

30 kDa Hollow Fiber Tangential Flow Filter Capsule

Description

SepraPor® XFC30 is a tangential flow filtration (TFF) capsule containing hollow fiber membranes. It is ideal for use in a variety of biopharmaceutical applications, including ultrafiltration, diafiltration, and purification. This TFF capsule is available in a range of sizes from bench through production scale for ease of scalability. The Seprapor® filter capsule is 100% integrity tested during manufacture and meets the critical demands of the pharmaceutical, biotechnology, and related industries.

Materials of Construction

All components of the filter are animal component free (ACF). These materials are listed for food contact use in the Code of Federal Regulations (CFR), Title 21, as below:

Polysulfone Hollow fibers: CFR Title 21, 177.1655 Fiber bundle netting: Polypropylene CFR Title 21, 177.1520 Capsule housing: Polysulfone CFR Title 21, 177.1655 Polysulfone Permeate ports: CFR Title 21, 177.1655 Retentate ports: Polysulfone CFR Title 21, 177.1655 Fiber encapsulation: Ероху CFR Title 21, 175.300

Retention Rating30 kDaTypical Fiber Lumen1.0 mm

Fluid Path Length (Nominal) 12 in (30 cm) 24 in (60 cm)

Maximum Diffusive Flow Rate 32 mL/min/m² @ 30 psig (2.07 bar), water with air

Normalized Clean Water Flux 80 - 265 LMH/bar (Liters/m²/h/bar) @ 25 °C

Operating Characteristics

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

Maximum operating temperature: 122 °F (50 °C) for short-term operations such as cleaning

Maximum feed pressure: 65 psig @ 77 °F (4.48 bar @ 25 °C)
Maximum transmembrane pressure: 50 psig @ 77 °F (3.45 bar @ 25 °C)

Sterilization

Autoclave: 121 °C to 123 °C (15 psi, 1.03 bar), 30 minutes.

Apply a validated autoclave sterilization cycle that increases and decreases temperature gradually. Consult autoclave sterilization instructions in the SepraPor® Green Doc document for further guidance. Capsules must not be in-line steam sterilized.

Biological Safety

SepraPor® filters meet the requirements as specified in the USP 43 Biological Reactivity Tests, in vitro <87> (cytotoxicity) and in vivo <88> (Class VI Plastics).

Quality Assurance

SepraPor® 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 SepraPor® filter is clearly marked with filter type, lot number, and unique serial number.



Ordering Guide

Part Number	Filter Diameter	Fluid Path Length	Effective Filtration Area	Retentate Ports ¹	Permeate Ports ¹
XFC030C012-7711	³ /8" (0.95 cm)	12" (30 cm)	0.12 ft ² (0.011 m ²)	3/4" sanitary flange	1/4" hose barb
XFC030C024-7711		24" (60 cm)	0.25 ft ² (0.023 m ²)		
XFC030C112-7722	³ / ₄ " (1.9 cm)	12" (30 cm)	0.45 ft ² (0.042 m ²)		³/8" hose barb
XFC030C124-7722		24" (60 cm)	0.91 ft ² (0.085 m ²)		
XFC030C212-00CC	1" (2.5 cm)	12" (30 cm)	0.81 ft ² (0.075 m ²)	1" sanitary flange	½" hose barb
XFC030C224-00CC		24" (60 cm)	1.6 ft ² (0.15 m ²)		
XFC030C312-FFCC	11/4" (3.2 cm)	12" (30 cm)	1.3 ft ² (0.12 m ²)	1½" sanitary flange	
XFC030C324-FFCC		24" (60 cm)	2.5 ft ² (0.23 m ²)		
XFC030C412-FFCC	2.0" (5.1 cm)	12" (30 cm)	3.9 ft ² (0.36 m ²)		
XFC030C424-FFCC		24" (60 cm)	9.0 ft ² (0.84 m ²)		
XFC030C512-FF00	3.0" (7.6 cm)	12" (30 cm)	9.9 ft ² (0.92 m ²)		1" sanitary flange
XFC030C524-FF00		24" (60 cm)	23 ft² (2.1 m²)		

¹ Retentate and Permeate port options are subject to capsule diameter size. For custom ports, please contact Meissner.

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