



0.1 μm *AseptiCap* KS

Sterilization Grade Hydrophilic Polyethersulfone (PES) Membrane Devices for Liquid Streams in Biopharmaceuticals

Data Sheet

Biopharmaceutical processing requires sterilizing grade microfiltration at multiple stages to meet specific process requirements.

Processes managers are continuously looking for microfiltration solutions to upstream, downstream, intermediate processes and final biological preparations. Since bio manufacturing is a multi stage process and bio molecules by nature are extremely sensitive, they are looking for:

- Minimizing protein losses due to adsorption to improve over all product yields
- Minimizing filter extracts which add up due to multiple points of use in a process
- High throughputs to achieve process economy
- Choice of filter end connections for easy and reliable quick connections
- Absolute retentions for higher sterility assurance

mdi produces a wide range of Sterilizing grade PES membrane devices to meet filtration requirements of biopharmaceutical processing. These filter devices are validated to meet compendia and regulatory requirements and are well characterized. They meet key process requirements such as high retention efficiency, very high protein recoveries, extremely low extractables, high throughputs, wide chemical compatibility and other important characteristics.

With the advantages of pre filtration layer built into the device for higher throughputs, linear scalability of filter area for smooth transitions from lab scale to pilot to process scale and widest range of end connections for quick and reliable connections to the existing fittings, **mdi** *AseptiCap* KS filters are a universal solution for process filtration.

AseptiCap KS

PES Membrane Devices for Biopharmaceuticals

Datasheet

AseptiCap KS 0.1 micron capsule filters uses **mdi** PES membrane in Polypropylene housing. No adhesives or glue are used in the manufacturing process and all bonding is done by heat welding.

The products are deeply validated for use in Biopharmaceutical applications and specially recommended for single use systems. *AseptiCap KS* are manufactured in class 10,000 clean rooms and ISO 9001 certified facilities.

Applications

Sterile Filtration of

- Cell culture media
- Cell culture media containing serum
- Media additives
- pH adjusters
- Final product concentrates

Key Features

- Absolute retention
- 100% integrity tested
- Low protein binding
- Very low hold up volume in filters
- High flow rates
- Serial construction with prefilter for higher throughput with fouling streams
- Bioburden maintained below 1000 cfu/device
- Endotoxin level certified to be <0.25 EU/ml
- Widest range of end connections
- Products available for total scalability from a few ml to thousands of liters
- Total traceability through unique serial number for each filter
- Individual certificate of quality for each device
- Sterilizable by EO gas or autoclaving

Validation Services

The regulatory requirements emphasize on the need to validate the efficacy of the 'Sterilizing Filter' with drug product under simulated worst-case conditions of use.

mdi provides validation services supported by customized validation protocols and world class test facilities to assist you in filter validations with your specific drug product.

mdi's quality management system emphasizes on quality by design rather by end product testing. Robust processes are developed for product manufacturing and are continuously monitored to ensure that the products meet their predetermined specifications and lot to lot reproducibility is ensured.

Certificate of Quality

Each capsule filter is accompanied by individual certificate of quality to ensure traceable documentation at user's end.

It certifies the product compliance to various regulatory as well as user requirements.

Validated for Microbial Retention

Integrity test data have been correlated to actual microbial retention with *Acholeplasma laidlawii* ATCC 23206 at a challenge level $\geq 10^7$ organisms per cm^2 to establish acceptable integrity test values. Also validated for retention of *B. diminuta* ATCC 19146 as per ASTM F838-05.

100% Integrity Tested

Each *AseptiCap KS* is tested for integrity to comply with validated Acceptable Integrity Test Specifications.

Flow Rate

Each lot is tested for clean water flow rates to ensure that flow rates are within the specifications.

Adsorption

AseptiCap KS filters are validated for low protein binding to ensure minimal active ingredient losses when used for filtration of high value proteins.

Pressure, Temperature Endurance

AseptiCap KS filters are validated to endure high operating pressure and temperature conditions which may be encountered during use.

These filters are also validated for high burst pressure to ensure user safety in case of inadvertent pressure build-up.

Extractables

Extractables/leachables from sterilizing filters, used at various stages of a biopharmaceutical manufacturing process, will add on and may impact the impurity profile of the desired product.

AseptiCap KS filters are validated to exhibit low extractables under harsh extraction conditions.

Bioburden Testing

Device bioburden is tested as per ISO 11737-1 and assured to be < 1000 cfu/device.

Endotoxin Testing

Aqueous extracts exhibit < 0.25 EU/ml as established by Limulus Amebocyte Lysate (LAL) Test

Total Traceability

AseptiCap KS filters come with completely traceable lot numbers and unique identification number to facilitate easy and fast retrieval of manufacturing and quality control data.

These unique lot and identification numbers are laser etched on each filter device and also printed on the labels of the box in which individual filter is packed.

Packaging Integrity

AseptiCap KS filters are fitted with vent caps and are packed in pouch to ensure package integrity during transit as well as to prevent particulate contamination while transferring to clean room assembly or process areas.

