



## AseptiCap NL/NS Nylon-66 Membrane Capsule Filters

### Data Sheet

**mdi** Nylon membrane capsule filters are ready to use, disposable, highly retentive filtration devices specially designed for sterilization of aqueous as well as organic solutions. Nylon-66 membrane, and polypropylene body used in these filters provide wide chemical compatibility. These capsule filters are heat resistant, biologically inert, autoclavable, and suitable for filtration and sterilization applications.

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 NL/NS filters are an ideal solution for pharmaceutical process filtration.

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, extremely low extractables, high throughputs, wide chemical compatibility and other important characteristics.

#### **Types Available**

- AseptiCap NS:
  - AseptiCap NL: Single Layer (without Prefilter)

Double Layer (with Prefilter)

### Applications

- Sterilizing filtration of stability batches in formulation development labs
- > Sterilization of compatible solvents and chemicals

#### **Key Features**

- Absolute retention
- > 100% integrity tested
- > 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

## Quality Assurance

### Datasheet

Asepticap NL/NS capsule filters use **mdi** Nylon 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 pharmaceutical applications. *Asepticap NL/NS* are manufactured in class 10,000 clean rooms and ISO 9001:2015 certified facilities.

**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 Brevundimonas diminuta ATCC 19146 as per ASTM F838-05 to establish acceptable integrity test values.

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

### 100% Integrity Tested

Each *AseptiCap NL/NS* 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.

#### Pressure, Temperature Endurance

AseptiCap NL/NS 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 may impact the impurity profile of the desired product.

*AseptiCap NL/NS* 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**

AseptiCap NL/NS 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**

AseptiCap NL/NS 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 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, invivo, USP <88> for class VI Plastics

### **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.

## **Easy Connect**

### Datasheet

1½" Sanitary Flange to ½"Barb Hose

#### Widest Range of End Connections

**mdi** AseptiCap NL/NS 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**

1<sup>1</sup>/<sub>2</sub>" Sanitary Flange to <sup>3</sup>/<sub>4</sub>" Sanitary Flange

**mdi** AseptiCap NL/NS 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 pharmaceutical 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.



Some end connections available with AseptiCap.

AseptiCap NL/NS with HighSecurity ½" 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 NL/NS* 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 is identical for all filter devices starting from 5 cm<sup>2</sup> to 18000cm<sup>2</sup> hence process scaling can be facilitated without triggering additional validation studies for given process conditions. **mdi** provides complete documentation for each of the *AseptiCap NL/NS* filters there by reducing the additional validation cost and time.



AseptiCap NL/NS 25mm, 5cm<sup>2</sup>



AseptiCap NL/NS 50mm, 20cm<sup>2</sup>



AseptiCap NL/NS 1", 250cm<sup>2</sup>/200cm<sup>2</sup>



AseptiCap NL/NS 2", 900cm<sup>2</sup>/700cm<sup>2</sup>



*AseptiCap NL/NS* 5", 1800cm<sup>2</sup>/1400cm<sup>2</sup>



AseptiCap NL/NS 8", 2700cm<sup>2</sup>/2100cm<sup>2</sup>

Filter Devices	Hold up Volume
AseptiCap NL/NS 25 mm	< 50µl
AseptiCap NL/NS 50 mm	< 300µl
AseptiCap NL/NS 1"	< 5ml
AseptiCap NL/NS 2"	< 25ml
AseptiCap NL/NS 5"	< 45ml
AseptiCap NL/NS 8"	< 60ml
AseptiCap NS 5"	< 80ml
AseptiCap NS 10"	< 150ml
AseptiCap NS 20"	< 250ml
AseptiCap NS 30"	< 350ml



