

STERILIZATION AND DISINFECTION

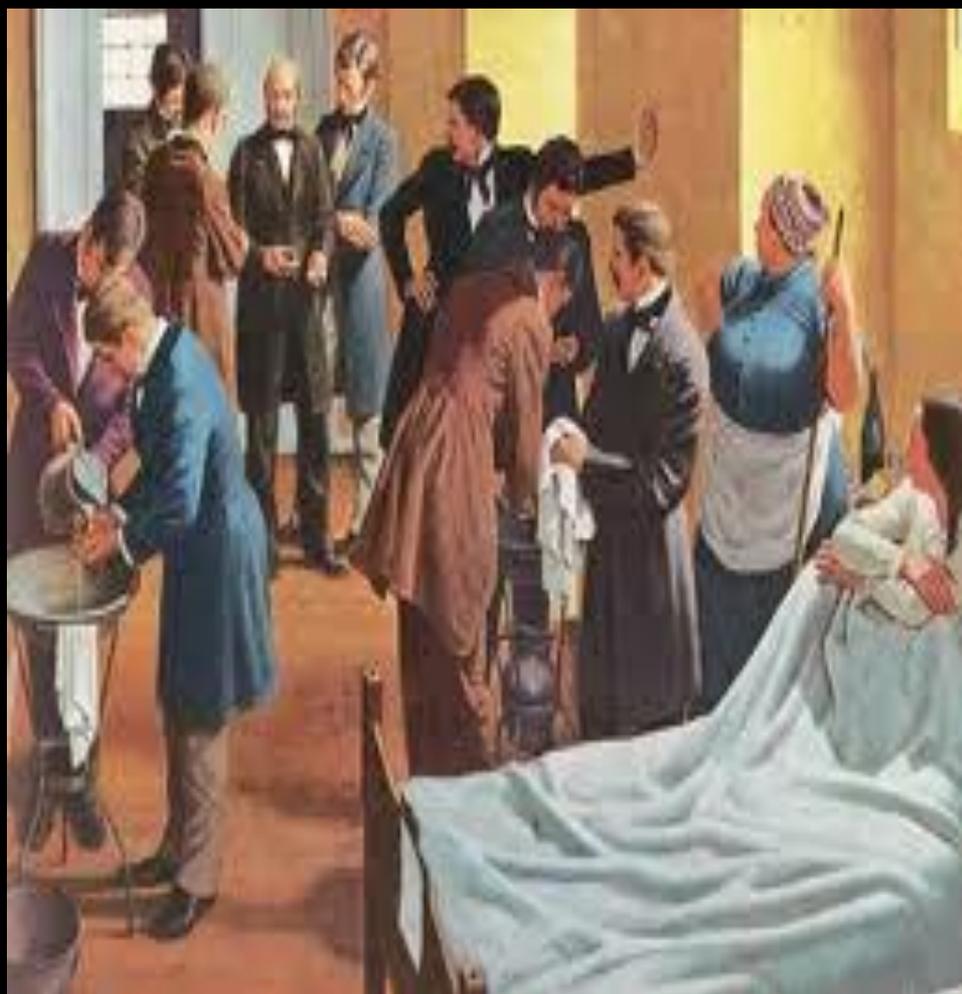
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INFECTIOUS DISEASES AND CLINICAL MICROBIOLOGY

History

- Heating, salting, drying in the sun
- Infected wounds in the middle ages: the use of mercury
- Phenol, wine, vinegar, chlorine : antiseptic
- Robert Boyle fermentation/ disease relationship (1663)
- Nicolas Appert production of canned (1810)
- Pasteur pasteurization and flame (1850)



