

EverLUX®

STW0.1 Cartridge

Description

The EverLUX® STW0.1 features two serially layered PES hydrophilic membranes with the coarser highly asymmetric upstream layer optimized for protection of the asymmetric 0.1 µm downstream layer. The filter provides the added benefit of certification that meets the critical demands of the pharmaceutical, biotechnology, and related industries.

Materials of Construction

All components of the EverLUX® filter are either animal free or in compliance with EMEA/410/01 Rev. 3 (EDQM 5.2.8 07/2011:50208), and US Code of Federal Regulations 9 CFR 94.18 and 21 CFR 189.5. These materials are listed for food contact use in the Code of Federal Regulations (CFR), Title 21, as below:

Membranes:	Polyethersulfone	CFR Title 21, 177.2440
Upstream support:	Polypropylene	CFR Title 21, 177.1520
Downstream support:	Polypropylene	CFR Title 21, 177.1520
Outer guard:	Polypropylene	CFR Title 21, 177.1520
Core:	Polypropylene	CFR Title 21, 177.1520
End caps:	Polypropylene	CFR Title 21, 177.1520
O-rings:	Typically Silicone	CFR Title 21, 177.2600
Sealing method:	Thermal bonding	

Pore Size 0.1 µm

Minimum Bubble Point 80 psi (5,5 bar), water
30 psi (2,1 bar), 60% IPA / 40% water
27 psi (1,9 bar), 70% IPA / 30% water

Maximum Diffusion Rate 25 mL/min per 10" @ 40 psi, water (25 mL/ min per 25 cm @ 2,8 bar, water)

Typical Water Flow Rate 0.7 psid/gpm per 10" (0,78 L/min at Δp 10 mbar per 25 cm)

Bacterial Retention >10⁷ per cm² removal of *Brevundimonas diminuta* per ASTM F838-05

Operating Characteristics

100 °F @ 80 psid (38 °C @ Δp 5,5 bar)	150 °F @ 60 psid (66 °C @ Δp 4,1 bar)	180 °F @ 30 psid (82 °C @ Δp 2,1 bar)
--	--	--

Sterilization

Autoclave: 121 to 135 °C (15 to 30 psi, 1 to 2 bar), 30 to 60 min, ≥ 3 cycles. Water wet membrane prior to autoclaving.
Steam in place: 121 to 135 °C (15 to 30 psi, 1 to 2 bar), 30 to 60 min, ≥ 3 cycles, water wet membrane first.

