# **EverLUX®** Filters





### Table of Contents - EverLUX<sup>®</sup> Filters



EverLUX <sup>®</sup> Filters	3
Materials of Construction	3
Components	3
Configurations	3
Dimensions	4
Operating Characteristics	5
Cartridge Installation Instructions	6
Wetting Best Practices	7
Water & Product	7
Isopropyl Alcohol	7
Flushing Guidelines	8
Integrity Testing	9
Manual Bubble Point Test of a Filter Housing System	9
Manual Bubble Point of a Capsule, UltraCap <sup>®</sup> & UltraCap <sup>®</sup> H.D.	10
EverLUX <sup>®</sup> Bubble Point Specifications	10
Manual Diffusive Flow Test of a Filter Housing System	11
EverLUX <sup>®</sup> Diffusive Flow Specifications <sup>1</sup>	12
Filter Integrity Test Preparations	13
Pre-Use/Post-Use FIT Considerations	14
Autoclave Instructions	15
Inline Steam Sterilization Procedure	16
Storage & Shelf Life	17
EverLUX <sup>®</sup> Membrane Grade Descriptions	18
Ordering Matrix	19
Cartridge	19
Small Flow Elements (SFE Filters)	20
Capsule (CA/CB)	21
Capsule (CM/CK)	22
Capsule (CF)	23
Capsule (CS/CL/CJ)	24
UltraCap <sup>®</sup> (T-Style & Inline)	25
UltraCap <sup>®</sup> H.D. (T-Style & Inline)	26



### **EverLUX®** Filters

EverLUX® is an advanced PES membrane filter that provides high contaminant capacity, extended service life, and high flow rates when filtering a wide range of aqueous and biological liquids. This hydrophilic filter is ideal for sterilization and bioburden reduction in a range of low-to-high contaminant liquids, including pharmaceutical preparations, biopharmaceuticals, parenterals, vaccines, complex biologicals, serum, cell and tissue culture media, buffers, media additives, supernatants, process intermediate, ophthalmic and other dilute preservative solutions, UPW, chemicals, alcohols, and sanitizing agents.

### **Materials of Construction**

The EverLUX® filter is manufactured using high quality components made from nontoxic and biologically inert raw materials. All components of the EverLUX<sup>®</sup> filter are FDA listed for food contact use in the Code of Federal Regulations (CFR), Title 21 as below:

#### **Components**

Media:	Polyethersulfone (PES)	CFR Title 21, 177.2440
Upstream/Downstream support:	Polypropylene	CFR Title 21, 177.1520
Core/Outer guard:	Polypropylene	CFR Title 21, 177.1520
End caps/Adaptors:	Polypropylene	CFR Title 21, 177.1520
Capsule housing:	Polypropylene	CFR Title 21, 177.1520
O-rings:	Buna, EPR or Silicone	CFR Title 21, 177.2600
	Teflon <sup>®</sup> over Silicone or	CFR Title 21, 177.1550
	Teflon <sup>®</sup> over Viton <sup>®</sup>	
Sealing method:	Thermal bonding	

The EverLUX® filter complies with European Commission Regulation No. 10/2011. The filter meets requirements as specified in the current USP Class VI plastics, pyrogen and cytotoxicity tests. No binders, adhesives, or surfactants are used in the construction of EverLUX<sup>®</sup> filters. The filters are non-fiber-releasing as defined in 21 CFR 210.3(b)(6) and 211.72.

### **Configurations**

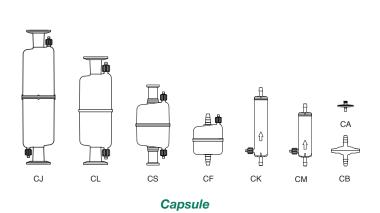
EverLUX<sup>®</sup> can be ordered in a variety of configurations from SFE filter cartridges through UltraCap<sup>®</sup> H.D. high capacity capsule filters.

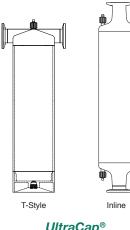


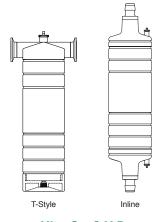


SFE Cartridge









**UltraCap®** 

UltraCap® H.D.



### **Dimensions**

Cartridge	Diameter		Length (nominal)	EFA (SM	,** IH, SLH, SPH - Grade)		FA TW, STS, STC - Grade)
	2.75" (7.0	cm)	10" (25 cm)	7.3	ft² (0.68 m²)	6.	8 ft² (0.63 m²)
	Υ.		20" (50 cm)		S ft <sup>2</sup> (1.36 m <sup>2</sup> )		3.6 ft <sup>2</sup> (1.26 m <sup>2</sup> )
			30" (75 cm)		9 ft² (2.03 m²)		).4 ft² (1.90 m²)
			40" (100 cm		2 ft² (2.71 m²)		7.2 ft² (2.53 m²)
SFE Cartridge	Diameter		Length (nominal)	EFA (SM	** IH, SLH, SPH - Grade)	El (S	FA TW, STS, STC - Grade)
	2.25" (5.7	′ cm)	2.5" (6.4 cm	n) 1.6	ft² (0.15 m²)	1.	2 ft² (0.11 m²)
			5" (12.7 cm)	) 3.3	ft² (0.31 m²)	2.	6 ft² (0.24 m²)
Capsule	Diameter		Length (nominal)	EFA (SM	". IH, SLH, SPH - Grade)		FA TW, STS, STC - Grade)
CA2	0.98" (25	mm)	1.18" (30 m	m) 0.00	04 ft² (3.6 cm²)	0.	004 ft² (3.6 cm²)
CB2	0.02 (50 m	nm	2.51 (64 mm	n) 0.02	2 ft² (19.6 cm²)	0.	02 ft² (19.6 cm²)
CK2	1.25" (3.2	cm)	6.25" (15.9 cm) 0.45 ft <sup>2</sup> (415 cm <sup>2</sup> )		0.	0.35 ft² (325 cm²)	
CM2	1.25" (3.2	cm)	5.50" (14.0	cm) 0.36	6 ft² (335 cm²)	0.	28 ft² (260 cm²)
CF-B/CF2-B	2.25" (5.7	cm)	3.3" (8.3 cm	ı) 0.70	) ft² (650 cm²)	_	
CL/CL2	2.75" (7.0	cm)	6.9" (17.5 c	m) 3.3	ft² (0.31 m²)	2.	6 ft² (0.24 m²)
CS/CS2	2.75" (7.0	cm)	4.5" (11.4 c	m) 1.6	ft² (0.15 m²)	1.	2 ft² (0.11 m²)
CJ/CJ2	2.75" (7.0	cm)	10.0" (25.4	cm) 5.2	ft² (0.48 m²)	4.	0 ft² (0.37 m²)
UltraCap®	Diameter	Length (nominal)		ule Dimension all)	EFA** (SMH, SLH, SPH - Gra	ade)	EFA (STW, STS, STC - Grade)
T-Style	3.25" (8 cm)	10" (25 cm	n) 12.3"	(31.2 cm)	7.3 ft² (0.68 m²)		6.8 ft² (0,63 m²)
		20" (50 cm	r) 22.3"	(56.6 cm)	14.6 ft <sup>2</sup> (1.36 m <sup>2</sup> )		13.6 ft² (1.26 m²)
		30" (75 cm	n) 32.3"	(82 cm)	21.9 ft² (2.03 m²)		20.4 ft² (1.90 m²)
Inline	3.25" (8 cm)	10" (25 cm	n) 14.8"	(37.6 cm)	7.3 ft <sup>2</sup> (0.68 m <sup>2</sup> )		6.8 ft <sup>2</sup> (0.63 m <sup>2</sup> )
		20" (50 cm	n) 24.9"	(63.2 cm)	14.6 ft² (1.36 m²)		13.6 ft <sup>2</sup> (1.26 m <sup>2</sup> )
		30" (75 cm	n) 34.9"	(88.6 cm)	21.9 ft² (2.03 m²)		20.4 ft² (1.90 m²)