AseptiCap NS 10", 6000cm<sup>2</sup>

# Specifications AseptiCap NL/NS

## Datasheet

		Construction							
Final Filter Pore	e Size	0.2 μm	0.45 μm						
Pre-filter Meml (in case of Asep		0.8 μm, 0.45μm	0.8 µm						
Membrane		Nylon- 66							
Plastic Parts		Polypropylene							
		Integrity Testing / Retention							
Bubble Point (with 50% IPA	Wetted)	> 17psi (1.19Kg/cm²)	> 11psi (0.77Kg/cm²)						
Microbial Rete Retention (LR\	ntioMicrobial Bacterial / >7 for)	Brevundimonas diminuta (ATCC 19146) per cm <sup>2</sup>	Serratia marcescens (ATCC 14756) per cm <sup>2</sup>						
		Size							
Size		25 mm	50 mm						
EFA (Effective	Filtration Area)	5cm <sup>2</sup>	20cm <sup>2</sup>						
Operational Ra	adius (with Vent/ Drain)	15 mm	28 mm						
		Operational							
Max. Operatin	g Temperature	55 ℃	60 °C						
Max. Different	ial Pressure	5Kg/cm <sup>2</sup> (75 Psi) @ 25° C	3Kg/cm <sup>2</sup> (42 Psi) @ 30° C						
Hold-up Volun	ne(with air purge)	<50µL	<300µL						
Burst Pressure		> 14 Kg/cm <sup>2</sup> > 8 Kg/cm <sup>2</sup>							
Sterilization By Gas By Autoclave		Sterilizable by Ethylene Oxide							
		Autoclavable at 125°C for 30 minutes. Can not be in-line steam sterilized							
Shelf Life		3 years after EO sterilization							
		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 ISO 11737-1: 2018							
Bacterial Endo	toxin	Aqueous extracts exhibit < 0.25 EU/ml as established by Limulus Amebocyte Lysate (LAL) Test as per USP <85>							
Non Fiber Rele	asing	Passes test as per USP and comply with USFDA 21 CFR Part 210.3(b)(6) for fiber release							
Extractables w	ith WFI	Passes NVR test as per USP <661>							
Particle Sheddi	ing	The filtrate complies with USP <788> test for partic	ulate matter in injections						
TOC/Conductiv	vity 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 WFI flush							
Indirect Food A	Additive	All Polypropylene components meet the FDA Indire 21 CFR 177.1520	ect Food Additive requirements cited in						
Good Manufac	turing Practice	These products are manufactured in a facility which adheres to Good Manufacturing Practices							
Oxidizable Sub	ostances	Passes test as per USP <1231>							
Quality Manag	ement System	ISO-9001 Certified							
USFDA		DMF No. 015554							

# Specifications AseptiCap NL/NS

### Datasheet

		Con	struction							
Final Filter Por	e Size	0.2 µ	ım	0.45 µm						
Pre-filter Mem (in case of Ase		0.8 μm, 0	).45µm	0.8 μm						
Membrane		Nylon- 66								
Support Layer			Polye	ester						
Body and Cor	e		Polypro	pylene						
		Integrity Te	sting / Retentior	า						
Bubble Point (with 50% IPA	Wetted)	> 17psi (1.19	9Kg/cm²)	> 11psi (0.77	/Kg/cm²)					
Microbial Rete Retention (LR)	entioMicrobial Bacterial V >7 for)	Brevundimor (ATCC 19146		Serratia ma (ATCC 14756						
			Size							
Size		1″	2″	5″	8″					
Effective Filtra	tion AseptiCap NL	250cm <sup>2</sup>	900cm <sup>2</sup>	1800cm <sup>2</sup>	2700cm <sup>2</sup>					
Area (Nomina	I) AseptiCap NS	200cm <sup>2</sup>	700cm <sup>2</sup>	1400cm <sup>2</sup>	2100cm <sup>2</sup>					
Operational R	adius (with Vent/ Drain)	200000								
•			30 mm 65 mm 65 mm 65 mm							
Vent and Drai	n	1/4" Hose Barb with Silicone "O" rings								
		Оро	erational							
Max. Operatin	ig Temperature	80 °C @ < 30 psi (2 Kg/cm²)								
Max. Different	ial Pressure	< 60 psi (4 Kg/cm²) @ 30 °C								
Sterilization	By Gas	Sterilizable by Ethylene Oxide								
Stermzation	By Autoclave	Autoclavable at 125°C for 30 minutes. Can not be in-line steam sterilized								
Shelf Life		3 years after EO sterilization								
		As	surance							
Toxicity		Passes Biological reactivity te	est, In Vivo, as per USP <	<88> for Class VI plastics						
Bioburden		Bioburden level is < 1000 cfu/filter device as per ISO 11737-1: 2018								
Bacterial Endo	otoxin	Aqueous extracts exhibit < 0.25 EU/ml as established by Limulus Amebocyte Lysate (LAL) Test as per USP <85>								
Non Fiber Rele	easing	Passes test as per USP and comply with USFDA 21 CFR Part 210.3(b)(6) for fiber release								
Extractables v	vith WFI	Passes NVR test as per USP <661>								
Particle Shedo	ling	The filtrate complies with USP <788> test for particulate matter in injections								
TOC/Conduct	ivity 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 WFI flush								
Indirect Food	Additive	All Polypropylene componer	nts meet the FDA Indire	ect Food Additive requirement	s cited in 21 CFR 177.1520					
Good Manufa	cturing Practice	These products are manufac	These products are manufactured in a facility which adheres to Good Manufacturing Practices.							
Oxidizable Su	bstances	Passes test as per USP <1231	>							
Quality Manag	gement System	ISO-9001 Certified								
USFDA		DMF No. 015554								

