

Wisconsin HAI Long-Term Care Education Series

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WISCONSIN DEPARTMENT
of HEALTH SERVICES



Wound infection prevention

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Objectives

1. Describe the four steps of wound hygiene and how they contribute to infection prevention
2. Compare treatment recommendations for wounds with suspected infection
3. List sequelae of wound infection
4. Review practical tips for infection prevention in the administration and delivery of care

Long term care stats – in context

- 65,600 regulated long-term care facilities (LTCF) in the US
- ~70% of people turning 65 are expected to need long-term care at some point in their life
- 18% of the older persons will spend over a year in a nursing facility
- 58% of adults receive nursing home care after age 50

Uneven distribution of risk

However, the lifetime risk of receiving paid care is not evenly distributed across the population.

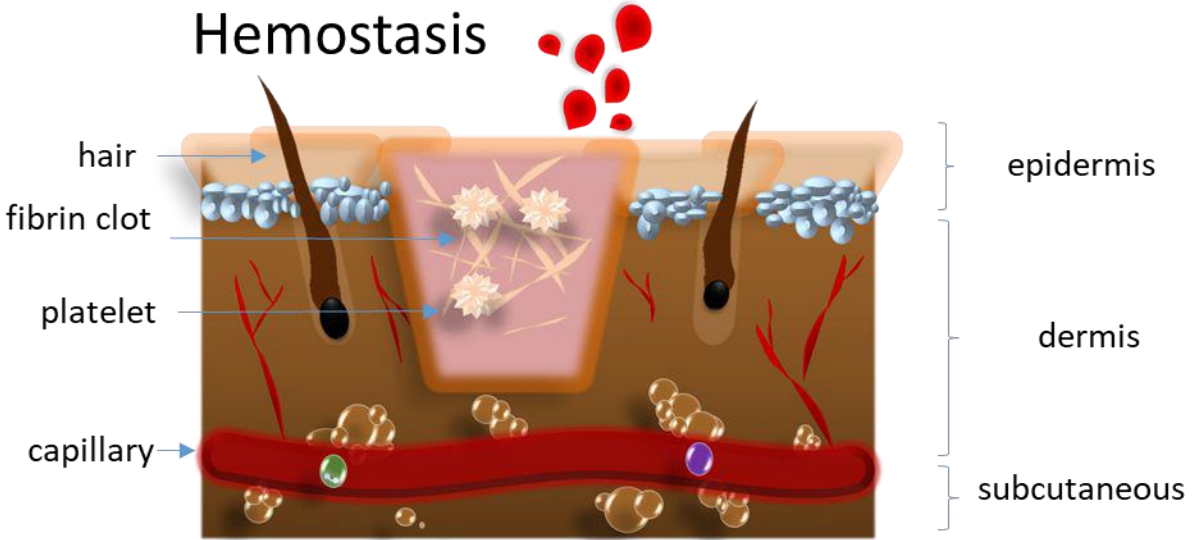
Lengthy spells of severe LTSS needs and paid care are much more common among older adults with few financial resources than their wealthier counterparts.

Infection prevention and control in LTC

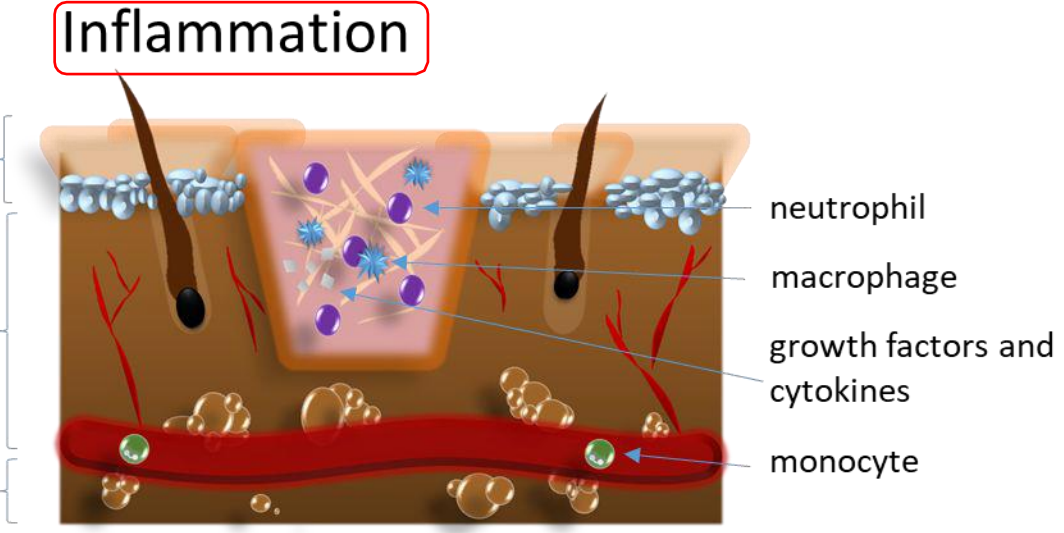
- Healthcare associated infections account for as many as 380,000 deaths annually
 - Acute care facilities estimates are 1.7 million infections and 99,000 associated deaths each year
- Infection prevention and control (IPC) guidelines are well-defined in the acute care setting, evidence of effectiveness for long-term care facilities (LTCF) is missing
- Both residents AND staff have increased risk of infection

Stages of Wound Healing

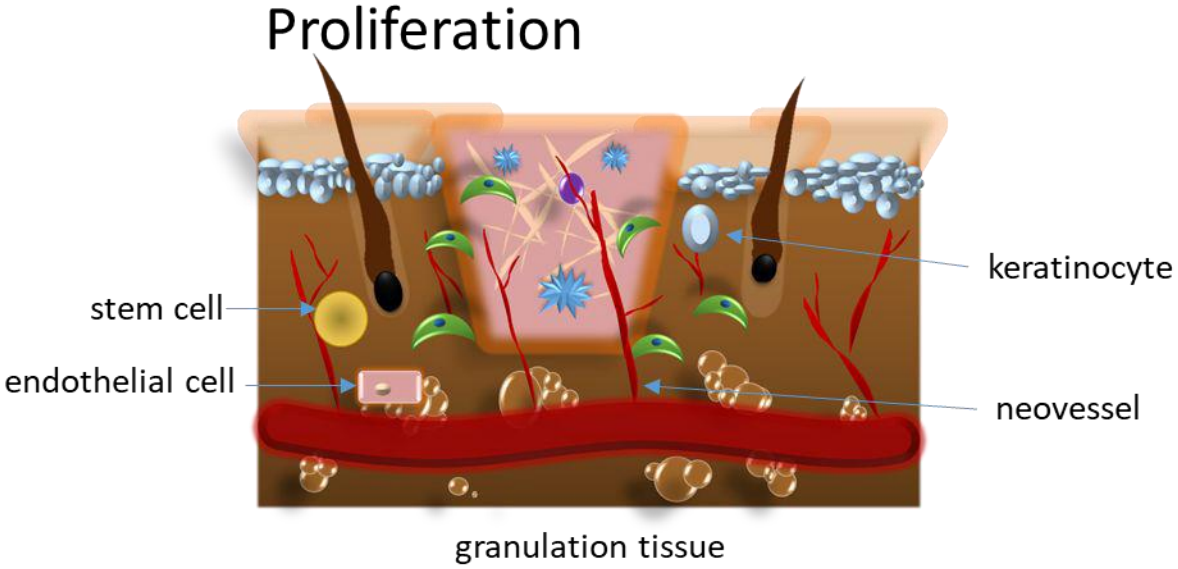
Hemostasis



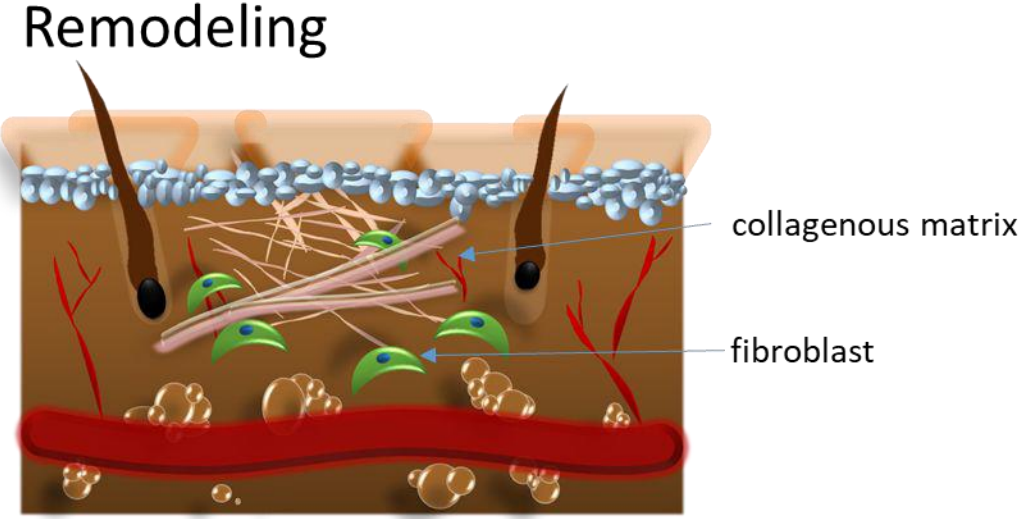
Inflammation



Proliferation



Remodeling



Spectrum of delayed wound healing

Modifiable and non-modifiable

- Comorbid conditions
- Nutritional adequacy
- Habits and physical capabilities
- Wound environment



Conditions that contribute to delayed wound healing

Systemic Factors affecting wound healing		
Nutrition and Hydration	Deficiencies, swallowing difficulties	Fluid restrictions
Medications	Steroids, anticoagulants, chemotherapy	
Systemic Infection	Vasopressors, increased metabolic demand	
Incontinence	Fecal, urinary, frequency	
Immobility	Use of calf muscles, repositioning	
Comorbid disease states	Endocrine disorders	Diabetes , thyroid disorder
	Hematologic	Anemia, systemic sclerosis, polycythemia, myeloproliferative disorders
	Cardiopulmonary problems	COPD, CHF
	Circulatory disease	Peripheral arterial disease, venous insufficiency , lymphedema, HTN, history of DVT or CVA
	Gastrointestinal	Inflammatory bowel disease, malnutrition, gastroparesis
	Autoimmune	Rheumatoid arthritis, lupus, inflammatory bowel disease
	History of radiation, sun exposure, smoking	

All patients undergo a comprehensive exam including:

- complete history
- physical assessment

Components of Wound Assessment

- Location
- Size (tunneling/undermining)
- Drainage
- Wound tissue
- Odor
- Peri-wound
- Edges/Margins
- Full thickness or partial thickness



Peri-wound



Maceration – softening of the tissues; is due to excess moisture and is considered an abnormal finding. It presents as white tissue at the edges of the wound

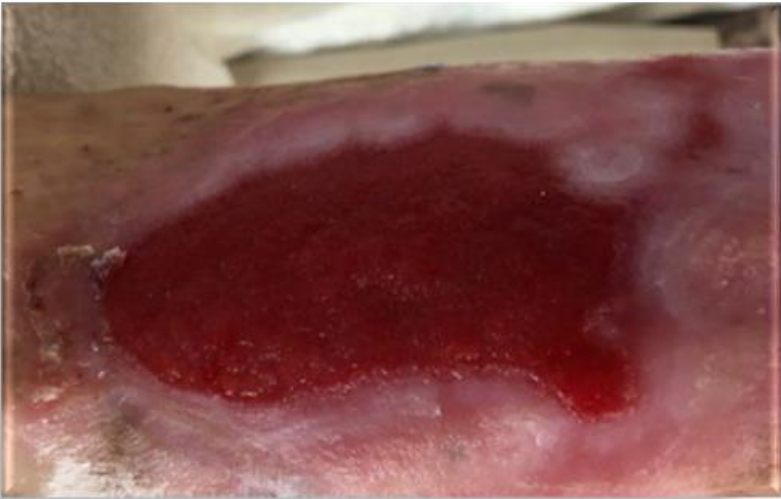


Erythema – redness; not necessarily from infection



Hyperpigmentation/hemosiderin staining
heme = blood
sid= iron
Permanent staining

Tissue Types



Granulation – tissue composed of new blood vessels, connective tissue, fibroblasts and inflammatory cells which fills an open wound when it starts to heal; typically appears deep pink or red with an irregular, granular surface



Epithelialization-Regeneration of epidermis across a wound surface. The color of the epithelium ranges from pearly to pink

Tissue Types continued



Slough – soft moist avascular (devitalized) tissue; may be loose or firmly adherent