Semmelweis Wien Medical Faculty

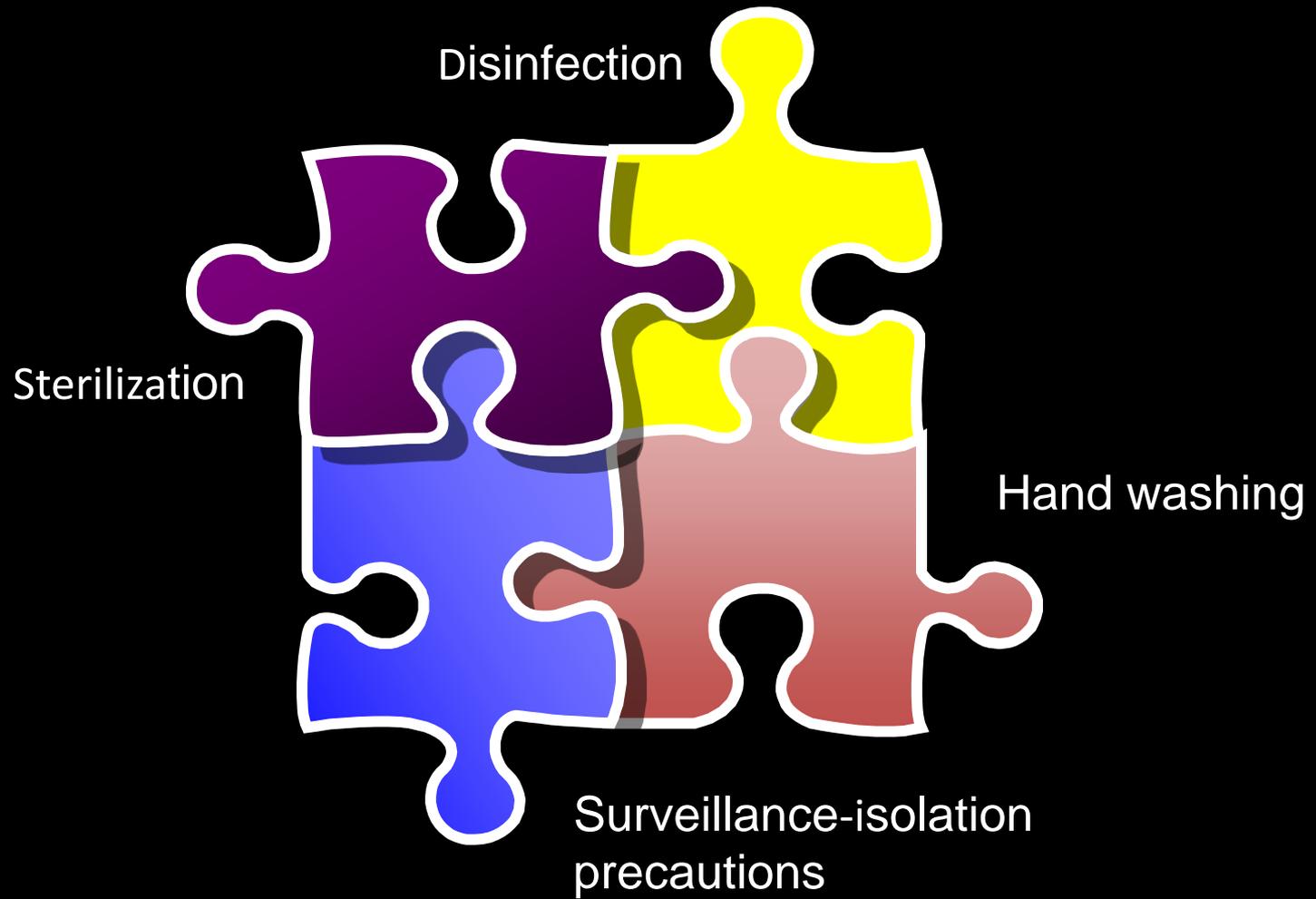
- Transport of microorganisms from autopsy room to the ward : first scientist



Semmelweis Wien Medical Faculty

- Hand Washing: The main behaviour in the clinics today





**S T E R I L
A L A N**

**T E M I Z
A L A N**

**K I R L I
A L A N**



Introduction

- Microorganisms are the agents of
 - Contamination
 - Infection
 - Decay

- Hence it becomes necessary to remove them from materials and areas.

Introduction

- In mid 18 century Lister developed Aseptic Techniques to prevent contamination of surgical wounds.
- Prior to this development:
 - Nosocomial infections caused death in >10% of surgeries
 - Up to 25% mothers delivering in hospitals died due to infection

Definitions

STERILIZATION is the total elimination of all microorganisms including spores

- Typically the last things to die are the highly heat- and chemical-resistant bacterial endospores
- Instruments used for invasive procedures must be sterilized prior to use
- Moist heat or steam, radiation, chemicals (e.g., glutaraldehyde), and ethylene oxide (a gas) are employed for sterilization
- Sterilization by autoclaving, which uses moist heat, is used in most hospital and microbiology laboratory settings

Definitions

DISINFECTION is the elimination of pathogens, except spores, from inanimate objects

- Disinfectants are chemical solutions used to clean inanimate objects

(physical processes, e.g., UV radiation, may also be employed to effect disinfection)

- Germicides are chemicals that can be applied to both animate (living) and inanimate objects for the purpose of eliminating pathogens
- Antiseptics are formulated for application to living tissue

Definitions

- **SEPSIS :**

Comes from Greek for decay or putrid.
Indicates bacterial contamination.

- **ASEPSIS :**

Absence of significant contamination on
inanimate surfaces

- **ANTISEPSIS :**

Reduction or Inhibition of microbes found
on living tissue

Definitions

- **Aseptic techniques** are used to prevent contamination of surgical instruments, medical personnel, and the patient during surgery.
- Aseptic techniques are also used to prevent bacterial contamination in food industry.

Definitions

- BACTERIOSTATIC AGENT :

An agent that inhibits the growth of bacteria, but does not necessarily kill them.

