



STERILE CAPSULE FILTER

These filters are designed for applications where sterility of the affluent is mandatory (Sterile Compounding and Pharmaceutical use). The product contains a pharmaceutical grade PTFE membrane (Sterilizing Grade) and the filters are built in a 10,000-class clean room, flushed, dried and 100% integrity tested prior to packaging. The product is factory sterilized by Ethylene Oxide gas (ETO) ensuring sterility.

Technical Data Sheet

Catalog Number: MKPF020LFLM-ETO



Materials of Construction

Filter Membrane: 0.2 um Pharmaceutical Grade PTFE Sterilizing Grade - Hydrophobic

Supports: Polypropylene

Core: Polypropylene

Capsule Body: Polypropylene

Filter Dimensions and Specifications

Outer Diameter: 59 mm (2.32")

Length, Fitting to Fitting: 92 mm (3.62")

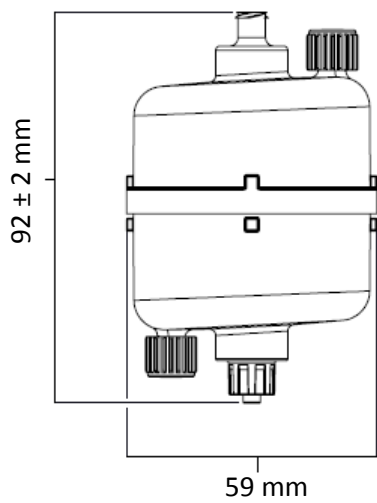
Filter Area: 480 cm² (74.4 in²)

Sterilization: The filter is factory sterilized by Ethylene Oxide Gas (ETO)

Inlet Fitting: Luer Loc Female

Outlet Fitting: Luer Loc Male

Volume Filtered: 8.8 L to 16.5 L



Integrity Data

Pore Size	Min, Bubble Point, 25°C	Challenge Microorganisms	CFU per cm ²
0.20 µm	≥3.5 bar (50 psi)	Brevundimonas Diminuta (ATCC 19146)	7

Operating Parameters

- Maximum Working Pressure: 5.5 bar (80 Psi) @ 25 °C (77 °F)
- Maximum differential Pressure (forward): 4.1 bar (60 Psi) @ 25 °C (77 °F)
- Maximum differential Pressure (reverse): 2.1 bar (30 Psi) @ 25 °C (77 °F)
- Maximum Operating Temperature: 80 °C (176 °F)
- Recommended Replacement Pressure: 2.4 bar (35.3 Psi)

Safety

- UPS Bacteria Endotoxins: ≤25 EU/ml
- Factory Sterilized: Ethylene Oxide Gas (ETO)
- Product was 100% Integrity Tested and flushed with filtered water @ 0.05 µm

Regulatory

- Complies with USP 797 Guidelines
- ASTM F838-05 - Bacterial Retention
- FDA 21 CFR 177.1655
- USP Class VI - Biological Reactivity
- ISO 10993-Part 1. 5 - Cytotoxicity
- ISO 14001; ISO 13485; OHSMS 18001 Certified.
- ASTM Hemolysis testing