Objectives

- Describe basic principles of cleaning, disinfection, sterilization
- Identify when to use cleaning, disinfection, or sterilization
- Describe how to monitor cleaning, disinfection and sterilization processes
Terminology

- **Cleaning**
  - general removal of debris (dirt, food, feces, blood, saliva and other body secretions)
  - reduces amount of organic matter that contributes to proliferation of bacteria and viruses

- **Disinfection**
  - removes most organisms present on surfaces that can cause infection or disease

- **Sterilization**
  - the killing or removal of all organisms
Cleaning, Disinfection and Sterilization in Healthcare Settings

- Practice standards are based on Spaulding’s Classification system
- Healthcare devices and equipment designated as
  - Critical
  - Semi-critical
  - Non-critical
- Categories define level of reprocessing required
Critical Items

• Require sterilization
• Includes items that enter sterile tissue or the vascular system
• Examples include surgical instruments and accessories, biopsy forceps, cardiac and urinary catheters, implants, needles
Semi-Critical Items

• Require minimum high level disinfection (or sterilization)
• Includes items in contact with non-intact skin or mucous membranes
• Examples include respiratory therapy equipment, anesthesia equipment, flexible and laryngoscopes, bronchoscopes, GI endoscopes, cystoscopes, vaginal ultrasonic probes
• Cleaning process must precede high-level disinfection
Non-Critical Items

- Require intermediate-level or low-level disinfection
- Includes items in contact only with intact skin
- Examples include BP cuffs, stethoscopes, durable mobile patient equipment
Environmental Cleaning

- Patient environment can facilitate transmission of bacteria and viruses
  - By direct contact
  - On hands of healthcare personnel
- Contaminated surfaces increase potential for transmission of bacteria and viruses between patients
- Items categorized as non-critical (intermediate or low disinfection) or require cleaning only

\[ \times \text{ represents VRE culture positive sites} \]
Policy Considerations

• Include in policy all surfaces and equipment that can reasonably be expected to be contaminated by bacteria (high touch surfaces)
• Define responsibility and frequency for cleaning and disinfecting patient care equipment and surfaces
• Monitor compliance with policy
• Staff should be able to answer question “How do you know whether this item has been cleaned and/or disinfected?”
• Cleaned/disinfected items should be labeled (date/time)
High Touch Surfaces in Patient Rooms

- Considered non-critical
- Must be cleaned *then* disinfected on a regular basis
- Examples include:
  - Bedrails
  - Call bell
  - Telephones
  - TV remote
  - IV pump
  - IV poles
  - Toilet, commode chair
  - Overbed table
  - Light switches
  - Doorknobs
  - Respiratory and other bedside equipment
  - Computer keyboard
  - Chairs
Increased acquisition risk from prior room occupant
6 studies as of January 2011

- Huang
  - MRSA
  - VRE
- Hardy
  - MRSA
- Dress
  - VRE
- Shaugnessy
  - C. difficile
- Datta
  - MRSA
- Nseir
  - Pseudomonas
- Nseir
  - Acinetobacter

Average = 120%

Increased Risk of Acquisition (%)

Items Requiring *only* Cleaning

- Floors, walls, and windows
- Chairs and other furniture used by individuals who are clothed
- Private offices and other non-public, non-patient care areas
- Bed curtains should be changed when soiled and w/ terminal cleaning

*Clarify in policy what needs to be cleaned and not necessarily disinfected*
Use Microfiber for Cleaning

- Densely constructed synthetic strands ~1/16\textsuperscript{th} the diameter of a human hair
- Attracts dust, cleans ~50% better than comparable cotton
- Easier to use, lighter, designed for repeat usage

HICPAC Disinfection & Sterilization Guideline 2008, Rutala
Monitor Environmental Cleaning Processes

• Bioluminescence (outcome measure)
  • Monitors for light emissions produced if organism present
  • Results difficult to interpret because it is unknown whether organism remains viable and thus transmissible
  • Expensive

• Fluorescence (process measure)
  • Monitors for chemical markers that fluoresce with ultraviolet (black) light if not removed during cleaning

• Culturing
  • Should *not* be done except during some outbreak investigations

• Visual inspection
  • Make routine rounds and provide feedback to frontline staff
Linens