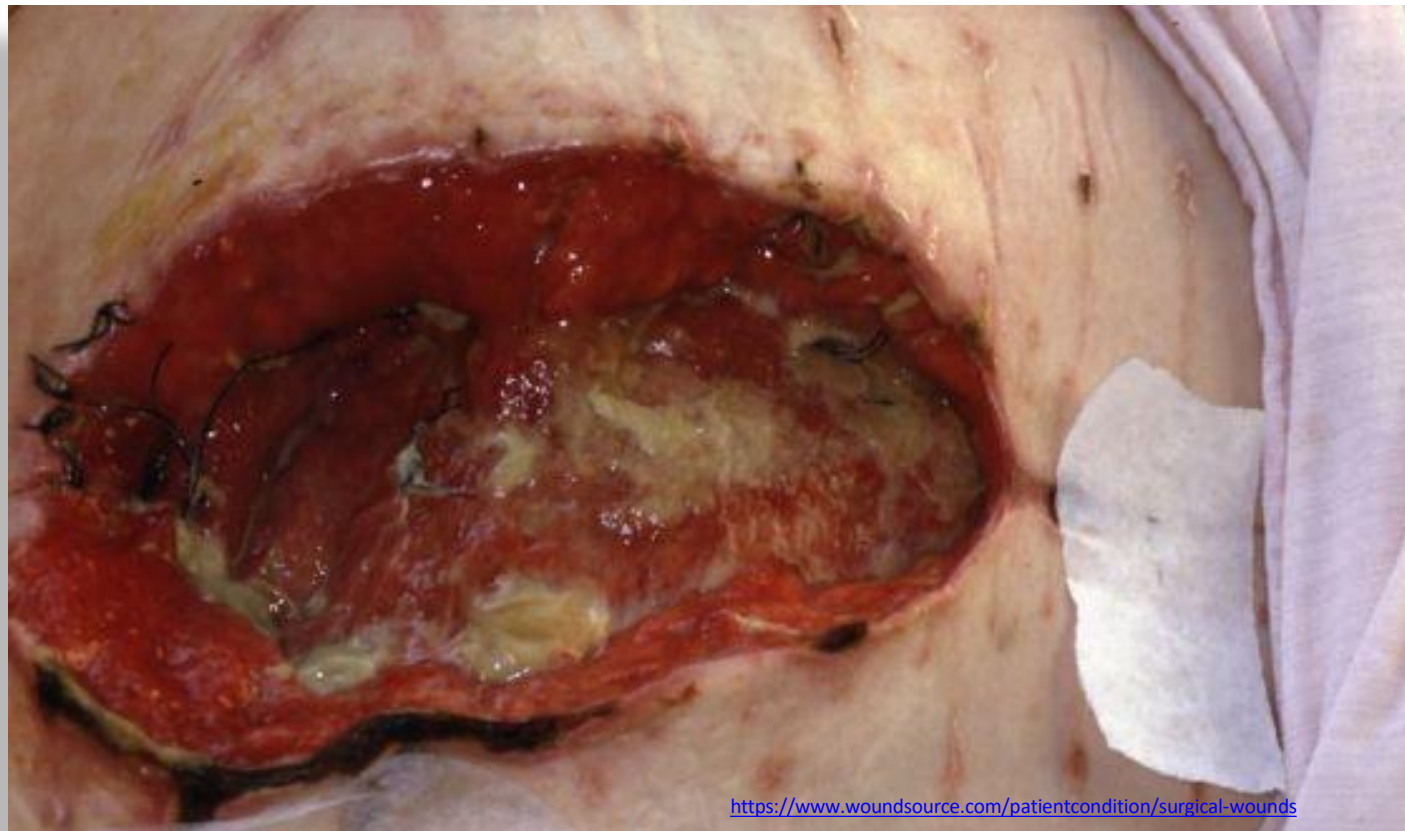


Eschar – Black necrotic devitalized tissue; tissue can be loose or firmly adherent, hard, soft or soggy

Tendon/ fascia



Organ and vessel



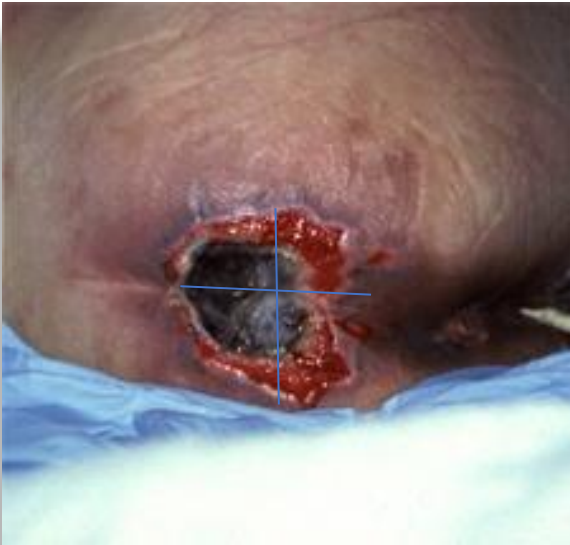
Bone



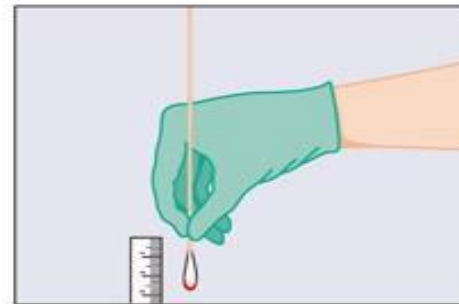
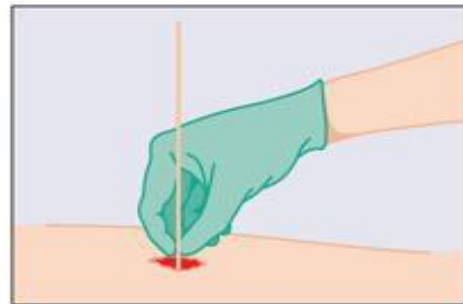
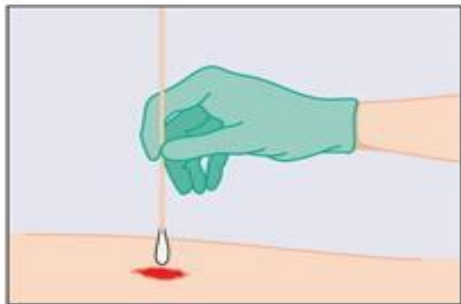
Probe to bone

- Using a sterile blunt metal tool gently search the base of the wound for hard, gritty surfaces
- Screening tool in conjunction with the patient's pretest probability
- Reliability may vary by the ulcer location and the expertise of the clinician performing the test
 - Best on the foot and evidence is on DFUs
- Systematic review evaluating the performance of the probe-to-bone test (using bone histopathology or culture as the reference standard):
 - pooled sensitivity 87%
 - specificity 83%

Wound Measurement



- Do not estimate size by comparing to an object
(e.g. wound is about the size of cheeseburger)
1. Length=longest length from patient head to toe position
 2. Widest=widest width from patient side to side position
 3. Depth (if $>0.1\text{cm}$) = gently place cotton tipped applicator into deepest part of wound



Undermining versus Tunneling

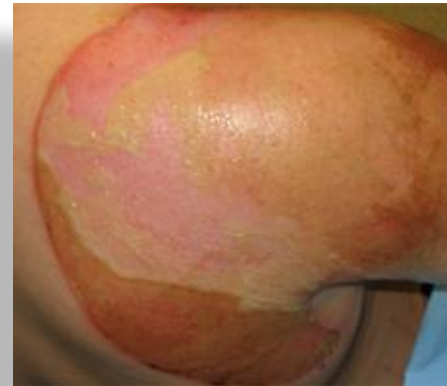


Undermining



Tunneling

Wound types



Wound types



Skin Tear



Perineal Dermatitis



Pressure Ulcer



Venous Ulcer



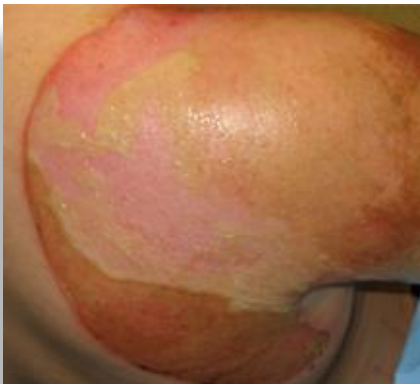
Diabetic Ulcer



Arterial Ulcer



Surgical Wound



Burn

Wound Hygiene

- A protocol of care that is delivered every time wound care is provided
- 4 simple steps
- Implemented world-wide

Wound Hygiene

- 1. Cleanse**
2. Debride
3. Refashion
4. Dress

Cleanse

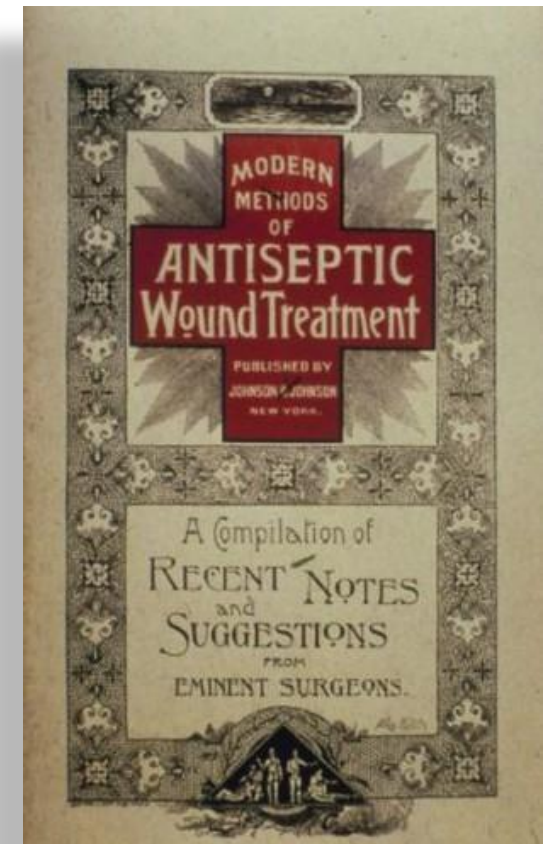
Non Cytotoxic Cleansers:

- Normal saline
- Hypochlorous acid
- Soap (mild) and water

Cytotoxic Cleansers:

- Non-dilute bleach solutions
- Povidone Iodine Solution
- High dose silver (silver sulfadiazene, certain contact layers)
- Chlorhexidine Gluconate
- Hydrogen Peroxide

Cytotoxicity: the dose makes the poison



Effective/safe antimicrobials

- Hypochlorous Acid (HClO)
- Same molecule secreted by neutrophils
- Non-toxic to fibroblasts
- A 15-second soak leads to 7 log reduction in pathogenic bacteria including:
 - *K pneumonia*;
 - *E faecalis*,
 - *S aureus* including MRSA
 - *S epidermis*
 - *A baumannii*
 - *P aeruginosa*
 - *E coli*
 - *C albicans*



Effective/Safe antimicrobials

- Methylene Blue/Gentian Violet
 - Medical-Grade Honey
 - Silver & Copper
 - DACC
 - PHMB
 - Cadexomer Iodine*
 - * betadine is potentially cytotoxic and should not be used on open wounds as better choices are usually available.
- All are available in advanced topical wound dressings



How to Cleanse Wounds

- ✓ Wash the wound, peri-wound, and the entire extremity
- ✓ Sterile water/normal saline = tap water
- ✓ Don't need to excessively scrub,
-but moderate pressure as tolerated
- ✓ Use gauze or clean wash cloth

How to Cleanse Wounds



- Decrease bacteria/fungus, optimize skin integrity
- Don't let patients soak their wounds!
- Moisturize (eczema, diabetic autonomic neuropathy)

Wound Hygiene

1. Cleanse
2. **Debride**
3. Refashion
4. Dress

Bioburden



the bacteria and inflammatory contents (proteases, pro-inflammatory cytokines) of a wound that potentiates the inflammatory stage and limits healing

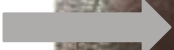
slough



eschar



hyperkeratotic plaques



* Debridement of hard-to-heal wounds is standard of care

granulation

Forms of debridement

Faster



Slower

- Sharp
- Biological
- Conservative sharp
- Mechanical
- Chemical
- Enzymatic
- Autolytic

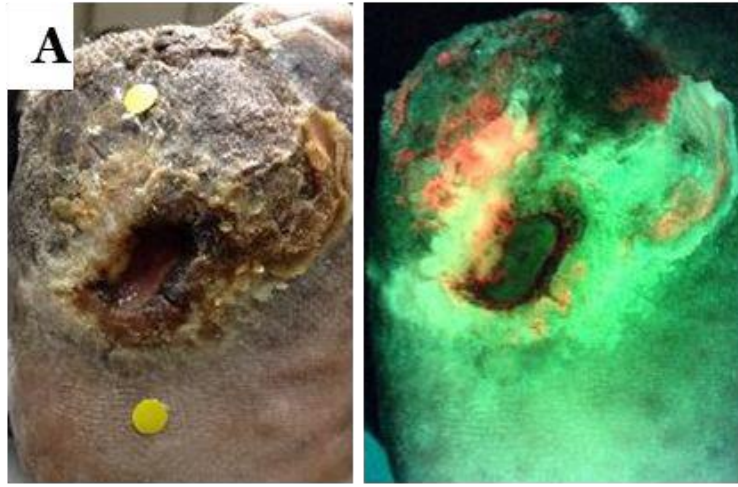


Debridement

- Removes necrotic tissue, which minimizes proteolytic enzymes
- Removes biofilm
- Decreases bacterial bioburden
- Decreases inflammation
- Promotes epithelial edge advancement



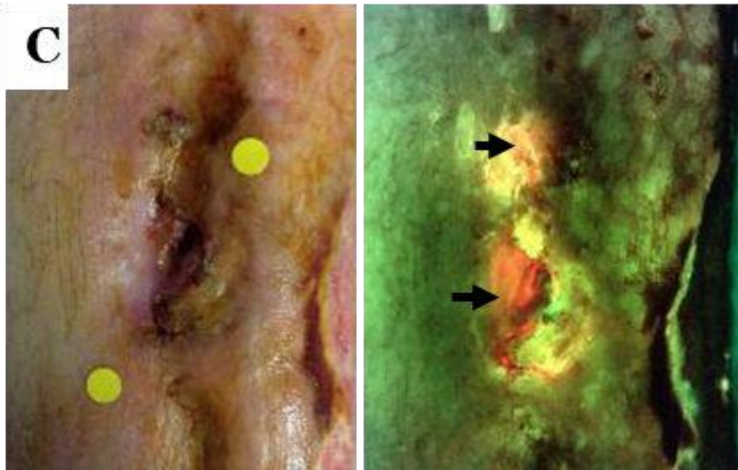
Fluorescence imaging reveals hidden load



Serratia marcescens and mixed anaerobes
(heavy growth)



Proteus mirabilis
(light growth)



Mixed bacteria (heavy growth)



Mixed bacteria (heavy growth)

Green = matrix components from tissues
Red/pink = bacteria ($>10^4$ CFU/g)