UltraCap®H.D.	Diameter	Length (nominal)	Capsule Dimension (overall)	EFA** (SMH, SLH, SPH - Grade)	EFA (STW, STS, STC - Grade)
T-Style	3.5" (9 cm))	10" (25 cm)	11.7" (29.7 cm)	7.3 ft² (0.68 m²)	6.8 ft² (0.63 m²)
		20" (50 cm)	21.1" (53.6 cm)	14.6 ft <sup>2</sup> (1.36 m <sup>2</sup> )	13.6 ft² (1.26 m²)
		30" (75 cm)	30.6" (77.7 cm)	21.9 ft <sup>2</sup> (2.03 m <sup>2</sup> )	20.4 ft² (1.90 m²)
		40" (100 cm)	40.0" (101.6 cm)	29.2 ft² (2.71 m²)	27.2 ft² (2.53 m²)
		50" (125 cm)	49.5" (125.7 cm)	36.5 ft <sup>2</sup> (3.39 m <sup>2</sup> )	34.0 ft² (3.16 m²)
Inline	3.5" (9 cm	10" (25 cm)	17.3" (43.9 cm)	7.3 ft <sup>2</sup> (0.68 m <sup>2</sup> )	6.8 ft <sup>2</sup> (0.63 m <sup>2</sup> )
		20" (50 cm)	26.8" (68.1 cm)	14.6 ft <sup>2</sup> (1.36 m <sup>2</sup> )	13.6 ft² (1.26 m²)
		30" (75 cm)	36.2" (91.9 cm)	21.9 ft <sup>2</sup> (2.03 m <sup>2</sup> )	20.4 ft <sup>2</sup> (1.90 m <sup>2</sup> )
		40" (100 cm)	45.7" (116.1 cm)	29.2 ft² (2.71 m²)	27.2 ft <sup>2</sup> (2.53 m <sup>2</sup> )
		50" (125 cm)	55.2" (140.2 cm)	36.5 ft <sup>2</sup> (3.39 m <sup>2</sup> )	34.0 ft <sup>2</sup> (3.16 m <sup>2</sup> )

\*inlet/outlet fittings extend beyond stated diameter | \*\*EFA = Effective Filtration Area



### **Operating Characteristics**

#### Cartridges and SFE (Small Flow Elements)

Maximum Operating Temperatures and Pressures 80 psid @ 32 °F to 100 °F (Δp 5.5 bar @ 0 °C to 38 °C)

60 psid @ 150 °F (Δp 4.1 bar @ 66 °C)

30 psid @ 180 °F (Δp 2.1 bar @ 82 °C)

#### Capsules - CA/CB Models

Maximum Operation Pressure & Temperature, Liquids 75 psig @ 32 °F to 100 °F (5.2 bar @ 0 °C to 38 °C)

#### Capsules - CM/CK Models

Maximum Operation Pressure & Temperature, Liquids 100 psig @ 32 °F to 122 °F (6.9 bar @ 0 °C to 50 °C)

#### Capsules - CF/CS/CL/CJ Models

Maximum Operating Pressure, Liquids 75 psig @ 32 °F to 100 °F (5.2 bar @ 0 °C to 38 °C)

Maximum Operating Temperature Rating 160 °F @ 35 psig (71 °C @ 2.4 bar)

#### UltraCap® Model

Maximum Operation Pressure & Temperature, Liquids 75 psig @ 32 °F to 100 °F (5.2 bar @ 0 °C to 38 °C)

45 psig @ 140 °F (3.1 bar @ 60 °C)

#### UltraCap® H.D. Model

Maximum Operation Pressure & Temperature, Liquids 90 psig @ 32 °F to 100 °F (6.2 bar @ 0 °C to 38 °C)

55 psig @ 140 °F (3.8 bar @ 60 °C) Maximum Operating Pressure & Temperature, Gas 50 psig @ 32 °F to 100 °F (3.4 bar @ 0 °C to 38 °C)

Maximum Operating Pressure & Temperature, Gas 100 psig @ 32 °F to 122 °F (6.9 bar @ 0 °C to 50 °C)

Maximum Operating Pressure, Gas 50 psig @ 32 °F to 100 °F (3.4 bar @ 0 °C to 38 °C)

Maximum Operating Pressure & Temperature, Gas 50 psig @ 32 °F to 100 °F (3.4 bar @ 0 °C to 38 °C)

30 psig @ 140 °F (2.1 bar @ 60 °C)

Maximum Operating Pressure & Temperature, Gas 60 psig @ 32 °F to 100 °F (4.1 bar @ 0 °C to 38 °C)

35 psig @ 140 °F (2.4 bar @ 60 °C)



### **Cartridge Installation Instructions**

Meissner filters are available in a number of different adapter and O-ring configurations designed to fit modern filter housings. The filter should fit snugly in the housing. Improper installation can impair filtration efficiency.

