

Data Sheet

AseptiVent® TF

Hydrophobic PTFE Membrane Devices for Sterile Filtration of Air/Gases

Pharmaceutical and Biopharmaceutical manufacturing involves sterile filtration of air/gases for a multitude of critical processes such as air sparging, bioreactor venting, fermentor exhaust, dry powder filling, WFI tank venting etc. The critical nature of these processes and associated high costs require the highest degree of reliability for the filter device with regard to its retention efficiency, flow rates, service life and mechanical and thermal stability.

mdi

mdi produces a wide range of PTFE membrane capsule filters to meet filtration requirements of biopharmaceutical and pharmaceutical processing.

These filters are validated for microbial retention with liquid bacterial challenge test as per ASTM F838-05 to provide a high degree of sterility assurance for critical applications involving sterilization of air/gases.

Applications

- Fermentor exhaust
- > Sterile air sparging in fermentors and bioreactors
- Sterile venting of cell factories, bioreactors and fermentors
- > Sterilization of environmental air in isolators

- > Venting of sterile collection vessels
- > Cleaning sterile surfaces
- WFI tank venting
- > Nitrogen blanketing
- Dry powder injectable filling
- > Sterile air for dryers and micronizers

Key Features

- Absolute Retention
- > Hydrophobic
- High heat stability
- Wide chemical compatibility
- Heat sealed to ensure 'no leaching'
- > 100% Integrity tested
- Bioburden maintained below 1000 cfu/device
- Endotoxin level certified to be <0.25 EU/ml</p>
- Widest range of end connections
- Total traceability through unique serial number for each filter
- > Individual certificate of quality for each device
- Sterilizable by EO gas or autoclaving

Quality Assurance

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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 *B. diminuta* ATCC 19146 as per ASTM F838 to establish acceptable integrity test values.

Samples from each lot are subjected to microbial challenge test before final lot release.

100% Integrity Tested

Each *AseptiVent*[®] *TF* is tested for integrity to comply with validated Acceptable Integrity Test Specifications.

Flow Rate

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

Pressure, Temperature Endurance

AseptiVent[®] *TF* 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.

AseptiVent[®]*TF* filters are validated to exhibit low extractables under harsh extraction conditions.

Bioburden Testing

Device bioburden is tested as per ISO 117 37-1 and assured to be <1000 cfu/device.

Endotoxin Testing

Aqeous extracts exhibit <0.25 EU/ml as established by Lumulus Amebocyte Lysate (LAL) test as per USP <85>.

Total Traceability

AseptiVent[®] TF filters come with completely traceable lot numbers and unique identification number to facilitate easy and fast retrieval of manufacturing and quality control data associated with each filter.

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

AseptiVent[®] *TF* 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 indirect food additives
- Materials of construction tested for toxicity as per Biological Reactivity Tests, In vitro, USP <87> for cytotoxicity and In vivo, USP <88> for class VI Plastics

Easy Connect

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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 AseptiVent[®] TF 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.

Customized Connectivity

mdi AseptiVent[®] TF filters are available in a wide range of end connections and are also customized to offer different inletoutlet 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.





³⁄₄″ Sanitary Flange



½″ HB



1⁄4″ SHB

1¹/₂" Sanitary Flange



Single Stepped Hose Barb



Quick Connector

Some end connections available with AseptiVent® TF Capsule Filters



AseptiVent® TF with HighSecurity 1/2" hose barb connection

Linear Upscaling from R&D to Production Process

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Scientists in process development labs working with cell factories or small bioreactors require small area hydrophobic filters for air/gas filtration or sterile venting.

A scale up of these processes for larger productions requires larger area devices.

mdi offers a wide range of *AseptiVent*[®] *TF* Hydrophobic PTFE capsule filters to provide linear scale up from lab scale to pilot scale to full scale biopharmaceutical manufacturing processes. The appropriate size filter can be selected on the basis of the bioreactor size and required flow rates.