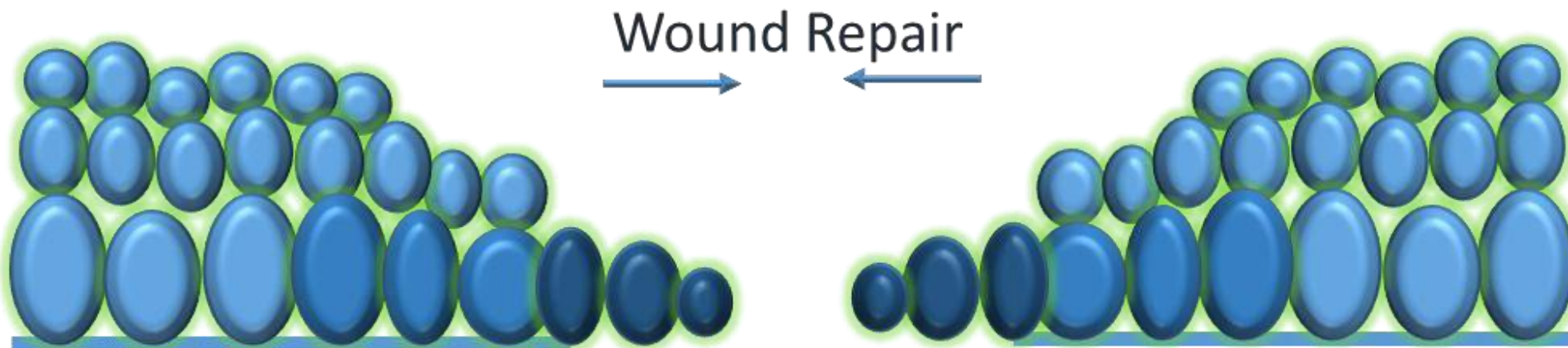
Wound Hygiene

1. Cleanse
2. Debride
- 3. Refashion**
4. Dress

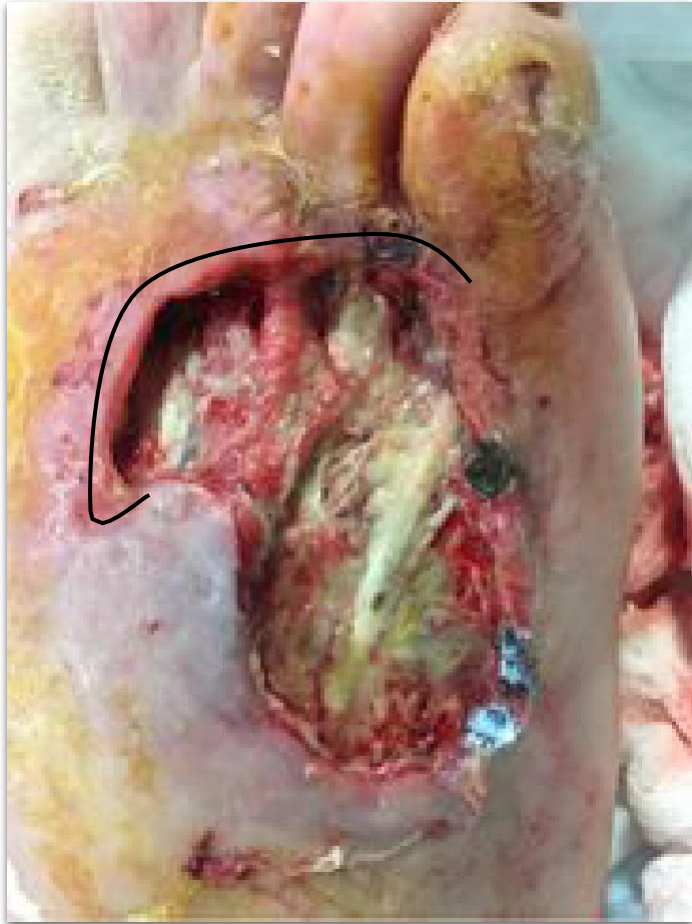
Refashion edges

Goals:

- attached edges
- “cereal bowl” shaped wound
- epithelial edge advancement



Refashion edges



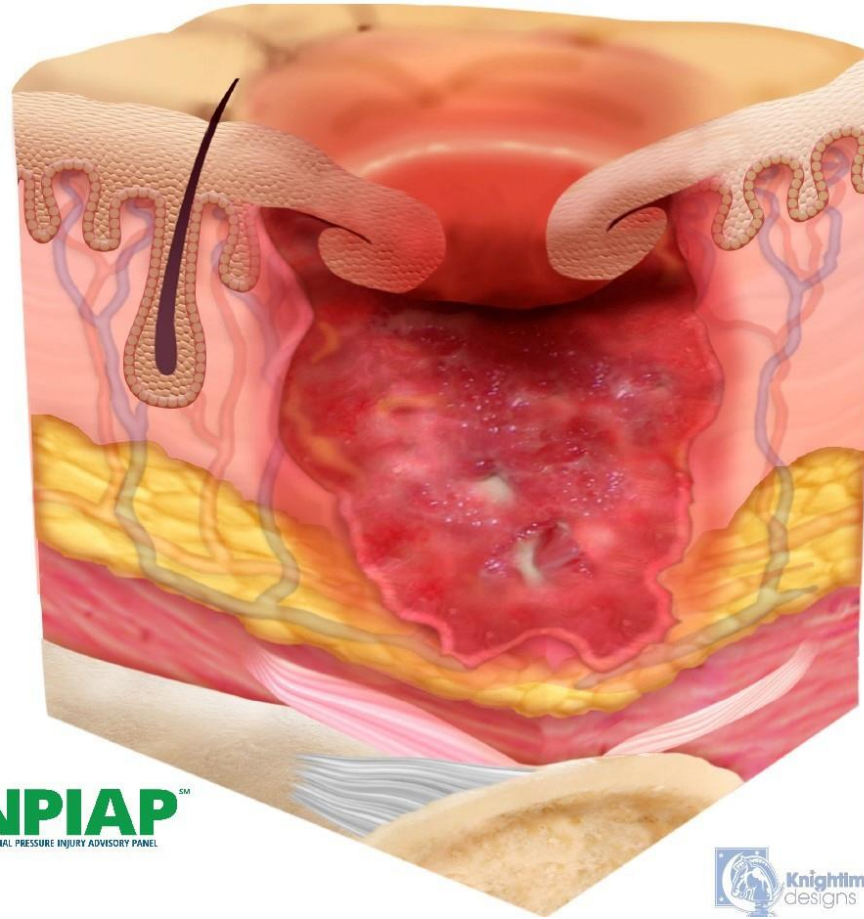
- Epibole
- Callus
- Hyperkeratosis
- Un-attached edges



Epibole



Area of Focus



Refashion Edges

- Diabetic foot ulcer presentation to wound clinic after 3x/week RN dressing changes and non-contact low frequency ultrasound.
- Presented with wound bed obscured by hyperkeratosis.
- Curette callus to go layer by layer - allows investigation of whether ulcer is present



Removal of blister and callus reveals wound base



- H/o neuropathic ulcer to site
- with new bullae
- Fluctuant area where we know there is a pocket of non-viable tissue and/or fluid
 - scalpel
 - curette
- Curette remaining callus to go layer by layer
 - allows investigation of whether ulcer is present

Wound Hygiene

1. Cleanse
2. Debride
3. Refashion
- 4. Dress**

Dress = Moist Wound Healing

Published in Lancet by Dr. Winter in 1962



Moist wounds heal:

- ✓ Faster
- ✓ With less scarring
- ✓ With fewer infections
- ✓ More cost effectively
- ✓ And less pain



- No soaking!
- Don't leave open to air
- Wounds don't need to "air out"
- Caveat: eschar on ischemic limbs



Moisture balance

- Wound drainage is irritating to surrounding skin = skin breakdown and dermatitis
- Too much drainage?
 - Step up dressing absorbency
 - Skin protectant
 - Increase dressing change frequency



Cover up!

Covering a wound with a protective dressing:

- ✓ Maintains a moist wound environment
- ✓ Minimizes bacterial contamination
- ✓ Maintains normothermia
- ✓ Protects wounds from trauma



1. Gauze doesn't protect the wound from bacterial contamination.

- bacteria can penetrate 60 layers of gauze!

2. Gauze potentiates inflammation

- foreign body reaction
- doesn't remove bacteria, cytokines, and proteases in drainage

3. Doesn't hold fluid away from wound

- has to be changed daily
- causes local hypothermia

4. Gauze is used in the lab to grow biofilm!



Typical Wound Regimen

1. Cleanse the wound, and the whole extremity (foot/leg/arm) with warm soapy water.
2. Spray with hypochlorous acid and let sit for 5 minutes.
3. Pat the skin around the wound dry.
4. Apply a foam bordered dressing
5. Change 2-3 times per week dependent on drainage.
6. Address etiology:
 - Diabetic/neuropathic foot ulcers: offload (don't walk on it, wear appropriate footwear, urinal, wheelchair, knee scooter)
 - Venous leg ulcers: compression, 30-40mmHg
 - Pressure Ulcer: offload (no donut cushions, use waffle/gel/roho cushions, reposition, micro reposition, limit time in chair, heel protectors)



Acute infection (planktonic) v chronic infection

This microscopic image shows a cross-section of plant tissue, likely a leaf, stained with a purple dye. The tissue structure is visible, including the epidermis and underlying cells. Numerous dark, circular spores are scattered throughout the tissue, indicating an acute infection. The spores are concentrated in certain areas, particularly near the surface, and are surrounded by a clear, uninfected-looking tissue. The overall appearance is that of a planktonic infection, where the pathogen is present in a free, non-replicative state.

Infection

Studies show that between 30% - 50% of in-patient antibiotic use is either unnecessary or inappropriate

16.4% of all antibiotic prescriptions are for wounds

53-71% of patients are prescribed at least one course per chronic wound

2.8 million antibiotic-resistant infections occur in the United States each year

More than 35,000 people die from these infections

Complications of wound infection

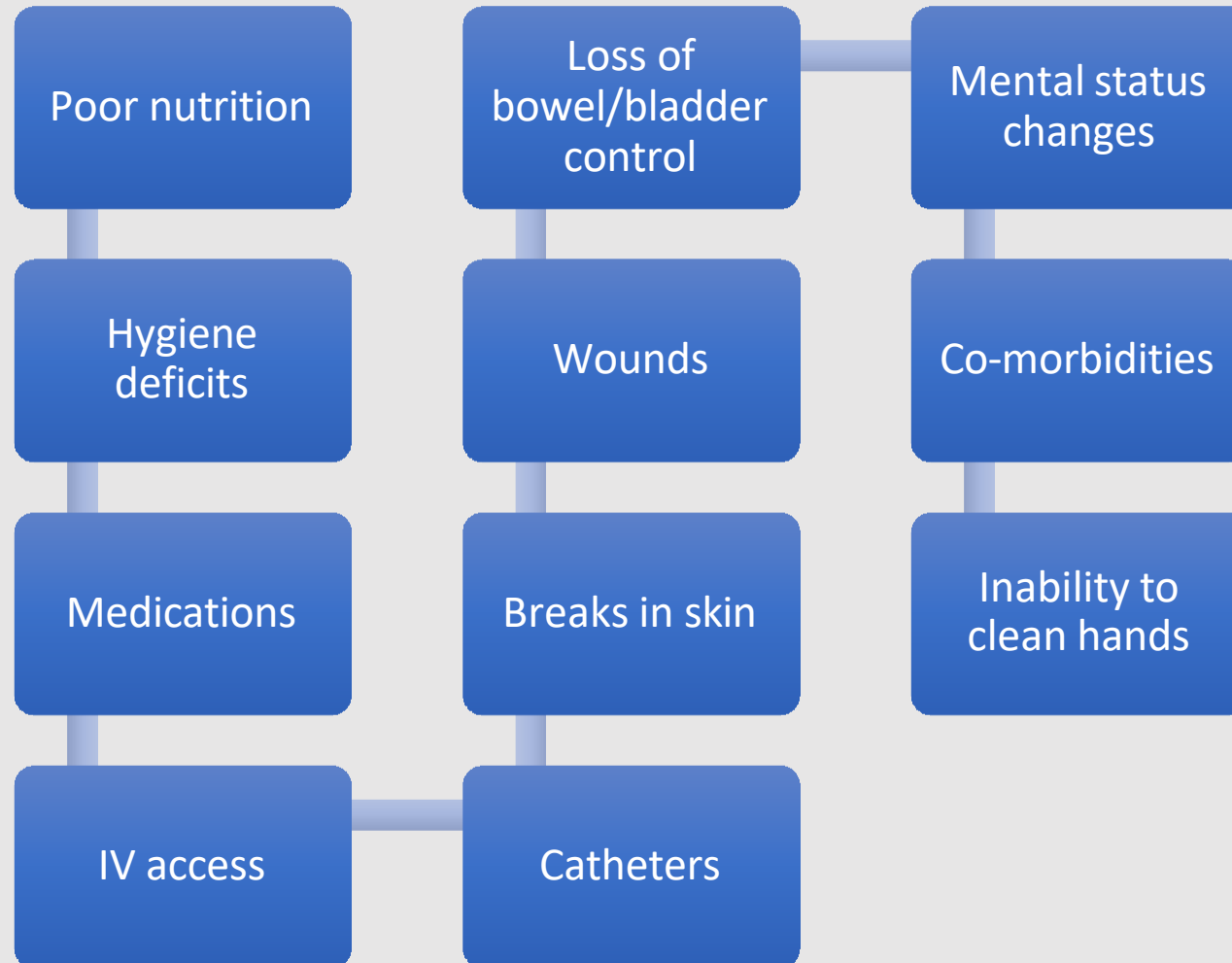


Wound infection risk with aging

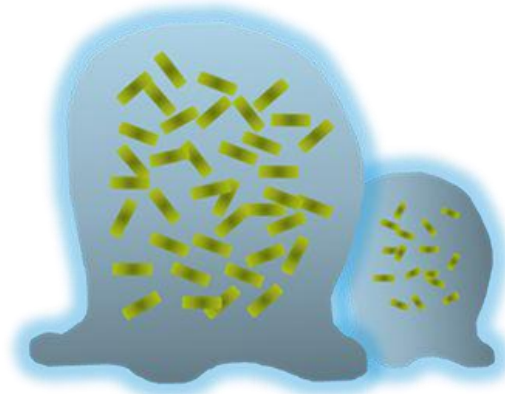
- Aging causes accumulation of disease and pharmaceuticals that affect our ability to heal and our immune system, increasing risk for infection
- People who live or work together are more likely to share microbes (eg long term care environment)
- Many infections can be prevented with basic infection prevention and control