- 1. Verify that the correct filter part number for the application has been chosen.
- 2. Keep the filter in its plastic bag to avoid contaminating the cartridge as long a possible. Cut open the bag at the O-ring end. While holding the bagged cartridge, lubricate the O-rings by dipping the O-rings into clean water or other suitable liquid compatible with the process fluid.
- 3. Line up the open end of the cartridge with the housing seat and install using a slight twisting motion while holding the bagged cartridge near the O-ring adapter. Verify that the O-rings are fully seated and not twisted. If the cartridge has locking tabs, rotate the tabs into place with a clockwise motion until engaged. Caution: always rotate cartridges while firmly grasping the O-ring end of the cartridge to prevent excessive torque damage to the filter.
- 4. Repeat with additional cartridges. Remove protective bags from the cartridges. If present, install cartridge retainer system (plate or spring). Reassemble housing.



### Wetting Best Practices

#### Water & Product

The following is a guide for proper wetting of the membrane.

- 1. Attach and secure filter via tubing to source of wetting fluid.
  - a. Users may use a pump to monitor flow rate or pressurized transfer to monitor pressure on the upstream.
- 2. Open any inlet vent valves to allow air to escape the housing/capsule and close the downstream valve.
- 3. Slowly fill the capsule/housing at the start to prevent turbulence (air entrapment); the capsule is oriented so the vent is on the highest position.
- 4. Close the high point vent valve when all air is released (identified by fluid exiting vent valve).
- 5. Following closure of a vent valve, open downstream to allow flow.
- 6. Re-vent the housing after about a minute to allow any accumulated air to escape.
- 7. Typical filter wetting times are listed in the table below. Longer wetting times can be used to ensure complete wetting of the filter. Recirculated systems may be used.
- 8. Application of back pressure should be utilized to help ensure a uniform flow distribution through the entire length of the filter.
  - a. Back pressure is not reverse pressure, it is a restriction placed on the outlet of the filter during forward flow by using a partially closed valve, smaller piping size, tubing clamp, or similar constriction.

#### Typical Filter Wetting Times

Configuration	Flow Rate L/min	Time
CM, CK, CF Capsules	1	1-2 minutes
CS Capsules, 2.5" SFE Cartridges	4	5-6 minutes
CL Capsules, 5" SFE Cartridges	4	5-6 minutes
CJ Capsules	8	5-10 minutes
10" UltraCap <sup>®</sup> , UltraCap <sup>®</sup> H.D., and Cartridges <sup>1</sup>	15	5-10 minutes <sup>1</sup>

<sup>1</sup> For >10" filters, the wetting time should be multiplied by the number of 10" modules. It is advised to increase the flow rate if process conditions allow.

Configuration	Wetting guidance
CA, CB Capsules	Due to the size of the CA & CB capsules, very small volumes are required to wet the filter. If no vent is present, introduce fluid into the capsule with the outlet point upwards so that air may escape. For capsules without a vent, airlock can occur after the membrane has been wetted, if more fluid is introduced and air enters the capsule. Slowly add fluid to the filter capsule until no air bubbles are visible in effluent downstream. If the filter has a vent, slowly introduce fluid into the capsule until all air has escaped through the vent on the upstream side.

#### Isopropyl Alcohol

Isopropyl Alcohol (IPA) is a low surface tension fluid. As a result, membranes are more easily wetted when IPA is used. It is common practice to simply submerge the filter in a container with the IPA wetting product and allow it to sit for 5-10 minutes with large filters leaning towards the 10-minute time. Following wetting, it is important to flush out any remaining product in the filter tubing, housing, or capsule filter prior to putting the filter into service.

Configuration	Time	
CA, CB Capsules	< 1 minute	
CM, CK, CF, CS, CL, CJ Capsules	2.5 minutes	
2.5", 5" SFE Cartridges	3-5 minutes	
10" UltraCap <sup>®</sup> , UltraCap <sup>®</sup> H.D., and Cartridges	10 minutes	



### **Flushing Guidelines**

Meissner recommends flushing the filter, prior to use or service, which can be used to wet the filter. This process is beneficial to filtrate quality and helps eliminate leachables by reducing extractables. Installation can generate contaminants in or on the filters. These contaminants are most evident during start-up but will dissipate after the initial few liters of fluid have passed through the filter. Without flushing prior to use, contaminants likely will be present downstream of the filter.

The following is a general guide for filter flushing. The flushing recommendations provided by Meissner are based on Rinse-Up TOC data using DI water passed through a filter following sterilization. When passing DI water through the filter, the effluent's TOC was analyzed at intervals until the effluent's TOC value met the United States Pharmacopeia (USP) Requirement for purified water (< 0.5 ppm). Since the following is based on DI water as a flushing fluid, suitable flushing volumes may differ from process fluid to process fluid. We offer these recommendations as a general guideline, and users commonly devise their own flushing method based on the configuration or set-up of their process. Please contact Meissner for Rinse-Up TOC data requests.

Configuration	Suggested Minimum Flow Rate L/min	Typical Total Flush Volume, Liters <sup>1</sup>
CA Capsules	See N	lote 3
CB Capsules	N/A	0.08
CM/CK Capsules	0.5-1	1-2
CF Capsules	2	2-3
CS Capsules	2-3	2-3
CL Capsules	4	3-8
CJ Capsules	8	5-15
10" UltraCap <sup>®</sup> , UltraCap <sup>®</sup> H.D. Capsules & Cartridges <sup>2</sup>	15	12-20

#### **EverLUX®** Flushing Guidance<sup>1</sup>

<sup>1</sup> Flushing volumes vary based on sterilization method.

<sup>2</sup> For filters larger than 10", multiply the value by the number of 10" units in the cartridge/capsule.