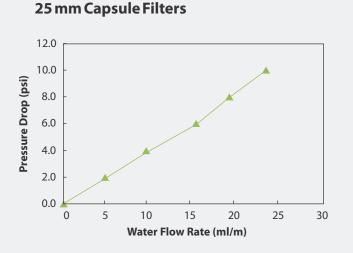
# Specifications AseptiCap NL/NS

### Datasheet

		Const	truction							
Final Filter Por	re Size	0.2 μm	1	0.45 μm	1					
Pre-filter Mem (in case of Ase		0.8 μm, 0.4	0.8 μm							
Membrane		Nylon- 66								
Support Layer			Polyester							
Body and Core	e		Polypropyle	ne						
		Integrity Tes	ting / Retention							
Air Diffusion Fl (water wetted)	low per 10″ Capsule Filter )	< 30ml/min @ 37 psi (2.	60 Kg/cm²)	<30ml/min @ 22 psi	(1.54 Kg/cm²)					
Microbial Bactor Retention (LRV		Brevundimona (ATCC 19146) ا		Serratia marc (ATCC 14756)						
		2	Size							
Size		5″	10″	20″	30″					
Effective Filtra	ition Area (Nominal)	3000 cm <sup>2</sup>	6000 cm <sup>2</sup>	12000 cm <sup>2</sup>	18000 cm <sup>2</sup>					
Operational R	adius (with Vent/Drain)	78 mm	78 mm	78 mm	78 mm					
Vent and Drain	n	<sup>1</sup> ⁄4" Hose Barb with Silicon	e "O" rings							
		Оре	ational							
Max. Operatin	ig Temperature	80 °C @ < 2 Kg/cm² (30 psi)								
Max. Differential Pressure		< 4 Kg/cm² (60 psi ) @ 30 °C								
	By Gas	Sterilizable by Ethylene Oxide								
Sterilization	By Autoclave	Autoclavable at 125 °C for 30 minutes. Can not be in-line steam sterilized								
Shelf Life		3 years after EO sterilization								
		Ass	urance							
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 11737-1:2018								
Bacterial Endo	otoxin	Aqueous extracts exhibit < 0.25 EU/ml as established by Limulus Amebocyte Lysate (LAL) Test as per USP <85>								
Non Fiber Rele	easing	Passes test as per USP and comply with USFDA 21 CFR Part 210.3(b)(6) for fiber release								
Extractables w	vith WFI	Passes NVR test as per USP	<661>							
Particle Shedd	ling	The filtrate complies with L	ISP <788> test for partice	ulate matter in injections						
TOC/Conduct	ivity at 25 ℃	Meets the WFI requirements of USP <643> for Total Organic Carbon and USP <645> for Water Conductivity after a specified volume of WFI flush								
Indirect Food	Additive	All Polypropylene components meet the FDA Indirect Food Additive requirements cited in 21 CFR 177.1520								
Good Manufa	cturing Practice	These products are manufactured in a facility which adheres to Good Manufacturing Practices.								
Oxidizable Sul	bstances	Passes test as per USP <123	1>							
Quality Manag	gement System	ISO-9001 Certified								
USFDA		DMF No. 015554								

