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## PRODUCT CARE INSTRUCTIONS: CLEANING METHOD 2

DEVICE(S): All Ocular MaxField® Glass and MaxLight® CR-39 Indirect Diagnostic/Laser Lenses\*

\*Note: See Cleaning Method 3 for OI-20A, MaxAC 20D Indirect and OI-28A, MaxAC 28D Indirect.

WARNINGS	<ul> <li>Read all instructions before use.</li> <li>Follow instructions and warnings as issued by manufacturers of any decontaminants, disinfectants and cleaning agents used.</li> <li>Wherever possible avoid the use of abrasive materials for cleaning and drying.</li> <li>Incorrect handling and care or misuse can lead to premature wear of these devices.</li> <li>Inspect these devices carefully for damage, cracks or malfunctions before each use.</li> <li>Do not use damaged devices.</li> <li>Use only approved disinfectant solutions (e.g., FDA, DGHM, CE Mark).</li> <li>Each device requires cleaning and disinfection before its first use and any subsequent use.</li> <li>Ensure cleaning and disinfected and dry state.</li> <li>Store devices in a cleaned, disinfected and dry state.</li> <li>Sterilize all devices before surgery.</li> <li>Never Steam Autoclave or Boil listed lenses.</li> <li>Never soak in Acetone or Other Solvents.</li> </ul>
Limitations on reprocessing	Repeated processing has minimal effect on these devices.  Rapid cooling may damage devices.

INSTRUCTIONS		
Point of Use:	Clean with alcohol wipe.	
Preparation for decontamination:	Reprocess all devices as soon as reasonably practical following use.	
Cleaning: Automated	Not recommended.	
Cleaning: Manual	Wipe: Clean with alcohol wipe.	
	Then: Proceed with either disinfection or sterilization instructions.	
	Caution: If fluid/gas exchange has occurred, wipe lens with alcohol to remove any trace of oil present. If lens is not promptly and properly cleaned, permanent damage may result.	
Disinfection:	Disinfectant solutions (e.g., Approved by FDA, DGHM, CE Mark) may be used in accordance with label instructions of the disinfectant manufacturer. Pay strict attention to disinfectant manufacturers recommended concentrations and contact durations. Ensure that disinfectant solution makes complete contact with all device surfaces and lumens.	
	After manual high level disinfection, soak and rinse lens in large volume of cool or tepid sterile water for 1 minute and thoroughly flush lumens. Repeat this procedure 2 times with fresh rinse water to ensure removal of disinfection solution.	
	Caution: To avoid damage to the lens, do not exceed recommended exposure time.	
Drying:	Dry devices carefully with lint free tissues or hospital grade compressed air and place in a dry storage case.	
Maintenance, Inspection and Testing:	Inspect these devices carefully for damage, cracks or malfunctions before each use. Do not use damaged devices.	
Packaging:	Standard biological peel packs ( <i>wrapped</i> ) may be used. The pack should be large enough to contain the device without stressing the seals. Biological peel packs ensure sterility after the sterilization process.	

	EO Minimum Time: 1 hour Temperature: 130°F (54°C) Aerations Time: 12 Hours  STEAM AUTOCLAVE NO
Sterilization:	STERRAD (See notes 1 & 2 below).  STERRAD 100NX System (Standard Cycle) <sup>1,2</sup>   STERRAD NX: Standard Cycle <sup>1,2</sup>   STERRAD 100S, 200: Short Cycle <sup>1,2</sup>   STERRAD 50 <sup>1,2</sup> Process product in STERRAD approved tray or container and wrap when applicable. Follow STERRAD instructions.  Not compatible with: MaxLight CR-39 Indirects.
	Steris SYTEM 1E Follow Steris insctructions Not compatible with MaxLight CR-39 Indirects  Steris V-Pro Models <sup>1</sup> No
	3M <sup>TM</sup> Optreoz <sup>TM</sup> 125-Z Low Temperature Sterilization System – Cycle 1 <sup>1</sup> Follow 3M <sup>TM</sup> Optreoz <sup>TM</sup> 125-Z Low Temperature Sterilization System instructions.  Notes: 1. Colored aluminum will fade to a natural aluminum color within 25 cycles.
	2. Polyacetal components (black or white plastic) may have limited life after repeated sterilization with this method. For information on compatibility with alternative product care methods, contact Customer Service.
Storage:	Ensure devices are cleaned, disinfected and dry before storage. Store in a clean and dry room temperature environment.

	Other forms of cleaning and sterilization equipment are available. Please consult instructions of the processing equipment or manufacturer for compatibility claims. All cleaning and sterilization processes require validation at the point of use.
Additional Information:	Note: These lenses are known to be compatible with Glutaraldehyde (2% or 3.4%), BLEACH (10% solution mixed at: 1 part bleach to 9 parts cool or tepid water, recommended exposure time = 10 minutes; Bleach is corrosive to metals, to avoid corrosion do not exceed recommended exposure times), Medical disinfectant wipes (Asepti-Wipe II, Cavicide, DisCide Ultra, Envirocide, Tristel Wipes System and Opti-Cide-3) and Medical disinfectant solutions such as Cidex and Cidex OPA. Compatible with Steris Resert except: OI-120M, OI-60M, OI-66M, OI-HM-78M, OI-78M, and OI-SP.
Manufacturer contact:	See brochure for telephone number and address of local representative.

The instructions contained herein have been validated as being CAPABLE of preparing a medical device for re-use. It remains the responsibility of the processor to ensure that the reprocessing as actually performed using equipment, material and personnel in the reprocessing facility achieve the desired result. This normally requires validation and routine monitoring of the process.

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