AseptiVent® TF 25mm 5 cm²



AseptiVent® TF 37mm 10 cm²



AseptiVent® TF 50mm 20 cm²



AseptiVent[®] TF 1" 250cm²



AseptiVent® TF 2" 500cm²



AseptiVent® TF 5" 1000cm²

Bioreactor Size	Filter Devices	EFA* (Nominal)
100 ml Shake Flasks	AseptiVent® TF 25mm	5cm²
Up to 1 liter Shake Flasks	AseptiVent® TF 37mm	10cm ²
Up to 50 liter	AseptiVent® TF 50mm	20cm ²
Up to 100 liter	AseptiVent® TF 1"	250cm ²
Upto 300 liter	AseptiVent® TF 2"	500cm ²
Upto 1000 liter	AseptiVent® TF 5"	1000cm ²
upto 5000 liter	AseptiVent® TF 8"	2000cm ²
more than 5000 liter	AseptiVent® TF 10"	6000cm ²

*Effective Filtration Area



AseptiVent® TF 8" 2000cm²



AseptiVent® TF 10" 6000cm²

Specifications

AseptiVent[®] TF- 25mm, 37mm, 50mm

		Constru	uction						
Pore Size		0.2 μm			0.45 μm				
Membrane		Hydrophobic PTFE							
Support Layers		Polypropylene							
Body and Core			Polypropy	lene					
		Integrity Test	ing/Retention						
Bubble Point		\geq 22 psi (1.54 Kg/cm ²) with 70%	IPA/Water Solution	<u>></u> 10 psi (0.7 Kg/	cm ²) with 70% IPA/Water Solution				
Microbial Retention		LRV >7 for Brevundimon (ATCC 19146) per			7 for Serratia marcescens TCC 14756) per cm²				
l			ize	-					
Size		25 mm	37 mr	n	50 mm				
Effective Filtration Are	a (Nominal)	5 cm ²	10 cm	2	20 cm ²				
Operational Radius (w	ith Vent/ Drain)	15 mm	23 mr	n	28 mm				
		Operat	tional						
Max. Operating Tempe	erature		60° C						
Max. Differential Press	ure	42 psi (3 Kg/cm²) @ 30 °C							
Burst Pressure		> 14 Kg/cm ²	> 8 Kg/	cm²	> 8 Kg/cm ²				
	By Gas	Sterilizable by Ethylene Oxide							
Sterilization	By Autoclave	Autoclavable at 125°C for 30 minutes, 30 cycles. Can not be in-line steam sterilized							
Shelf Life		3	years after Ethylene	Oxide sterilizati	on				
		Assur	ance						
Microbial Bacterial R	etention	Validated as per ASTM F 838							
Cytotoxicity		Passes Biological reactivity test, In Vitro, as per USP <87> for cytotoxicity							
Toxicity		Passes Biological reactivity test, I	n Vivo, as per USP <8	8> for Class VI plas	tics				
Bioburden		Bioburden level is < 1000 cfu/filter device as per ISO 11737-1							
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 comp	ly with USFDA 21 CFF	Part 211.72 and 2	10.3 (b)(6) for fiber release				
Oxidizable Substance	25	Passes test as per USP <1231>							
Particle Shedding		The filtrate complies with USP <7	788> test for particula	ite matter in inject	ions				
Indirect Food Additiv	e	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							

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Specifications AseptiVent®TF-1", 2", 5", 8"

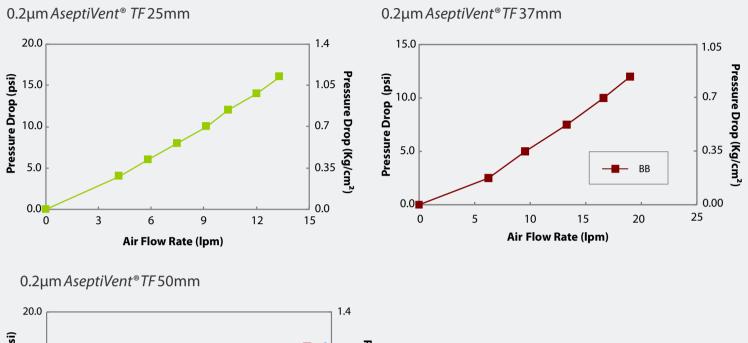
		Cons	struction						
Pore Size		0.2 μn	n	0.45	um				
Membrane		Hydrophobic PTFE							
Support Layers		Polypropylene							
Body and Core			Polypropyle	ene					
		Integrity T	esting/Retention						
Bubble Point		\geq 22 psi (1.55 Kg/cm ²) with 2	70% IPA/Water Solution	\geq 10 psi (0.7 Kg/cm ²) with	70% IPA/Water Solution				
Microbial Retention		LRV >7 for Brevundir (ATCC 19146)		LRV >7 for Serrat ATCC 14756					
			Size						
Size		1″	2″	5″	8″				
Effective Filtration Area	a (Nominal)	250 cm ²	500 cm ²	1000 cm ²	2000 cm ²				
Operational Radius (wi	ith Vent/ Drain)	30 mm	65 mm	65 mm	65 mm				
		Оре	erational						
Max. Operating Tempe	erature	80 °C @ < 30 psi (2 Kg/cm²)							
Max. Differential Press	ure	< 60 psi (4 Kg/cm²) @ 30 °C							
Chaviliantian	By Gas								
Sterilization	By Autoclave	Autoclavable at 125°C for 30 minutes, 50 cycles. Can not be in-line steam sterilized							
Shelf Life		3 years after Ethylene Oxide sterilization							
		Ass	surance						
Microbial Bacterial R	etention	Validated as per ASTM F 838							
Cytotoxicity		Passes Biological reactivity test, In Vitro, as per USP <87> for cytotoxicity							
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 ISO 11737-1							
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 co	omply with USFDA 21 CFR	Part 211.72 and 210.3 (b)(6)) for fiber release				
Oxidizable Substance	25	Passes test as per USP <1231	>						
Particle Shedding		The filtrate complies with US	SP <788> test for particulat	e matter in injections					
Indirect Food Additiv	re	All Polypropylene componer	nts meet the FDA Indirect F	ood 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							
		DMF No. 015554							

