STyLUX® is a hydrophilic PES membrane filter compatible with a wide range of liquids. It withstands a wide pH range (1-14) and is suited for removal of microorganisms and particulates in high-purity liquids and aqueous chemicals. The filter’s asymmetric membrane provides absolute retention but also superior flow rates and contaminant capacity. The STyLUX® filter delivers reliable and consistent high-quality performance and is ideal for prefiltration or final filtration when sterility assurance is not required.

Materials of Construction
All components of the STyLUX® mini capsule are animal component free. These materials are listed for food contact use in the Code of Federal Regulations (CFR), Title 21, as below:

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<tbody>
<tr>
<td>Membrane:</td>
<td>Polyethersulfone</td>
<td>Polypropylene</td>
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<td>Polypropylene</td>
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<td>Polypropylene</td>
<td>Thermal bonding</td>
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<tr>
<td>CFR Title 21, 177.2440</td>
<td>CFR Title 21, 177.1520</td>
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Pore Size
0.1 µm

Operating Characteristics
- Operating temperature range: 32 °F to 122 °F (0 °C to 50 °C)
- Maximum operating pressure: 100 psig @ 122 °F (6.9 bar @ 50 °C)
- Maximum temperature rating: 160 °F @ 35 psig (72 °C @ 2.4 bar)
- Maximum reverse pressure: 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, ≥ 3 cycles.
- Gamma irradiation: 25 to 40 kGy once. Do not autoclave irradiated capsules.
- Capsules must not be in-line steam sterilized.

Biological Safety
STyLUX® filters meet the requirements as specified in the current USP <88> Class VI plastics, physicochemical, oxidizable substances, and USP <87> cytotoxicity tests. Bacterial endotoxin levels in aqueous extracts of STyLUX® filters are less than 0.5 EU/mL, as determined using the Limulus amebocyte lysate (LAL) test USP <85>. No binders, adhesives or surfactants are used in the construction of STyLUX® filters. Filters comply with European Commission Regulation No 10/2011.

Quality Assurance
STyLUX® 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 STyLUX® filter is clearly marked with filter type, lot number and serial number.
Ordering Guide

C K 2 SL 0.1 - 11 2

Capsule Options
- Standard capsule: C
- Sterile capsule: G

Effective Filtration Area
- 0.45 ft² (415 cm²): K
- 0.36 ft² (334 cm²): M

Vent/Drain Ports
- 0: No vent/drain port
- 1: One luer port with cap, inlet side
- 2: Standard; two luer ports with caps
- 4: Two sanitary valves with hose barbs
- 5: One sanitary valve with hose barb, inlet side
- 6: One sanitary valve with hose barb, outlet side

Inlet/Outlet Connections
- 11: 1/4" hose barb
- 1B: 1/4" hose barb w/filling-bell
- 22: 3/8" hose barb
- 2B: 3/8" hose barb w/filling-bell
- 41: 1/4" MNPT, 1/4" hose barb outlet
- 44: 1/4" MNPT
- 71: 3/4" sanitary flange inlet, 1/4" hose barb outlet
- 72: 3/4" sanitary flange inlet, 3/8" hose barb outlet
- 77: 3/4" sanitary flange

Housing Material Designator
- Gamma-stable polypropylene, animal component free (ACF): 2

Product
- STyLUX® hydrophilic PES membrane

Grade Retention Rating
- SL: 0.1 µm

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

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