Wound infection risk with aging



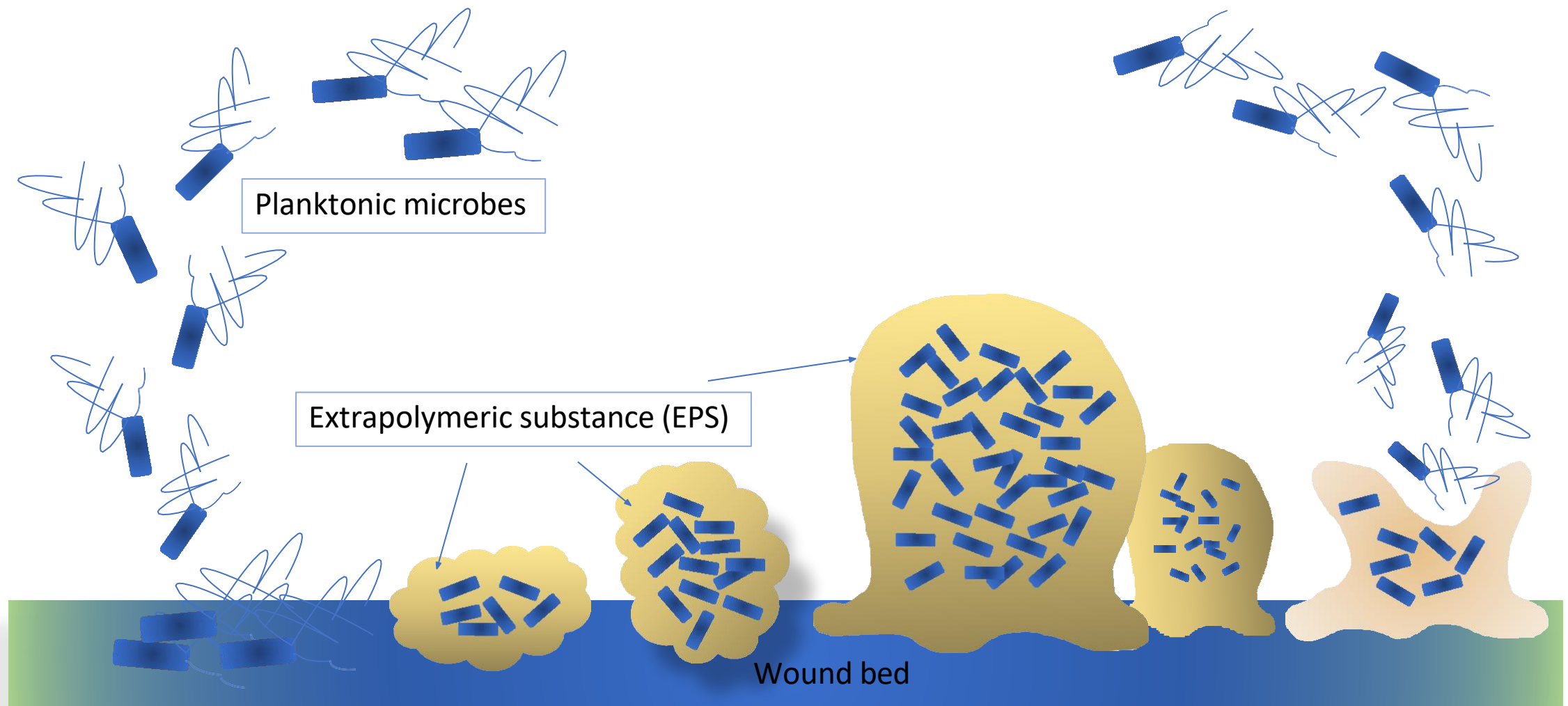
What is colonization?



Carrying microbes without causing disease / causing undetectable disease



When you're colonized you can still spread the microbes to others and cause disease



Reversible attachment

Irreversible attachment

Maturation

Dispersion

Biofilm is a complex, often polymicrobial, structure that can form on an array of surfaces including: teeth, the hull of ships, implanted medical devices, showers and bathtubs, and notably chronic and hard-to-heal wounds

Biofilm

*You can't see it; it's present in most wounds; best removed through sharp debridement



Protects Bacteria and Fungus in a EPS (usually polysaccharide)



Begins to reform in 24-48 hours even after surgical removal



Is tolerant to most topical antibiotics and antimicrobials



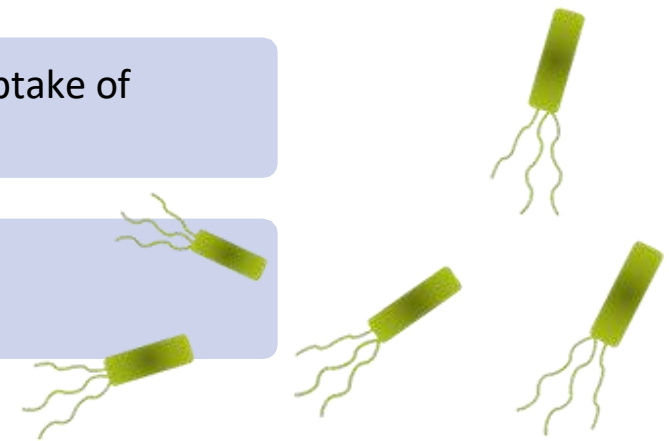
Allows bacteria to share resistance genes/nutrients through quorum sensing



In sessile/biofilm phenotype not metabolically active = Decreased uptake of bacteriocidal mechanisms



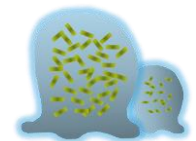
Delays wound healing



Chronic Inhibitory Bacterial Load (CIBL)

Bacteria in biofilm
delay wound healing
even when not causing
acute infection

Treating with oral or systemic ATB and NOT addressing biofilm is not enough



Wound infection s/s



Sign – what you can observe

Increased drainage
Wound getting larger
Unhealthy tissue
Redness
Swelling
Pus
Odor



Symptom- what the resident tells you

Pain
Subjective fevers
Subjective size or appearance
Reported odor

Inflammation

Limitations: differentiating infectious inflammation from other forms

- autoimmune diseases
- reperfusion injuries
- pressure
- venous dermatitis
- DVT
- Burn, expected healing rim
- Contact or irritant dermatitis
 - Eg. Moisture/ drainage on periwound



Limitations of clinical s/s of infection

- Substantial variability in how wounds are assessed and deemed infected or not
- Disease states limit clinical s/s: diabetes, PAD,
- Skin tone limitations
 - Erythema



Fluorescence Guided culturing

Violet light with filter causes bacteria at greater than 10^4 CFU/g) that produce porphyrins to glow red

Green = matrix components from tissues

Culture to specific location based on imaging

More sensitive than clinical s/s alone



Culturing Wounds

Levine technique

How to culture:

- Clean the wound with soap and water or normal saline
- Do not culture pus or frank necrotic tissue
- Culture intact tissue
- Push down in one place to express fluid from the wound bed

When to culture:

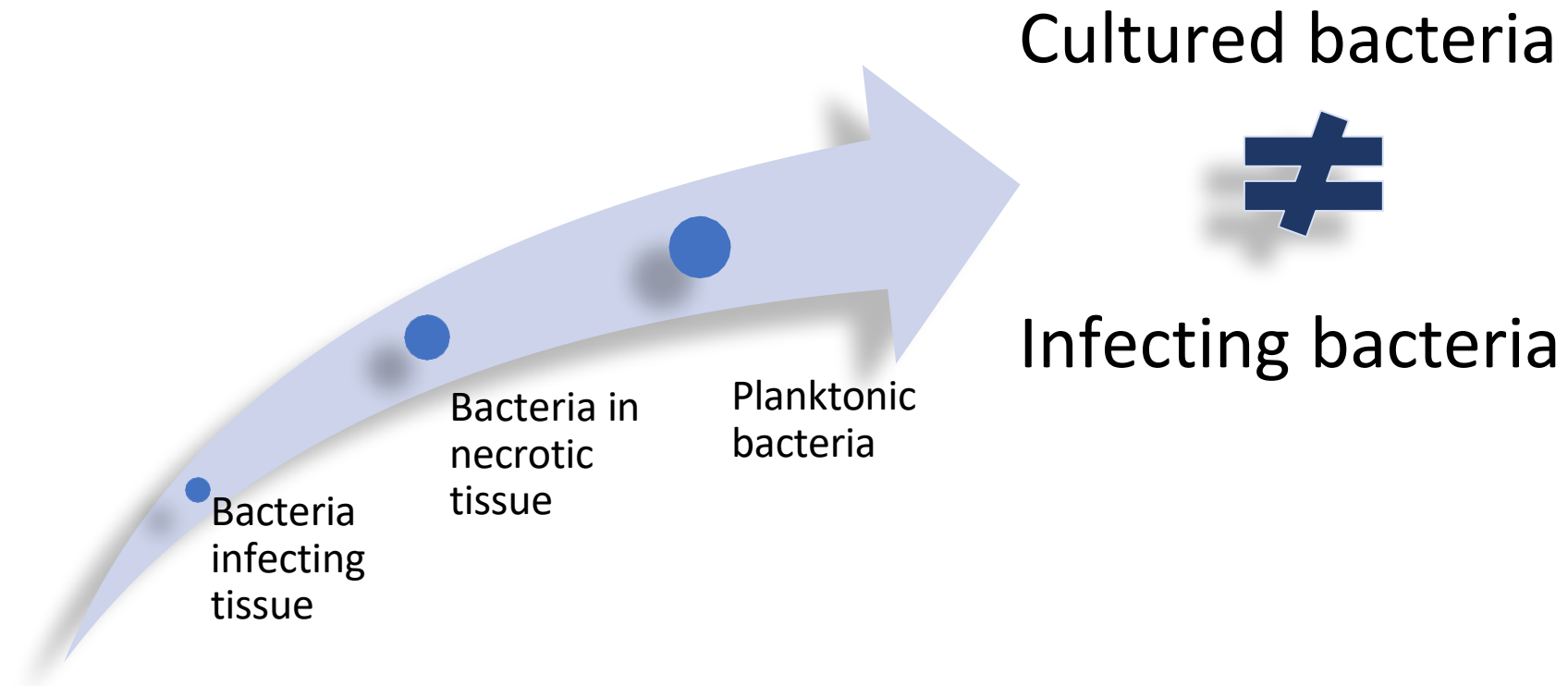
- When you suspect wound infection and culture would guide antibiotic selection

When NOT to culture:

- If you don't suspect infection



Bacteria in Necrotic Tissue v Healthy Tissue:





Topical antibiotics in wound care

Topical antibiotics = antibiotic resistance + allergic skin reactions

More likely to develop an allergy when used on skin

Former allergens of the year:

- bacitracin
- neomycin

Allergies and resistance can apply to oral and parenteral form

Does not penetrate biofilm

❖ *Planktonic bacteria that develop resistance can share resistance genes into biofilm via quorum sensing*



Burns and silversulfadiazene (SSD) cream

- 450,000 people receive medical attention for burns annually
- 45,000 people are hospitalized for burns
- Systematic reviews of SSD in burns show it can actually increase infection rates and hospital length of stay
- Burn wounds don't need prophylactic antibiotics
- Good dressings for burns are hydrogels and hydrogel sheets, which can reduce pain



Antimicrobial stewardship

Antibiotic resistance is one of the biggest public health challenges of our time.

It is on the rise worldwide.

- killing 1.27 million globally
- associated with nearly 5 million more deaths

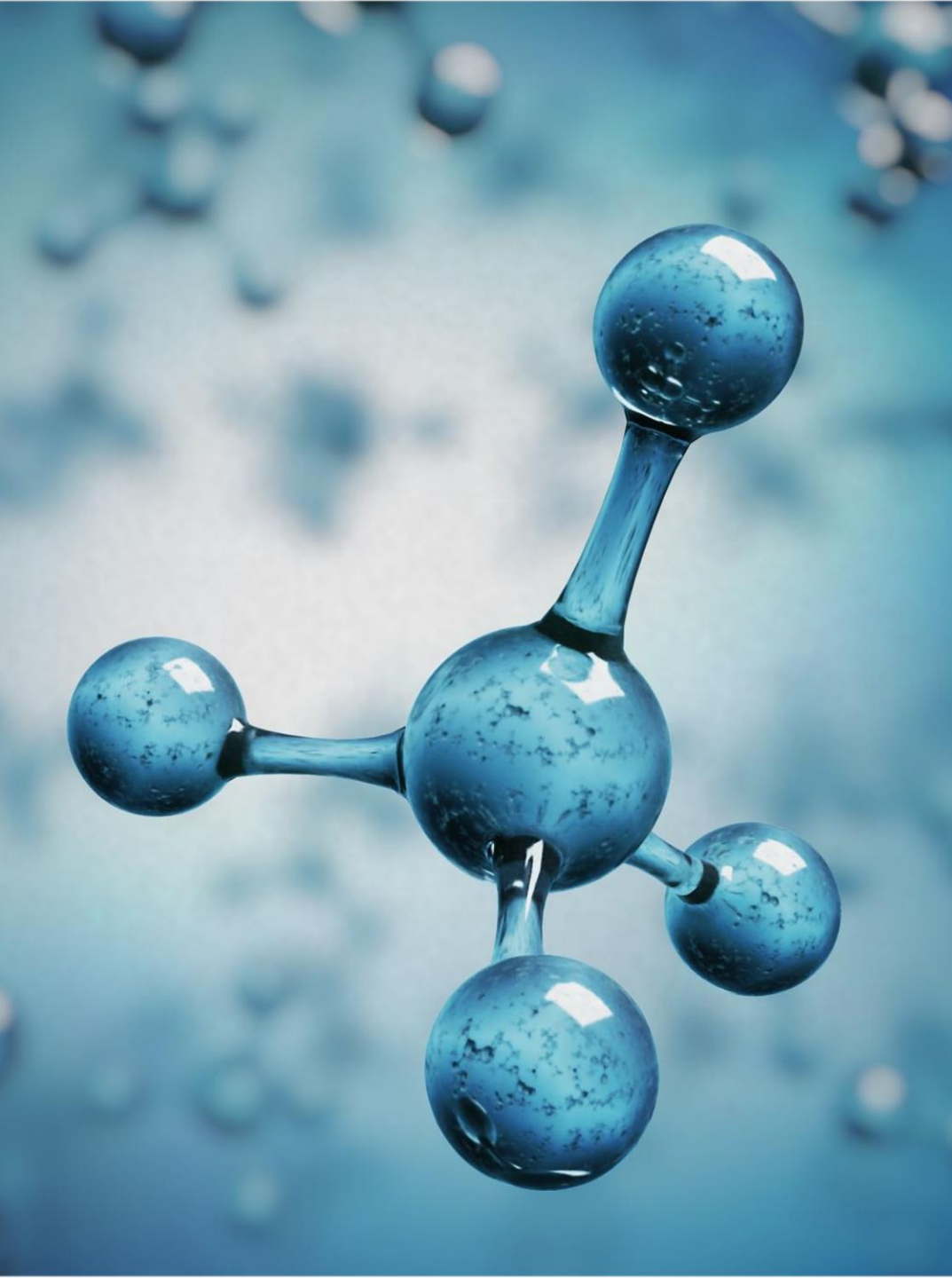


Antibiotic use eliminates non-resistant bacteria,
increasing the proportion of resistant bacteria that remain.