## **Typical Water Flow Rates**

### Datasheet



### 0.2µm AseptiCap NS, 1" Capsule Filters

12

10

8

6

4

2

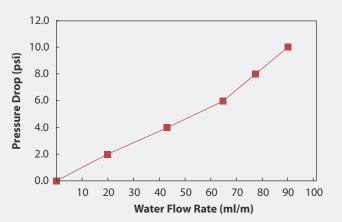
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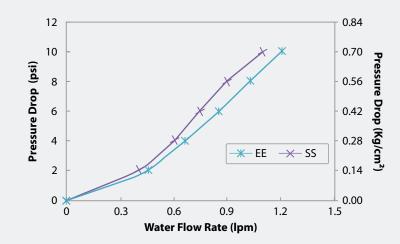
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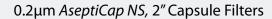
0.3

Pressure Drop (psi)

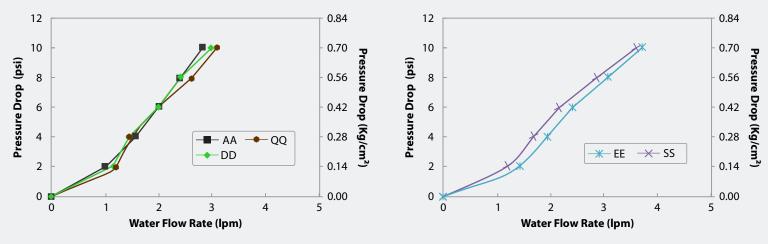
50 mm Capsule Filters







0.6



0.84

0.70

0.56

0.42

0.28

0.14

0.00

1.5

DD

1.2

- AA

0.9

Water Flow Rate (lpm)

Pressure Drop (Kg/cm<sup>2</sup>)

### **End Connection Type:**

A: <sup>1</sup>/<sub>4</sub>" Stepped Hose Barb Q: <sup>1</sup>/<sub>2</sub>" Single Step Hose Barb

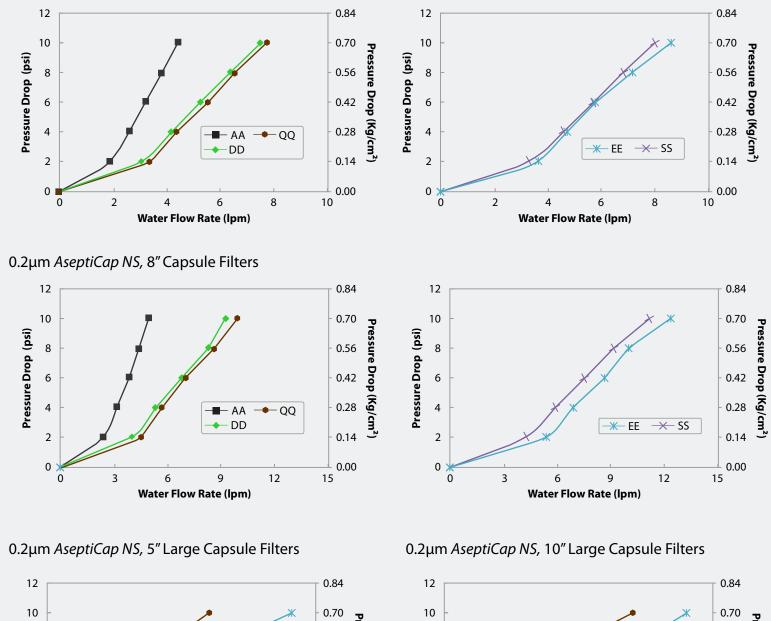
E: 1<sup>1</sup>/<sub>2</sub>" Sanitary Flange