- BACTERICIDE AGENT :

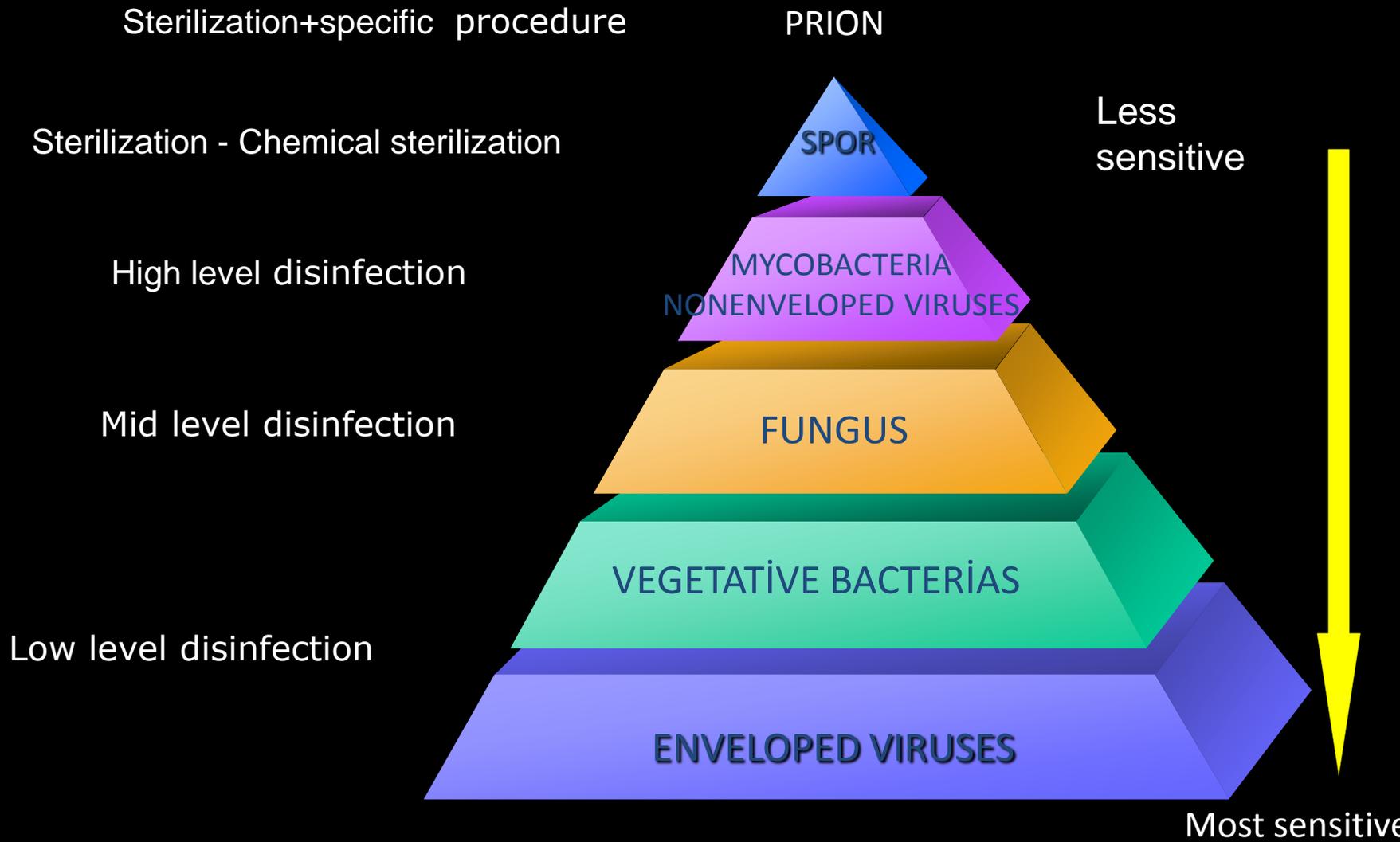
An agent that kills bacteria. Most do not kill Endospores

- SPOROCIDE AGENT :

An agent that kills spores

Definitions

- **SANITIZATION** : Lowering of microbial counts to prevent transmission in public setting (e.g., restaurants & public rest rooms)
- **DEGERMING** : Mechanical removal of microbes, e.g., from hands with washing



SPAULDING classifications (1960)

- Medical and surgical equipments can be divided 3 group : depends on the make an infection capability
 - CRITICAL
 - SEMICRITICAL
 - NONCRITICAL

SPAULDING classifications

- Critical devices : Penetration to the steril tissues, must be sterilized
- Critical objects which enter normally
 - Sterile tissue
 - Vascular system
 - Blood flows

SPAULDING classifications

- Semicritical devices : objects that touch mucous membranes or skin
- Devices is not intact with steril tissue
- Sterilization preferred
- A disinfection process (High level disinfection- HLD) that kills all microorganisms but HLD alternative

SPAULDING classifications

- Noncritical devices: object that touch only intact skin
- Low level disinfection (LLD) enough

