

EC REP

■ OCULAR INSTRUMENTS

 $2255\ 116^{\mathrm{TH}}$ Ave NE, Bellevue, Washington 98004-3039 USA

T: 425-455-5200 or 800-888-6616 F: 425-462-6669

E: ocular@ocularinc.com I: www.ocularinc.com

EM ERGO EUROPE: Molenstraat 15 2513 BH, The Hague The Netherlands

T: +31.70.345.8570; F: +31.70.346.7299

PRODUCT CARE INSTRUCTIONS: CLEANING METHOD 3 DEVICE(S): ALL Ocular Surgical Lenses and Rings* and OI-20A, OI-28A, O4MAC, O4MAC-LR, O4MAC-H, OG3MAC-10.

*Note: See Cleaning Method 1 for OLIV-EQNA, Landers NA Equatorial Vitrectomy Lens, OLIV-WFNA, Landers NA Wide Field Vitrectomy Lens, OLTK-7.2 or 8.2, Landers WF Temporary Keratoprosthesis, OTSG, Thorpe Surgical Gonioscope, OUV-132-2, Peyman-Wessels-Landers 132D Upright Vitr. Lens. OWIV-HMNA. Woldoff NA High Mag Vitrectomy Lens. OKSG. Khaw Surgical Gonioprism.

Upright vitr. Lens, Owiv-Himiya, woldoff iya High Mag vitrectomy Lens, OksG, Khaw Surgical Gontoprism.					
	Read all instructions before use.				
	Follow instructions and warnings as issued by manufacturers of any decontaminants, disinfectants and				
	cleaning agents used.				
	Wherever possible avoid the use of abrasive materials for cleaning and drying.				
WARNINGS	Incorrect handling and care or misuse can lead to premature wear of these devices.				
	Inspect these devices carefully for damage, cracks or malfunctions before each use.				
	Do not use damaged devices.				
	• Use only approved disinfectant solutions (e.g., FDA, DGHM, CE Mark).				
	Each device requires cleaning and disinfection before its first use and any subsequent use.				
	Ensure cleaning and disinfection solutions fully contact all device surfaces and lumens.				
	Store devices in a cleaned, disinfected and dry state.				
	Sterilize all devices before surgery.				
	Allow devices to air cool to room temperature before handling and use.				
	Repeated processing has minimal effect on the performance on these devices ^{1, 2, 3} .				
	End of a product's service life is therefore determined by wear and tear and by damage due to use, such as				
Limitations on	scratches caused by mechanical cleaning (e.g. by hard brushes), or calciferous residues (e.g. hard water used in the				
reprocessing	sterilizer) that impair the optical quality. Thus end of a product's service life varies and is therefore to be				
	determined by the user.				
	Rapid cooling may damage devices.				

INSTRUCTIONS					
Point of Use:	Rinse: Immediately upon removal from patient's eye, thoroughly rinse in cool or tepid water to avoid soil drying on surfaces or lumens.				
Preparation for decontamination:	Reprocess all devices as soon as reasonably practical following use. Disassemble devices only where intended.				
Cleaning: Automated	Not recommended.				
Cleaning: Manual	Wash: Place a few drops of low foaming mild soap (i.e., neutral pH (7.0) detergent formulated for medical instruments) on a moistened cotton ball. Gently clean with a circular motion until all soil has been removed. Flush all lumens with detergent solution to remove soil.				
	Rinse: Thoroughly rinse lens and flush lumens in cool or tepid high purity water, then dry carefully with a <i>non-linting</i> tissue or hospital grade compressed air.				
	Inspect: Visually inspect all surfaces, crevices, joints, holes and lumens for complete removal of soil and fluid. If any soil or fluid is visible, then repeat cleaning.				
	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. Caution: If used on an ulcerated cornea, lens must be STERILIZED before next procedure.				
Drying:	Dry devices carefully with lint free tissues or hospital grade compressed air and place in a dry storage case.				
Maintenance,	Inspect these devices carefully for damage, cracks or malfunctions before each use. Do not use damaged devices.				
Inspection and Testing:					

