

mdi AseptiSure KR Cartridge filters incorporate a low protein binding PES membrane with polypropylene drainage layers to ensure pH compatibility from 1-14 making these ideal for alkaline fluid streams.

Special Features

- Low protein binding
- Very wide chemical compatibility, even with very high alkaline solutions.
- Large filtration area
- High throughputs
- pre-flushed to minimize particulate release after installation
- Non-toxic material of construction
- Multiple steam sterilizable
- Heat sealed, no glues or adhesives



Applications

- Sterile filtration of alkaline solutions for pH control

Microbially Validated as per ASTM F 838-05

Complies with USDFA 21 CFR 210.3(b)(6)

Meets and Exceeds USDFA 21 CFR 177.1520

Specifications

Construction				
Final Filter Pore Size	0.2µm		0.45µm	
Membrane	Hydrophilic PES			
Support Layers	Polypropylene			
Body and Core	Polypropylene			
Integrity Testing / Retention				
Bubble Point	> 50psi (3.52Kg/cm ²) with Water		> 30psi (2.11Kg/cm ²) with Water	
Air Diffusion Flow (10")	< 40ml/min @ 37 psi (2.6Kg/cm ²) with Water		< 45ml/min @ 22 psi (1.54Kg/cm ²) with Water	
Microbial Retention	LRV >7 for Brevundimonas diminuta (ATCC 19146) per cm ²		LRV >7 for Serratia marcescens (ATCC 14756) per cm ²	
Size				
Size	5"	10"	20"	30"
Effective Filtration Area (Nominal)	3000cm ²	6000cm ²	12000cm ²	18000cm ²
Operational				
Max. Operating Temperature	80 °C @ < 2 Kg/cm ² (30 psi)			
Max. Differential Pressure	3.5 Kg/cm ² (50 psi) @ 25 °C			
Reverse Pressure	< 0.7 Kg/cm ² (10 psi) @ 25 °C			
Typical Water Flow Rates (10")	40 lpm @ 0.70 Kg/cm ² @ 27 °C		55 lpm @ 0.70 Kg/cm ² @ 27 °C	
Sterilization	Autoclavable/In-line steam sterilizable at 121 °C for 30 minutes, 25 cycles			

Assurance	
Toxicity	Passes Biological reactivity test, In Vivo, as per USP <88> for Class VI plastics
Bioburden	Bioburden level is < 1000 cfu/filter device as per ANSI/AAMI/ISO 11737-1 : 1995
Bacterial Endotoxin	Aqueous extracts exhibit < 0.25 EU/ml as established by Limulus Amebocyte Lysate (LAL) Test as per USP <85>
Non Fiber Releasing	Passes test as per USP and comply with USFDA 21 CFR Part 210.3(b)(6) for fiber release
pH Compatibility	Compatible with pH range of 1 - 14
Extractables with WFI	Passes NVR test as per USP <661>
Oxidizable Substances	Passes test as per USP <1231>
Particle Shedding	Complies with USP <788> test for particulate matter in injections
TOC/Conductivity at 25 °C	Meets the WFI requirements of USP <643> for Total Organic Carbon and USP <645> for Water Conductivity after a specified volume of purified water flush
Indirect Food Additive	All Polypropylene components meet the FDA Indirect Food Additive requirements cited in 21 CFR 177.1520
Good Manufacturing Practice	These products are manufactured in a facility which adheres to Good Manufacturing Practices.
Quality Management System	ISO-9001 Certified
USFDA	DMF No. 015554

Ordering Information

Type		Size		Pore Size		Adaptor		Elastomer		Sterility		Pack Size	
	Code	Length and filtration Area	Code		Code		Code		Code		Code		Code
AseptiSure KR	CPKR	5" (3000 cm ²)**	53	0.2 µm	01	7P	A0	Silicone	SS	Non Sterile	1	1	01
		10" (6000 cm ²)	54	0.45 µm	02	7P without fin	A1	EPDM	SE				
		20" (12000 cm ²)	55			28 with Fin	C0	Viton	SV				
		30" (18000 cm ²)	56			'0'	D0	FEP Encapsulated Viton	FV*				

Example

CPKR	55	01	A0	SS	1	01
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* FV is available in Adapter Code A0 (7P) only

** Size 5" are available in Code A0 (7P) and A1 (7P without fin) only

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