Wounds without evidence of soft tissue or bone infection do not require antibiotic therapy.

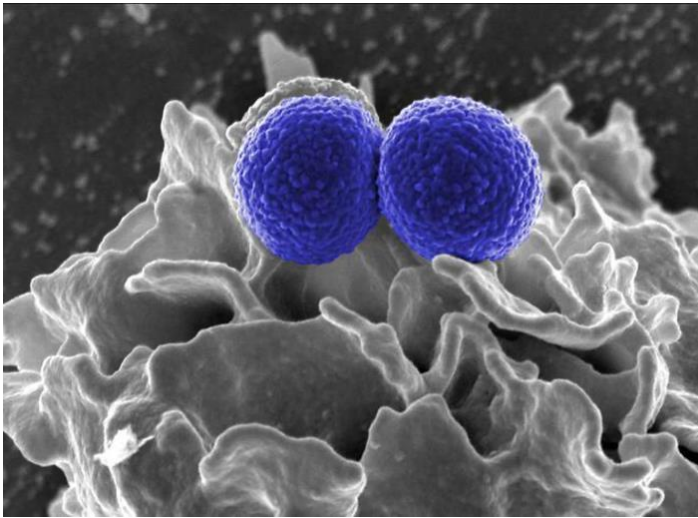




Safe antimicrobials

- Silver in dressings and gels
 - Next gen silver dressings have BEC and EDTA
- Copper dressings
- Manuka honey
- Methylene blue and gentian violet dressings
- Cadexomer iodine
- Biofilm disruptors
- Hypochlorous acid

Considerations in the use of antimicrobials



Cytotoxicity

Lack of resistance

Works via multiple mechanisms

Allows for early intervention and avoidance of atb's and antifungals



Multi-Drug Resistant Organisms:

Microorganisms, predominantly bacteria, that are resistant to one or more classes of antimicrobial agents

Bacteria have adapted and are no longer killed by an antibiotic

Make it more difficult to treat an infection

Examples of MDRO

MRSA (methicillin-resistant staphylococcus aureus)

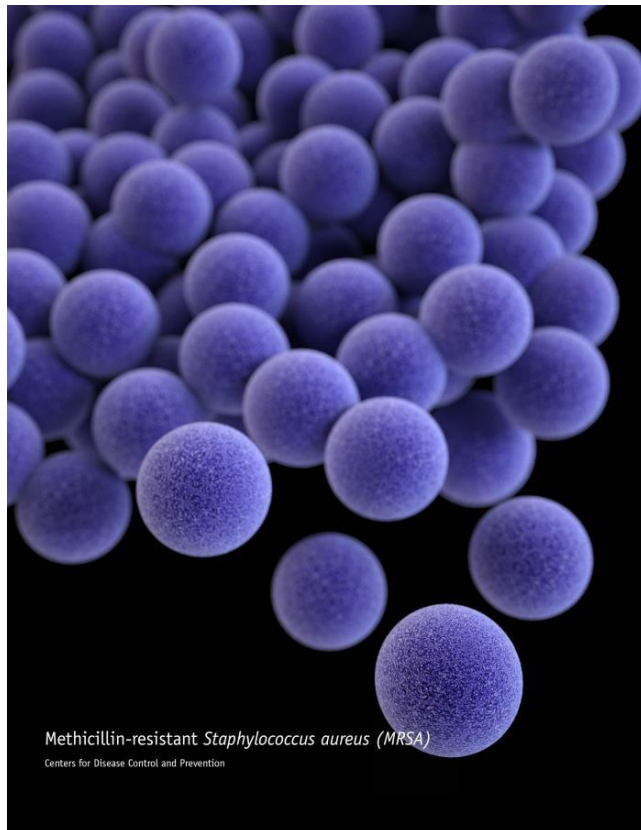
VRE (vancomycin-resistant Enterococcus)

C. Difficile (Clostridium difficile)

ESBL (extended spectrum beta lactamase) bacteria

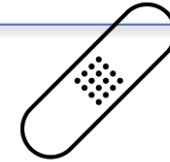
CRE (carbapenem-resistant Enterobacteriaceae)

Highest risk of MDRO:



Indwelling medical devices and/or lines.

Presence or history of chronic wounds.



History of residing in congregate living settings.

History of out-of-state or international health care.

History of frequent surgeries or procedures.

Frequent or prolonged stays in hospitals or long-term care facilities.

Underlying chronic medical conditions.

MDRO characteristics



In most instances, MDRO infections have clinical manifestations that are similar to infections caused by susceptible pathogens



Antimicrobial resistance rates are also strongly correlated with hospital size, tertiary-level care, and facility type (e.g., LTCF)



Options for treating often extremely limited



Increased LOS



Cost



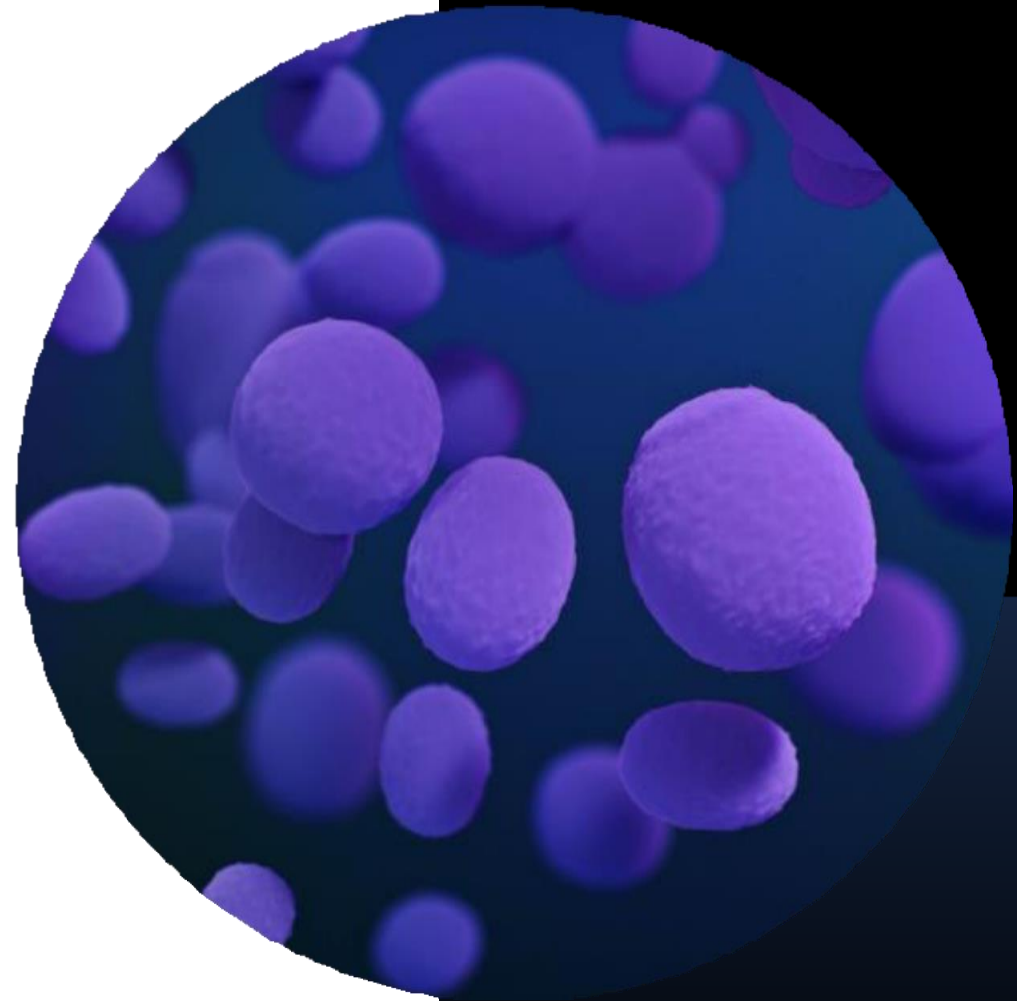
Mortality

MDR Candida Auris

Cases have more than tripled from 2022 to 2023

Drivers:

- overall antifungal use,
- subtherapeutic drug levels at sites of infection/colonization,
- drug sequestration in the biofilm matrix,
- in the setting of outbreaks, suboptimal infection control.





MDRO Prevention & Control

Nearly all studies reporting successful MDRO control employed a median of 7 to 8 different interventions concurrently or sequentially

Prevent and control MDRO spread

Follow infection prevention best practices

Appropriate clinical practices incorporated into routine patient care

Accurate diagnosis of infectious etiologies

Judicious antimicrobial selection and utilization

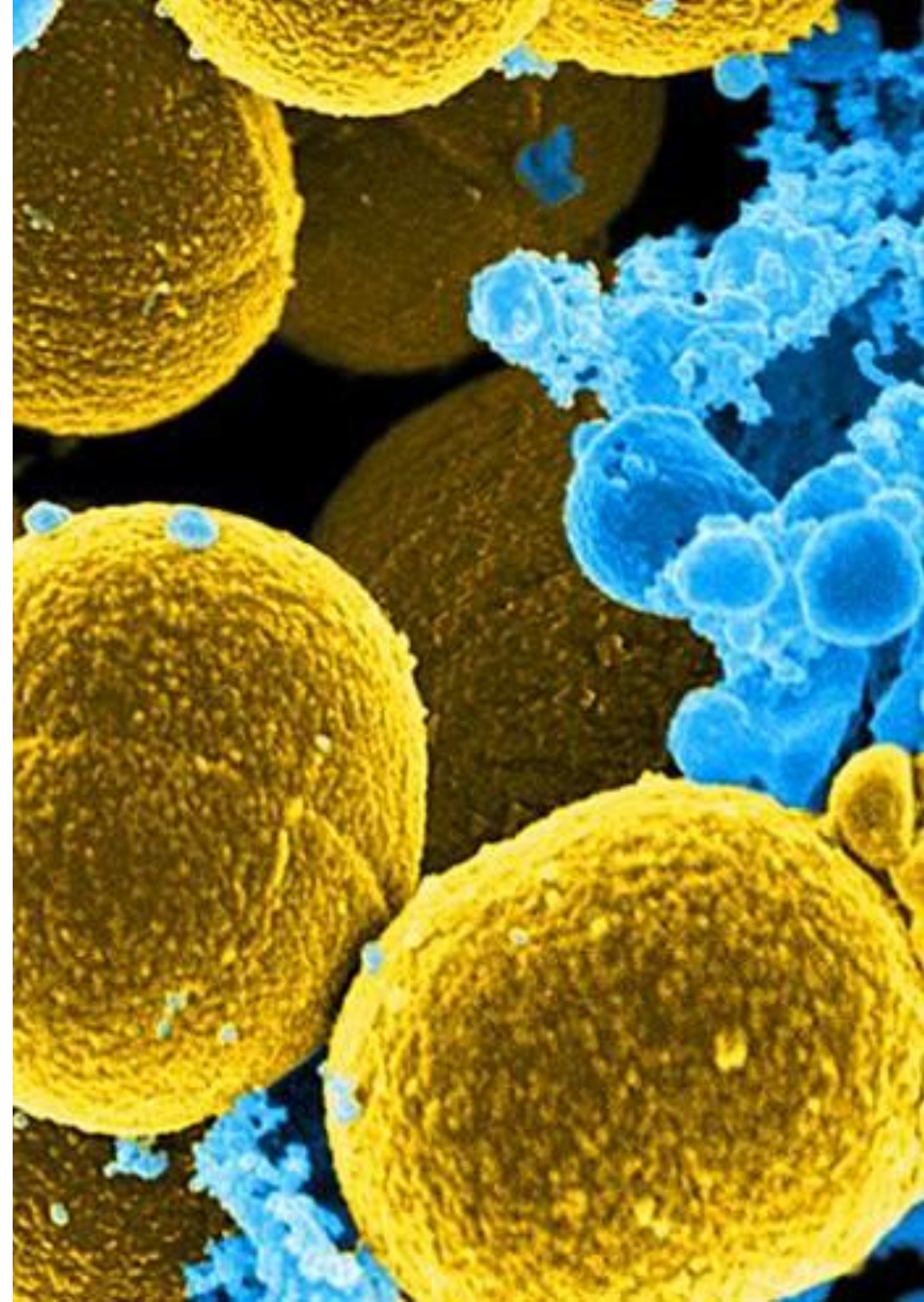
There is ample epidemiologic evidence to suggest that MDROs are carried from one person to another via the hands of HCP

Facilities must display hand hygiene posters in heavily trafficked areas.



