins and receptacles (stored wet).
- ✓ Antiseptics and disinfectants decanted or 'topped up' from bulk containers.
- ✓ Intravenous solutions contaminated by repeated access or used as diluent for multiple patients.
- ✓ Humidifiers in ventilators and incubators.
- ✓ Haemodialysis effluent points.
- ✓ Reusable suction bottles.
- ✓ Chronic wounds, especially those with high exudate levels (e.g., deep burn injuries, and pressure and venous stasis ulcers).



## PATHOGENESIS AND VIRULENCE OF *Pseudomonas aeruginosa* <sup>3,6</sup>

Its nutritional needs are simple, and it is tolerant of a wide variety of physical conditions, including ambient temperature. Although viable at temperatures ranging from 4°C - 42°C, the optimum temperature for the growth of *P. aeruginosa* is 37°C (body temperature). *P. aeruginosa* is also a 'facultative anaerobe', which means it can adapt to, and proliferate in, conditions of partial or total oxygen depletion.

**The first step in the infectious process is adherence of the microbe to host tissue, which enhances the capacity of the organism to produce disease.**

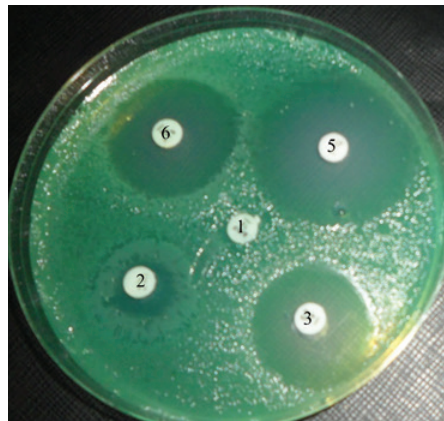
- The ability of *P. aeruginosa* to invade tissue is derived from the production of extracellular enzymes and cytotoxins which break down physical barriers and damage host cells, while its **bacterial capsule** or **slime layer** effectively protects it from the host's immune defences, such as antibodies or phagocytosis by leukocytes.

- Another important virulence strategy used by *P. aeruginosa* is the production of **biofilm**, which facilitates the attachment of these bacteria to plastic surfaces, medical devices and wounds. The 3-dimensional structure of biofilm provides protection from the immune response, antimicrobial agents (e.g., antibiotics) and adverse environmental conditions.
- ‘**Quorum sensing**’ is a revolutionary discovery, whereby bacteria use a complex series of chemical signals to communicate with each other to coordinate the formation of biofilm, invasive infective processes, their motility, and density in comparison to other species! *Pseudomonas* species in particular are known to use quorum sensing to gain a competitive advantage over other microbial species.<sup>6</sup>

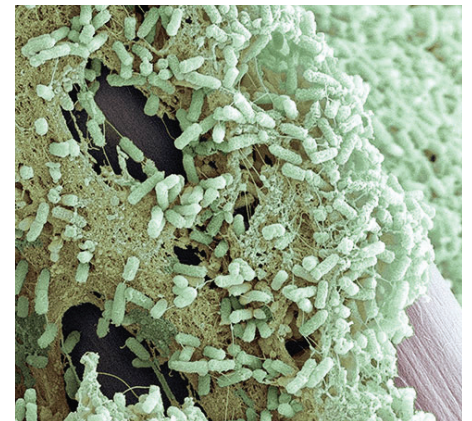
*Pseudomonas* is a large and diverse species... ‘*aeruginosa*’ is derived from the Latin word meaning ‘copper rust’, describing the characteristic blue-green colour of *P. aeruginosa* colonies, and sickly-sweet odour from the production of the metabolite ‘pyocyanin’. ‘Pyo’ is also a reference to pus.