<sup>3</sup> Due to the size of these capsules, please refer to the Wetting Guidance.



### Integrity Test: Manual Bubble Point of a Filter Housing System

#### (Consult Meissner for Pressure Hold Procedure)

#### Wetting

Please refer to wetting instructions on Page 7 under Wetting Best Practices.

#### Filter and Housings - Manual Bubble Point Test

The manual bubble point test relies on a wetted, microporous membrane filter, a housing or holder to contain the wetted filter, a calibrated pressure gauge, a regulated gas pressure source (usually compressed air or nitrogen), narrow diameter downstream tubing, and a beaker containing water in which the tubing is immersed. Pressure is increased gradually until a steady stream of bubbles is observed to come from the tubing. This pressure is referred to as the bubble point. The following highlights the basic steps required to perform a bubble point test on a filter.

#### Procedure (Figure 1)

- 1. Close all valves.
- 2. Open valves  $V_1$  and  $V_3$  fill the housing with water or appropriate wetting fluid. Close  $V_3$  once fluid escapes. Open  $V_6$  to wet the cartridge.
- 3. Close  $V_1$  after the cartridge is wetted.
- 4. Open V<sub>2</sub> and apply regulated air pressure to the inlet side of the system.
- 5. If necessary, open  $V_4$  to drain the downstream volume of water.
- 6. Close V<sub>4</sub>.
- 7. Open  $V_5$  and close  $V_6$ .
- 8. While observing for continuous bubbling from the bubble point tube, slowly increase the air pressure. When a rapid, steady stream of bubbles is observed, the bubble point pressure has been reached. (Do not confuse the diffusive flow for the bulk gas flow of the bubble point.) Record the pressure where this occurs and compare it to the chart on Page 10.
- 9. A bubble point value lower than the specification is an indication of one of the following:
  - Fluid with different surface tension than the recommended test fluid
  - Integral filter but wrong pore size
  - High temperature
  - Incompletely wetted membrane
  - Non-integral membrane or seal

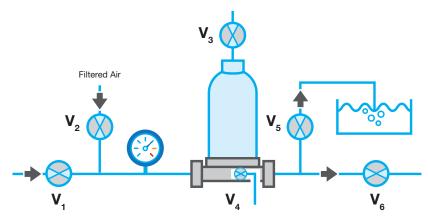


Figure 1



### Integrity Test: Manual Bubble Point of a Capsule, UltraCap<sup>®</sup> & UltraCap<sup>®</sup> H.D.

- 1. Wet the filter capsule well. Drain the capsule housing.
- Connect regulated compressed air and a calibrated gauge to the inlet and a narrow ID hose to the filter outlet and immerse the open hose end in a beaker of water. Alternatively, connect an automated integrity tester to the inlet and follow manufacturer's directions.
- 3. Apply air pressure to the filter to approximately 10 psi less than the minimum bubble point and observe the hose in the water beaker. The initial pleat pack compression will produce a large amount of bubbling that will decrease rapidly and be replaced by regularly spaced bubbles produced by air diffusing through the filter membrane.
- 4. Slowly increase the air pressure and observe the steady stream of bubbles in the water. The bubble point is the pressure at which a marked change in the rate of bubbling occurs, accompanied by an increase in sound volume. Record the pressure where this occurs and compare it to the chart below.

#### **Quality Awareness - All Integrity Test Methods**

- 1. While increasing the air pressure, do not confuse the additional bubbles produced by a rapid increase in air pressure with the bubble point. Observe the bubbles when the air pressure has stabilized.
- 2. If the integrity test fails, rewet the filter with additional water and repeat the test. If the filter has been used to filter product, it may take several minutes of high volume flush to remove the product and return to the water integrity test value. It is not appropriate to recirculate the water in this case.
- 3. To troubleshoot membrane wetting challenges, the filter may be rewet with a low surface tension fluid such as 60% IPA and integrity tested in that fluid when applicable. Please see further guidance on Page 7 under Wetting Best Practices, Isopropyl Alcohol, as well as on Pages 13 and 14 under Troubleshooting Guidance and Pre-Use/Post-Use FIT Considerations.

Grade & Pore Size	DI Water	60% IPA PSI (bar)	70% IPA PSI (bar)	
STW0.1	80 psi (5.52 bar)	30 psi (2.06 bar)	27 psi (1.86 bar)	
STC0.1	110 psi (7.68 bar) <sup>1</sup>	36 psi (2.48 bar)	35 psi (2.41 bar)	
STS0.2	50 psi (3.45 bar)	16.3 psi (1.12 bar)	15.7 psi (1.08 bar)	
SMH0.4	40 psi (2.76 bar)			
SMH0.6	22 psi (1.52 bar)	Please contact Meissner if you require these values.		
SPH0.2	42 psi (2.90 bar)	i you require these values.		
STW0.2	The recommended test method for these filters is diffusive flow.			

#### **EverLUX®** Bubble Point Specifications

<sup>1</sup> Bubble point value exceeds capsule pressure rating and is only recommended for filter cartridges in stainless steel housings.



### Integrity Test: Manual Diffusive Flow Test (Filter Housing System)

The diffusion test may be performed on EverLUX<sup>®</sup> STW0.2 filter cartridges and capsules. The filter is wetted, drained, and a constant air pressure is applied. Diffusional air flow through the membrane is measured.

The manual diffusive flow test relies on a wetted membrane filter of sterilizing grade, a housing or holder to contain the wetted filter, a calibrated pressure gauge, a regulated gas-pressure source (usually compressed air or nitrogen), downstream tubing, and an inverted, water-filled graduated cylinder suspended in a container of water. Test pressure is increased to the manufacturer's recommended test pressure. The following highlights the basic steps required to perform a diffusive flow test on a single cartridge filter.