MDRO spread

- Wash hands!
- Cover cuts and wounds with a dressing
- Follow isolation precautions when in place
- Complete the full round of antibiotics



Infection Control Assessment and Response (ICAR) tool for General Infection Prevention and Control (IPC) across settings

- Level of detail not sufficient for specialty areas like burn units
- Module 3 – observation form- wound care
- Module 8 – wound care facilitator guide

Prevention of cross transmission

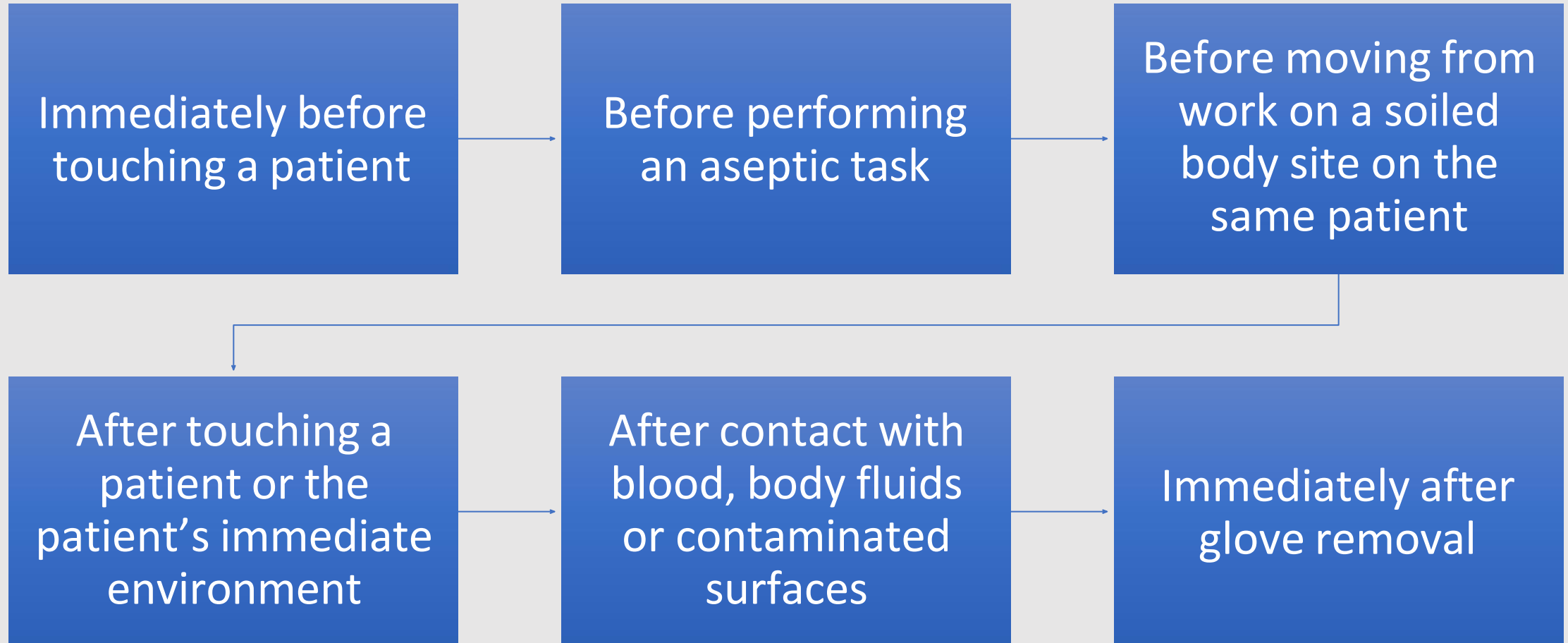
Must be performed
wherever wound care
is performed

Many facilities have
wound care teams but
nursing personnel will
still perform dressing
changes

Practical hygiene tips for big impact

- Alcohol-based hand sanitizer dispensers easily accessible to HCP while performing wound care?
- Do HCP perform hand hygiene before performing wound care?
- Do HCP don clean gloves and other recommended PPE?
- Gloves changed and hand hygiene performed when moving from dirty to clean tasks
 - Moving amongst wounds
- Maintain separation between clean and dirty supplies

Hand hygiene occurs



PPE recommendations

- Gloves should be worn during wound care procedures
- Gowns should be worn when wound care requires significant contact with resident or their immediate environment, such as when turning or positioning a resident for wound care or if the procedure could generate splashes or sprays (eg during irrigation)
- Face protection such as goggles or a face mask/face shield should be worn during wound care procedures that may generate splashes or aerosols such as irrigation, pulse lavage, and handling equipment such as vacuum assisted closure devices
- Additional PPE may be warranted if the patient/resident is on precautions
- Wear gloves when it can be reasonably anticipated that contact with blood or other potentially infectious materials could occur
- Wear a gown appropriate to the task during activities that could cause contact with body fluids
- Use protective eyewear and a mask or face shield during procedures and activities that could generate splashes or sprays

Prior to starting procedure

- Clean supplies gathered and placed on a clean surface in the room?
- Dressing materials and equipment should be selected and gathered prior to entering the patient/resident area to avoid accessing the supply cart/clean storage during the procedure
- Only the materials needed for an individual patient should be brought into the patient's room or treatment area
 - Placed on a clean surface
 - Away from potential sources of contamination (away from sink splash zones)
 - Brought prior to beginning wound care

Where is the
wound care
performed?

- Patient/resident room
- Procedure room
- Operating room
- Unknown

Where are
clean wound
care supplies
stored?

- Patient/resident room
- Procedure room
- Wound care cart
- Clean supply closet

Does the wound care
clean supply cart remain
outside the patient
immediate care area?

Where are wound supplies stored?

- Maintain separation between clean and soiled equipment to prevent cross contamination
- If wound care cart is used, it should not enter the patient/resident's immediate care area (room)

What happens to
unused disposable
supplies that enter
the patient/resident
area?

- Discarded
- Returned to clean supply storage (eg cart, closet, bin) for use on other patients/residents
- Dedicated to the patient/resident
 - How/where these supplies are stored and how the facility ensures they remain dedicated to the patient/resident

What happens to unused disposable supplies that enter the patient/resident area?

- Maintain separation between clean and soiled equipment to prevent cross contamination
- Dressings/supplies that enter the patients care area are now soiled
 - must either be discarded or dedicated to the patient

What happens to unused disposable supplies that enter the patient/resident area?

If supplies dedicated to individual patient/resident:

- Should be properly labeled
- Stored in a manner to prevent cross-contamination or use on another patient
 - Eg. designated cabinet in the patient's room

Is any wound care
equipment used for more
than one patient?

Bandage scissors

Bandage scissors should not be transported in pockets

- r/o cross contamination

WOUND CARE equipment and supplies

- If fresh bandages are cut for the resident, it should be done with clean scissors, not with scissors used to cut off soiled bandages
- Wound care dressings can be disposed of in the regular trash unless they are dripping or saturated with blood or other regulated body fluids.
- Dedicate tape, sprays, creams, and all wound care products to an individual resident and do not store used sprays with clean wound care supplies.

WOUND CARE equipment and supplies

- Clean and disinfect the surface (e.g., over bed table) where wound care supplies will be placed prior to setting down wound care supplies in resident room.
- Store wound care supplies in a clean area of resident room.

Is re-useable
equipment cleaned
and disinfected
after each use?

- Re-usable wound care equipment should be cleaned and disinfected after each use
- Level of disinfection depends on type of equipment
- Eg. bandage scissors only require low or intermediate level disinfection v debridement supplies must be sterile

Who is responsible for
cleaning/disinfecting
the equipment before
use on another
patient?

How/where is cleaning and disinfection performed?

- Dedicated (in-house) wound care team
- Dedicated (external/consultant) wound care team
- Nursing personnel
- Unknown

Who is responsible
for
cleaning/disinfecting
the equipment before
use on another
patient?

If device reprocessing
performed elsewhere
devices must be contained
and transported to prevent
cross contamination
(soaking in
detergent/cleaner in a
biohazard container)

Is topical medication either dedicated to an individual patient or aliquoted for individual patient/resident use prior to entering the patient/resident room?

- Includes creams/sprays/ointments
- Dedicated containers must be properly labeled and stored
- Once enters the patient care area must be dedicated to patient or discarded

Dedicate multi dose vials to a single patient whenever possible.

If multi dose vials are used for more than one patient, restrict the medication vials to a centralized medication area and do not bring them into the immediate patient treatment area (ie OR, patient room, cubicle)

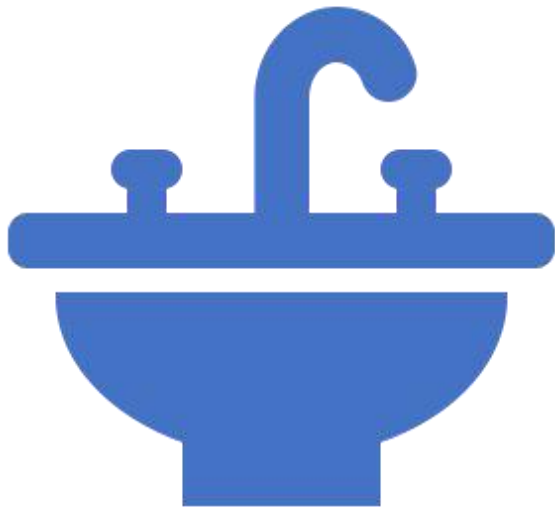
Are potentially contaminated surfaces cleaned and disinfected after wound care completed?

- You must discard PPE and perform hand hygiene after completing wound procedures
- Routine and targeted cleaning of the environment is required!
 - Close proximity to the patient
 - Frequently touched surfaces

Infected Roomies?

Consider how long an infected resident has been their roommate before moving them

- When resident colonized or infected they may need to be moved to single room to prevent transmission
- If cannot be in single room, cohort with same germ
- If cannot cohort, place infected residents with low risk residents



Patient considerations with MDRO wound

- Have infected residents shower last
- A patient with MDRO should not be moved to room with resident dependent on staff for ADLs
- A patient with MDRO should not be moved to room with urinary Catheter, IV catheter, or an open wound

Administrative controls

1. Facility has current, evidence-based policies and procedures readily available regarding wound detection, assessment, and management, which are reviewed and updated on an annual basis
2. Resources about wound care are available for staff to utilize should questions or concerns arise (i.e., nursing reference book with checklists).
3. The facility has a competency-based program for training all personnel who provide wound care upon hire and annually thereafter. Education is provided when new equipment or protocols are introduced.
4. The facility audits (monitors and documents) adherence to wound care policies and procedures and provides feedback to health care workers (HCWs), including contracted staff, regarding their performance of wound care. Audits should be conducted with a standardized tool on a routine basis. HCWs will receive education focused on gaps identified during audits.
5. The facility keeps a record of all types of wound and skin infections identified in residents receiving wound care. When necessary, transmission-based precautions or Enhanced Barrier Precautions (EBP) are implemented based on CDC guidance.

If you're going to audit its recommended to do so on at least 2 different staff, when direct observations can't be used ask staff

Wound hygiene in practice



- ✓ Cleanse
- ✓ Debride
- ✓ Refashion
- ✓ Dress

Presentation to wound clinic of traumatic foot ulcer

Wound hygiene in practice



Week 1



Week 2



Week 3

Wound hygiene in practice



Week 4



Week 5



Epithelialized

Conclusion

- Moist wounds heal quicker with less infection
- AVOID topical antibiotic ointments, unless needed based on culture results
- AVOID cytotoxic cleansers and wound dressings
- When needed, refer to a wound specialist

Typical Wound Regimen

1. Cleanse the wound, and the whole extremity (foot/leg/arm) with warm soapy water.
2. Spray with hypochlorous acid and let sit for 5 minutes.
3. Pat the skin around the wound dry.
4. Apply a foam bordered dressing
5. Change 2-3 times per week dependent on drainage.
6. Address etiology:
 - Diabetic/neuropathic foot ulcers: offload (don't walk on it, wear appropriate footwear, urinal, wheelchair, knee scooter)
 - Venous leg ulcers: compression, 30-40mmHg
 - Pressure Ulcer: offload (no donut cushions, use waffle/gel/roho cushions, reposition, micro reposition, limit time in chair, heel protectors)

Conclusions

1. Wound supplies that enter the patient area must be dedicated to the patient, discarded, or properly sanitized.
2. Wound carts cannot enter the patient area
3. Sanitize following procedures
4. Use clean supplies