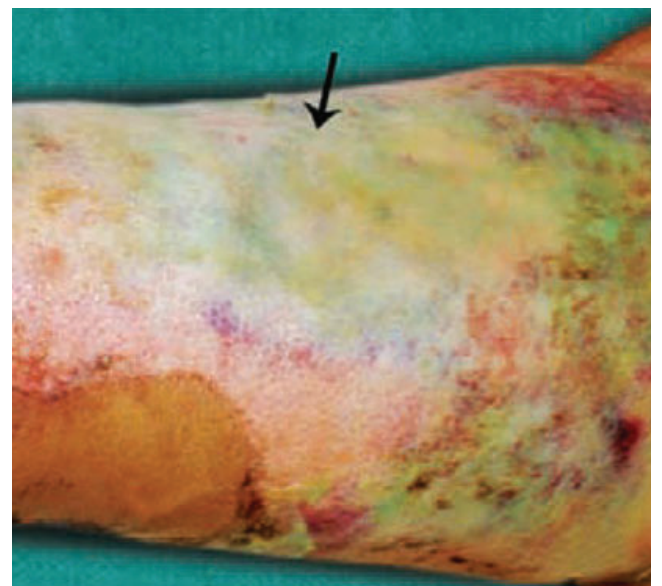
Scanning electron microscopy image of *P. aeruginosa*’s ‘whip-like’ flagellae which aid its attachment, motility and escape from phagocytic neutrophils.



Laboratory culture plate illustrating the characteristic slimy blue-green colonies of *Pseudomonas aeruginosa*.



Scanning electron microscopy image of *P. aeruginosa* biofilm on the fibres of a gauze dressing.



**Chronic and heavily exuding wounds (such as venous stasis ulcers and full thickness burns) are prone to infection with *Pseudomonas aeruginosa*.**

**Note the characteristic blue-green exudate (caused by the pigment pyocyanin).**



### A significant outbreak investigation finding

A spot check on the sluice room in the ICU revealed that the plastic bathing basins were not correctly stored or inverted after cleaning to promote air-drying; i.e., they were stacked inside one another and were therefore still 'wet' when reused.

### Infection prevention and control actions implemented

Stainless steel basins were purchased and clearly marked with bed numbers. Twice as many basins available vs the total number of ICU beds enabled the storage of a basin at its designated bed, whilst the spare basins were stored (inverted) on a designated stainless-steel rack in the sluice room. The use of plastic basins was immediately discontinued.



## DID YOU KNOW?

**Mupirocin** (Pseudomonic acid A) is an antibiotic produced by *Pseudomonas fluorescens*, which has a high level of activity against staphylococci and streptococci bacteria.

**Mupirocin** is used as an intranasal antibiotic to treat patients and healthcare workers who are colonised with methicillin-resistant *Staphylococcus aureus* (MRSA).<sup>7</sup>

## THE BIOFILM-BASED APPROACH TO WOUND INFECTIONS CAUSED BY *P. aeruginosa*<sup>8,9</sup>



***Pseudomonas* species thrive in moisture-rich environments; therefore, venous stasis ulcers, stage 2-4 pressure injuries and deep burn wounds are vulnerable to infection if slough and exudate levels are not actively managed.**

**Polymicrobial** (multiple species) **biofilms are present in chronic wounds** – microscopic structures which cannot be seen with the naked eye. With advances in molecular technology, however, it is now widely recognised that biofilm structures have a negative impact on healing, so the active management of biofilms should be integral to any wound infection treatment plan.

### Clinical signs of biofilm activity:

- ✓ Delayed healing despite optimal wound management and support of underlying patient comorbidities
- ✓ Failure or recalcitrance to appropriate topical and/or systemic antimicrobial treatment
- ✓ Increased wound exudate/moisture
- ✓ Low-level chronic inflammation
- ✓ Low-level erythema
- ✓ Poor granulation or friable, bleeding hypergranulation
- ✓ Covert signs of infection