Other Regulatory Compliance

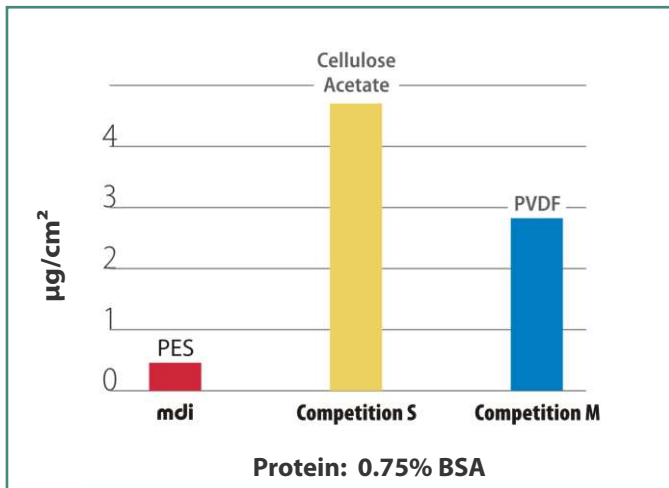
- Complies with USFDA 21 CFR 210.3(b)(6) for fiber release
- Complies with USFDA 21 CFR 177.1520 for fractional dissolution
- Materials of construction tested for toxicity as per Biological Reactivity Tests, In vivo, USP <88> for class VI Plastics
- Complete filter devices tested for cytotoxicity as per Biological Reactivity Tests, In vitro, USP <87>

Low Protein Binding

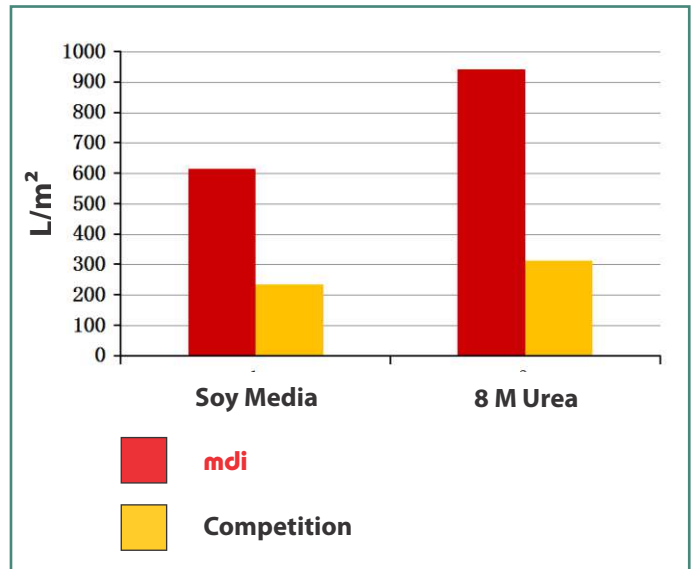
A comparative study on **mdi** PES membrane exhibits much lower protein adsorption than other competing membranes of Cellulose Acetate and PVDF.

The low protein binding results in increased overall product yield and higher throughputs with biological streams.

Protein Binding ($\mu\text{g}/\text{cm}^2$)



High Throughputs



The high throughput translates to lower filtration costs, less number of filter changes and overall economy of operations.

Very Low Hold-Up Volumes

mdi PES membrane capsule filters are designed to offer very low hold-up volumes to minimize filtration losses and maximize product recovery.

0.1 μm AseptiCap Filters	Protein Binding
25 mm, 5 cm^2	1.7 μg
50 mm, 20 cm^2	7 μg
1", 250 cm^2	88 μg
2", 500 cm^2	187 μg
10", 6000 cm^2	2275 μg

Filter Devices	EFA* (Nominal)	Hold up Volume
AseptiCap KS, 25mm	5 cm^2	< 50 μl
AseptiCap KS, 50mm	20 cm^2	< 200 μl
AseptiCap KS, 1"	250 cm^2	< 5ml
AseptiCap KS, 2"	500 cm^2	< 25ml
AseptiCap KS, 5"	1000 cm^2	< 45ml
AseptiCap KS, 8"	2000 cm^2	< 60ml

*EFA: Effective Filtration Area

Extractables

It is useful to evaluate extractables that may be leached out of the filter and enter the process stream. **mdi** filters give low extractables under harsh preconditioning and extraction conditions.

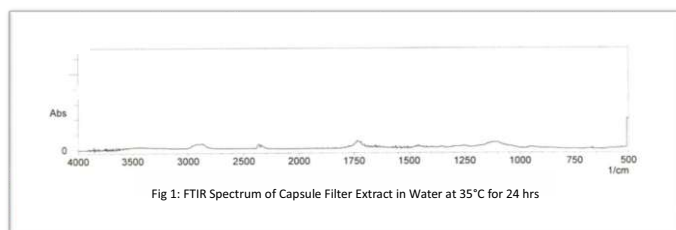
Low extractables mean less addition to impurity profile of the biological product from the filters.