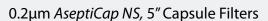
D: 1/2" Hose Barb

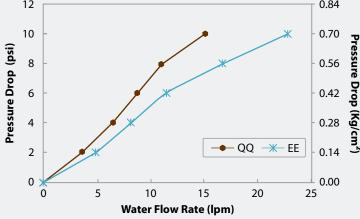
S: 34" Sanitary Flange

## **Typical Water Flow Rates**

### Datasheet







### **End Connection Type:**

A: <sup>1</sup>/<sub>4</sub>" Stepped Hose Barb Q: <sup>1</sup>/<sub>2</sub>" Single Step Hose Barb

E: 11/2" Sanitary Flange

Pressure Drop (psi)

8

6

4

2

0

0

v Flange D:

10

D: 1/2" Hose Barb

Water Flow Rate (Ipm)

20

S: 3/4" Sanitary Flange

40

QQ

30

**DST DNXLNXX1403A** 

0.56

0.42

0.28

0.14

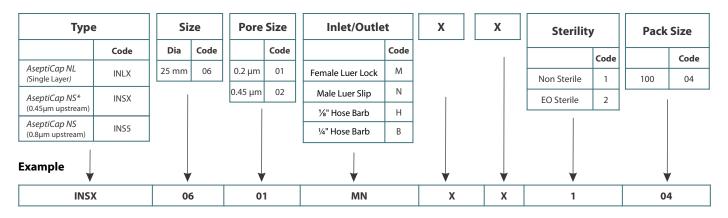
0.00

50

## **Ordering Information**

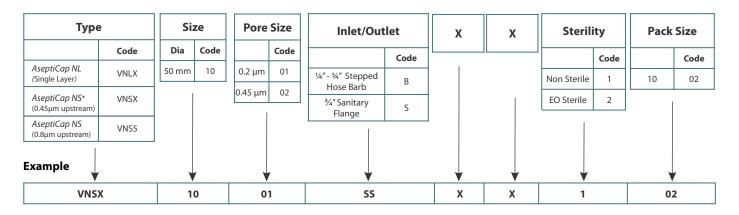
## Datasheet

### AseptiCap NL/NS 25mm



\*0.45µm Upstream is only available in 0.2µm Pore Size

### AseptiCap NL/NS 50mm



\*0.45  $\mu m$  Upstream is only available in 0.2  $\mu m$  Pore Size

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

Connections Available								
Inlet/Outlet	25mm	50mm						
¼" - ¾" Stepped Hose Barb	х	$\checkmark$						
<sup>3</sup> ⁄ <sub>4</sub> " Sanitary Flange	х	$\checkmark$						
Female Luer Lock	Inlet Only	х						
Male Luer Slip	Outlet Only	Х						
⅓″ Hose Barb	$\checkmark$	х						
Male Luer Lock	Outlet Only	х						
1/4" Hose Barb	$\checkmark$	х						

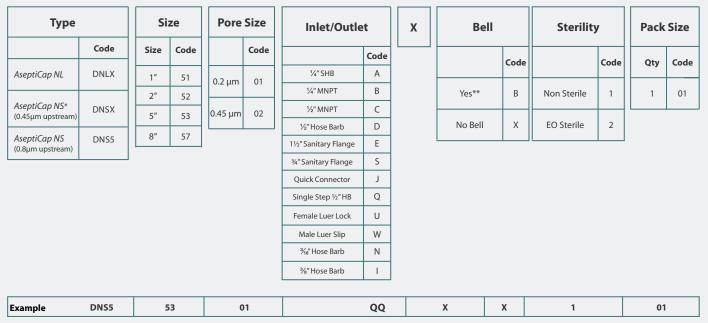
Dimension (in mm)	Inline Capsule Filters					
Inlet/ Outlet	25mm	50mm				
¼″ - ¾ <b>″</b> Stepped Hose Barb I/O	-	79				
¼" Single Step Hose Barb I/O	38	-				
¾" Sanitary Flange I/O	-	51				
Female Luer Lock Inlet/ Male Luer Slip Outlet	23	-				
1⁄8" Hose Barb I/O	36	-				
Operational Radius	15	28				