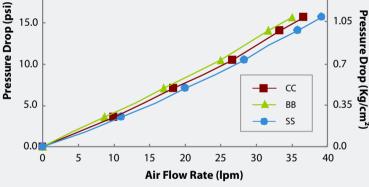
Specifications AseptiVent® TF- 5", 10", 20", 30"

		Cons	struction						
Pore Size		0.2 μn	n	0.45	μm				
Membrane		Hydrophobic PTFE							
Support Layers		Polypropylene							
Body and Core			Polypropy	ene					
		Integrity T	esting/Retention						
Air Diffusion Flow (709) (10" Capsule Filter)	6 IPA Wetted)	<u><</u> 45 ml/min @ 16 ps	ii (1.12 Kg/cm²)	<u><</u> 45 ml/min @ 8	psi (0.56 Kg/cm²)				
Microbial Retention		LRV >7 for Brevundir (ATCC 19146)			atia marcescens 56) per cm ²				
			Size						
Size		5″	10″	20″	30″				
Effective Filtration Area	a (Nominal)	3000 cm ²	6000 cm ²	12000 cm ²	18000 cm ²				
Operational Radius (w	ith Vent/ Drain)	78 mm	78 mm	78 mm	78 mm				
		Оре	erational						
Max. Operating Tempe	erature	80 °C @ < 30 psi (2 Kg/cm²)							
Max. Differential Press	ure	60 psi (4 Kg/cm²) @ 30 ℃							
	By Gas	Sterilizable by Ethylene Oxide							
Sterilization	By Autoclave	Autoclavable at 125°C for 30 minutes, 30 cycles. Can not be in-line steam sterilized							
Shelf Life			3 years after Ethylene C	xide sterilization					
		Ass	urance						
Microbial Bacterial R	etention	Validated as per ASTM F 838							
Cytotoxicity		Passes Biological reactivity test, In Vitro, as per USP <87> for cytotoxicity							
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 ISO 11737-1							
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 co	omply with USFDA 21 CFR	Part 211.72 and 210.3 (b)(6	5) for fiber release				
Oxidizable Substance	es	Passes test as per USP <1231	>						
Particle Shedding		The filtrate complies with US	P <788> test for particula	te matter in injections					
Indirect Food Additiv	re	All Polypropylene componer	nts meet the FDA Indirect	Food Additive requirement	s cited in 21 CFR 177.1520				
Good Manufacturing	Practice	These products are manufactured in a facility which adheres to Good Manufacturing Practices							
Quality Management	t System	ISO-9001 Certified							
USFDA		DMF No. 015554							

Air Flow Rates

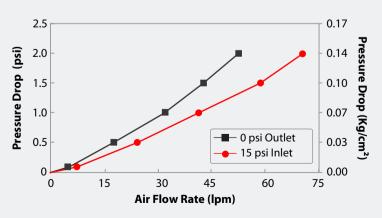
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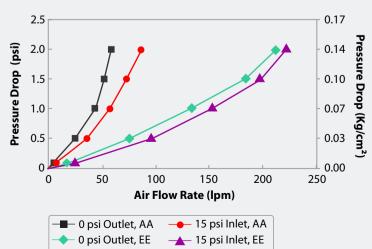