Critical objects



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Processing Critical Objects

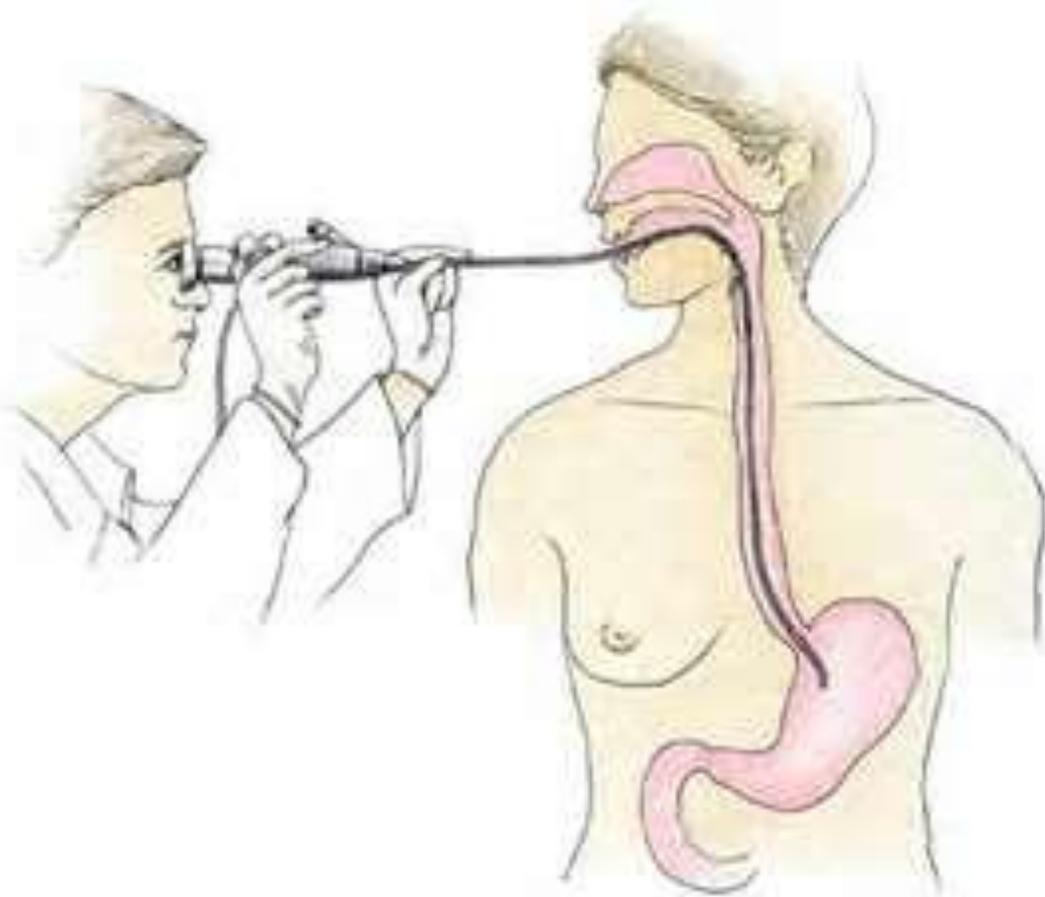
- Classification : enter sterile tissue
- Object : must sterilized
- Level germicidal action: kill all m.o. and spores
- Examples : surgical instruments and devices
cardiac catheters, implants, etc
- Method : Steam, ETO, Hydrogen peroxide
plasma, chemical sterilization

Chemical sterilization of critical objects

- Glutaraldehyde ($\geq 2.0\%$)
- Hydrogen peroxide-HP (7.5%)
- Peracetic acid-PA (0.2%)
- HP(1%) and PA (0.08%)
- HP (7.5%) and Phenol (1.93%)

- Exposure time per manufacturers recommendations

Semicritical objects



Processing semicritical objects

- Classification : Contact with mucous membranes or skin that is not intact
- Object: Free of all microorganisms
- Level germicidal action: Kill all microorganisms except high numbers of bacterial spores
- Examples: Respiratory-anesthesia equipment, GI endoscopes, thermometer, etc
- Method: Sterilization or high level disinfection

HLD of semicritical objects

- Glutaraldehyde $\geq 2\%$
- Ortho-phthalaldehyde 0.55%
- Hydrogen peroxide 7.5 %
- HP and peracetic acid 7.5% - 0.23%
- Hypochlorite 650-675 ppm
- Glutaraldehyde and phenol 1.21% - 1.93%