Extraction Time: 24 hours

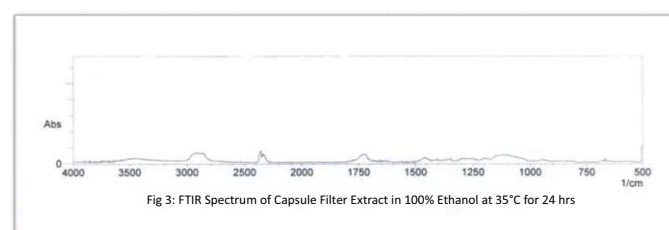
Model Solvent	Non Volatile Residue	
	<i>AseptiCap KS 1"</i> (250 cm ²)	<i>AseptiCap KS 10"</i> (6000 cm ²)
Water @ 35 °C	1.6 mg	38.26 mg
Water @ 80 °C	1.8 mg	43.04 mg

Model Solvent	Non Volatile Residue	
	<i>AseptiCap KS 1"</i> (250 cm ²)	<i>AseptiCap KS 10"</i> (6000 cm ²)
100% Ethanol @ 35 °C	13.4 mg	320.43 mg

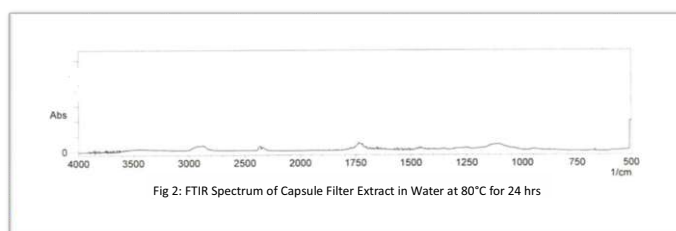
FTIR Analysis of Extractables From *AseptiCap KS 1"* Capsule Filter with Water @ 35 °C



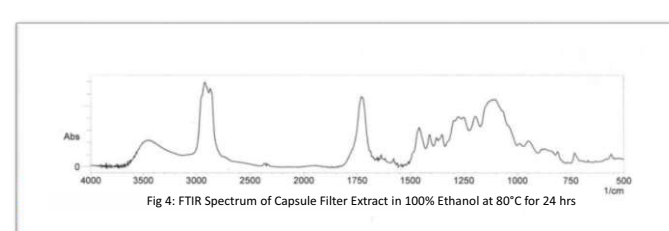
FTIR Analysis of Extractables From *AseptiCap KS 1"* Capsule Filter with 100% Ethanol @ 35 °C



FTIR Analysis of Extractables From *AseptiCap KS 1"* Capsule Filter with Water @ 80 °C



FTIR Analysis of Extractables From *AseptiCap KS 1"* Capsule Filter with 100% Ethanol @ 80 °C



The Spectrum of extracts from *AseptiCap KS* capsule filters with 100% ethanol under extreme extraction conditions show presence of various components used in the manufacture of **mdi** PES membrane capsule filters.

Easy Connect

Widest Range of End Connections

Biopharmaceutical processes involve transfer of high value fluids through multiple process steps. Making high quality, reliable, flexible and functionally convenient connectivity with filters is of utmost value to the bio-processors.

mdi *AseptiCap* KS filters offer a wide range of reliable end connections for functional convenience and customized connectivity.

Validated for Performance

These end connections are manufactured with tight dimension tolerance and are validated for strength and connection integrity under extreme use conditions as well as for their ability to withstand prevalent sterilization methods including EO sterilization and autoclaving.



1/2" HB



1/2" Single Stepped
Hose Barb



1/4" MNPT



1/4" SHB



Quick Connector



Male Luer Slip



3/8" Hose Barb



Female Luer Lock



1 1/2" Sanitary Flange



3/4" Sanitary Flange



1/2" MNPT



1" Hose Barb

Variety of end connections

Customized Connectivity

mdi *AseptiCap* KS filters are available in a wide range of end connections and are also customized to offer different inlet-outlet combinations to meet the unique connectivity needs in biopharmaceutical process assemblies where, for example, stainless steel components with sanitary flange connections are sometimes required to be connected to single use disposable systems through quick-connectors or hose barb connections.



1 1/2" Sanitary Flange
to 1/2" Barb Hose



1 1/2" Sanitary Flange
to 3/4" Sanitary Flange



AseptiCap with HighSecurity
1/2" hose barb connection

Linear Upscaling from R&D to Production Process

Datasheet

Scientists are concerned about filter fluid interaction impacting the stability, purity, strength etc. of the drug product, and they take a keen interest in filter selection at the formulation development stage itself. Although preliminary compatibility data support initial filter selection, for stability studies detailed filter validations are required to provide enough documented evidence to justify specific filter use.

A critical requirement that needs to be addressed at this stage is of scalability from R&D to pilot scale to full scale production processes.

mdi offers a wide range of *AseptiCap KS* filters to provide linear scale up from lab scale to production process. While scaling up the process, the appropriate size filter can be selected by increasing the effective filtration area of filter proportionate to the process fluid volumes.

All Materials of construction as well as manufacturing process are identical for all filter devices starting from 5 cm² to 18000cm² hence process scaling can be facilitated without triggering additional validation studies for given process conditions. **mdi** provides complete documentation for each of the *AseptiCap KS* filters there by reducing the additional validation cost and time.



