SteriLUX®

0.2 µm VMH-grade Standard Capsule Filter — CS(2), CL(2), CJ2 Model

Description

The SteriLUX[®] VMH0.2 capsule filter is a hydrophilic PVDF membrane filter with high flow rates, high throughputs, low protein binding properties and broad chemical compatibility. It is recommended for filtration of pharmaceutical preparations, active ingredients, biopharmaceuticals, parenterals, vaccines, biologicals including dilute protein solutions, cell and tissue culture media, media additives, ophthalmic and other dilute preservative solutions, UPW, chemicals, alcohols, and sanitizing agents.

Materials of Construction

All components of the filter capsule are either animal free (ACF) or in compliance with EMA/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:

Membrane: Upstream support: Downstream support: Outer guard: Core: End caps: Capsule housing: Sealing method:	Polyvinylidene fluoride (PVDF) Polypropylene Polypropylene Polypropylene Polypropylene Polypropylene Polypropylene Thermal bonding	CFR Title 21, 177.2510 CFR Title 21, 177.1520 CFR Title 21, 177.1520
Pore Size	0.2 μm	
Minimum Bubble Point	50 psi (3.44 bar), water 18 psi (1.24 bar), 60% IPA/40% water 17 psi (1.17 bar), 70% IPA/30% water	
Maximum Diffusion Rate	CS: 2.7 mL/min @ 40 psi (2.76 bar), water CL: 5.7 mL/min @ 40 psi (2.76 bar), water CJ: 8.8 mL/min @ 40 psi (2.76 bar), water	
Typical Water Flow Rate	CS: 0.24 gpm @ 1 psid (0.13 L min ⁻¹ at Δp 10 mbar) CL: 0.51 gpm @ 1 psid (0.28 L min ⁻¹ at Δp 10 mbar) CJ: 0.88 gpm @ 1 psid (0.48 L min ⁻¹ at Δp 10 mbar) Capsule inlet/outlet selection may affect pressure drop and flow rate.	
Bacterial Retention	$>10^7$ per cm ² removal of <i>Brevundimonas diminuta</i> per ASTM F838	
Operating Characteristics Operating temperature range: Maximum temperature rating: Maximum operating pressure: Maximum operating pressure: Reverse operating pressure:	32 °F to 100 °F (0 °C to 38 °C) 160 °F @ 35 psig (72 °C @ 2.4 bar) 75 psig @ 100 °F (5.2 bar @ 38 °C), liquid service 50 psig @ 100 °F (3.4 bar @ 38 °C), gas service 15 psig @ 100 °F (1.0 bar @ 38 °C)	

Sterilization

Autoclave: 121° C to 135° C (15 to 30 psi, 1 to 2 bar), 30 to 60 minutes, \geq 3 cycles. Irradiation: 25 to 40 kGy once. Do not autoclave irradiated capsules. Capsules must not be in-line steam sterilized.

Biological Safety

SteriLUX[®] filters meet the requirements as specified in the current USP <88> Class VI plastics, <87> cytotoxicity and physicochemical tests; after flush, filters comply with USP 43 oxidizable substances test. Bacterial endotoxin levels in aqueous extracts of SteriLUX[®] filters are less than 0.5 EU/mL, as determined using the *Limulus* amebocyte lysate (LAL) test. No binders, adhesives, or surfactants are used in the construction of SteriLUX[®] filters. Filters comply with Commission Regulation (EU) No. 10/2011.



Quality Assurance

SteriLUX[®] 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 SteriLUX[®] filter is clearly marked with filter type, lot number, and serial number.



*CJ filters are animal component free (ACF) and only available with a gamma-stable polypropylene housing. **3/4 sanitary flange fitting option is only available for CS2, CL2, and CJ2 capsules.

Additional information about this filter product is available in the SteriLUX® Green Docs document at www.meissner.com/green-docs.

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