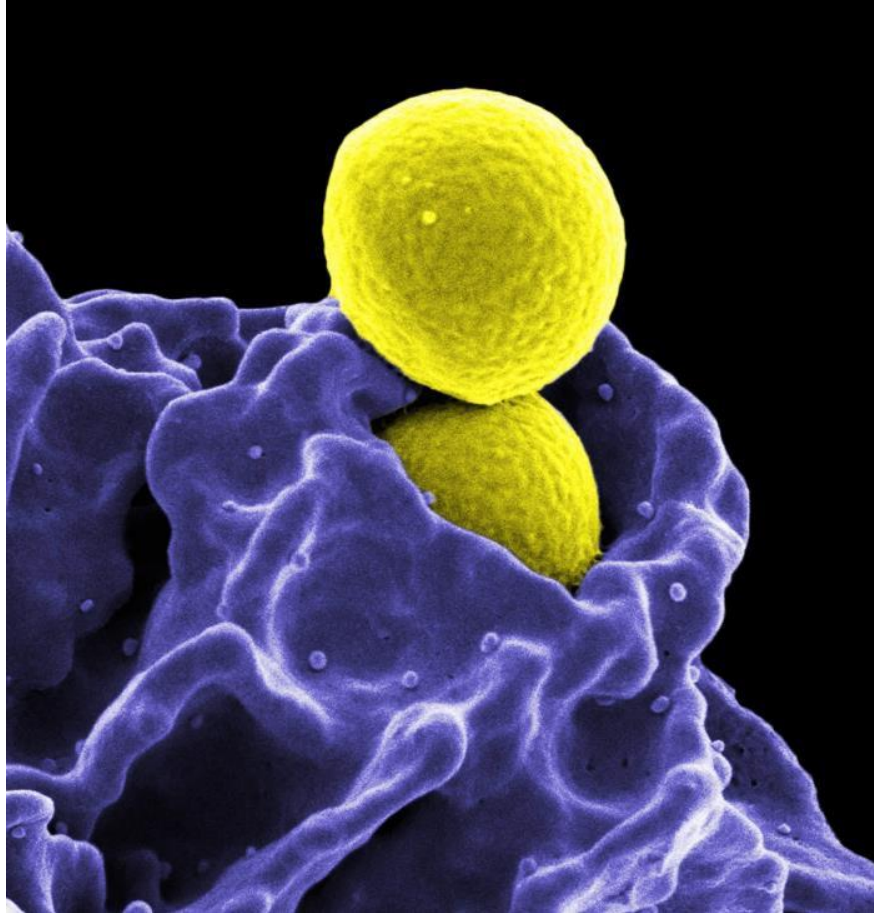
References

1. What is the Lifetime Risk of Needing and Receiving Long-term Services and Supports? ASPE Research Brief from the HHS Office of the Assistant Secretary for Planning and Evaluation; Office of disability, aging, and Long-term Care Policy. 2019, April. Retrieved 3/22/24 from https://aspe.hhs.gov/sites/default/files/migrated_legacy_files//188046/LifetimeRisk.pdf
2. Burdsall D, Schweon W, Collier S. A Unit Guide to Infection Prevention for Long-Term Care Staff. Agency for Healthcare Research & Quality (AHRQ); Healthcare-Associated Infections Program Toolkit. 2017, March. Retrieved 3/21/24 from ahrq.gov/hai/quality/tools/cauti-ltc/modules/resources/guides/infection-prevent.html
3. U.S. Department of Health and Human Services. National Action Plan To Prevent Healthcare-Associated Infections: Road Map to Elimination. Chapter 8, Long-Term Care Facilities. April 2013. <http://health.gov/hcgp/pdfs/hai-action-plan-ltcf.pdf>. Accessed September 23, 2015.
4. Wagner LM, Roup BJ, Castle NG. Impact of infection preventionists on Centers for Medicare and Medicaid quality measures in Maryland nursing homes. *Am J Infect Control*. 2014 Jan;42(1):2-6. PMID: 24388467.
5. Siegel JD, Rhinehart E, Jackson M, Chiarello L, and the Healthcare Infection Control Practices Advisory Committee. Management of Multidrug-Resistant Organisms in Healthcare Settings, 2006. Centers for Disease Control and Prevention. <http://www.cdc.gov/hicpac/pdf/guidelines/MDROGuideline2006.pdf>. Accessed May 5, 2016.
6. Siegel JD, Rhinehart E, Jackson M, Chiarello L, and the Healthcare Infection Control Practices Advisory Committee. 2007 Guideline for Isolation Precautions: Preventing Transmission of Infectious Agents in Healthcare Settings. Centers for Disease Control and Prevention. <http://www.cdc.gov/hicpac/pdf/isolation/Isolation2007.pdf>. Accessed May 5, 2016.
7. Get Smart: Know When Antibiotics Work. Centers for Disease Control and Prevention. <http://www.cdc.gov/getsmart/community/index.html>. Accessed November 19, 2015. 6. Bloodborne Pathogens. Occupational Safety and Health Administration. https://www.osha.gov/pls/oshaweb/owadisp.show_document?p_table=STANDARDS&p_id=10051. Accessed November 19, 2015.
8. Bloch N, Männer J, Gardiol C, Kohler P, Kuhn J, Münzer T, Schlegel M, Kuster SP, Flury D. Effective infection prevention and control measures in long-term care facilities in non-outbreak and outbreak settings: a systematic literature review. *Antimicrob Resist Infect Control*. 2023 Oct 18;12(1):113. doi: 10.1186/s13756-023-01318-9. PMID: 37853477; PMCID: PMC10585745.
9. Hurd, Michael D., Pierre-Carl Michaud, and Susann Rohwedder. 2014. "The Lifetime Risk of Nursing Home Use." In *Discoveries in the Economics of Aging*, edited by David A. Wise (81- 109). Chicago, IL: University of Chicago Press.
10. Wound Care Infection Prevention Recommendations for Long-Term Care Facilities. Minnesota department of health. 2022, October. Retrieved 3/22/24 from <https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/woundcare.pdf>
11. McGuire J, Love E, Vlahovic TC, et al. The ABCSS System for Chronic Wound Management: A New Acronym for Lower Extremity Wound Management. *Wounds*. 2020;32(suppl 11):S1-S25.
12. Lipsky BA, Berendt AR, Deery HG, et al. Diagnosis and treatment of diabetic foot infections. *Clin Infect Dis* 2004; 39:885.
13. Butalia S, Palda VA, Sargeant RJ, et al. Does this patient with diabetes have osteomyelitis of the lower extremity? *JAMA* 2008; 299:806.
14. Grayson ML, Gibbons GW, Balogh K, et al. Probing to bone in infected pedal ulcers. A clinical sign of underlying osteomyelitis in diabetic patients. *JAMA* 1995; 273:721.
15. Álvaro-Afonso FJ, Lázaro-Martínez JL, Aragón-Sánchez FJ, et al. Does the location of the ulcer affect the interpretation of the probe-to-bone test in the diagnosis of osteomyelitis in diabetic foot ulcers? *Diabet Med* 2014; 31:112.
16. García Morales E, Lázaro-Martínez JL, Aragón-Sánchez FJ, et al. Inter-observer reproducibility of probing to bone in the diagnosis of diabetic foot osteomyelitis. *Diabet Med* 2011; 28:1238.
17. Lam K, van Asten SA, Nguyen T, et al. Diagnostic Accuracy of Probe to Bone to Detect Osteomyelitis in the Diabetic Foot: A Systematic Review. *Clin Infect Dis* 2016; 63:944.
18. Senneville E. Editorial Commentary: Probe-to-Bone Test for Detecting Diabetic Foot Osteomyelitis: Rapid, Safe, and Accurate-but for Which Patients? *Clin Infect Dis* 2016; 63:949.
19. Murphy C, Atkin L, Swanson T, Tachi M, Tan YK, Vega de Ceniga M, Weir D, Wolcott R. International consensus document. Defying hard-to-heal wounds with an early antibiotic intervention strategy: wound hygiene. *J Wound Care* 2020; 29(Suppl 3b):S1–28.
20. Punjateawakupt A, Napavichayanun S, Aramwit P. The downside of antimicrobial agents for wound healing. *Eur J Clin Microbiol Infect Dis*. 2019;38(1):39-54. doi:10.1007/s10096-018-3393-51.
21. Chlorhexidine (2016) In: Aronson JK (ed) *Meyler's side effects of drugs*, Sixteenth edn. Elsevier, Oxford, pp 239–248. <https://doi.org/10.1016/B978-0-444-53717-1.00474-1>
22. Karpinski TM, Szkaradkiewicz AK (2015) Chlorhexidine—pharmaco-biological activity and application. *Eur Rev Med Pharmacol Sci* 19(7):1321–1326
23. Giannelli M, Chellini F, Margheri M, Tonelli P, Tani A (2008) Effect of chlorhexidine digluconate on different cell types: a molecular and ultrastructural investigation. *Toxicol in Vitro* 22(2):308–317. <https://doi.org/10.1016/j.tiv.2007.09.012>
24. Hidalgo E, Dominguez C (2001) Mechanisms underlying chlorhexidine-induced cytotoxicity. *Toxicol in Vitro* 15(4–5):271–276
25. Hildago E, Bartolome R, Dominguez C: Cytotoxicity mechanisms of sodium hypochlorite in cultured human dermal fibroblasts and its bactericidal effectiveness. *Chemico-Biological Interactions* 2002; 139: 265-282.
26. Sakarya S, Gunay N, Karakulak M, Ozturk B, Ertugrul B. Hypochlorous Acid: an ideal wound care agent with powerful microbicidal, antibiofilm, and wound healing potency. *Wounds*. 2014;26(12):342-350.
27. Chindera K, Mahato M, Sharma AK, et al. The antimicrobial polymer PHMB enters cells and selectively condenses bacterial chromosomes. *Sci Rep*. 2016;6:23121. Published 2016 Mar 21. doi:10.1038/srep23121
28. Bishop AJ (2018) Using antimicrobial dressing to treat infected wounds. *The Diabetic Foot Journal* 21(3): 168–71
29. Jonas SK, Riley PA, Willson RL. Hydrogen peroxide cytotoxicity. Low-temperature enhancement by ascorbate or reduced lipotease. *Biochem J*. 1989;264(3):651-655. doi:10.1042/bj2640651
30. Pilcher BK, Wang M, Qin XJ, Parks WC, Senior RM, Welgus HG. Role of matrix metalloproteinases and their inhibition in cutaneous wound healing and allergic contact hypersensitivity. *Ann N Y Acad Sci*. 1999 Jun 30;878:12-24. doi: 10.1111/j.1749-6632.1999.tb07671.x. PMID: 10415717.
31. Doughty, DB & McNichol LL (editors). *Wound, Ostomy and Continence Nurses Society™ core curriculum: Wound management*. Philadelphia, PA: Wolters Kluwer; 2016. ISBN:9781496343604
32. Wickline S. Wounds Heal Better When Debrided Often. *MedPageToday*. 2013. Accessed online: www.medpagetoday.com/dermatology/generaldermatology/40692
33. Wilcox JR, Carter MJ, Covington S. Frequency of debridements and time to heal: a retrospective cohort study of 312 744 wounds. *JAMA Dermatol*. 2013;149(9):1050-8.
34. Bryant, R, & Nix, D. *Acute & Chronic Wounds: Current management concepts*, fourth edition. St Louis, MO: Elsevier. 2012. ISBN: 9780323069434
35. Shakeri H, Lemmens K, Gevaert AB, De Meyer GRY, Segers VFM. Cellular senescence links aging and diabetes in cardiovascular disease. *Am J Physiol Heart Circ Physiol*. 2018 Sep 1;315(3):H448-H462. doi: 10.1152/ajpheart.00287.2018. Epub 2018 May 11. PMID: 29750567.
36. Regulski, M. Understanding Diabetic Induction of Cellular Senescence: A concise review. 2018. *WOUNDS*, 30(4), 96-101.
37. Winter GD. Formation of the scab and the rate of epithelization of superficial wounds in the skin of the young domestic pig. *Nature*. 1962;193:293-294. doi:10.1038/193293a0
38. Bryant, R, & Nix, D. *Acute & Chronic Wounds: Current management concepts*, fourth edition. St Louis, MO: Elsevier. 2012. ISBN: 9780323069434
39. Ovington LG. Hanging wet-to-dry dressings out to dry. *Home Healthc Nurse*. 2001;19(8):477-484. doi:10.1097/00004045-200108000-00007
40. de Wert LA, Schoonhoven L, Stegen JHCH, et al. Improving the effect of shear on skin viability with wound dressings. *J Mech Behav Biomed Mater*. 2016;60:505-514. doi:10.1016/j.jmbbm.2016.03.006
41. Cordrey R. Gauze, Impregnated Gauzes, and Contact Layers. *Adv Skin Wnd Care*. 2012. 1:120-125.
42. Lawrence JC. Dressings and wound infection. *Am J Surg*. 1994;167(1A):215-245. doi:10.1016/0002-9610(94)90006-x
43. Rippon M, Davies P, White R. Taking the trauma out of wound care: the importance of undisturbed healing. *J Wound Care*. 2012;21(8):359-368. doi:10.12968/jowc.2012.21.8.359
44. *Biggest Threats & Data*. (2020, June 18). Center for Disease Control & Prevention. <https://www.cdc.gov/drugresistance/biggest-threats.html>
45. Dolk FCK, Pouwels KB, Smith DRM, Robotham JV, Smieszek T. Antibiotics in primary care in England: which antibiotics are prescribed and for which conditions?. *J Antimicrob Chemother*. 2018;73(suppl_2):ii2-ii10. doi:10.1093/jac/dkx504
46. Siddiqui AR, Bernstein JM. Chronic wound infection: facts and controversies. *Clin Dermatol*. 2010;28:519–526
47. Caputo WJ, Monterosa P, Beggs D. Antibiotic Misuse in Wound Care: Can Bacterial Localization through Fluorescence Imaging Help?. *Diagnostics (Basel)*. 2022;12(12):3207. Published 2022 Dec 17. doi:10.3390/diagnostics12123207

