

# Quick Reference for Reprocessing in the Clinical Office Setting

## 1. Policies and Procedures:

- Ensure there are written policies and procedures for all aspects of reprocessing
- Single-use medical instruments are preferred and must not be reprocessed
- Equipment used to clean, disinfect or sterilize must meet Health Canada/PHAC and CSA standards

## 2. Education and Training for Reprocessing Staff

- It is strongly recommended that any staff responsible for reprocessing complete the Provincial Infectious Disease Advisory Committee's 'Reprocessing in the Community Online Learning Course' (see reverse for link to this resource)
- Written, device specific reprocessing instructions are available to reprocessing staff

## 3. Physical Reprocessing Space:

- There is a designated area that is physically separate from direct care areas and from where clean items are handled and stored
- There is a dedicated hand washing sink and/or ABHR for hand hygiene, eye washing station, point-of-use sharps container, and an appropriate supply of personal protective equipment is available for staff

## 4. Reprocessing Level:

| Class                | Use   | Minimum Level of Reprocessing   | Examples  |
|----------------------|---|---|---|
| <b>Critical</b>      | Enters sterile body site, including the vascular system                               | Cleaning followed by sterilization  | <ul style="list-style-type: none"><li>• Surgical instruments</li><li>• Uterine sounds</li><li>• Tenaculum</li><li>• Forceps</li><li>• Biopsy instruments</li><li>• Foot care/podiatry equipment</li></ul> |
| <b>Semi-Critical</b> | Comes in contact with non-intact skin or mucous membranes but does not penetrate them | Cleaning followed by high-level disinfection. Sterilization is preferred                  | <ul style="list-style-type: none"><li>• Vaginal specula</li><li>• Endoscopes</li><li>• Biopsy instruments</li><li>• Anaesthesia equipment</li><li>• Tonometer</li></ul>                                   |
| <b>Non-Critical</b>  | Touches only intact skin and non mucous membranes, or does not directly touch patient | Cleaning followed by low-level disinfection (in some cases, cleaning alone is acceptable) | <ul style="list-style-type: none"><li>• ECG machines</li><li>• Oximeters</li><li>• Stethoscopes</li><li>• Blood pressure cuffs</li></ul>  |

*\* Spaulding's classification of medical equipment and required level of reprocessing (Adapted from Best Practices for Cleaning, Disinfection and Sterilization of Medical Equipment/Devices, Public Health Ontario)*

## 5. Instrument Cleaning:

- Immediately after use, instruments are immersed in an appropriate diluted cleaning solution (e.g. enzymatic cleaner) or are treated with an agent that prevents drying of secretions/body fluids
- Dirty and clean instruments remain separate in covered containers
- Instruments are cleaned manually with a detergent or enzymatic solution, or in an ultrasonic washer, or in an automated washer-disinfector
- Cleaning equipment (e.g. brushes, sponges) is disposable or is cleaned and disinfected with a high-level disinfectant between uses
- Instruments are rinsed after cleaning, and dried prior to high-level disinfectant or sterilization
- Detergent or enzymatic cleaning solution is discarded after each use

## 6. High-Level Disinfection:

- Disinfection logs are required, including test strip monitoring, concentration and exposure time, and disinfectant temperature for automated endoscope reprocessors
- Disinfectant has not passed the manufacturer's expiry date
- Chemical test strips are used to determine whether an effective concentration is present
- Instrument are thoroughly rinsed with sterile, filtered or tap water, depending on intended use
- Instrument are dried after disinfection before storing in a dry, covered container (e.g. dried with a lint free cloth)

## 7. Quality Control of Sterilization Process:

- Equipment/devices are wrapped prior to sterilization using approved sterilization packaging (wrap or pouch)
- Chemical indicators (CIs) are placed appropriately in and/or on each package (if not already included as part of the pouch/pack wrap)
- Date of sterilization is recorded on the wrapper prior to sterilization

## 8. Sterilization:

**Instruments are placed in the sterilizer according to manufacturer's instructions. Do not overload!**

- Sterilization logs are required for all sterilizers/autoclaves
- Sterilizer is tested with biological indicators (BIs) to verify that bacterial spores are killed after cycle (e.g. spore testing)
- After each cycle, mechanical printout (if available) is reviewed in order to verify cycle time, temperature and pressure
- Medical instruments are not used if the chemical and/or biological indicators fail

## 9. Storage of Disinfected/Sterile Medical Equipment/Devices:

- Sterile items are stored in their sterile packaging until time of use
- Sterile items are stored in a clean, dry, dust-free area (e.g. closed shelves or containers), not at floor level, away from debris, drains, moisture, sinks and vermin in order to prevent contamination and maintain disinfection/sterility until time of use
- Prior to use, ensure sterility of the packaging has not been compromised

\*All information based on the document "Infection Prevention and Control for Clinical Office Practice (PIDAC, 2013)

\*\*Please note: This is not an exhaustive list. **Please see resources for more detail.**

## Available Resources for Healthcare Professionals:

- Reprocessing in Community Health Care Settings (\*\*Online Learning Course)  
<https://secure.publichealthontario.ca/en/LearningAndDevelopment/OnlineLearning/InfectiousDiseases/Reprocessing/Pages/default.aspx>
- Infection Prevention and Control for Clinical Office Practice (PIDAC)  
[https://secure.publichealthontario.ca/en/eRepository/IPAC\\_Clinical\\_Office\\_Practice\\_2013.pdf](https://secure.publichealthontario.ca/en/eRepository/IPAC_Clinical_Office_Practice_2013.pdf)
- Best Practices for Cleaning, Disinfection and Sterilization of Medical Equipment/Devices  
[https://www.publichealthontario.ca/en/eRepository/PIDAC\\_Cleaning\\_Disinfection\\_and\\_Sterilization\\_2013.pdf](https://www.publichealthontario.ca/en/eRepository/PIDAC_Cleaning_Disinfection_and_Sterilization_2013.pdf)
- Best Practices for Infection Prevention and Control Programs in Ontario  
[https://www.publichealthontario.ca/en/eRepository/BP\\_IPAC\\_Ontario\\_HCSettings\\_2012.pdf](https://www.publichealthontario.ca/en/eRepository/BP_IPAC_Ontario_HCSettings_2012.pdf)
- Checklist for Reprocessing Endoscopy Equipment (Public Health Ontario)  
[https://secure.publichealthontario.ca/en/eRepository/IPAC\\_Clinical\\_Office\\_Practice\\_Checklist\\_Endoscopy\\_Equipment\\_2013.pdf](https://secure.publichealthontario.ca/en/eRepository/IPAC_Clinical_Office_Practice_Checklist_Endoscopy_Equipment_2013.pdf)
- Checklist for Reprocessing (Public Health Ontario)  
[https://secure.publichealthontario.ca/en/eRepository/IPAC\\_Clinical\\_Office\\_Practice\\_Checklist\\_Reprocessing\\_2013.pdf](https://secure.publichealthontario.ca/en/eRepository/IPAC_Clinical_Office_Practice_Checklist_Reprocessing_2013.pdf)



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