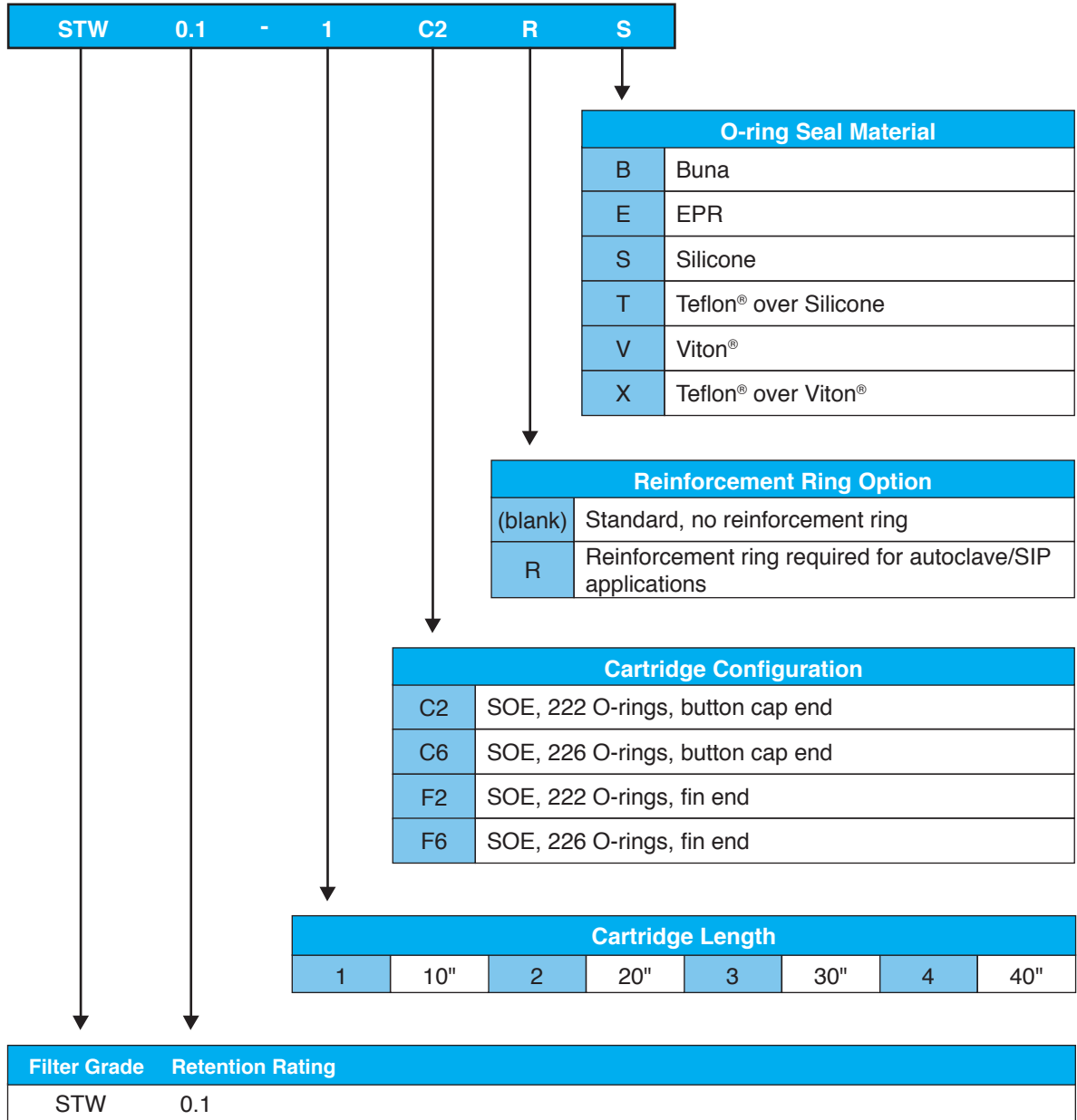
Biological Safety

EverLUX® filters meet the requirements as specified in the current USP Class VI plastics, physicochemical, oxidizable substances, and cytotoxicity tests. Bacterial endotoxin levels in aqueous extracts of EverLUX® filters are less than 0.5 EU/mL, as determined using the *Limulus* amoebocyte lysate (LAL) test. No binders, adhesives or surfactants are used in the construction of EverLUX® filters. Filters comply with Commission Regulation (EU) No 10/2011.

Quality Assurance

Each EverLUX® STW0.1 is supplied with a Certificate of Quality verifying the high standards and superior performance of the product. EverLUX® filters comply with the Food and Drug Administration Code of Federal Regulations, Title 21, Parts 210 and 211. Product is manufactured and packaged in a cleanroom facility that, through voluntary compliance, meets or exceeds FDA Good Manufacturing Practice Standards. To ensure product reliability, Meissner's Quality Assurance staff continually audits the manufacturing process for conformance to its Quality Management System. Each EverLUX® filter is integrity tested during manufacture and is clearly marked with filter type and lot number.

Ordering Guide



Additional information about EverLUX® filter products is available in the Green Docs document which is viewable at <https://www.meissner.com/downloads/everlux-gd002-2.1.pdf>

Viton and Teflon are registered trademarks of DuPont Dow Elastomers L.L.C. & The DuPont Company. EverLUX is a registered trademark of Meissner Filtration Products.