

SUREPORE PES MEMBRANE FILTER CARTRIDGE

PES Membrane - Sterile Liquid Filter

SurePore PES Filter Cartridges are specially designed to provide a reliable sterilizing solution at an economical cost. Hydrophilic PES membrane cartridges require no prewetting and are ready to use. In addition, these filters provide excellent performance in pharmaceutical applications.



FEATURES AND BENEFITS

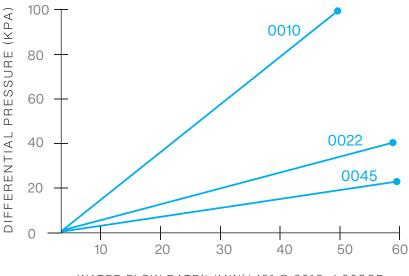
- Low diffusion flow
- Inherently hydrophilic PES membrane
- High surface area provides excellent flow rates and extended service life while maintaining high bacteria removal efficiency
- · Low protein binding



QUALITY STANDARDS

- Bacterial quantitative retention of 107 CFU/ cm2 Brevundimonas Diminuta (ATCC 19146) according to ASTM F838 methodology.
- 100% Integrity testing in manufacturing.
- Each filter is fully traceable with unique serial number.
- Manufactured in a facility which adheres to ISO 9001:2015 Practices.
- Full Regulatory Compliance with followign:
- Bacterial Endotoxin: Aqueous extraction of autocalved filter contains
 <0.25 EU/ml as determined by Limulus Amebcyte Lysate (LAL), USP
 <85>.
- Non-fiber Releasing: Component materials meet the criteria for a "Non-fiber-releasing filter" as defined in 21 CFR 210.3(b)(6).
- Component Material Toxicity:
- Meet the requirement of USP <87> In Vitro Cytotoxicity Test;
- Meet the Criteria of USP <88> Biological Reactivity Test for Class VI-121°C plastics.
- TOC/Conductivity at 25°C: Autoclaved filter effluent meet the USP <643> for Total Organic Carbon and USP <645> for Water Conductivity per WFI requirements after a UPW flush of specified volume.
- Particle Shedding: Autoclaved filter effluent meet the USP <788> for large volume injections.
- Indirect Food Additive: All component materials meet the FDA indirect Food Additive requirements cited in 21 CFR 177-182, and EU framework regulation [1935/2004/EC]





WATER FLOW RATE(L/MIN)/ 10" @ 20°C, 1.00SCP

MATERIALS OF CONSTRUCTION				
Filter Media	PES Membrane			
Cage/Support	Polypropoylene			
Core/End Caps	Polypropoylene			

	STERILIZATION
Inline Steam Sterilization	up to 100 cycles (135°C for 30min < 0.3 bar per cycle)
Autoclave	up to 200 cycles (130°C for 30min per cycle)

INTEGRITY TEST DATA			
Bubble Point	BP: ≥ 0.32 MPa (water), 0.22 µm BP: ≥ 0.20 MPa (water), 0.45 µm		
Diffusion Flow	DF: ≤ 25 ml/min/10" @ 0.275 Mpa, 0.22 µm DF: ≤ 25 ml/min/10" @ 0.15 Mpa, 0.45 µm		

OPERATING CONDITIONS					
Max. Operating Pressure		6.9 bar (100 psi) at 25°C			
		4.0 bar (58 psi) at 60°C			
		2.4 bar (35 psi) at 80°C			
Max. Differential Pressure	Forward	6.9 bar (100 psi) at 25°C			
		4.0 bar (58 psi) at 60°C			
		2.4 bar (35 psi) at 80°C			
	Reverse	3.0 bar (44 psi) at 25°C			
		1.0 bar (15 psi) at 80°C			
Effective Filtration Area		0.58m²/φ 69-10inch			

ORDERING INFORMATION						
PFSHR	Removal Ratings	End Cap	Minimal Length	Seal Material	-P	
	0010 =0.10μm	HSF = 226/Fin (PBT Insert)	05 = 5"	S = Silicone		
	0022 =0.22µm	HSC = 226/Flat (PBT Insert)	10 = 10"	E =EPDM		
	0045 =0.45µm	HTF = 222/Fin (PST Insert)	20 = 20"	V = Viton		
		HTC = 222/Flat (PBT Insert)	30 = 20"	P = PFA/Viton		
		DOE = Double Open End	40 = 20"			

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