SAFETY, RELIABILITY and PERFORMANCE

Transforming low temperature sterilization



SAFETY



V-PRO™ Sterilizers are SAFE for the User, the Patient, the Medical Devices and the Environment*

No toxic residues are released to the environment, only water and oxygen vapor

- Hydrogen Peroxide emissions from V-PRO™ Sterilizers are well below the established Permissible Exposure Limits (PEL) for all countries where applicable
- The 15-minute Short-Term Exposure Limit (STEL) levels of Hydrogen Peroxide are below all applicable standards

*V-PRO[™] Low Temperature Sterilization System, Environmental H202 Safety Profile. Independent testing, using a fully validated OSHA test method, demonstrates industry-leading levels of Safety for Healthcare employees and the environment. For further information and scientific studies visit www.steris.com

Designed to meet parametric release criteria in accordance to the requirements of EN ISO 14937

The performance of the V-PRO™ maX Sterilizer is ensured through rigorous control and independent monitoring of various critical parameters during the cycle. If any process parameter is outside of the established ranges, the cycle will abort

"Testing was confirmed in vitro and in vivo with multiple strains of prions (including vCJD/BSE) under worst case test conditions and in the presence/absence of cleaning. Fichet, G. E. Comoy, K. Antioga, C. Duval, G. McDonnell, J.P. Desys, 2004. Novel methods for disinfection of prion-contaminated medical devices. The Lancet 364: 521-526. Fichet G., K. Antioga, E. Comoy, J.P. Deslys, G. McDonnell (2007) ripoi inactivation using a new gaseous hydrogen peroxide sterilisation process. J. Hosp. Infect. 67: 278-386. McDonnell G., C. Dehen, A. Perrin, V. Thomas, A. Igel-Egalon, P.A. Burke, J.P. Deslys, E. Comoy (2013) Cleaning, Disinfection and Sterilization of Surface Prion Contamination. J. Hosp. Infection 85: 268-273. NOTE: These claims and this brochure are for Use only in Europe, Middle East and Africa.

SAFETY

The STERIS Device Testing team has the knowledge, expertise and tools to provide you confidence when reprocessing validated devices. This team works closely with medical device manufacturers to test devices for material compatibility and sterilization efficacy.

The validated list of medical devices for the V-PRO™ Sterilizers is growing. Examples of currently validated devices include:



Broad device manufacturers endorsements















Endoscope	Arthroscopes Choledochoscopes Bronchoscopes	Bronchoscopes Intubation Endoscopes	Cystoscopes Laparoscopes	¹da Vinci® Endoscopes Rhinolaryngoscopes	Hysteroscopes Ureteroscopes
Endoscope Accessories	Cameras	Sheaths/Adapters/Bridges	Cannulas	Fibre Optic Light Cables	Obturators
Powered Instruments	Batteries	Chuck	Drills	Saws	Handpieces
General Surgical Instruments	Curettes	Forceps	Scissors	Retractors	
Packaging	Wraps	Rigid Containers	Trays	Pouches	
Other Devices	Bougies Transducers	Defibrillation Paddles Ultrasound Probes	Laryngoscope Bl Gonio/Vitrectomy	ades and Handles Lens	Flow probes



Discover our online compatibility matrix!

We constantly update a long list of instruments validated for use in V-PRO™ Sterilizers

¹da Vinci endoscopes is a registered trademark of Intuitive Surgical, Inc.

and many more.

RELIABILITY



Large colour touchscreen and cycle countdown

Colour-coded display screens for easy identification of cycle status



Plug and play: only a switch on/off button! You just need an electrical outlet

- No special ventilation
- No drain
- No special utility requirement

Easy and safe handling of VAPROX™ HC Sterilant cartridge

- The teardrop shape of the VAPROX™ HC Sterilant cartridge ensures that only one option of insertion is possible
- No need for medical waste containers. The empty cartridge of VAPROX™ HC Sterilant can be safely discarded as a normal waste





Easy loading and unloading

2 removable sliding shelves for better ergonomics and ample space for bulky loads

PERFORMANCE

Largest Chamber Capacity = Fewer Cycles to Run



is the most productive on the market!

Non-Lumen Cycle

Up to 23 kg (both shelves) Non-lumened flexible endoscopes.

