Pharmsteri[™] PES Disposable Filter

Cost-effective, safe, sterile filtration for pharmaceutical applications

PharmsteriTM PES disposable filters are designed for provide smart, disposable, safe filtration for pharmaceutical applications. Constructed of polypropylene (PP) components and a hydrophilic polyethersulfone (PES) membrane, these disposable filters can help eliminate cleaning problems and extra validation costs.

Our portfolio of Pharmsteri PES filters is 100% tested during manufacturing to ensure high quality performance, providing sterility assurance. The T-style configuration is ideal for quicker and easier installation.



Allows high flow and low hold-up volume of liquids.

APPLICATIONS

- · Bioburden reduction or sterile filtration for injections
- · Bulk chemicals
- Buffers

FEATURES & BENEFITS

Polypropylene and PES construction	Allows use in a broad range of applications
1.5" Tri-Clamp connector	Allows easier, user-friendly installation
Optimized form factor	Allows high flow and low pressure drop, lowering cost of ownership
All components meet USP <88> Class VI Plastics Test	Provides biosafety insurance and global regulatory compliance



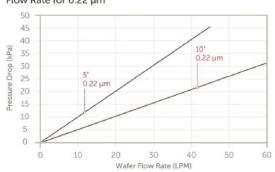


SPECIFICATIONS

Retention rating	0.22,0.45,0.45+0.22µm			
Materials	Membrane	Polyethersulfone (PES)		
	Housing, support	Polypropylene		
	Sleeve, core, top, end caps	Polypropylene		
	O-ring	Silicone		
Sealing technology	Thermal bonding without adhesives			
Filtration area	Single layer filter,10"	≥0.6 m²		
	Dual layer filter,10"	≥0.53 m²		
Integrity test (0.22 µm)	Minimum bubble point at 25°C (77°F)	≥3500 mbar		
	Maximum diffusion flow at 2800 mbar @ 25°C (77°F)	5": ≤15 mL/min		
		10": ≤30 mL/min		
Maximum operating conditions	Maximum forward differential pressure	3.5 bar @ 25° \pm 2°C (77°F), 3.0 bar @ 60° \pm 2°C (140°F)		
Sterilization cycles	Maximum 5 cycles of autoclave @ 127°C (261°F), 30 min			
Endotoxin releasing	<0.25 EU/mL according to the USP test methodology			
TOC	<0.5 mg/L after a water flush of 11 L/10" filter			
Conductivity	≤1.3 µS/cm after water flush of 11 L/10" filter			
Particle release	Meets the requirements of WFI set by USP			
NVR	After autoclaving and 24-hours in 50% ethanol water with vibration (contact Entegris for details)			
Material toxicity	All components meet the requirements of the USP <88> Reactivity Test for Class VI Plastics			
Bacterial challenge test	Passed the bacterial challenge test using <code>Brevundimonas diminuta</code> (ATCC 19146) at a minimum challenge concentration $1x10^7$ CFU/cm² (0.22 μ m) per ASTM methodology			
Indirect food additive	Meets the FDA Indirect Food Additive requirements cited in 21 CFR 177-182			
Food contact material	(EU) No. 1935/2004, (EU) No. 10/2011			
Quality	Manufactured in accordance with ISO 9001 and ISO 13485 certified quality management system			

PERFORMANCE DATA

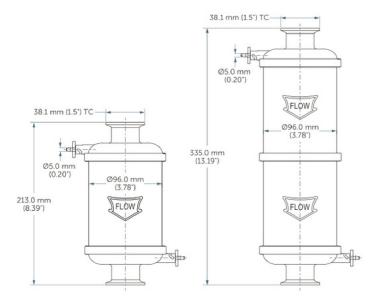
Flow Rate for 0.22 μm



Flow Rate for 0.45 μm 25 20 10° 0.45 μm 5 0.45 μm 6 0.45 μm 7 0.45 μm 8 0.45 μm 9 0.45 μm 9 0.45 μm 9 0.45 μm 9 0.45 μm



DIMENSIONS



ORDERING INFORMATION

PART NUMBER	RETENTION RATING	LENGTH	
PSTSUS022FS05-A2	0.22 µm	5"	
PSTSUS022FS10-A2	0.22 µm	10"	
PSTSUS045FS05-A2	0.45 µm	5"	
PSTSUS045FS10-A2	0.45 µm	10"	
PSTSUSD42FS5-A2	0.45+0.22µm	5″	
PSTSUSD42FS10-A2	0.45+0.22µm	10"	

FOR MORE INFORMATION

Please call your Regional Customer Service Center today to learn what Entegris can do for you. Visit <u>Argenpore.com</u> and select the <u>Contact Us</u> link to find the customer service center nearest you.

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