AseptiCap KS
25mm, 5cm²



AseptiCap KS
50mm, 20cm²



AseptiCap KS
1", 250cm²



AseptiCap KS
2", 500cm²



AseptiCap KS
5", 1000cm²



AseptiCap KS
8", 2000cm²

Filter Devices	EFA* (Nominal)	Hold up Volume
<i>AseptiCap KS</i> , 25 mm	5cm ²	< 50µl
<i>AseptiCap KS</i> , 50 mm	20cm ²	< 200µl
<i>AseptiCap KS</i> , 1"	250cm ²	< 5ml
<i>AseptiCap KS</i> , 2"	500cm ²	< 25ml
<i>AseptiCap KS</i> , 5"	1000cm ²	< 45ml
<i>AseptiCap KS</i> , 8"	2000cm ²	< 60ml
<i>AseptiCap KS</i> , 5"	3000cm ²	< 80ml
<i>AseptiCap KS</i> , 10"	6000cm ²	< 150ml
<i>AseptiCap KS</i> , 20"	12000cm ²	< 250ml
<i>AseptiCap KS</i> , 30"	18000cm ²	< 350ml



AseptiCap KS
10", 6000cm²

0.1 µm AseptiCap KS (with Prefilter)

Construction

Membrane	0.1 µm Hydrophilic PES
Prefilter Membrane	0.2 µm or 0.45 µm Hydrophilic PES
Plastic parts	Polypropylene

Integrity Testing/ Retention

Bubble Point	≥ 31 psi (2.18 Kg/cm ²) with 50% IPA
Bacterial Retention	LRV> 7 for <i>Acholeplasma laidlawii</i> ATCC 23206 per cm ² of filter area
	LRV> 7 for <i>Brevudimonas diminuta</i> ATCC 19146 per cm ² of filter area as per ASTM F 838-05

Size

Size	25mm	50mm
Effective Filtration Area (Nominal)	5 cm ²	20 cm ²
Operational Radius (with Vent/ Drain)	15 mm	28 mm

Operational

Max. Operating Temperature	55 °C	60 °C
Max. Differential Pressure	75 psi (5 Kg/cm ²) @ 25 °C	42 psi (3 Kg/cm ²) @ 30 °C
Sterilization	By Gas	Sterilizable by Ethylene Oxide
	By Autoclave	Autoclavable at 125 °C for 30 minutes, 25 Cycles. Can not be in-line steam sterilized
Shelf Life	3 year after EO sterilization	

Assurance

Toxicity	Passes Bioreactivity test, In Vivo, as per USP <88> for Class VI plastics
Cytotoxicity	Passes Biological Reactivity Tests, In vitro, USP <87> for cytotoxicity
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
TOC and Conductivity	Meets the WFI requirements of USP for TOC <643> and Conductivity <645> after a 500ml flush
pH Compatibility	Compatible with pH range of 1 - 10
Extractables with WFI	Passes NVR test as per USP <661>
Indirect Food Additives	Comply with USFDA 21 CFR Part 177.1520
Oxidizable Substances	Passes test as per USP <1231>
Quality Management System	ISO-9001 Certified
USFDA	DMF No. 015554

0.1 µm AseptiCap KS (with Prefilter)

Construction				
Membrane	0.1 µm Hydrophilic PES			
Upstream Membrane (in case of AseptiCap KS-γ)	0.2 µm or 0.45 µm Hydrophilic PES			
Support Layers	Polyester			
Plastic parts	Polypropylene			
Integrity Testing/ Retention				
Bubble Point	≥ 31psi with 50% IPA/ Water solution			
Bacterial Retention	LRV> 7 for <i>Acholeplasma laidlawii</i> ATCC 23206 per cm ² of filter area			
	LRV> 7 for <i>Brevudimonas diminuta</i> ATCC 19146 per cm ² of filter area as per ASTM F 838-05			
Size				
Size	1"	2"	5"	8"
Effective Filtration Area (Nominal)	250cm ²	500cm ²	1000cm ²	2000 cm ²
Operational Radius (with Vent/ Drain)	40 mm	65 mm	65 mm	65 mm
Vent and Drain	¼" Hose Barb with Silicone "O" rings			
Operational				
Max. Operating Temperature	80 °C @ < 30 psi (2 Kg/cm ²)			
Max. Differential Pressure	60 psi (4 Kg/cm ²) @ 30 °C			
Sterilization	By Gas	Sterilizable by Ethylene Oxide		
	By Autoclave	Autoclavable at 125 °C for 30 minutes, 25 Cycles. Can not be in-line steam sterilized		
Shelf Life	3 year after EO sterilization			
Assurance				
Toxicity	Passes Bioreactivity test, In Vivo, as per USP <88> for Class VI plastics			
Cytotoxicity	Passes Biological Reactivity Tests, In vitro, USP <87> for cytotoxicity			
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			
TOC and Conductivity	Meets the WFI requirements of USP for TOC <643> and Conductivity <645> after a 3 liter flush			
pH Compatibility	Compatible with pH range of 1 - 10			
Extractables with WFI	Passes NVR test as per USP <661>			
Indirect Food Additives	Comply with USFDA 21 CFR Part 177.1520			
Oxidizable Substances	Passes test as per USP <1231>			
Quality Management System	ISO-9001 Certified			
USFDA	DMF No. 015554			

0.1 µm AseptiCap KS (with Prefilter)

