

Reprocessing Instructions for Reusable Medical Devices



Device(s): The following instructions are for all reusable medical devices supplied by PADGETT INSTRUMENTS unless stated otherwise with the packaging of the product. These instructions are intended for use only by persons with the required specialist knowledge and training.

WARNINGS	- Follow instructions and warnings as issued by manufacturers of any disinfectants, disinfectants and cleaning agents used. Whenever possible avoid use of mineral acids and bleach, abrasive agents. - No part of the process shall exceed 140° C. - Some sensitive materials (e.g. Aluminium) are damaged by high alkaline solutions (pH > 10). - Devices with long, narrow cannula, hinges and blind holes require particular attention during cleaning. Note: when reprocessing medical devices, always handle with care, wearing protective clothing, gloves and eyewear in accordance with local Health & Safety procedures.
LIMITATIONS ON REPROCESSING	- Repeated processing has minimal effect on these instruments. - End of life is normally determined by wear and damage in use. - Any specific limitations on the number of reprocessing cycles shall be made available with the instrument.
INSTRUCTIONS	
FROM POINT OF USE	Whenever possible, do not allow blood, debris or bodily fluids to dry on instruments. For best results and to prolong the life of the medical device reprocess immediately after use. If they cannot be reprocessed immediately, use an enzymatic foam spray-cleaner to help prevent soil from drying.
PREPARATION FOR DECONTAMINATION	- Reprocess all instruments as soon as it is reasonably practical following use. - Disassemble only where intended, without the use of tools unless specifically provided by the manufacturer. Where instructions for disassembly are required, these are available with the product.
CLEANING AUTHORITY	- Use only either CE marked or validated washer-disinfector machines and low-foaming, non-bleaching cleaning agents and detergents following the manufacturers' instructions for use, warnings, concentrations and recommended cycles. - Load instruments carefully, with any hose joints and hinges open and so that any dimensions in instruments can drain. - Place heavy instruments with care in the bottom of containers, taking care not to overload wash baskets. - Place instruments with concave surfaces facing down to prevent pooling of water. - Where available, use appropriate attachments to flush inside nozzles and devices with lumens or cannula. - Ensure that soft, high purity water which is controlled for bacterial endotoxins is used in the final rinse stage. Note: automatic cleaning may not be suitable for all lumens and cannula, in which case clean manually with a water jet gun, if available, and an appropriate brush (and saline if provided) that reaches the depth of the lumen. After manually cleaning, pass all devices through an automatic cleaning cycle to achieve disinfection. Note: these instructions have been validated using a washer-disinfector cycle validated to include two cold rinses at <13°C, a detergent cycle and a rinse cycle both at >50°C, a disinfection cycle operating at a temperature of between 80°C and 87°C for a minimum holding time of 3 minute (actual holding time in excess of 2 minutes 50 seconds) and a 20 minute drying cycle. The detergent used was a low-foaming, non-bleaching spray wash detergent cleaner (type 1,2 pH), and the rinse and a neutral pH low-foaming, non-ionic surfactant with integrated alcohol.



Manual For Reprocessing Medical Devices Csao

David Baud



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Announcements & Index ,1985-12 **Medical Device Reprocessing Manual** Rick Bauer,2017 **The Joint Commission Guide to Reprocessing Reusable Medical Devices** ,2023 **User Handbook for Medical Device Reprocessing in Community Health Care Settings** CSA Group,2014 This handbook is intended to be a comprehensive guideline for medical device reprocessing MDR in a variety of community health care settings e g dental surgi centre and physicians offices However it is not meant to replace the use of available MDR standards and guidelines see Annex B Page 1

Government reports annual index ,199? Electrical & Electronics Abstracts ,1997 *International Congress Calendar* ,1990 Who's who in the West ,2000 Reprocessing of Single-use Medical Devices United States. Congress. Senate. Committee on Health, Education, Labor, and Pensions,2000 *Reprocessing of Reusable Medical Devices in Health Service Organizations* Standards Australia (Organization),Standards New Zealand,2014 **Compatibility of Medical Devices and Reprocessing Equipment with Decontamination Agents** Medical Devices Agency,2001 *Medical Devices Reprocessing Workbook* Tammy Bennardo,2017 **Sterile Medical Devices** ,1984 **Import of Medical Devices** ,1988

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