Packaging:	0 1	packs (wrapped) may be uals. Biological peel packs er			enough to contain the device			
	EO Minimum Time: Temperature: Aerations Time:	1 hour 130°F (54°C) 12 Hours	isure steri	inty arter the stermza	ation process.			
	STEAM AUTOCLAVE ³ Prep: Rinse devices with sterile water. Place product in sterilization case.							
	Gravity Cycle (wrappe Temperature: Time: Dry Time:	270°F (132°C) min. 15 minutes min. 15 minutes min.	or	Temperature: Time: Dry Time:	250°F (121°C) min. 30 minutes min. 15 minutes min.			
	Pre-Vacuum Cycle (w. Temperature: Time: Dry Time:	rapped) 270°F (132°C) min. 4 minutes min. 20 minutes min.	or	Temperature: Time: Dry Time:	273°F (134°C) min. 3 minutes min. 20 minutes min.			
	FOR IMMEDIATE USE ONLY -FLASH AUTOCLAVE Gravity Cycle (umwrapped) Temperature: 270°F (132°C) min. Time: 10 minutes min.							
	Pre-Vacuum Cycle (un Temperature: Time:	nwrapped) 270°F (132°C) min. 4 minutes min.	or	Temperature: Time:	273°F (134°C) min. 3 minutes min.			
	could leave a cloudy file	water in steam sterilizer is recommon the lens. The deposits cate that of a new lens. Allow Vite	n only be r	emoved by regrinding	1 0			
Sterilization:	STERRAD 100NX: Standard Cycle ^{1, 2} Process product in STERRAD approved tray or container and wrap when applicable. Follow STERRAD instructions. Not compatible with: OCTK-6.5, OLTA, OLTA-2, Silicone tubing and Luer adapters supplied with products (i.e., lumens are less than 0.7mm ID). STERRAD NX: Standard Cycle ^{1, 2} STERRAD 100S, 200: Short Cycle ^{1, 2} STERRAD 50 ^{1, 2} Process product in STERRAD approved tray or container and wrap when applicable. Follow STERRAD instructions. Not compatible with: OLV-1-IN, OLV-1-IR, OBVI, OFVI, OPFVI, OMVI, OPGVI, OPVI-3, OPGVI, OPVI-3, OCTK-6.5, OLTA, OLTA-2, Silicone tubing and Luer adapters supplied with products (i.e., lumens are less than 1.00 mm ID)							
	Steris V-Pro models ¹ Follow Steris instructions. All models not compatible with: OHBVE, OHFVE, OHMVE, OHWVE, OCTK-6.5 V-Pro 1 is not compatible with: OPVI-3, OPGVI, OMVI, OPFVI, OBVI, Silicone tubing and Luer adapters supplied with products (i.e., lumens are less than 1.00 mm ID)							
	3M TM Optreoz TM 125-Z Low Temperature Sterilization System – Cycle 1 ¹ Follow 3M TM Optreoz TM 125-Z Low Temperature Sterilization System instructions. Not compatible with: OCTK-6.5, OHFVE, OHMVE, OHBVE, OHWVE, OBVI, OFVI, OPFVI, OMVI, OPGVI, OPVI-3, OLTA, OLTA-2 Silicone tubing and Luer adapters supplied with products (lumens smaller than 1.0mm diameter are not compatible)							
	3M TM Optreoz TM 125-Z Low Temperature Sterilization System − Cycle 2 ¹ Follow 3M TM Optreoz TM 125-Z Low Temperature Sterilization System instructions. Not compatible with: OCTK-6.5							
	Note: 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. 3. Devices containing coated mirrored surfaces may exhibit minor accumulative changes with repeated cycling.							
	For information on compatibility with alternative product care methods, contact Customer Service.							
Storage:	Ensure devices are cleane	ed, disinfected and dry before	storage. S	Store in a clean and dr	y room temperature environment.			
					cessing equipment or the manufacturer for			
Additional Information:	compatibility claims. All cleaning and sterilization processes require validation at the point of use. 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 coor tepid water, recommended exposure time = 10 minutes; Bleach is corrosive to metals, to avoid corrosion do not exceed recommended exposure times), Medical disinfectant wipes (e.g. 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: OBVI, OHBVE, OHFVE, OHMVE, OHWVE, OIV-							
Manufacturer contact:	132, and all HRI lenses.	one number and address of l	local repre	esentative				

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.