Noncritical objects



Processing noncritical objects

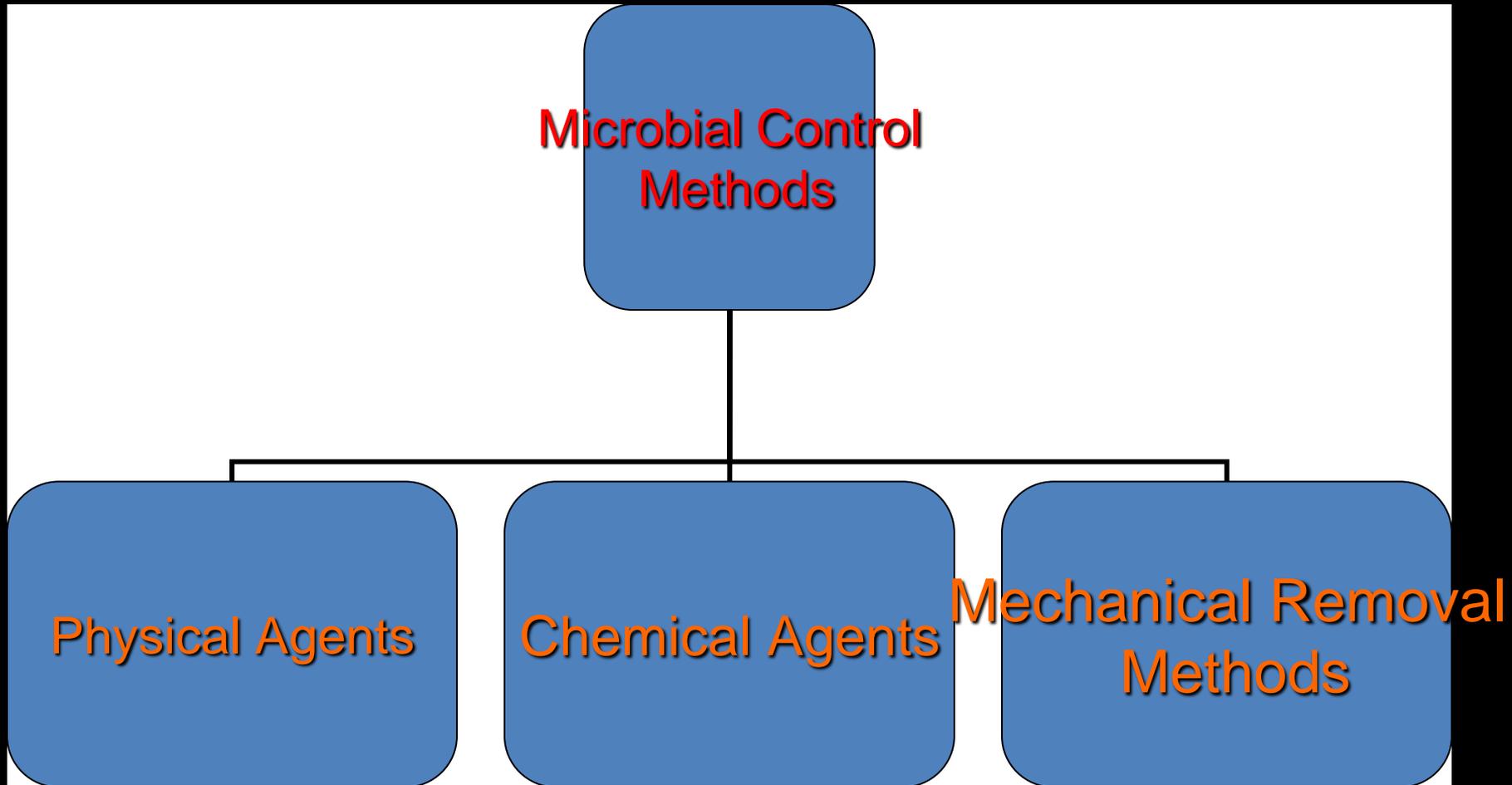
- Classification: will not come in contact with mucous membranes and skin
- Object: can be expected to be contaminated with some organisms
- Level germicidal action: kill vegetative bacteria, fungi and viruses
- Examples: bedpans, bed, EKG leads, walls, floors, etc
- Method : low level disinfection

Low level disinfection for noncritical objects

- Ethyl or isopropyl alcohol 70-09 %
- Chlorine 100 ppm
- Phenol ud
- Iodophor ud
- Quaternary ammonium ud

- Ud: manufacturers recommended use dilution

Methods of sterilization



Physical Agents

Heat

Radiation

Dry

Moist

Ionizing

Non Ionizing

Incineration

Steam Under Pressure

X Ray, Cathode, Gamma

UV

Dry Oven

Sterilization

Boiling Water/Hot Water Pasteurization

Sterilization

Disinfection

Disinfection

Chemical Agent

Gas

Liquids

Sterilization

Disinfection

Animate

Inanimate

Chemotherapy

Antiseptics

Sterilization

Disinfection

Mechanical Removal Methods

Filtration

Air

Liquids

Disinfection

Sterilization

Sterilization

- Steam sterilization
- Dry heat sterilization
- Gas sterilization
 - Ethylene Oxide
 - Formaldehyde
 - Gas plasma
 - Ozon
 - Chlorindioxide
- Radiation
- Liquid chemical sterilization

Contaminated equipments-concept of sterilization

- Decontamination**
 - Transportation to the Sterilization unit**
 - washing, rinsing, Packaging**
 - Make sterilization**
 - Sterilization protection**
-
- Records must be kept under the each phase**

Steam sterilization

- Advantages

- Non-toxic, Cycle easy to control and monitor
- Inexpensive, Rapidly microbicidal
- Least affected by organic/inorganic soils
- Rapid cycle time
- Penetrates medical packing, device lumens

- Disadvantages

- Deleterious for heat labile instruments
- Potential for burns

Steam sterilization

HEAT (C)	PRESSURE (PSi)	TIME (MIN)
121	15	15
126	20	10
132	27	4
134	30	3
FOR PRIONS 134 C 18 MINUTE REQUIRE		

Dry Heat Sterilization

- ❑ Works under the high heat
- ❑ At 140 degree 4 hour,
- ❑ At 160 degree 2.5 hour,
- ❑ At 170 degree 1 hour,
- ❑ At 180 degree 30 minute need for sterilization period