- All linen handled as if contaminated with blood or body fluids (Standard Precautions)
  - Bag linen at point of use
  - Wear PPE when sorting and agitate minimally
- Laundry equipment must be maintained to prevent microbial contamination*
- New laundry technologies allow linen washing without requirements for hot water and chlorine
  - Hot water - 160°F x 25 min
  - Cold water - 71-77°F with 125 ppm chlorine bleach rinse or equivalent detergent
- Detergents not required to have stated anti-microbial claims*

*Manufacturer’s instructions for use must be followed
Cleaning, Disinfection, and Sterilization of Medical Instruments and Devices

You CANNOT achieve disinfection or sterilization without pre-cleaning
  • As organic material dilutes disinfectants, bioburden must be reduced for processes to be effective

Clean all medical instruments and devices as a first step
  • Remove visible soil
  • May need to disconnect or separate instrument parts
  • Avoid organic material drying on equipment by rinsing or soaking in an enzymatic solution
Personal Protection

When cleaning soiled medical instruments, wear

- Long sleeved impervious gown
- Eyewear
- Mask or mask with face shield
- Gloves
- Cap
- Chemical goggles (when mixing or changing solution)
Disinfection

- Eliminates or kills most bacteria, many virus types, some fungi (not prions)
- Cannot be accomplished without first cleaning
- Time-dependent process
- Levels of disinfection - high, intermediate, or low
- Hospitals must use EPA-approved product for desired level of disinfection
  - Has minimally a tuberculocidal label claim
Disinfection - continued

• Follow manufacturer’s recommendations to achieve disinfection and to avoid medical device damage method
  ▫ Use correct dilution – more is not better!
  ▫ Use correct contact time
  ▫ Use correct temperature

• Understand employee and environmental safety issues
  ▫ Do not exceed exposure limits
  ▫ Know permissible exposure levels
  ▫ Assess compatibility with gloves, basins, other products
EPA Registration of Disinfectants

- Labeled as high level vs. intermediate vs. low level
- May include degrees of approval
  - Limited approval, e.g. kills Hepatitis B and HIV but not approved for spores
- Select disinfectant based on what you are trying to accomplish
  - Environmental vs. medical device disinfection
- Can search EPA website by product name
  
  www.epa.gov/oppad001/chemregindex.htm
High-level Disinfection - Glutaraldehyde

- Ensure achievement of temperature requirements
- Test product prior to each use
  - Can get diluted with frequent use
  - Follow facility policy
  - Test strips expire; monitor dates
- Change product as indicated by test and as manufacturer requires
- Maintain log records
- Ensure competency of staff
Endoscopes/Bronchoscopes

- United States
  - Infection: 1/1.8 million procedures
- Professional organization guidelines
  - Minimum high-level disinfection
- Ensure competency of personnel performing process
- Outbreaks associated with failure to comply with guidelines for disinfection/sterilization

Ambulatory Surgery in the United States, 2006. NHSR Number 11.26pp
Endoscopy/Bronchoscopy Associated Infections

- **Endoscopy**
  - >280 Infections transmitted, some fatal
  - >70%: *Salmonella* and *Pseudomonas aeruginosa* (others: HBV, *Strongyloides stercoralis*, H. *pylori*, *Trichosporan*)

- **Bronchoscopy**
  - >90 documented infections transmitted
  - Mycobacteria, *Pseudomonas aeruginosa*
  - Mycobacteria are resistant to many disinfectants
  - High level disinfectants
    - 2% glutaraldehyde at 20°C for 20min is most common

Weber, DJ, Rutala WA. ICHE 2001: 22:403-408
The 5 Steps of Endoscope Re-Processing

1. Clean
   • Remove debris/tissue which can impede disinfection process, flush all lumens (water & enzymatic cleaner)

2. High Level Disinfection
   • Perfuse through ALL channels with disinfectant

3. Rinse
   • Sterile or filtered water/tap water followed by alcohol rinse

4. Dry
   • Forced air

5. Store
   • Hang vertically – Promote drying & Avoid recontamination
The 5 steps of Endoscope Re-Processing - continued