The growth and multiplication of *P. aeruginosa* is likely to induce an alkaline wound pH and promote biofilm formation – especially in the presence of high exudate levels and occlusive dressings. **The pH of the wound directly influences all biochemical reactions which take place during the process of healing.** Research has proven that a slightly acidic pH helps to rebalance wound flora, increases antimicrobial activity, oxygen release, protease activity and angiogenesis. <sup>8,9</sup>

### Biofilm-busting and pH reducing strategies <sup>8,9</sup>

- 1. Debridement** converts a chronic or ‘stalled’ wound to an acute wound by removing non-viable tissue which stimulates inflammation and microbial growth. It reduces the ‘barrier to healing’ caused by biofilm and a high bioburden, and helps the wound to progress out of a prolonged inflammatory phase and into the **proliferative phase**. Senescent cells (which have stopped dividing and are non-responsive to growth factors and cytokines) are also removed, making way for healthy, responsive cells.
- 2. Polyhexamethylene biguanide** (a.k.a. PHMB) combined with a surfactant detergent cleanser is a non-cytotoxic antiseptic which disrupts lipoproteins in biofilm, interferes with chemical signalling (‘quorum sensing’) and slows biofilm regrowth.
- 3. Acetic acid 1% solution** (the active component in vinegar) is widely used as a cleanser and wound ‘soak’ to reduce wound pH and eradicate *P. aeruginosa*.
- 4. Hypochlorous acid (HOCl)** exerts a microbicidal effect against a wide spectrum of bacterial, viral and fungal pathogens. Its action mimics the impact of normal neutrophil activity in the wound whereby hydrogen peroxide is produced and converted to HOCl in the presence of chlorine and hydrogen ions in the exudate.
- 5. Dressings** should be selected for their proven fluid handling capacity (even under compression), and ideally, be able to sequester (contain or isolate) microorganisms and harmful proteases away from the wound bed, to hinder the cyclical inflammatory process which delays healing.



During March this year, the US Centers for Disease Control and Prevention (CDC) identified a multistate outbreak of extensively drug-resistant (XDR) *P. aeruginosa* eye infections associated with ‘artificial tears’ in multi-use eye drop bottles.

This is the first time the carbapenem-resistant outbreak strain (i.e., Verona integron-mediated [VIM] metallo- $\beta$ -lactamase and Guiana extended-spectrum- $\beta$ -lactamase [VIM-GES-CRPA]), has been reported. <sup>10</sup>



### ANTIMICROBIAL STEWARDSHIP (AMS)

The emergence of MDR/XDR bacteria has led to the revival of older antibiotics, with colistin being a good example. In pandrug-resistant (PDR) *P. aeruginosa* infections, colistin is often used in synergistic combinations with other antibiotics (i.e., their combined effect is greater than the sum of the effects seen when each drug is given alone).

**Pandrug-resistance (PDR) is defined as non-susceptibility to all agents in all antimicrobial categories** (i.e., the bacterial isolates are not susceptible to any clinically available drug). The prevalence of PDR bacteria is difficult to judge as isolates are rarely assessed against all possible antibiotics; however, PDR Gram-negative bacteria have been reported in more than 20 countries across all continents of the world. The mortality rate in patients infected with these bacteria is considered to be at least 50-70%. <sup>1,11</sup>

**Liaison with a medical microbiologist is recommended when prescribing antimicrobial agents for drug-resistant isolates.**