- ❑ Is not preferred nowadays

Sterilization with chemical agent

- ❑ If equipment is not resistance to the high heat, sterilization can make with chemical agent
- ❑ In this method, germicidal effect depends on the nucleic acid alkanization.
- ❑ Ethyleneoxide (EO), Formaldehyde

Sterilization with chemical agent

- Ethylene oxide (EO) a gas
 - Flammable
 - Explosive
 - Toxic
 - Carcinogenic
- Before using must wait 3-14 days
- Can be use implant sterilization

Sterilization with chemical agent

- Formaldehyde a gas
 - Toxic
 - Carcinogenic
- Is not suitable for implant devices
- Does not have permission in Canada and USA

Sterilization with chemical agent

- ❑ In this method (gas plasma) does not need heat and moisture
- ❑ Cold and dry sterilization
- ❑ Sensitive materials (optical, electronic devices) can make sterilized

Sterilization with chemical agent

- ❑ Hydrogen peroxide (HP)
 - ❑ Steel
 - ❑ Aluminum
 - ❑ Titanium
 - ❑ Ceramics
 - ❑ Plastic
 - ❑ Glass materials for suitable to use

Sterilization with liquid chemical agent

- Dipped into the liquid to the method of sterilization
- Can not protect the sterility of the material after the procedure
- That's way not use routinely

Sterilization with liquid chemical agent

Chemical agent	time	heat
Glutaraldehyde (% > 2.0)	10 hour	20-25°C
Hydrogen peroxide-HP (% 7.5)	5 hour	20-25°C
Peracetic acid-PA (% 0.2)	12 min	50-56°C
HP (1.0%) + PA (% 0.08)	8 hour	20°C
HP (7.5%) + PA (% 0.23)	180min	20° C
HP (8.3%) + PA (% 7)	5 hour	25°C
Glut (% 1.12) + Phenol (%1.93)	12 hour	25°C
Glut (% 3.4) + Isopropanol (% 26)	10 hour	- 20° C

Sterilization control methods

- ❑ Maintenance and calibration should be done on a regular basis
- ❑ Devices, heat, pressure and time indicators reflect the correct measurements
- ❑ Temperature, pressure, gas concentration and the time given to these graphs

Sterilization control methods

- ❑ Applied to two types of indicators are used to test the validity of the sterilization
- ❑ Chemical indicators / Biological indicators
- ❑ If there is a change in the expected conversion of sterilization indicator to understand that the Company has requested.

Chemical indicators

- ISO standards are divided into six classes according to their usage according to their capacities and evaluation.

CLASS 1-PROCESS INDICATOR

CLASS 2-SPECIAL TEST indicator

CLASS 3-ONE PARAMETER INDICATORS

CLASS 4-MULTI-PARAMETER INDICATORS

CLASS 5-Integrator

CLASS 6-Emulator

Chemical indicators

- ❑ 1 - Process indicator: Package up indicator. History tells us that the sterilization process of the material. Adhesive tapes or labels can use
- ❑ 2 - Special test indicators: Bowie-Dick type test packs are used
- ❑ 3 - Single-parameter indicators: Sterilization process indicates that a particular variable to the desired value has been reached.

Chemical indicators

- ❑ 4 - Multi-parameter indicators: Two or more critical variables to show the achievement of the desired values
- ❑ 5 - Integrator: Sterilization process checks all critical variables
 - ❑ Is equivalent to the effectiveness of biological indicators.

Chemical indicators

- 6 - Emulator (control cycle indicator): Method is set to the values that are critical to the achievement of the device helps to control all the variables

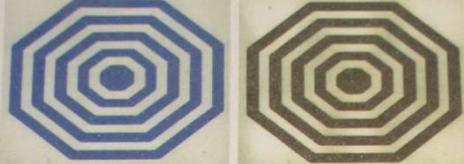
Chemical indicators

- ❑ The effectiveness of the vacuum system used to control the steam saturation and Bowie-Dick test should be done every day before the first operation

Sterilization control

4A MEDICAL **BOWIE & DICK TEST PACK RE**

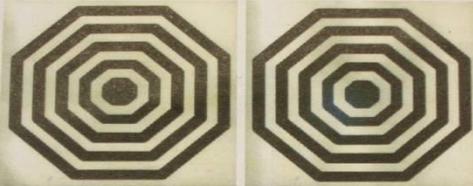
Before exposure **After exposure** **Successful Process:**



Unused 4A Bowie & Dick Pack
The colourchange has been occur
the Bowie & Dick Test Pack.

All the surface of test card turns into green colour,
which verifies the conditions of Bowie & Dick Test.