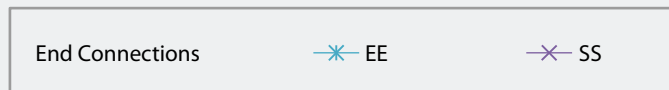
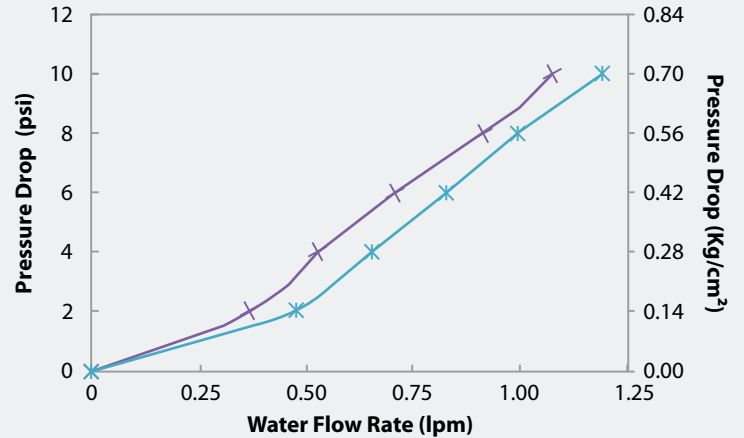
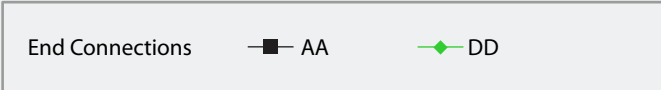
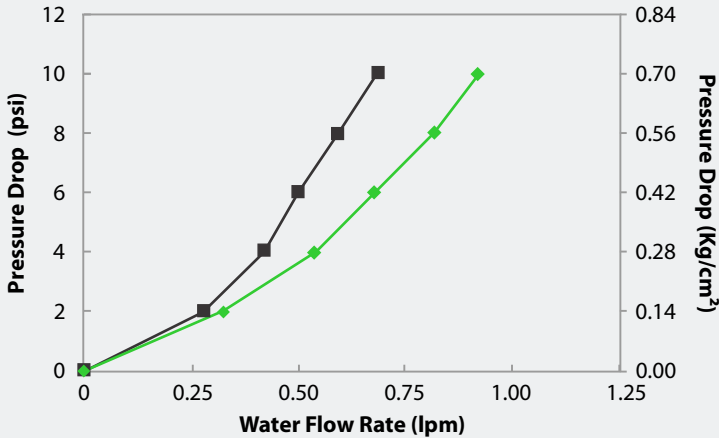
Construction		
Membrane	0.1 µm Hydrophilic PES	
Upstream Membrane (in case of AseptiCap KS-γ)	0.2 µm or 0.45 µm Hydrophilic PES	
Support Layers	Polyester	
Plastic parts	Polypropylene	
Integrity Testing/ Retention		
Bubble Point	≥ 31psi with 50% IPA/ Water solution	
Bacterial Retention	LRV> 7 for <i>Acholeplasma laidlawii</i> ATCC 23206 per cm ² of filter area	
	LRV> 7 for <i>Brevudimonas diminuta</i> ATCC 19146 per cm ² of filter area as per ASTM F 838-05	
Size		
Size	5" 10" 20" 30"	
Effective Filtration Area (Nominal)	3000cm ² 6000cm ² 12000cm ² 18000 cm ²	
Max. Air Diffusion Flow (@ 50psi (3.51 Kg/cm ²) with water)	≤ 15 ml/min ≤ 29 ml/min ≤ 58 ml/min ≤ 87 ml/min	
Operational Radius (with Vent/ Drain)	80 mm 80 mm 80 mm 80 mm	
Vent and Drain	¼" Hose Barb with Silicone "O" rings	
Operational		
Max. Operating Temperature	80 °C @ < 30 psi (2 Kg/cm ²)	
Max. Differential Pressure	60 psi (4 Kg/cm ²) @ 30 °C	
Sterilization	By Gas	Sterilizable by Ethylene Oxide
	By Autoclave	Autoclavable at 125 °C for 30 minutes, 25 Cycles. Can not be in-line steam sterilized
Shelf Life	3 year after EO sterilization	
Assurance		
Toxicity	Passes Bioreactivity test, In Vivo, as per USP <88> for Class VI plastics	
Cytotoxicity	Passes Biological Reactivity Tests, In vitro, USP <87> for cytotoxicity	
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	
TOC and Conductivity	Meets the WFI requirements of USP for TOC <643> and Conductivity <645> after a 10 liter flush for 5" capsule filters and 20 liter flush for 10" capsule filters	
pH Compatibility	Compatible with pH range of 1 - 10	
Extractables with WFI	Passes NVR test as per USP <661>	
Indirect Food Additives	Comply with USFDA 21 CFR Part 177.1520	
Oxidizable Substances	Passes test as per USP <1231>	
Quality Management System	ISO-9001 Certified	
USFDA	DMF No. 015554	