References

48. Metcalf DG, Bowler PG. Biofilm delays wound healing: A review of the evidence. *Burns Trauma*. 2013;1(1):5-12. Published 2013 Jun 18. doi:10.4103/2321-3868.113329
49. Rodrigues CF, Kaushik KS, Light C. Biofilms in Wounds: New Advances in Therapy and in Healing Management. *Biomedicines*. 2021;9(2):193. Published 2021 Feb 16. doi:10.3390/biomedicines9020193
50. Swoboda, L. Biofilm-Based Wound Management: Implications for Clinical Practice. *Wound Management & Prevention*. 2022;68(8):5,6.
51. Metcalf DG, Bowler PG. Biofilm delays wound healing: A review of the evidence. *Burns Trauma*. 2013;1(1):5-12. Published 2013 Jun 18. doi:10.4103/2321-3868.113329
52. Warrior A, Satyamoorthy K, Murali TS. Quorum-sensing regulation of virulence factors in bacterial biofilm. *Future Microbiol*. 2021;16:1003-1021. doi:10.2217/fmb-2020-0301
53. Ciaccia L. Fundamentals of Inflammation. *Yale J Biol Med*. 2011;84(1):64-65.
54. Haalboom M, Blokhuis-Arkes MHE, Beuk RJ, et al. Culture results from wound biopsy versus wound swab: does it matter for the assessment of wound infection?. *Clin Microbiol Infect*. 2019;25(5):629.e7-629.e12. doi:10.1016/j.cmi.2018.08.012
55. Stallard Y. When and How to Perform Cultures on Chronic Wounds?. *J Wound Ostomy Continence Nurs*. 2018;45(2):179-186. doi:10.1097/WON.0000000000000414
56. Copeland-Halperin LR, Kaminsky AJ, Bluefeld N, Miraliakbari R. Sample procurement for cultures of infected wounds: a systematic review. *J Wound Care*. 2016;25(4):S4-S10. doi:10.12968/jowc.2016.25.Sup4.S4
57. James GA, Swogger E, Wolcott R, et al. Biofilms in chronic wounds. *Wound Repair Regen*. 2008;16(1):37-44. doi:10.1111/j.1524-475X.2007.00321.x
58. Armstrong DG, Edmonds ME, Serena TE. Point-of-care fluorescence imaging reveals extent of bacterial load in diabetic foot ulcers. *Int Wound J*. 2023;20(2):554-566. doi:10.1111/iwj.14080
59. Biggest Threats & Data. (2020, June 18). Center for Disease Control & Prevention. <https://www.cdc.gov/drugresistance/biggest-threats.html>
60. Dolk FCK, Pouwels KB, Smith DRM, Robotham JV, Smieszek T. Antibiotics in primary care in England: which antibiotics are prescribed and for which conditions?. *J Antimicrob Chemother*. 2018;73(suppl_2):ii2-ii10. doi:10.1093/jac/dkx504
61. Sasseville, Denis Neomycin, Dermatitis: January-February 2010 - Volume 21 - Issue 1 - p 3-7. doi: 10.2310/6620.2009.09073
62. Biggest Threats & Data. (2020, June 18). Center for Disease Control & Prevention. <https://www.cdc.gov/drugresistance/biggest-threats.html>
63. Jacob SE, Nijhawan RI. (2003). Focus On: Bacitracin Allergen of the Year 2003. *The Dermatologist*. <http://www.the-dermatologist.com/content/focus-on-bacitracin-allergen-year-2003>
64. Neomycin Sulfate. (2020). Contact Dermatitis Institute. <https://www.contactdermatitisinstitute.com/neomycin-sulphate.php>
65. Dire DJ, Coppola M, Dwyer DA, Lorette JJ, Karr JL. Prospective evaluation of topical antibiotics for preventing infections in uncomplicated soft-tissue wounds repaired in the ED. *Acad Emerg Med* 1995;2:4-10.
66. Langford JH, Artemi P, Benrimoj SI. Topical antimicrobial prophylaxis in minor wounds. *Ann Pharmacotherapy* 1997;31:559-563.
67. Delves, P. J., & Roitt, I. M. (1998). *Encyclopedia of immunology*. Academic Press.
68. Ramos G, Cornistein W, Cerino GT, Nacif G. Systemic antimicrobial prophylaxis in burn patients: systematic review. *J Hosp Infect*. 2017;97(2):105-114. doi:10.1016/j.jhin.2017.06.015
69. Barajas-Nava LA, López-Alcalde J, Roqué i Figuls M, Solà I, Bonfill Cosp X. Antibiotic prophylaxis for preventing burn wound infection. *Cochrane Database Syst Rev*. 2013;(6):CD008738. Published 2013 Jun 6. doi:10.1002/14651858.CD008738.pub2
70. Norman G, Christie J, Liu Z, et al. Antiseptics for burns. *Cochrane Database Syst Rev*. 2017;7(7):CD011821. Published 2017 Jul 12. doi:10.1002/14651858.CD011821.pub2
71. Nímia HH, Carvalho VF, Isaac C, Souza FÁ, Gemperli R, Paggiaro AO. Comparative study of Silver Sulfadiazine with other materials for healing and infection prevention in burns: A systematic review and meta-analysis. *Burns*. 2019;45(2):282-292. doi:10.1016/j.burns.2018.05.014
72. Nešporová K, Pavlík V, Safránková B, et al. Effects of wound dressings containing silver on skin and immune cells [published correction in *Sci Rep*. 2021 Feb 17;11(1):4369]. *Sci Rep*. 2020;10(1):15216. Published 2020 Sep 16. doi:10.1038/s41598-020-72249-3
73. McNally L, Brown SP. Visualizing evolution as it happens. *Science*. 2016;353(6304):1096-1097. doi:10.1126/science.aah5641
74. Werth, B. Merck Manual Professional Version. Overview of Antibacterial Drugs. Reviewed/Revised May 2022. Modified Sep 2022. Retrieved 2/13/24 from <https://www.merckmanuals.com/professional/infectious-diseases/bacteria-and-antibacterial-drugs/overview-of-antibacterial-drugs#v1002013>
75. Centers for Disease Control and Prevention, National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Division of Healthcare Quality Promotion (DHQP). Last Reviewed November 5, 2015. Edited October 2017 and October 2022. Retrieved 2/11/24 from <https://www.cdc.gov/infectioncontrol/guidelines/mdro/prevention-control.html>
76. Arendrup MC, Patterson TF. Multidrug-Resistant Candida: Epidemiology, Molecular Mechanisms, and Treatment. *J Infect Dis*. 2017;216(suppl_3):S445-S451. doi:10.1093/infdis/jix131
77. Wound Care Infection Prevention Recommendations for Long-Term Care Facilities. Minnesota department of health. 2022, October. Retrieved 3/22/24 from <https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/woundcare.pdf>
78. Minnesota Pollution Control Agency: Infectious Waste Management Guidance for Transporters (www.pca.state.mn.us/sites/default/files/w-sw4-31.pdf)

Resources: Infection Prevention Basics

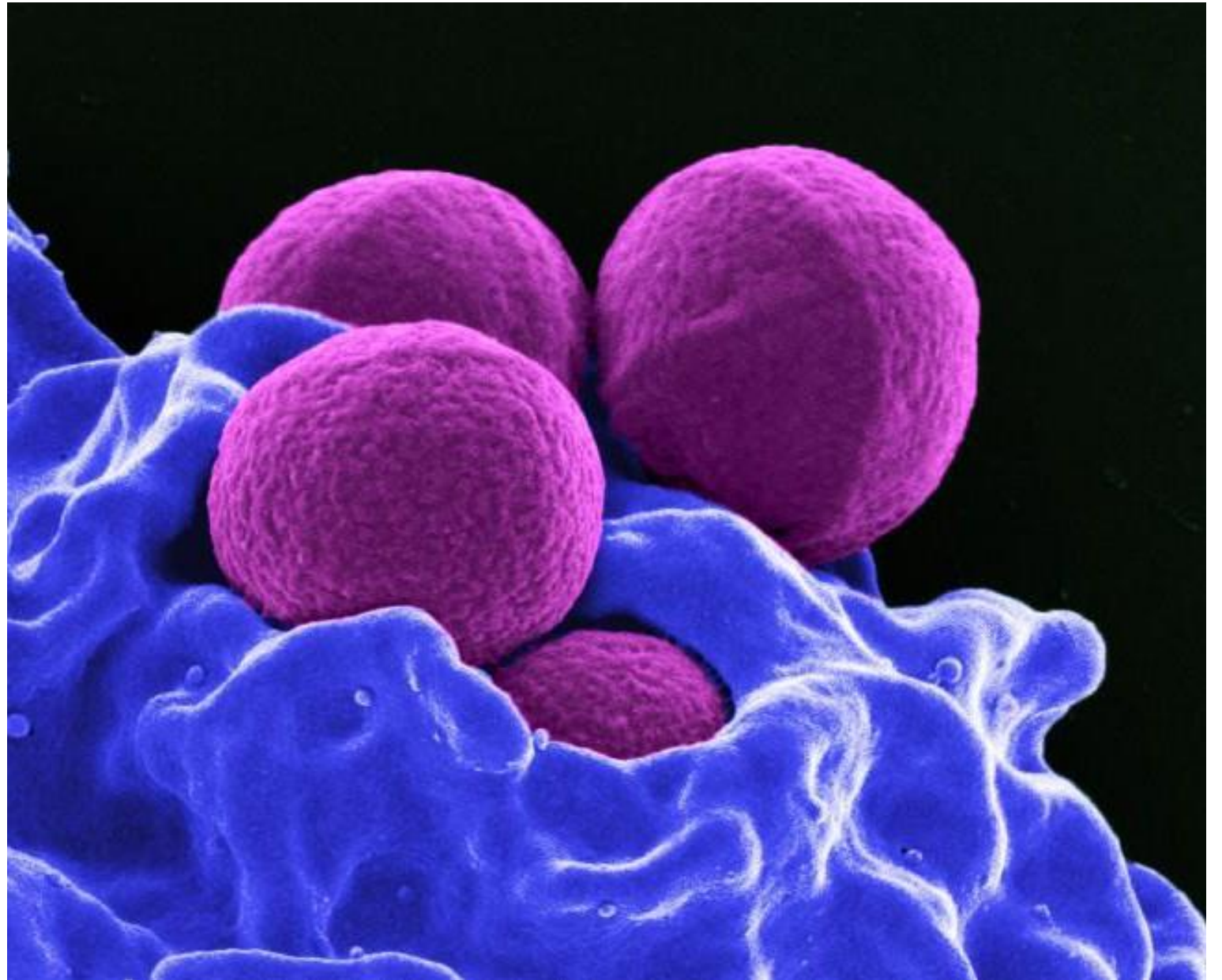


- CDC: Guideline for Isolation Precautions: Preventing Transmission of Infectious Agents in Healthcare Settings (www.cdc.gov/infectioncontrol/pdf/guidelines/isolation-guidelines-H.pdf)
- CDC: Implementation of Personal Protective Equipment (PPE) in Nursing homes to Prevent the Spread of Multidrug- Resistant Organisms (MDROs) (www.cdc.gov/hai/containment/PPEinNursingHomes.html)
- Wound Care Infection Prevention Recommendations for Long-Term Care Facilities. Minnesota department of health. 2022, October. Retrieved 3/22/24 from <https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/woundcare.pdf>
- CDC: Guideline for Disinfection and Sterilization in Healthcare Facilities (www.cdc.gov/infectioncontrol/pdf/guidelines/disinfection-guidelines-H.pdf)
- CDC: Healthcare Providers | Hand Hygiene (www.cdc.gov/handhygiene/providers/index.html)

Thank you! Questions?

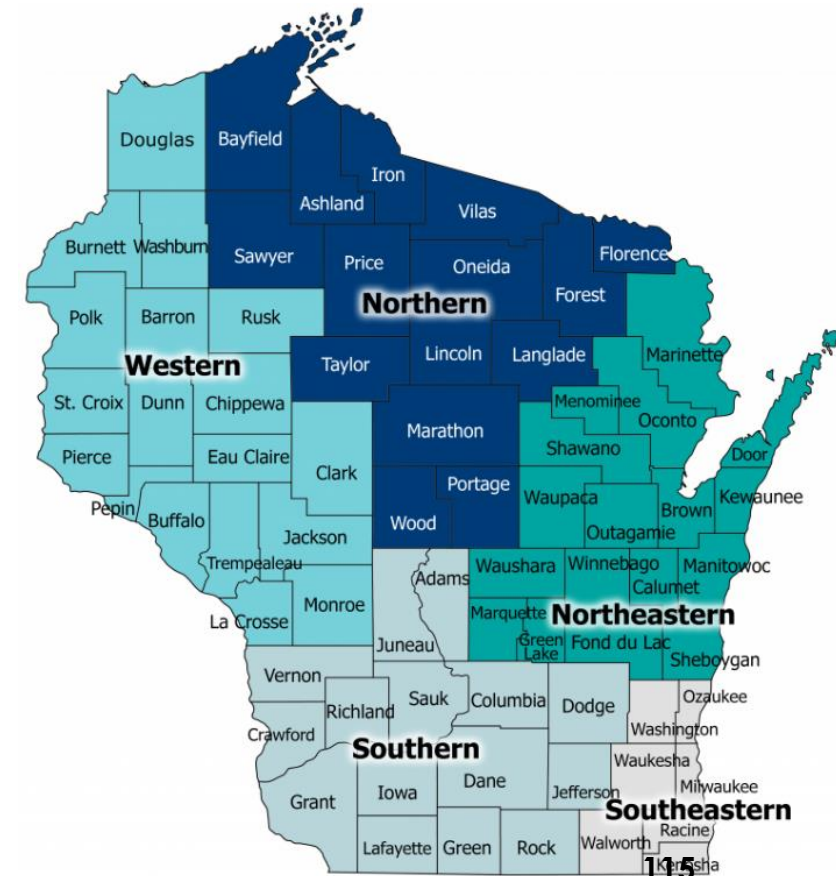
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HAI Infection Prevention Education webpage

HAI Infection Prevention Education

The resources below are intended to connect health care facility infection preventionists (IP) with education materials to support their role in preventing, detecting, and responding to healthcare-associated infections.

IPs play an essential role in facility infection prevention policy development, surveillance, and risk assessment.

IPs serve as a resource to other staff and programs within their facilities.

In addition to the state in-person trainings and online references below, there are a number of links to trusted education resources, including the CDC (Centers for Disease Prevention and Control), the Centers for Medicare and Medicaid Services (CMS), and the Association for Professionals in Infection Control and Epidemiology (APIC).



The [IP Starter Kit](#) provides Infection Preventionists a brief background and resources for some of the many infection prevention-related responsibilities within health care facilities.

Resources for infection preventionists Long-Term Care Education series

The long-term care (LTC) education series provides education presentations on topics that include infection prevention, HAIs, antibiotic stewardship, disease surveillance, and outbreak response for staff at skilled nursing facilities, assisted living facilities, local health departments, and other LTC stakeholders. Each session features a new, timely topic presented by the Department of Health Services (DHS) program staff, HAI Infection Preventionists, partner organizations, or other external subject matter experts.

View the [full library](#) of education sessions. **Note:** All 2021 and 2022 education sessions can be found by visiting the full library

Have a topic request?

Send topic ideas or requests that you have for the long-term care education series or the IP lunch and learn series to DHSWIHAIPreventionProgram@dhs.wi.gov

Upcoming LTC Education Session

Date: April 18, 2024

**Topic: Multidrug-Resistant Organisms in
Wisconsin**



**WISCONSIN DEPARTMENT
of HEALTH SERVICES**