Failure **Failure** **Low Vacuum**



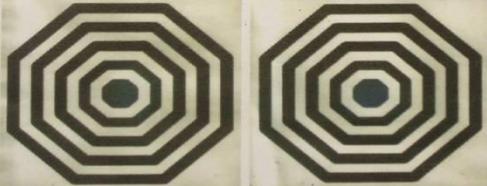
Problems:

- 1) Not enough vacuum power
- 2) Over heated water which vacuum pump decreases the
- 3) Pressure valve or pressure functioning properly gives fau

Solutions:

- 1) The problem in the vacu
- 2) The resistance of autoclave
- 3) The vacuum display in the

Failure **Failure** **Air Leakage**



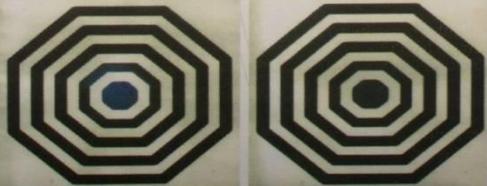
Problems:

- 1) the door joint in bad conditi
- 2) There can be leak in valves
- 3) Check up, incase of leak in t

Solutions:

- 1) Replace the door joint of the a
- 2) Contacting to the authorised s

Failure **Failure** **Non-condensed Gases**



Problems:

During the condensation of steam, completely.

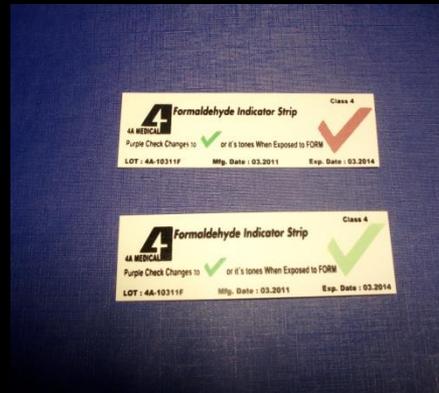
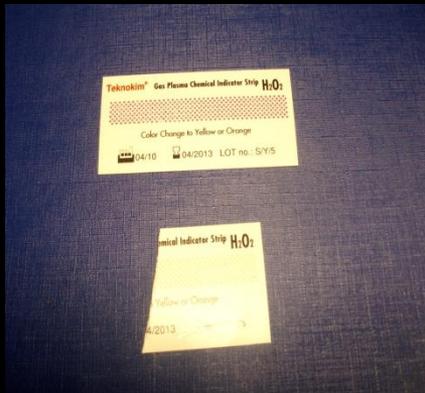
This is because of conditions of its The environment which is compos the failure in the result of Bowie&D

Solutions:

- 1) Deionized water should be used
- 2) The water should be heated at lea
- 3) gases can be prevented.
- 2) Check all joints incase of air le

Wet Steamy **Overheated s**

Sterilization control



Chemical indicators

- ❑ Process set sterilizer include class 1 indicator on each package
- ❑ If you are prompted single-parameter indicators in package (class 3) can be used
- ❑ **ESSENTIAL STERILIZATION CONTROL:** For each load sterilizer include indicator of at least one ISO grade 4.5 or 6
- ❑ This indicators are evaluated by placing all materials in the load can be controlled

Chemical indicators

- ❑ The use of chemical indicator is important to select the correct parameters
- ❑ If 121 C to 134 C sterilizer indicator is left in a set, or vice versa, the results are interpreted incorrectly.

Biologic indicators

- ❑ Measures microbicidal effectiveness of the sterilization process
- ❑ Contain a certain number of bacteria spores
- ❑ Steam sterilization:
Bacillus stearothermophilus
- ❑ Dry heat, EO, gas plasma sterilization
Bacillus subtilis is used
- ❑ After the procedure, control is made by culture of bacteria spores

Biologic indicators

- Pressure steam sterilization

 - Those with graphic print feature once a week,

 - Used every day if is not have graphic feature

 - To be implanted materials should be used in each conversion

- EO sterilization used for each conversion

 - Cabin volume <300 liters and 2,

 - > 300 liters, at least 3

 - use of biological indicators.

Biologic indicators

- Biological indicators of sterilization by using quality should be checked
 - After repairs
 - Packaging material change
 - Package size changes

Biologic indicators

- ❑ The disadvantage of biological indicators requires 48 hours after the procedure for the control of spore culture method
- ❑ The evaluation by the method of measuring the enzymatic activity of bacteria spores results can be obtained in 4 hours.

Biologic indicators



Sterilization record

- Protocol number
 - Applied method of sterilization
 - The selected program
 - Load the content
 - Critical variables measuring records
 - The official name / surname
-
- These registration forms must be kept for 5 years

Protection of steril materials

- Only health staff to the entry into room
- When you enter the store clean apron, bonnet, hand washing
- Which are protected from dust and insects
- Protect from sunlight
- Not Dust-binding surfaces
- Easy to clean the floor
- Room temperature: 22-24 C
- Humidity: 35-70% level

Unsuitableness materials

- Packaged with not suitable materials and methods
- Package integrity is corrupted
- Non-related information on the sterilization
- Stained, wet materials
- Improper storage conditions
- When the humid after autoclaves
- Contacting contaminated surfaces
- MATERIALS ARE NOT ACCEPT STERILE



Disinfection

- The removal of unwanted microorganisms from media
- Less effective than sterilization
- Bacterial spores can not disappear from media
- Is performed using heat or chemicals

Disinfection with heat

- ❑ Hot water (pasteurization), a simple, harmless, highly effective
- ❑ Plasma fraction preparations for about 6 hours at 60 ° C, vaccine and sera may be used
- ❑ 4 log reduction in microbial load endoscope washing with hot water is provided.