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## **Ordering Information**

## Datasheet

### AseptiCap NL/NS 1", 2", 5", 8"



\*0.45µm Upstream is only available in 0.2µm Pore Size

Inlet/Outlet	Size/Length								
	1″	2″	5″	8″					
<sup>1</sup> /4" Stepped Hose Barb	$\checkmark$	$\checkmark$	$\checkmark$	$\checkmark$					
½" Single Step Hose Barb	x	$\checkmark$	$\checkmark$	$\checkmark$					
½"Hose Barb	$\checkmark$		$\checkmark$	$\checkmark$					
1½" Sanitary Flange	$\checkmark$		$\checkmark$	$\checkmark$					
¾" Sanitary Flange	$\checkmark$		$\checkmark$	$\checkmark$					
Quick Connector	$\checkmark$								
½″ MNPT	х								
¼″ MNPT (18TPI)	$\checkmark$		$\checkmark$						
Female Luer Lock	$\checkmark$		$\checkmark$	$\checkmark$					
Male Luer Slip	Outlet Only	х	х	х					
3/16" Hose Barb	V		Outlet Only	х					
3/8" Hose Barb	х								

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

Dimensions (in mm)		Small Capsule Filters								
End Connections	1″	2″	5″	8″						
1/4" 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						
1⁄2" 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						
3/8" Hose Barb I/O	-	115	167	217						
Operational Radius	40	65	65	65						

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

## **Ordering Information**

## Datasheet

### AseptiCap NS 5", 10", 20", 30"

Туре		Si	Size Pore Size			Inlet/Outlet	x	Inline/T-line		Sterili	ty	Pack Size		
	Code	Size	Code		Code		Code			Code		Code	Qty	Code
AccentiCon NC*		5″	53	0.2 μm	01		couc							
(0.45µm upstream)	ptiCap NS* LNSX μm upstream)		54	0.2 µm		1½" Sanitary Flange	E		Inline	Х	Non Sterile	1	1	01
AseptiCap NS		20″	55	0.45 μm	02	<sup>3</sup> 4" Sanitary Flange	S		T-line	т	EO Sterile	2	·	
(0.8µm upstream)	LNS5	30" 56 Single Step ½" Hose Ba		Single Step ½" Hose Barb	Q									
						³∕₃" Hose Barb	I							
						1" Hose Barb	Z							
Example														
LNS5		5	6	C	01	EE		х	х		1		01	

\* 0.45μm Upstream is only available in 0.2μm Pore Size

#### 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		10″	20″	30″	10″	20″	30″
1/2" Single Step Hose Barb	1	1	1	1	x		x x		1½" Sanitary Flange I/O	205	330	600	855	340	580	840
<sup>72</sup> Single Step Hose Barb		V	N	N	×	^			<sup>3</sup> ⁄4″ Sanitary Flange I/O	214	335	х	х	х	х	х
1½" Sanitary Flange	$\checkmark$	Ī	1/2" Single Step Hose Barb I/O	218	336	630	890	x	x	х						
¾" Sanitary Flange		$\checkmark$	х	х	х	x	х		1½" Sanitary Flange Inlet ½" Hose Barb Outlet	212	334	620	870	x	х	x
³∕a″ Hose Barb					х	x	х		¾″ Hose Barb I/O	211	332	634	878	х	х	х
							x x		1" Hose Barb I/O	х	405	635	895	х	х	х
1" Hose Barb	Х	$\checkmark$	$\checkmark$		Х	X		Operational Radius	80	80	80	80	80	80	80	

### Advanced Microdevices Pvt. Ltd.

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