Typical Water Flow Rates

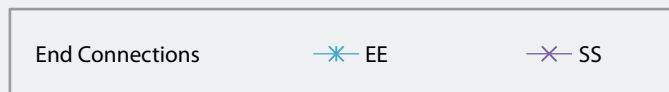
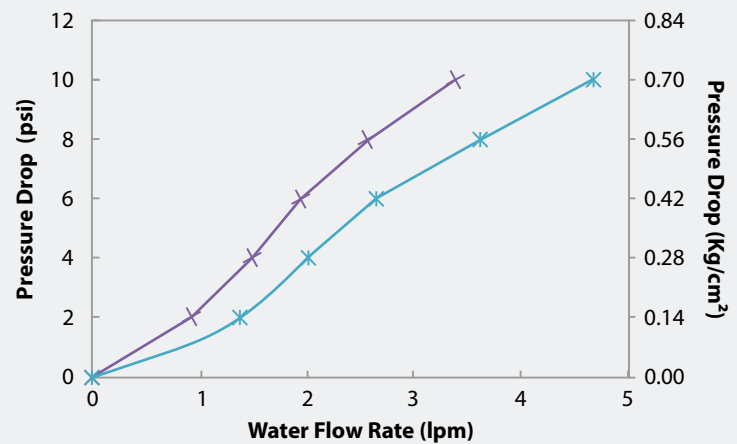
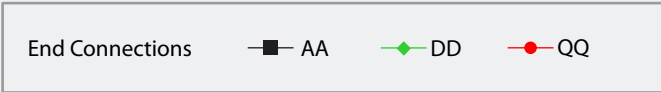
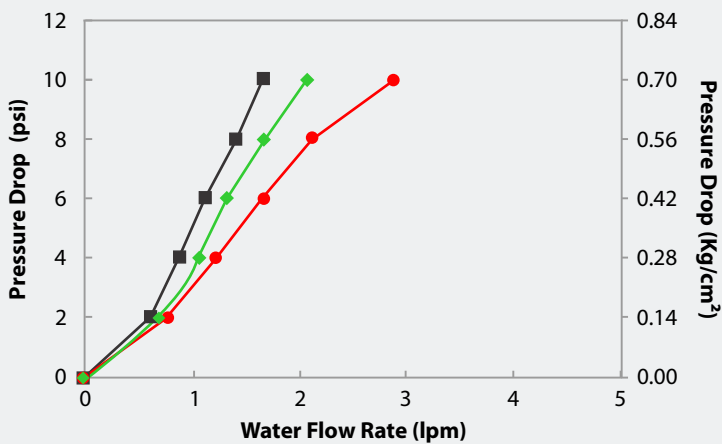
0.1 μm AseptiCap KS

Datasheet

AseptiCap KS, 1" Capsule Filter



AseptiCap KS, 2" Capsule Filter



End Connection Type:

A: 1/4" Stepped Hose Barb

E: 1/2" Sanitary Flange

D: 1/2" Hose Barb

S: 3/4" Sanitary Flange

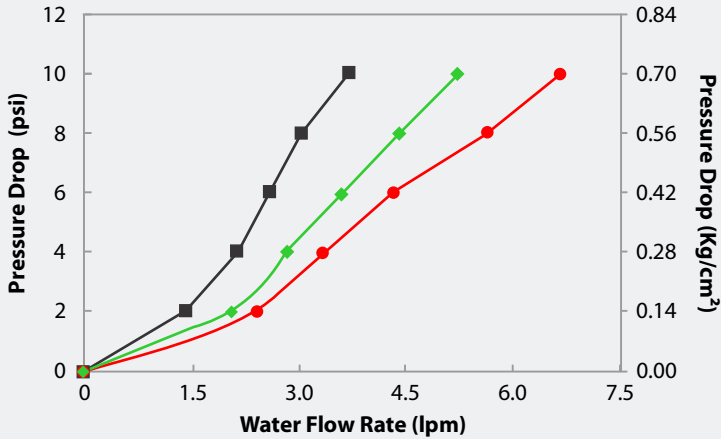
Q: 1/2" Single Step Hose Barb

Typical Water Flow Rates

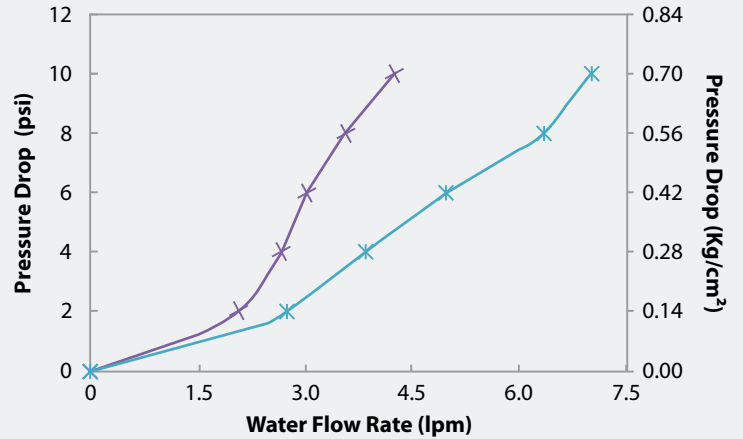
0.1 μm AseptiCap KS

Datasheet

AseptiCap KS, 5" Capsule Filter

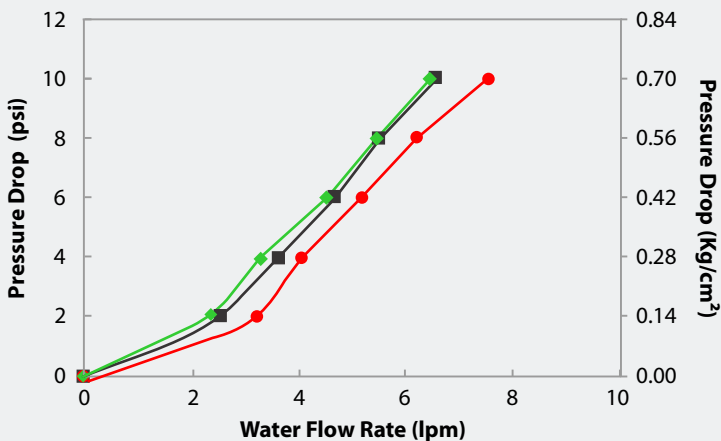


End Connections ■ AA ◆ DD ● QQ

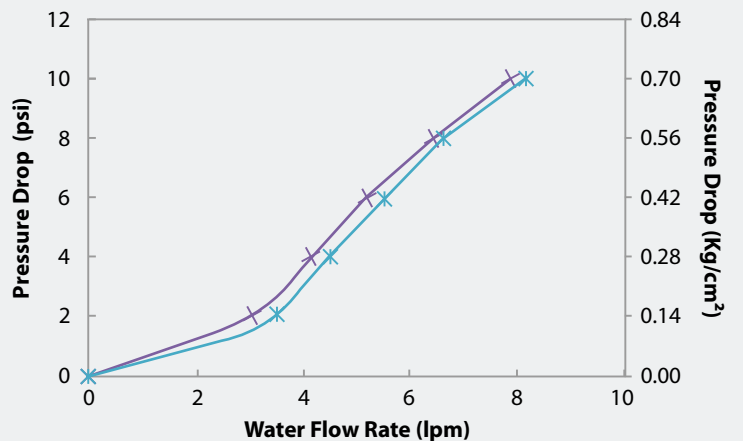


End Connections * EE * SS

AseptiCap KS, 8" Capsule Filter



End Connections ■ AA ◆ DD ● QQ



End Connections * EE * SS

End Connection Type:

A: 1/4" Stepped Hose Barb

E: 1/2" Sanitary Flange

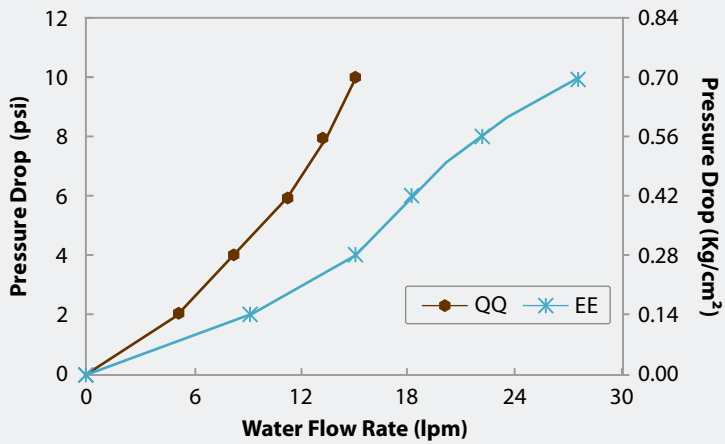
D: 1/2" Hose Barb

S: 3/4" Sanitary Flange

Q: 1/2" Single Step Hose Barb

Typical Water Flow Rates 0.1 μm AseptiCap KS

AseptiCap KS, 10" Capsule Filter



End Connection Type:

E: 1½" Sanitary Flange

Q: ½" Single Step Hose Barb

0.1 µm AseptiCap KS 25mm PES Membrane Capsule filter

Type		Size		Pore Size		Inlet/Outlet		X	X	Sterility		Pack Size	
	Code		Code		Code		Code				Code		Code
AseptiCap KS (0.45 µm Upstream)	IKSX	25mm	06	0.1µm	36	Female Luer Lock	M			Non Sterile	1	100	04
AseptiCap KS (0.2 µm Upstream)	IKS1					Male Luer Slip	N			EO Sterile	2		
						1/8" Hose Barb	H						
						1/4" Hose Barb	B						
Example:													
IKSX		06		36		MN		X	X	1		04	

0.1 µm AseptiCap KS 50mm PES Membrane Capsule filter

Type		Size		Pore Size		Inlet/Outlet		X	X	Sterility		Pack Size	
	Code		Code		Code		Code				Code		Code
AseptiCap KS (0.45 µm Upstream)	VKSX	50mm	10	0.1µm	36	1/4" SHB	B			Non Sterile	1	12	08
AseptiCap KS (0.2 µm Upstream)	VKS1					3/4" Sanitary Flange	S			EO Sterile	2		
Example:													
VKSX		10		36		BB		X	X	1		08	

Note: Inlet/Outlet Connections and Dimensions available with different diameter filters as follows:

Connections Available			Dimension (in mm)		
Inlet/Outlet	25mm	50mm	Inlet/ Outlet	25mm	50mm
1/4" - 3/4" Stepped Hose Barb	X	√	1/4" - 3/8" Stepped Hose Barb I/O	-	79
3/4" Sanitary Flange	X	√	1/4" Single Step Hose Barb I/O	38	-
Female Luer Lock	Inlet Only	X	3/4" Sanitary Flange I/O	-	51
Male Luer Slip	Outlet Only	X	Female Luer Lock Inlet/ Male Luer Slip Outlet	23	-
1/8" Hose Barb	√	X	1/8" Hose Barb I/O	36	-
Male Luer Lock	Outlet Only	X	Operational Radius	15	28
1/4" Hose Barb	√	X			

0.1 µm AseptiCap KS PES Membrane Capsule filter

Type		Size		Pore Size		Inlet/Outlet		X	Bell		Sterility		Pack Size	
	Code		Code		Code		Code			Code		Code		Code
AseptiCap KS (0.45 µm Upstream)	DKSX	1"	51	0.1 µm	36	¼" SHB	A		Yes*	B	Non Sterile	1	1	01
		2"	52			½" Hose Barb	D		No Bell	X	EO Sterile	2		
		5"	53			1½" Sanitary Flange	E		Bell with Cover	C				
8"	57	¾" Sanitary Flange	S											
AseptiCap KS (0.2 µm Upstream)	DKS1					Quick Connector	J							
						½" Single Step Hose Barb	Q							
						Female luer lock	U							
						Male luer slip	W							
						⅜" Hose Barb	N							
						⅝" Hose Barb	I							

*Bell is available with
½" HB outlet connections in 1", 2", 5" and 8" capsule filters
¼" SHB outlet connection in 1" capsule filters only

Example:

DKSX	57	36	DD	X	X	1	01
------	----	----	----	---	---	---	----

Inlet/Outlet Connections available with different Sizes/Length as follows:

Inlet/Outlet	Size/Length			
	1"	2"	5"	8"
¼" Stepped Hose Barb	√	√	√	√
½" Single Step Hose Barb	X	√	√	√
½" Hose Barb	√	√	√	√
1½" Sanitary Flange	√	√	√	√
¾" Sanitary Flange	√	√	√	√
Quick Connector	√	√	√	√
Female Luer Lock	√	√	√	√
Male Luer Slip	Outlet Only	X	X	X
⅜" Hose Barb	√	√	Outlet Only	X
⅝" Hose Barb	X	√	√	√

Bell at Outlet Available with (Size/Outlet)
1" / ¼" SHB
1", 2", 5", 8" / ½" HB

Dimensions (in mm)	Small Capsule Filters			
End Connections	1"	2"	5"	8"
¼" SHB I/O	94	122	172	223
¾" Sanitary Flange Inlet I/O	85	104	155	206
Quick Connector	100	113	164	218
1½" Sanitary Flange I/O	92	112	164	216
½" Hose Barb I/O	90	112	162	214
½" Single Step Hose Barb I/O	-	115	165	218
1½" Sanitary Flange Inlet ½" Single Step Hose Barb Outlet	-	112	165	216
⅜" Hose Barb I/O	-	115	167	217
Operational Radius	40	65	65	65

0.1 µm AseptiCap KS PES Membrane Large Capsule filter

Type		Size		Pore Size		Inlet/Outlet		X	Inline/ T-Line		Sterility		Pack Size			
	Code		Code		Code		Code			Code		Code		Code		
AseptiCap KS (0.45 µm Upstream)	LKSX	5"	53	0.1µm	36	½" Single Step Hose Barb	Q		Inline	X	Non Sterile	1	1	01		
		10"	54						T-Line	T					EO Sterile	2
		20"	55													
AseptiCap KS (0.2 µm Upstream)	LKS1	30"	56			1½" Sanitary Flange	E									
						¾" Sanitary Flange	S									
						⅜" Hose Barb	I									
						1" Hose Barb	Z									

Example:

LKSX	54	36	EE	X	T	1	01
------	----	----	----	---	---	---	----

Inlet/Outlet Connections available with different Sizes/Length as follows:

Inlet/Outlet	Inline				T-Line		
	5"	10"	20"	30"	10"	20"	30"
½" Single Step Hose Barb	√	√	√	√	X	X	X
1½" Sanitary Flange	√	√	√	√	√	√	√
¾" Sanitary Flange	√	√	X	X	X	X	X
⅜" Hose Barb	√	√	√	√	X	X	X
1" Hose Barb	X	√	√	√	X	X	X

Dimensions (in mm)	Inline Capsule Filters				T-line Capsule Filters		
End Connections	5"	10"	20"	30"	10"	20"	30"
1½" Sanitary Flange I/O	205	330	600	855	340	580	840
¾" Sanitary Flange I/O	214	335	x	x	x	x	x
½" Single Step Hose Barb I/O	218	336	630	890	x	x	x
1½" Sanitary Flange Inlet ½" Hose Barb Outlet	212	334	620	870	x	x	x
⅜" Hose Barb I/O	211	332	634	878	x	x	x
1" Hose Barb I/O	x	405	635	895	x	x	x
Operational Radius	80	80	80	80	80	80	80

Advanced Microdevices Pvt. Ltd.

20-21, Industrial Area, Ambala Cantt-133 006, INDIA

Tel : +91-171-2699290, 2699471

E-mail : info@mdimembrane.com

Website : www.mdimembrane.com