#### Procedure (Figure 2)

- 1. Close all valves.
- 2. Open  $V_1$  and  $V_6$ . Thoroughly wet the filter with water, opening vent  $V_3$  to allow trapped air to escape. The complete wetting of the filter is crucial to the accuracy of the test. An incompletely wet filter will fail. Close  $V_1$  and  $V_3$ .
- 3. Open V<sub>2</sub> and apply the specified air pressure to the inlet side of the system. This will push the upstream volume of water through the filter.
- 4. If necessary, open V<sub>4</sub> to drain the downstream volume of water. Water remaining downstream of the filter may cause an inaccurate diffusion reading by interfering with the air flow in the outlet tube. Close V<sub>4</sub>.
- 5. Open  $V_5$  and close  $V_6$ .
- 6. Verify the test pressure and adjust as necessary.
- 7. Fill a graduated cylinder with water and place the outlet tube under the inverted opening. Record the volume of air diffused per minute. The recorded flow rate must not exceed the flow rate specified for the filter per the chart on Page 12.

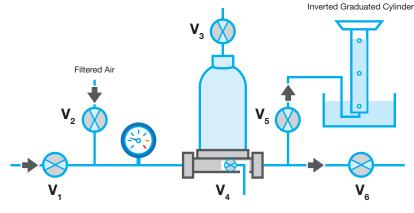


Figure 2

### **EverLUX® Diffusive Flow Specifications1**

Grade	Nominal Filter Size	Air psi (bar)	Water mL/min	
SMH0.4, SMH0.6, SPH0.2	All Sizes	Meissner recommends usi filter integrity testing meth		
	CM Capsules		1.2	
	CK Capsules		1.9	
	CS Capsules, 2.5" SFE Cartridges		6	
STW0.2	CL Capsules, 5" SFE Cartridges	30 psi (2.07 bar)	13	
	CJ Capsules		17	
	10" UltraCap <sup>®</sup> , UltraCap <sup>®</sup> H.D., Capsules & Cartridges <sup>2</sup>		30	
	CA, CB Capsules	Meissner recommends usi filter integrity testing met		
	CM Capsules		1.2	
	CK Capsules		1.4	
STS0.2	CS Capsules, 2.5" SFE Cartridges		5	
	CL Capsules, 5"SFE Cartridges	35 psi (2.41 bar)	11	
	CJ Capsules		16	
	10" UltraCap <sup>®</sup> , UltraCap <sup>®</sup> H.D., Capsules & Cartridges <sup>2</sup>		28	
	CA, CB, CM, CK Capsules	Meissner recommends using bubble point as a filter integrity testing method for these sizes.		
	CS Capsules, 2.5" SFE Cartridges		4.4	
STW0.1	CL Capsules, 5" SFE Cartridges	40 psi (2.76)	9.6	
	CJ Capsules		15	
	10" UltraCap <sup>®</sup> , UltraCap <sup>®</sup> H.D. Capsules & Cartridges <sup>2</sup>		25	
	CA, CB Capsules	Meissner recommends usi filter integrity testing met		
	CM Capsules		1.4	
	CK Capsules		1.2	
STC0.1	CS Capsules, 2.5" SFE Cartridges		5.3	
	CL Capsules, 5" SFE Cartridges	40 psi (2.76)	11	
	CJ Capsules		15	
	10" UltraCap <sup>®</sup> , UltraCap <sup>®</sup> H.D. Capsules & Cartridges <sup>2</sup>		25	

<sup>1</sup> Meissner does not publish IPA diffusive flow values for EverLUX<sup>®</sup>. <sup>2</sup> For >10" filters, maintain the pressure and multiply the mL/min by the number of 10" units to get the diffusive flow specification.

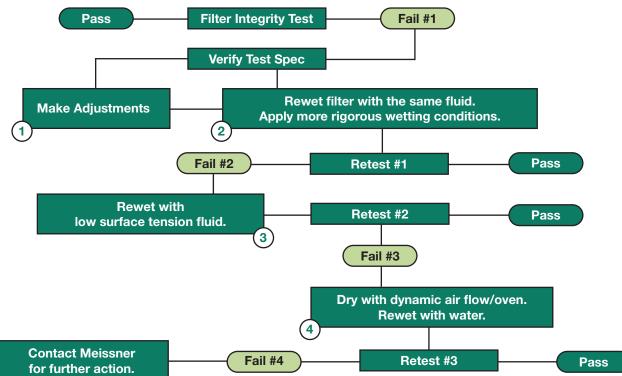


### **Filter Integrity Test Preparations**

- 1. Verify filter P/N matches specification, SOP, and filter to be installed.
- 2. Visually inspect filter to ensure that it is not damaged. Note damages, if applicable.
- 3. Verify the wetting fluid required, as well as note any preparation needed to execute the test with the specified wetting fluid (additional flushing, tubing, housing flush, etc.).
- 4. Verify the specific test gas (air, nitrogen, etc.).
- 5. Verify the source pressure.

**Troubleshooting Guidance** 

- 6. Verify that the filter assembly temperature is stable. Temperature changes can affect testing gas and distort diffusive flow values if not stable.
- 7. Verify testing set-up is without leaks. Leaks within the testing set up affect the FIT results by disturbing applied inlet pressure and by distorting the measured gas flow in the membrane.



Use the list on this page to verify testing specs. Make any adjustments needed and retest. If no adjustments are needed, rewet the filter in the same fluid and retest.

More rigorous wetting conditions can be enacted by increasing the flush time and volume, increasing pressure differential by increasing upstream pressure, or modifying downstream pressure. In addition to assisting with filter wetting, back pressure also facilitates the removal of air entrapped in the membrane pleats by further solubilizing the air (due to increase pressure in the system) and by compressing air bubbles to a size where they can freely pass through the membrane.

Rewet with a low surface tension fluid and retest. Meissner does not publish IPA integrity test values for EverLUX<sup>®</sup> STW 0.2. When using IPA to facilitate wetting for EverLUX<sup>®</sup>, it is advised that the IPA is flushed out with water thoroughly prior to retesting.

Dry with dynamic air flow or over drying at 50° C for PES for 8 hours, let cool, and retest with water. If the filter fails, contact Meissner for further action. If the filter passes, failure root cause is likely air lock. Air lock is the result of air being trapped between the downstream wetted membrane and the upstream wetted membrane and can be very difficult to remove with flushing alone.