Disinfection with heat

- ❑ Washing with hot water and detergent washing machines are used widely
- ❑ After disinfection, the instruments must be stored in suitable conditions
- ❑ The biggest disadvantage of thermal disinfection is not have standardization of safe control methods

Disinfection with heat

Thermal Disinfection heat and time	
Heat of surface (°C)	Disinfection time (min.)
≥ 80	2
75	10
70	15

Disinfection with chemical agent

- ❑ Used in many places within the hospital environment
- ❑ The efficacy of disinfection
 - ❑ Disinfectants used in external ambient conditions (pH, humidity, ambient temperature, water hardness)
 - ❑ Material that will be disinfected
 - ❑ Germicidal activity
 - ❑ Concentrations of use
 - ❑ Operating time DETERMINED

Disinfection with chemical agent

Disinfectants

Is inactivated by organic matter

There is no penetration properties

Affect the surface they come into contact

Disinfection with chemical agent

- ❑ External ambient conditions
 - ❑ Chlorinated compounds are negatively affected by the increase of heat
 - ❑ At Alkaline pH; glutaraldehyde, quaternary ammonium compounds enhancing the effect of , phenolic compounds, hypochloride, iodine effect decreases
 - ❑ Hard water; disinfectant inactivates and leaves on the material permanent precipitates

Disinfection

- ❑ Laparoscope-arthroscope-cystoscope
- ❑ Must be sterile when they enter the sterile space
- ❑ Between High-level disinfection and sterilization , there is no significant difference of infection
- ❑ After disinfection, should be rinse with sterile water

Disinfection

- Semi-critical instruments
 - Flexible Endoscopes
 - Laryngoscopes
 - Endotracheal tubes
 - anesthesia equipment
 - Pulmonary circulation equipment
 - Nasal and vaginal speculum and vaginal probes
 - Nebulizer cups, some ophthalmic instruments
 - Ear Syringe hose
 - Thermometers
 - hydrotherapy tanks

Disinfection

- ❑ Endoscopes
Heat sensitive endoscopes should be sterile or high-level disinfection
- ❑ Flexible endoscopes are contaminated with bacteria, according to entered the cavity (10^5 - 10^9 cfu / mL)
- ❑ Clear up the equipments reduce the bacterial load (log 4-6)
- ❑ Completely removed by cleaning with HIV virus contaminated endoscopes are shown

Disinfection

- ❑ Endocavitary probes
 - ❑ Echocardiography, vaginal / rectal probes
 - ❑ In addition to high-level disinfection, guidelines recommends that using condom or probe wrapper for each patient
 - ❑ Condoms are more secure than probe wrapper in terms of perforation leakage, 1.7% condom, 8.3% coating

Disinfection / Glutaraldehyde

- ADVANTAGES

- Does not corrosion of metals
- Effective in the presence of organic material
- Many materials required 10 hours for sterilization

- DISADVANTAGES

- Very irritant to mucous membranes and the skin
- After reconstitution, short shelf half-life (14-30 days)
- Expensive
- Re-used solutions, the concentration should be monitored

Disinfection / ortho-phytoaldehyde (OPA)

- ❑ Mechanism of action
 - ❑ Rapid-acting high-level disinfectant
 - ❑ 0.55% contains 1.2 benzendikarboxialdehyde
 - ❑ An alkylating agent
 - ❑ Continued effectiveness of acid and alkali environment (pH 3-9)

Disinfection / ortho-fytoaldehyde (OPA)

- ❑ According to glutaraldehyde mikobacterisidal activity begins earlier and stronger (6 min; 6x \log^9 decrease)
- ❑ OPA is effective to the Gluteraldehyde-resistant mycobacteria
- ❑ 0.5% OPA is not sporicidal, if pH increased sporicidal effect can be seen

Disinfection / ortho-phytoaldehyde (OPA)

- There is FDA-approved only for HLD
- High-level disinfection contact time of 12 min at 20 ° C is sufficient to
- Material compatibility is good
- When prepared solutions effectiveness of 14 days

Disinfection / ortho-phytoaldehyde (OPA)

- If swallowed, irritation of the digestive tract
- If contact with the skin, paint gray color
- Contact with eyes; itching, tearing and redness
- Long-term or repeated skin contact resulted with dermatitis
- Irritation of the respiratory system
- Expensive

Disinfection

- Test strips should be used for the pursuit of MEC
- After the useful life test strips should not be used (note date test strips is opened)
- The testing frequency, which depends on the often used solutions (daily use; should be tested at least once every day)
- Results must be recorded

Semicritical equipments

- Stethoscopes
- Sphygomanometer cuff
- Bed linen
- ECG electrodes
- Urine pan
- Food containers

Disinfection / protection of contamination

- Carrier containers must be cleaned, before re using safely
- After opening a sterile solution should not be considered as no longer sterile
- The expiration date must be on each product should be carefully monitored

Precautions

- ❑ Disinfection of critical and semi-critical devices never use low-level disinfectants
- ❑ Do not use HLD for cleaning noncritical materials and environment
- ❑ Use the recommended concentrations and contact time
- ❑ Disinfectant sometimes it can be toxic to the user ,always take safety precautions