Flexible Cycle

2 flexible endoscopes; or 1 endoscope and non-lumened load (total of 11 kg combined)

Lumen Cycle

20 stainless steel lumens (single, dual or triple channel devices)



Up to 5.4 kg (both shelves) Non-lumened flexible endoscopes. 1 flexible scope up to 990 mm 12 stainless steel lumens (single, dual or triple channel devices)

Exceptional conditioning phase and smart reservoir in the V-PRO™ Sterilizer

- Moisture check to verify the load is dry*
- If moisture is detected, the V-PROTM
 Sterilizer will attempt to remove residual moisture from the load and packaging
- Built-in reservoir in the V-PRO™ Sterilizer stores the VAPROX™ HC sterilant before injection during the conditioning phase and prevents the use of VAPROX™ HC Sterilant in case of aborted cycles

*Users must always ensure the load is thoroughly dried prior to placing the load in the sterilizer



Mixed loads: Process 2 flexible scopes in the same load or a flexible scope + a non-lumen load



Consistently Fast Cycle Times = Devices Ready When You Need Them

Non-Lumen Cycle

Flexible Cycle

Lumen Cycle















Mixed loads









All cycle times include conditioning phase

DISCOVER

Different sizes, different needs

Discover our range of V-PRO™ Sterilizers



Mobile and Compact



	V-PRO™ 600 Low Temperature Sterilization System	V-PRO™aX Low Temperature Sterilization System		
	Counter top version: 787 x 787 x 711 mm	Single Door Recessed: 838 x 962 x 1908 mm		
Overall dimensions (WxDxH)	Cart version: 787 x 787 x 1626 mm	Single Door Cabinet: 838 x 973 x 1908 mm		
		Double Door Recessed: 838 x 1010 x 1908 mm		
Chamber size	330W x 711D x 254H mm	432 x 381 x 826 mm		
Chamber usable volume	60 litres	136 litres		
Installation requirements	System installation requires no plumbing, ventilation or air supply – only a dedicated electrical connection is needed. A power cord is supplied for this connection			
Operating temperature	≤ 51.5 °C			
Chamber	Chamber and door assembly are aluminium equipped with a silicone rubber gasket for the door and a welded backhead for the chamber			
Door locking mechanism	Automatic door locking mechanism keeps the sterilization system door locked during the entire Sterilization Cycle. After cycle completion, the door is electrically unlocked			
Sterilant cup interface	Only accepts VAPROX™ HC Sterilant Cups. The system control automatically tracks the amount of VAPROX™ HC Sterilant used and the Sterilant expiration date			
Catalytic converter	Catalytic converter receives outflow from chamber during all cycle phases and converts hydrogen peroxide into water vapor and oxygen			
Control display panel	Control display panel is located on the front of the Sterilization unit. This colour touch panel provides user information and allows user inputs			
Control display panel	The display is a 640 x 480 pixel resolution, 5.7" screen	The display is a 640 x 480 pixel resolution, 10.4" screen		
Printer	Printer is located on the front of the Sterilization unit. This alphanumeric impact printer provides an easy-to-read permanent record of the Sterilization cycle. Printer provides a 5.7 mm, 24-character-wide cycle tape and paper take-up			



Faster, more Versatile

962 mm Recessed Single Door 973 mm Cabinet Single Door 1010 mm Recessed Double Door



838 mm

V-PRO™ STERILIZER DEDICATED SOLUTIONS

Enjoy the full range of accessories and indicators

Sterilant and accessories



VAPROX™ HC Sterilant



Trays and Organisers



Tray Mats



Tyvek® Pouches and Reels



Tape



ZENPACK SMMS Polywraps

Indicators



VERIFY® V24 Self-contained Biological Indicators



VERIFY® Biological Indicator Incubators



VERIFY® Biological Indicator Challenge Pack for VH2O2 Sterilization Processes



VERIFY® VH2O2 Vaporized Process Indicator Strips



VERIFY® VH2O2 Vaporized Process Label



VERDOC® VH2O2 Vaporized Record Cards with Process Indicator



VERIFY™ VH2O2 Indicator Tape

V-PRO™ Sterilizers - Transforming low temperature sterilization through:

SAFETY for the User, the Patient, the Medical Devices and the Environment. No toxic residues, only Oxygen and Water

RELIABILITY Broad range of device manufacturer endorsements, fewer moisture-related aborts increasing productivity

PERFORMANCE Faster cycle times and industry leading cycle claims will allow for a quick device turnaround and maximizing the number of devices per load



Get connected to STERIS University, encompassing training disciplines and quality Clinical Education, delivered by a team of Expert Industry Professionals.

For more details, ask **educationdesk@steris.com** or visit **university.steris.com**







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