### **Pre-Use/Post-Use FIT Considerations**

Wetting Fluid	Pre-Use	Post-Use
Water	Ensure filter, any associated tubing, and housing are clean before wetting.	It is important to completely flush any remaining product from the filter, tubing, and housing prior to integrity testing. Any diluted product remaining can impact the FIT results.
60%/70% IPA	If using alcohol for a wetting fluid, the filter must be dried or flushed after FIT and before putting the filter into service.	It is important to completely flush any remaining product from the filter, tubing, and housing prior to integrity testing. Any product remaining can impact the FIT results.
Product	Flushing the filter (not just wetting) may be required and is recommended if using process product as a testing fluid.	The filter membrane should already be sufficiently wet, and no further action should need to be taken prior to conducting the filter integrity test. Note: If the FIT indicates failure and retesting is desired with other wetting fluids, ensure all product remaining in the filter is completely flushed as it can impact the FIT results of other wetting fluids used.



### **Autoclave Instructions**

Meissner filters may be autoclaved repeatedly without loss of integrity. Note: EverLUX<sup>®</sup> filters must be wet with water prior to autoclaving if the filters are to be integrity tested post-sterilization. The filter may be fully immersed in water, open side up, until the core fills with water to wet the filter enough for autoclaving. Alternatively, the filter may be integrity tested prior to sterilization.

#### Capsule, UltraCap<sup>®</sup> & UltraCap<sup>®</sup> H.D. Filters

The following outlines the steps recommended in the autoclave sterilization of Meissner filter capsules. Gamma irradiated filters should not be autoclaved before use due to increased extractables and brittleness after both sterilization methods are employed.

- 1. Wet filter, drain.
- 2. Loosely cover the capsule inlet and outlet with autoclave wrap. All capsule vents are on the upstream side of the filter and should be loosened or removed to facilitate steam penetration. Hose barb vent valves must be opened at least two full turns to prevent valve leakage post autoclaving.
- 3. The weight of clamps or fittings attached to the capsule must be supported to avoid damaging the adapters. Sanitary flanges may have clamps and gaskets loosely attached to the filter. If fittings must be attached to flanges, tri-clamps are preferable to bi-clamps and should be tightened after the assembly has cooled.
- 4. Autoclave the capsule at a minimum of 121°C for 60 minutes or 125°C for at least 45 minutes with the capsule in a horizontal position using a slow exhaust or liquid cycle. T-style UltraCap<sup>®</sup> capsules may be autoclaved horizontally or with the outlet oriented downward to facilitate the removal of condensate from the downstream side of the filter. As autoclave systems vary, sterilization cycles should be validated under actual system or autoclave loading conditions. Downstream attachments can significantly increase the time required to sterilize the filter core.
- Allow the capsule to cool. Gently close vents finger tight. Excessive tightening of vent valves will damage the sealing surfaces. Integrity test if desired. Install filter into system aseptically.

### Cartridge & SFE (Small Flow Elements)

The following outlines the steps required to autoclave a Meissner filter cartridge and housing assembly. A stainless steel reinforcement ring is required for filter configurations with 222 or 226 O-rings.

- 1. Immerse the cartridge in water, drain, and install the filter into the housing. Loosely cover the inlet and outlet with autoclave wrap. Vent and drain valves should be fully open.
- 2. Autoclave the cartridge and housing assembly at a minimum of 121°C for 30 minutes with the filter outlets in an outlet down or horizontal position using a slow exhaust or liquid cycle. As autoclave systems vary, sterilization cycles should be validated under actual system or autoclave loading conditions. Assemblies attached to the outlet can increase the required sterilization times.
- 3. Allow the housing assembly to cool. Integrity test if desired.
- 4. Install the sterile filter assembly into the system using aseptic techniques.

Different autoclave temperature and time combinations may be used to sterilize the filters, but the combination should be validated to ensure that sterilization occurs under those conditions. Temperatures above 135°C are not recommended.





### **Inline Steam Sterilization Procedure**

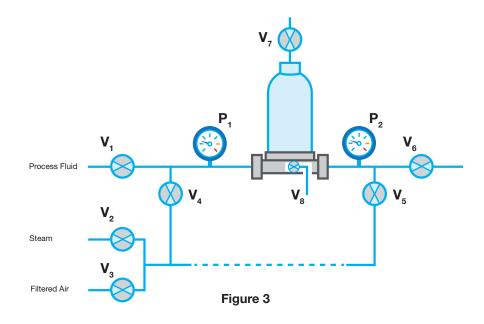
Steaming in place (SIP) is frequently used in critical applications where a sterile effluent is desired. To prevent damage to the filter cartridge's O-ring adapter, cartridges with 222 or 226 O-ring adapters must be reinforced with a stainless steel ring. Meissner filter cartridges with reinforced O-ring adapters are capable of repeated sterilization cycles without loss of integrity. The steps required to steam sterilize the EverLUX<sup>®</sup> filter cartridge and system using saturated steam are outlined in the procedure below.

The steam should be free of rust and other particulates. The housing should be clean before the cartridge is installed. EverLUX<sup>®</sup> filters must be wetted before steaming if they are to be integrity tested after use or post steaming. When steam sterilizing a wetted cartridge, upstream and downstream gauges must be provided to verify that the differential pressure across the membrane does not exceed 5 psi (0.3 bar). To assure sterilization, steam pressure in the assembly must not be allowed to fall below 15 psi (1 bar) or 121°C. Condensate should be drained from the system during sterilization. A typical piping schematic is outlined in Figure 3.

#### Caution: Capsules, UltraCap<sup>®</sup> and UltraCap<sup>®</sup> H.D. are not designed for inline steam sterilization!