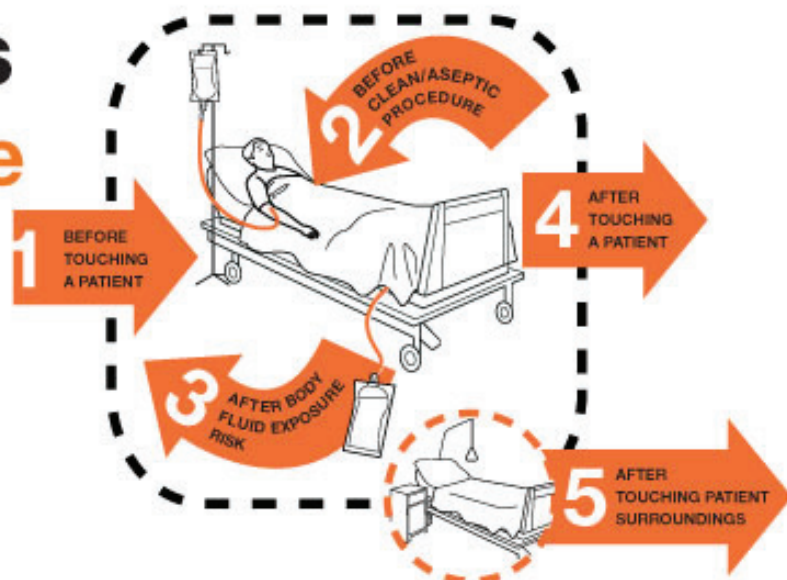


# THE BOTTOM LINE...

- ✓ **Standard Precautions** apply to all patient care, regardless of the patient's suspected or confirmed infectious state, and apply to all settings where care is delivered (these practices protect healthcare personnel and prevent healthcare personnel or the environment from transmitting infections to other patients). <sup>4,5</sup>
- ✓ **Hand hygiene is the single most important measure to prevent the spread of ALL types of microbes.**
  - > Jewellery and watches get in the way of performing hand hygiene correctly.
  - > Wearing rings also increases the carriage rate of Gram-negative bacteria on the hands of HCWs.
  - > HCWs who wear nail polish, acrylic and/or gel nails are more likely to harbour Gram-negative pathogens such as *P. aeruginosa* on their fingertips, both before and after hand-washing.
  - > An alcohol-based (60-80%) hand sanitiser is the preferred method for sanitising your hands when they are not visibly dirty. <sup>4,5,12</sup>
- ✓ **Glove use is NOT a substitute for hand hygiene** – dirty gloves contaminate your hands when they are removed. Always wash or sanitise your hands after removing gloves. <sup>4,5,12,13</sup>
- ✓ Practice **strict aseptic technique** for invasive procedures such as vascular and urinary catheterisation, injections and intravenous therapy, and wound care. <sup>4,5,10,12</sup>
- ✓ **Prioritise contact transmission prevention measures** with ongoing orientation, training and constant supervision of hand hygiene, the correct donning and doffing of PPE, environmental cleaning, and used linen and waste management. <sup>4,5,12,13</sup>
- ✓ Actively manage heavily exuding wounds with biofilm-active wound cleansers and antimicrobial (not antibiotic) dressings that have good fluid handling capacity, and change dressings timeously. <sup>8,9</sup>
- ✓ Detergent-based sodium hypochlorite disinfectants with proven efficacy against Gram-negative pathogens should be used for the routine cleaning of non-critical care equipment and environmental surfaces; and 'high touch' points should be disinfected at least twice daily. <sup>4,5</sup>

# Your 5 Moments for Hand Hygiene

- 1 BEFORE TOUCHING A PATIENT
- 2 BEFORE CLEAN / ASEPTIC PROCEDURE
- 3 AFTER BODY FLUID EXPOSURE RISK
- 4 AFTER TOUCHING A PATIENT
- 5 AFTER TOUCHING PATIENT SURROUNDINGS



## Supply the correct answer!

1. The term used to describe pathogens which exploit breaks in patients' immune defences is \_\_\_\_\_.
2. Common reservoirs which support the growth of *Pseudomonas aeruginosa* are those where \_\_\_\_\_ is present.
3. Biofilm enhances the ability of pathogens to \_\_\_\_\_ to host tissue and cause disease.
4. Chronic and complicated wounds with high levels of \_\_\_\_\_ are prone to infection with *P. aeruginosa*.
5. 'Bare below the \_\_\_\_\_' is an important infection prevention concept to facilitate effective hand hygiene.

ANSWERS: 1. Opportunistic 2. Moisture 3. Adhere 4. Exudate 5. Elbows



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