End Connection Type B: ¼" Stepped Hose Barb C: 1/8" MNPT S: ¾" Sanitary Flange

0.2µm AseptiVent® TF, 1" Capsule Filter

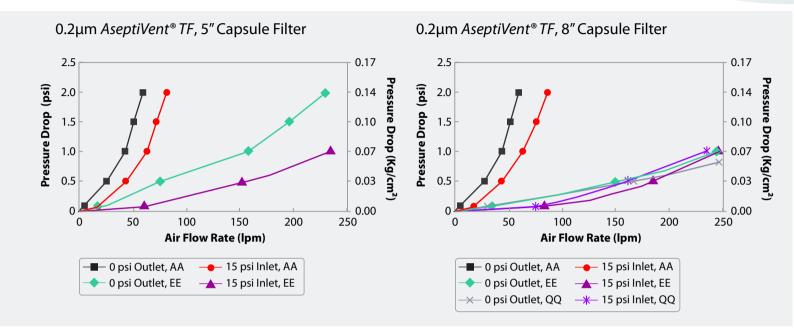


0.2µm AseptiVent® TF, 2" Capsule Filter

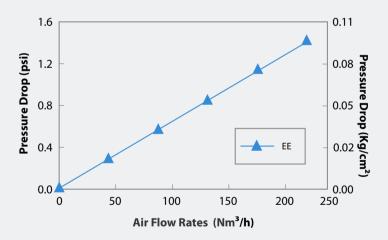


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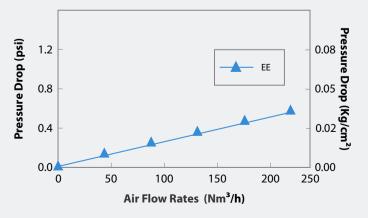
Air Flow Rates AseptiVent® TF



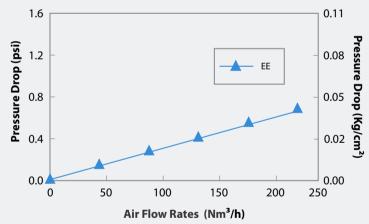
0.2µm AseptiVent® TF, 10" Capsule Filter



0.2µm *AseptiVent*[®] *TF*, 30" Capsule Filter



0.2µm AseptiVent®TF, 20" Capsule Filter



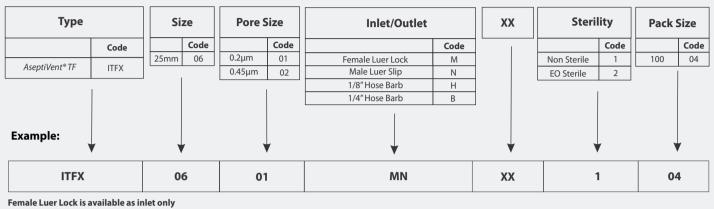
End Connection Type

A: 1/4" Stepped Hose Barb E: 11/2" Sanitary Flange Q: Single step 1/2" Hose Barb

Ordering Information

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AseptiVent® TF-25mm



Male Luer Slip is available as outlet only

AseptiVent®TF-37mm/50mm

Туре		pe Size P		Pore S	Pore Size Inlet/Outlet		X	х	Sterility		Pack Size		
	Code		Code		Code		Code		L		Code		Code
AseptiVent® TF	ITFX	37mm	08	0.2µm	01	" MNPT	С			Non Sterile	1	20 (37 mm)	09
	I	50mm	10	0.45µm	02	1⁄4″ SHB	В			EO Sterile	2	10 (50 mm)	02
						¾″ Sanitary Flange	S						
Example:			1	,	7	\downarrow		V	↓ ▼		1	•	
ITFX		1	0	01		SS		x	X	1		02	

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

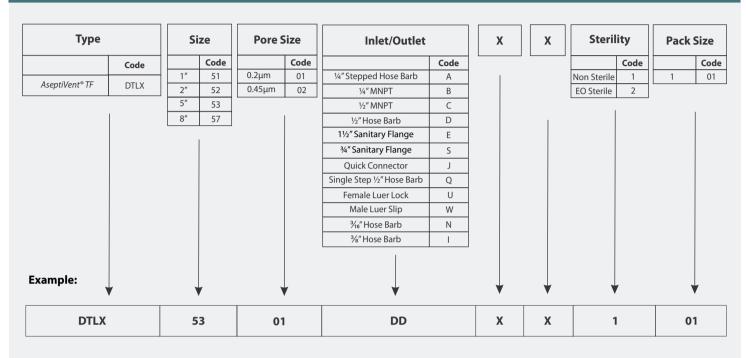
Connections Available											
Inlet/Outlet	25mm	37mm	50mm								
1/4" - 3/4" Stepped Hose Barb	х	\checkmark	\checkmark								
¾" Sanitary Flange	х	х	\checkmark								
Female Luer Lock	Inlet Only	х	х								
Male Luer Slip	Outlet Only	х	х								
1⁄8" Hose Barb	\checkmark	х	х								
Male Luer Lock	Outlet Only	х	х								
¹ ⁄4" Hose Barb		Х	х								

