Pleated Polyethersulfone (PES) Membrane Cartridge Filters



The Nucart® PES cartridge filters are available in $0.1\mu m$, $0.22\mu m$ & $0.45\mu m$ ratings and are especially designed & developed for critical applications such as sterile filtration, high product recovery, biological samples filtration in the Pharmaceuticals, Biologicals, Beverages & Allied industries. Available in various size options (3"-40"), these filters offer better throughput and are best suited as final filters or point-of-use filters in diversified applications.

Special Features & Benefits

- 100% integrity tested
- Reusable after multiple sterilization cycles
- Broad Chemical Compatibility across organic and aqueous sample; 1-14 pH
- Offered with High-flux hydrophilic PES membrane

Key Applications

- Buffer Filtration
- Vaccine Filtration
- Final Sterile Grade Filtration
- LVPs

- Water for Injections (WFIs)
- Sterilization of Hormonal Injectables
- Microbial Retention & Stabilization

Our Nucart Cartridge filters are quality assured for retention efficiency, integrity test and flow rate and validated for Heat Stability, Beta ratio test, fiber particle release, extractables and biosafety

In Complaince with Global Standards

Bacterial Endotoxin	The filtrate/Aqueous extraction from downstream of the filter exhibited endotoxin result < 0.25 EU/mL when tested as per USP <85> methodology
Oxidizable Substances	Oxidizable matter in filtered water meets the USP <1231> Oxidizable Substance Test requirements
Non-fiber Releasing	Meets the criteria for a "non-fiber releasing" filter as defined in 21 CFR 210.3(b)(6).
Particle Shedding	Meets Cleanliness per USP <788> for Particulates in Injectables
Extractable with water	Extractable passes within limit as specified by USP <661>
TOC/	Meets the USP <643> for Total Organic Carbon
Conductivity	Meets the USP <645> for Water Conductivity

- Manufactured in an ISO Class 8 Cleanroom Environment
- Complete Qualification Guide Available
- Critical raw material used for manufacturing are Compliant with FDA Indirect Food Additive requirements cited in 21 CFR 177.1520 & 21 CFR 177.2440
- Comply with USP <88> Reactivity
 Test for Class VI plastics
- Wide Chemical Compatibility
- 100% Integrity Tested

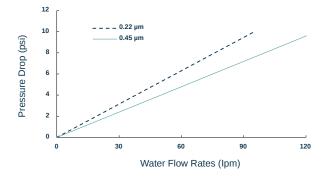
TECHNICAL SPECIFICATIONS



CONSTRUCTION MATERIALS

Filter Media: Polyethersulfone PES **Supporting Media:** Polyester

Core & Cage: Polypropylene



MAXIMUM OPERATING DIFFERENTIAL PRESSURE AND TEMPERATURE

Max Temp 80 °C @ ≤ 2 Kg/cm^2 Max Pressure 3.5 Kg/cm^2 @ 25 °C

Autoclavable 30 autoclave/steam sterilization cycles of

30 minutes at 121 °C/ 135 °C

INTEGRITY TEST DATA

Bubble Point:

0.10 μ m: \geq 1931 mbar (28 psi) (70% IPA/ water wetted) 0.22 μ m: \geq 3447 mbar (50 psi) (with water wetted) 0.45 μ m: \geq 2206 mbar (32 psi) (with water wetted)

Max Air Diffusion Flow (for 10" Cartridge):

0.10 μ m: \leq 30mL/min @3447 mbar (50 psi) (with water wetted)

0.22μm: ≤ 30 mL/min @ 2551 mbar (37 psi) 0.45μm: ≤ 35 mL/min @ 1655 mbar (24 psi)

Microbial Retention:

0.22μm: LRV > 7 for Brevundimonas diminuta 0.45μm: LRV > 7 for Serratia marcescens

ORDERING INFORMATION CODES:

Туре		Size			
Туре	Code		Length	EFA	Code
Single Layer	CFPE		3"	0.2m ²	03
CFPE	CFPER2		5"	0.3m ²	05
0.8µm upstream			9.75"	0.7m ²	9.75
CFPE	CFPER1		10"	0.7m ²	10
0.65µm upstream			20"	1.4m ²	20
CFPE	CFPEQ		30"	2.1m ²	30
0.45µm upstream	CFFEQ		40"	2.8m ²	40
0.2µm upstream	CFPEP				
High temp. PES	CFPH				

Pore Size		Ada	ptor	Rings / Gaskets		
Micron	Code	Type	Code	Ring	Code	
$0.10\mu m$	010	7P	U	Silicone	SS	
$0.22\mu m$	020	BEO	V	Viton	SV	
0.45 µm	045	K SEAL	K	EPDM	SE	
		Optiseal	X	Encapsulated	FV	
		4463	Υ	PTFE	ΓV	
		4463B	Z	Synthetic		
		M Disc	Q	Rubber SR		
		222	R	No Ring	XX	
		4440	W			

EXAMPLE: CFPE10045USS

Note: Other Specification are available on request



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