

Ultradyn[®]

TA0.2 Capsule

Description

The Ultradyn[®] TA sterilizing grade 0.2 µm PTFE hydrophobic filter capsule is manufactured using high quality components that are nontoxic and biologically inert.

Materials of Construction

All components of the Ultradyn[®] filter capsule 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:

Membrane:	Polytetrafluoroethylene	CFR Title 21, 177.1550
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
Capsule housing:	Polypropylene	CFR Title 21, 177.1520
Sealing method:	Thermal bonding	

Pore Size 0.2 µm

Minimum Bubble Point 16 psi (1,1 bar), 60% IPA

Maximum Diffusion Rate 1.2 ft²: 2.0 mL/min @ 10 psi, 60% IPA (2,0 mL/min @ 0,7 bar, 60% IPA)
2.5 ft²: 4.2 mL/min @ 10 psi, 60% IPA (4,2 mL/min @ 0,7 bar, 60% IPA)

Maximum WIT Rate 1.2 ft²: 0.12 mL/min water @ 36 psi (0,12 mL/min water @ 2,48 bar)
2.5 ft²: 0.25 mL/min water @ 36 psi (0,25 mL/min water @ 2,48 bar)
Specification can vary by instrumentation; consult factory.

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

Typical Air Flow Rate 1.2 ft²: 7 scfm @ 1 psid (8,62 Nm³h⁻¹ @ Δp 50 mbar)
2.5 ft²: 12.5 scfm @ 1 psid (15,4 Nm³h⁻¹ @ Δp 50 mbar)

Operating Characteristics

Normal operating temperature range: 32 °F to 100 °F (0 °C to 38 °C)

Maximum operating temperature rating: 160 °F @ 35 psig (71 °C @ 2,4 bar)

Maximum operating pressure (liquid service): 75 psig @ 100 °F (5,2 bar @ 38 °C)

Maximum operating pressure (gas service): 50 psig @ 100 °F (3,4 bar @ 38 °C)

Sterilization

Autoclave: 121 to 135 °C (15 to 30 psi, 1 to 2 bar), 30 to 60 min, ≥ 3 cycles.

Capsules must not be steamed in place (SIP). Gamma irradiation is not recommended.

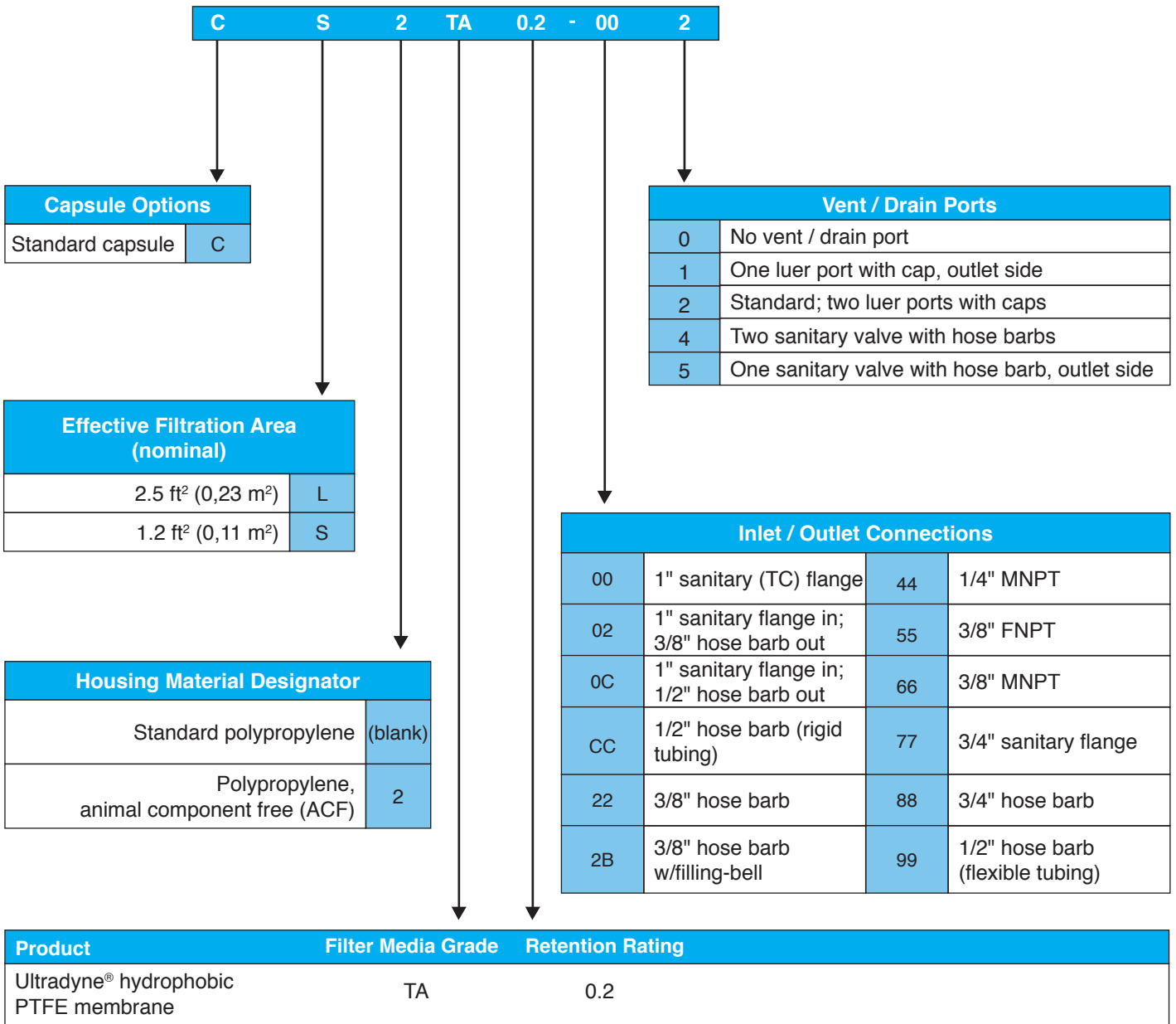
Biological Safety

Ultradyn[®] filters meet the requirements as specified in the current USP Class VI plastics, cytotoxicity and pyrogenicity tests. No binders, adhesives or surfactants are used in its construction. Filters comply with Commission Regulation (EU) No 10/2011.

Quality Assurance

Ultradyn[®] 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 Ultradyn[®] filter is integrity tested during manufacture and is clearly marked with filter type, lot number and serial number.

Ordering Guide



Additional information about Ultradyne® filter products is available in the Green Docs document which is viewable at <https://www.meissner.com/downloads/ultradyne-gd006.pdf>

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