- To avoid problems, the 5 steps must be performed in sequence
- Do not skip, bypass, shortcut any of the 5 steps
Factors Leading to Outbreaks from Endoscope/Bronchoscope Contamination

- Contaminated water supply
- Contaminated brushes for cleaning scope lumens
- Improper manual cleaning prior to disinfection
- Biofilm inside automatic washer
- Improper use of automatic washer
- Contaminated or expired disinfection reagent
- Inability or neglect to clean the suction channel
- Mechanical or design issues related to the endoscope/bronchoscope
Environmental Disinfectants

- Phenolics
  - “Gold Standard” in healthcare
  - Toxicity concerns prohibit use in nurseries, NICU
  - Does not kill spores

- Quaternary ammonium compounds
  - Approved for specific pathogens (read the label!)
  - Affected by water hardness
  - Affected by bioburden

Correct dilution is critical to effectiveness.
Environmental Disinfectants - continued

- Iodophors
  - Can be used in food preparation areas
  - Inactivated by organic materials, e.g. blood
  - Can stain surfaces

- Chlorine (bleach)
  - Inactivated by organic materials, e.g. blood
  - Kills spores, e.g. *C. difficile*
  - Corrosive
  - Highly toxic (deadly) if combined with ammonia
Environmental Disinfectants - continued

• Disinfectant spray-fog techniques for antimicrobial control in hospital rooms
  • Unsatisfactory method of decontaminating air and surfaces
  • Not recommended for general infection control in routine patient-care areas

• Ultraviolet Radiation
  • Dependent on strength and duration of exposure to light, ‘line of sight’, how well microorganism can withstand UV
  • Limited to destruction of airborne organisms, inactivation of microorganisms on surfaces, and water purification
Sterilization

Achieved by

• Steam
• Dry Heat
• Ethylene Oxide
• Peracetic Acid
• Plasma Gas (vaporized hydrogen peroxide)
• Glutaraldehyde (using higher concentrations and exposure times than for high-level disinfection)
Steam Sterilization - Autoclave

- Achieves rapid heating and penetration
  - Short exposure times (<20 minutes) but temperature must be maintained throughout
  - No toxicity to workers
  - Inexpensive
  - Can damage delicate instruments

- Items to be sterilized must be
  - Clean and free of protein (blood) or other organic material
  - Packaged so that the steam can penetrate

- Autoclave must be loaded correctly
Rapid Cycle or Flash Sterilization

- “Unwrapped” steam sterilization
- Should only be used when necessary
  - Do not flash whole trays of instruments
  - Items must be used immediately
  - Avoid by keeping adequate supply of frequently dropped items
- Maintain records or “flash logs”
  - Include all implants
  - Requires same monitoring processes as routine steam sterilization in hospital
  - Use to support need for additional instruments
Monitoring Sterilization

- **Mechanical Indicators**
  - Gauges, displays, printouts
  - Indicates if device working properly
  - Not indicator of sterility

- **Chemical Indicators**
  - Change color with timed exposure to heat, steam
  - Not indicator of sterility
  - Used to show items have gone through sterilization process

- **Biological Indicators**
  - Indicator of sterility
  - Demonstrates bacterial spores on test strips or in vials/containers have all been killed
  - Results can be available in 1 hour
Storage of Sterile Items

- Protect sterility until ready to use
  - **Store to protect packages from dust, moisture, falling on floor**
  - **Transport only covered, dry packages**
  - **Handle to protect package integrity**
- Rotate sterile items first in, first out
- Store and label for effective recall system
- Expiration date vs. Event-related sterilization
  - **Needs a program flex from L&C**
IP Role in Cleaning, Disinfection, and Sterilization

- Know the processes; update the policies
- Know directors of environmental services, sterile processing, operating room, endoscope services
- Know where all sterilization and disinfection is being done
  - May include
    - Radiology
    - GI dept
    - Cardiac cath lab
    - Wound care center
    - Outpatient clinics
    - Emergency room
    - Same day procedures
    - Ambulatory surgery
- Ensure staff know and follow contact times for products
  - Per manufacturer guidelines; on labels
Questions?

For more information, please contact any HAI Liaison Team member

Thank you