

Surface (Non-woven)



MICRO-PURE

АМРАтуре

Major Applications

Pre-filtration application

Pharmaceutical chemicals, Pharmaceutical ingredients, Drug substance, Manufacturing water

Quality standards

- Manufactured in ISO 9001 certified plant
- FDA 21 CFR compliant
- USP Class VI plastic biological safety testing compliant
- Certificate of quality is attached to the product
- Traceability by lot number

Fe	atu	res

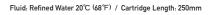
- 100% Polypropylene
- Pleated type with wide filtration area
- Variety of micron ratings

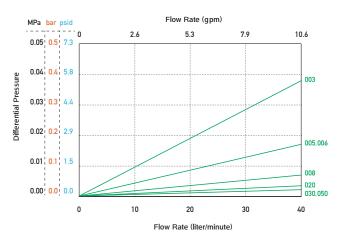
Advantages

- Wide range of fluid compatibility
- Excellent differential pressure vs. flow rate
- Optimal micron ratings selection according to the purpose

			Spec	ification	S			
	Grades	003	005	006	008	020	030	050
Mic	ron Ratings (μ m)	0.25	0.45	0.6	0.8	2.0	3.0	5.0
8	F.A. (m²/250L)	0.45	0.45	0.45	0.45	0.50	0.52	0.55
	Media		-		Polypropylene			
Materials	Core/Cage/Support				Polypropylene			
	End Cap	Polypropylene						
	Maximum ∆P			0.49MF	Pa at 20℃ (71psi	at 68°F)		
Maxin	num Operating Temp				80°C (176°F)			
	Length			125,	/250/500/750	mm		
Dimen- sions	0.D.				70.0mm			
	I. D.			26.1 (for F)/	26.9 (for 3, 4)/3	0.0 (for 7) mm		
Availabl	e sterilization methods		Inline stear	m, Autoclave *Ap	oplicable to only co	de 3, 4, 7 with Silic	one O-rings.	
Inline	steam sterilization	135°C (275°F) x 30 minutes x 10 cycles *Applicable to only code 3, 4, 7 with Silicone O-rings.						

Differential Pressure vs Flow Rate





Pa	Particle Removal Efficiency						
Grades		Pa	article Re	moval Ef	ficiency (%)	
Particle Size (μ m)	003	005	006	008	020	030	050
2.5	> 99.9	> 98	> 98				
3.5		>99.9	> 99.9				
4.5				> 98			
5.5				>99.9			
6.0					> 99	> 98	>98
8.5					> 99.9	>99.9	> 99.9
					•		
	G	est Co	nditions				
	F	Equipme Filtration Fluid	: 9	Liquid P Single P Refined	ass	Counter	

Flow Rate	: 10 liter/minute	

Dust : ACFTD+LATEX Beads

Validation items

Items	Evaluation criteria			
Endotoxin (LAL)	Extraction volume with water is less than 0.25 EU/mL and complies with USP \langle 85 $ angle$ requirements.			
Evaporation residues	Less than 5 mg of evaporation residue per 250 mm cartridge after 24 hours in ultrapure water following autoclave sterilization			
Potassium permanganate consumption	Meets the requirements of the USP Oxidizable Substance Test by flushing with at least 5,000 mL of ultrapure water after autoclaving			
Filter/component toxicity	USP (88) Biological Reactivity Tests For Class VI Plastics compliant			

*Please refer to the Validation Guide for detailed testing information.

Ordering Information



Code F Code 3 Code 4 Code 7 Image: Imag

*The contents of the catalog are subject to change without notice. *The performance data listed in the catalog are Typical values obtained under specific conditions based on our tests.

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Manufacturing is based on our Quality Management Systems that meet ISO9001 standards. Scope Design, Development, manufacture, and sales of filter cartridges, housings and filtration equipment.



5th Issue

Particle Removal Efficiency