#### Procedure (Figure 3)

- 1. Close all valves.
- 2. Open valve V<sub>4</sub>.
- 3. If cartridge is wet, open V<sub>5</sub>. (Use if filter will be integrity tested later.)
  - a. Slowly open V<sub>2</sub>. This will connect both sides of the filter to steam pressure.
  - b. Crack open  $V_7$  to vent trapped air.
  - c. Crack open V<sub>6</sub> allowing steam to flow through the system.
  - d. Slowly close V<sub>5</sub> but do not allow the differential pressure across the cartridge to exceed 5 psi (0.3 bar).
  - e. Leave drain V<sub>8</sub> cracked during sterilization to drain condensate.
- 4. If sterilizing a dry cartridge, slowly open V<sub>2</sub>. (Use if filter will not be integrity tested later.)
  - a. Crack open  $V_7$  to vent trapped air.
  - b. Crack open  $V_6$  to allow steam to flow through the system. Do not allow the differential pressure across the cartridge to exceed 5 psi (0.3 bar).
  - c. Leave drain V<sub>8</sub> cracked during sterilization to drain condensate.
- 5. Steam sterilize for 30 to 60 minutes at 15 to 20 psig (1.0 to 1.4 bar), or as long as experience dictates.
- 6. When sterilization is complete, close V<sub>2</sub>.
- 7. Open V<sub>3</sub> and introduce sterile air or nitrogen regulated to the same pressure as the steam.
- 8. Close  $V_{8}$  once steam and condensate flow stops.
- 9. Allow the system to cool to room temperature. Do not allow the differential pressure across the cartridge to exceed 5 psi (0.3 bar). Then close  $V_3$ ,  $V_7$ , and  $V_6$ . Keep the system under pressure until ready for use.
- 10. Crack vent  $V_7$  and allow the system pressure to equalize. The filtration process may now be started.





### Storage & Shelf Life

Meissner Filtration Products, Inc. manufactures a complete line of filter products and One-Touch<sup>®</sup> single-use assemblies. Products are suitably bagged and boxed for shipping and may be stored in the original packaging in a clean, dry area between 0°C and 40 °C (32 °F to 100 °F). The following gives the minimum shelf life expectancies for EverLUX<sup>®</sup> products.

#### **Filters**

The EverLUX<sup>®</sup> filter has an expected shelf life greater than 5 years in the cartridge, small flow, non-irradiated large and small capsule lines, and disc configurations. Filters may be used beyond their minimum expected shelf life if they were stored in their original packaging and are integrity tested prior to use and found to be within specification. Filter product age can be determined from the date on the original Certificate of Quality or Conformance.

Gamma irradiated filters, which include capsule, UltraCap<sup>®,</sup> and UltraCap<sup>®</sup>, H.D. capsules have a shelf life of at least 2 years from the date of irradiation. Gamma irradiated capsules are distinguished with part numbers beginning with the letter "G".

#### **One-Touch® Products**

The One-Touch<sup>®</sup> product line of single-use systems, including but not limited to biocontainers, tubing, and/or filter assemblies, has labeling that identifies the product specific expiration date. The standard shelf life of nonsterile and gamma irradiated One-Touch<sup>®</sup> products is 2 years from the date of manufacture. These standard time periods may be amended to reflect the various components included in a specific configuration, a change that will be indicated on the product label. Filters that are part of a One-Touch<sup>®</sup> single-use system begin with "C" and are irradiated once with the One-Touch<sup>®</sup> assembly.





### **EverLUX® Membrane Grade Descriptions**

- SLH = This single-layer PES membrane features a highly asymmetric pore structure but is not 100% integrity tested or flushed during manufacture. It is offered as an economical pr-filter or final filter when sterility assurance is not required. A Certificate of Conformance is provided on a lot basis.
- **SMH** = This standard, single PES membrane features a highly asymmetric pore structure. It is 100% integrity tested and flushed with DI water during manufacture. It is suited for critical applications when regulatory documentation requirements are minimal. A Certificate of Conformance is provided on a lot basis.
- **SPH** = This is an absolute, particulate rated filter. It is 100% integrity tested and DI flushed during manufacture. A Certificate of Conformance is provided on a lot basis.
- STW = This pharmaceutical validated, sterilizing grade filter features two serially layered, highly asymmetric PES membranes with the coarser upstream layer optimized for prefiltration. This filter meets full traceability requirements for the pharmaceutical industry. It is 100% integrity tested and flushed with DI water during manufacture. Each STW-grade filter is shipped with a Certificate of Quality stating exact quality control criteria and test performance results.
- STS = This pharmaceutical validated, sterilizing grade filter features a highly asymmetric PES membrane layered over an asymmetric PES membrane for an optimized pre and final filtration combination. This filter meets full traceability requirements for the pharmaceutical industry. It is 100% integrity tested and flushed with DI water during manufacture. Each STS grade filter is shipped with a Certificate of Quality stating exact quality control criteria and test performance results.
- STC = This pharmaceutical validated, sterilizing grade filter features two serially layered PES membranes, one highly asymmetric and one asymmetric, designed for 100% removal of mycoplasma. The coarser upstream layer is optimized for prefiltration. The filter is 100% integrity tested and DI flushed during manufacture, and it has the added benefit of certification that meets the critical needs of the pharmaceutical, biotechnology, and related industries.

Filter Grade	Sterilizing Applications	Integrity Tested & Flushed	Certificate of Conformance per Lot	Certificate of Quality per Filter	cGMP Traceability Conformance	Bacterial Rating Requirements
SLH	No	No	Yes	No	No	P, BR
SMH0.4	Yes	Yes	Yes	No	No	М
SMH0.6	Yes	Yes	Yes	No	No	М
SPH0.2	No	Yes	Yes	No	No	P, BR
STS0.2	Yes	Yes	No	Yes	Yes	M, S
STW0.2	Yes	Yes	No	Yes	Yes	M, S
STW0.1	Yes	Yes	No	Yes	Yes	S, MR*
STC0.1	Yes	Yes	No	Yes	Yes	S, AMR

#### **Application Documentation & Ratings Requirements**

#### **Bacterial Rating Requirements Key**

M: Microbially Rated

**BR:** Bioburden Reduction

P: Particle Rated S: Sterilizing Grade MR: Mycoplasma Reduction AMR: Absolute Mycoplasma Retention

\*Absolute Mycoplasma Retention is achievable with two or more STW 0.1 filters in series, or alternatively, one STC 0.1 filter.