	-		
Dimension (in mm)	Inlin	e Capsule Fi	ilters
Inlet/ Outlet	25mm	37mm	50mm
1/4" - 3/8" Stepped Hose Barb I/O	-	64	79
¼" Single Step Hose Barb I/O	38	-	-
¾" Sanitary Flange I/O	-	-	51
Female Luer Lock Inlet/ Male Luer Slip Outlet	23	-	-
¹ ∕ ₈ " Hose Barb I/O	36	-	-
Operational Radius	15	23	28

Ordering Information

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AseptiVent® TF Small Capsule Filters



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

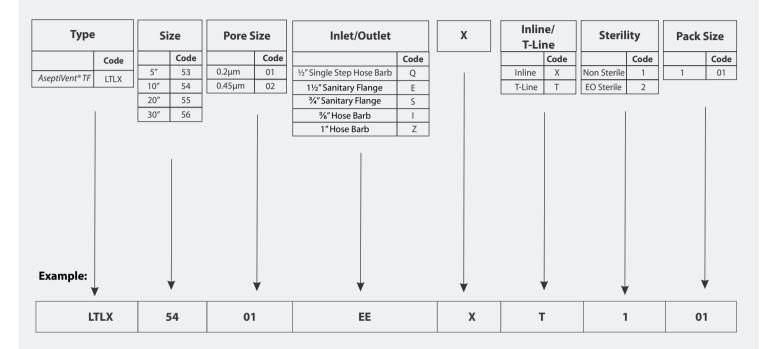
inlet/Outlet		Size/Length								
	1″	2″	5″	8″						
1/4" Stepped Hose Barb	\checkmark	\checkmark	\checkmark	\checkmark						
1⁄2"Hose Barb		\checkmark	\checkmark	\checkmark						
1½ ″ Sanitary Flange	\checkmark		\checkmark	\checkmark						
¾" Sanitary Flange	x	\checkmark	V	\checkmark						
Quick Connector				\checkmark						
½" Single Step Hose Barb	х		V	\checkmark						
1/4" MNPT	\checkmark			\checkmark						
1/2" MNPT	х		\checkmark	\checkmark						
Female Luer Lock	\checkmark		V	\checkmark						
Male Luer Slip	Outlet Only	х	х	х						
¾6″ Hose Barb			Outlet Only	х						
¾″ Hose Barb	х	\checkmark	\checkmark	\checkmark						

Dimensions (in mm)	Small Capsule Filters						
End Connections	1″	2″	5″	8″			
¼″ SHB I/O	94	122	172	223			
³ ⁄4" Sanitary Flange Inlet I/O	85	104	155	206			
Quick Connector	100	113	164	218			
1½" Sanitary Flange I/O	92	112	164	216			
1/2" Hose Barb I/O	90	112	162	214			
1/2" Single Step Hose Barb I/O	-	115	165	218			
1½" Sanitary Flange Inlet ½" Single Step Hose Barb Outlet	-	112	165	216			
3/8" Hose Barb I/O	-	115	167	217			
Operational Radius	40	65	65	65			

Ordering Information

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AseptiVent® TF Large Capsule Filters



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

	Inline				T-Line			Dimensions (in mm)	Inline Capsule Filters			ers	T-line Capsule Filters		
Inlet/Outlet	5″	10″	20″	30″	10″	20″	30″	End Connections	5″	10″	20″	30″	10″	20″	30″
1/2" Single Step Hose Barb	1	1	1		v	V	v	1½" Sanitary Flange I/O	205	330	600	855	340	580	840
⁷² Single Step Hose Barb	V	V	N	N	Х	^	X X	³ ⁄ ₄ " Sanitary Flange I/O	214	335	х	х	x	х	x
1½" Sanitary Flange	\checkmark	1/2" Single Step Hose Barb I/O	218	336	630	890	x	х	x						
¾" Sanitary Flange	\checkmark	\checkmark	х	х	х	x	х	1½" Sanitary Flange Inlet ½" Hose Barb Outlet	212	334	620	870	x	x	x
³∕₀″ Hose Barb		V	1		х	x	х	¾" Hose Barb I/O	211	332	634	878	x	х	х
							1″ Hose Barb I/O	х	405	635	895	х	х	х	
1" Hose Barb	Х				Х	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

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