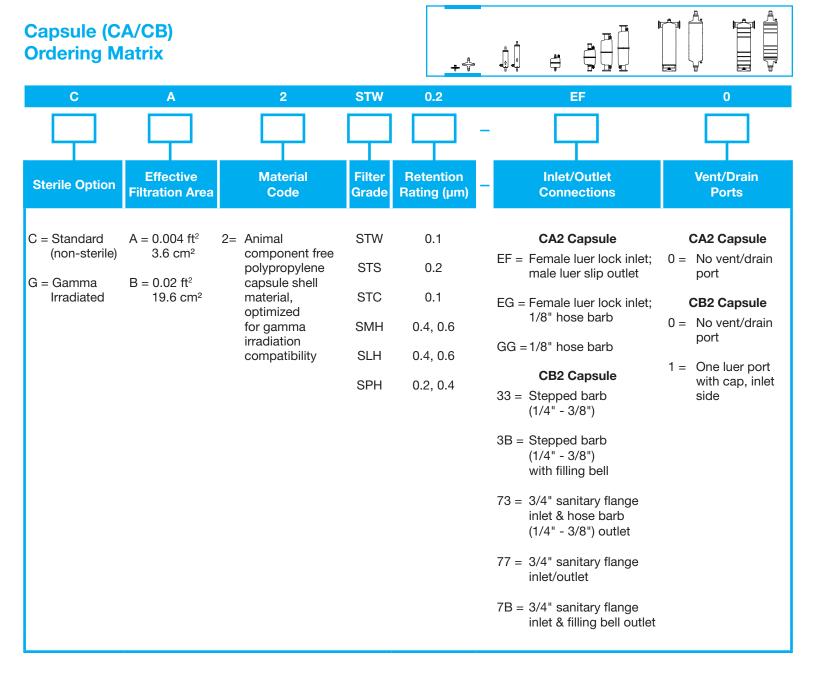
## Cartridge Ordering Matrix

STW	0.1	- 2	C6	R	S
	$\square$	- 🖵	$\square$	$\square$	$\square$
Filter Grade	Absolute Rating (µm)	Cartridge Length	End Cap Configurations	Reinforcement Ring Option	Seal Material (O-ring or Gasket)
Grade SMH STW STS STC SLH SPH	Rating (μm) 0.4, 0.6 0.1, 0.2 0.2 0.1 0.4, 0.6 0.2, 0.4	Length 1 = 10" 2 = 20" 3 = 30" 4 = 40"	Configurations $GS = DOE; flat gaskets (9.75", 19.5", 29.25", 39" length filters)GL = DOE; flat gaskets (20", 30", 40" length filters)C1 = SOE; 222 nO-Ring^{\circledast}, button cap endC2 = SOE; 222 O-rings, button cap endF1 = SOE; 222 nO-Ring^{\circledast}, fin endF2 = SOE; 222 O-rings, fin endF2 = SOE; 222 O-rings, fin endC5 = SOE; 226 nO-Ring^{\circledast}, button cap endC6 = SOE; 226 O-rings, button cap endF5 = SOE; 226 nO-Ring^{\circledast}, fin endF5 = SOE; 226 nO-Ring^{\circledast}, fin endF5 = SOE; 226 nO-Ring^{\circledast}, fin endF6 = SOE; 226 O-rings, fin end$	(Blank) = Standard - no reinforcement ring R = Reinforcement ring - required only for the 222 and 226 adapter	(O-ring or Gasket) <b>D-ring Seal</b> B = Buna E = EPR S = Silicone T = Teflon® over silicone V = Viton® X = Teflon® over Viton® <b>Gasket Seal</b> B = Buna E = EPR P = Polyethylene S = Silicone T = Teflon® V = Viton®
			DN = DOE; internal 120 O-rings RN = SOE; internal 120 O-ring, recessed cap end		
			DA = DOE; internal 213 O-rings RA = SOE; internal 213 O-ring, recessedcap end		

### Small Flow Elements (SFE Filters) Ordering Matrix

	OTW	0.1		<b></b>		<b>6</b>		D	<b>C</b>
L	STW	0.1		5		6		R	S
L			_						
								<u> </u>	
Model	Filter Grade	Absolute Rating (µm)	-	Length/Area		Adapter Selection		inforcement ling Option	O-ring Material
L	SMH	0.4, 0.6		For SMH, SLH, SPH	P =	Standard internal	(Blank) =	Standard - no	B = Buna
	•	,		grade filters:	•	116 O-ring	(,	reinforcement	E = EPR
	STW	0.1, 0.2		2 = 2.5"/1.6 ft <sup>2</sup> (0.15 m <sup>2</sup> )	2 =			ring	S = Silicone
	OTO	0.0		5 = 5.0"/3.3 ft <sup>2</sup> (0.31 m <sup>2</sup> )		autoclave/SIP applications,	R =	Reinforcement ring - required	T = Teflon <sup>®</sup> over Silicone
	STS	0.2		For STW, STS, STC		select "R" under "Reinforcement		only for the 222 and 226 adapter	V = Viton <sup>®</sup>
	STC	0.1		grade filters:	Ring Option")		when autoclaving	X = Teflon <sup>®</sup> over	
				2 = 2.5"/1.2 ft² (0.11 m²)	6 =	226 O-ring style		or steam sterilizing	Viton®
	SLH	0.4, 0.6		5 = 5.0"/2.6 ft <sup>2</sup> (0.24 m <sup>2</sup> )		locking adapter (for autoclave/		-	
	SPH	0.2, 0.4				(for autoclave/ SIP applications, select "R" under "Reinforcement Ring Option")			
					SK =	Skirt-flange adapter (no reinforcement or O-ring options available)			
					L =	116 O-ring with Mini Lock			





Capsule (CI Ordering Ma				+ 🕆					
С	К	2	STW	0.1	- 77	4			
$\square$	$\square$	2		$\square$	- 🖵	$\square$			
Sterile Option	Effective Filtration Area	Material Code	Filter Grade	Absolute Rating (µm)	Inlet/Outlet Connections	Vent/Drain Ports			
C = Standard (non-sterile)	For SMH, SLH, SPH grade filters:	2= Animal component	SMH	0.4, 0.6		0 = No vent/drain port			
G = Gamma irradiated	M = 0.36 ft² (335 cm²) K = 0.45 ft² (415 cm²)	free polypropylene capsule shell	STW	0.1, 0.2	1B = 1/4" hose barb with filling bell	1 = 1 luer port with cap, inlet side			
		material, optimized	STS	0.2		2 = Standard - 2 luer ports with caps			
	For STW, STS, STC grade filters:	for gamma irradiation compatibilityl	STC	0.1	2B = 3/8" hose barb with filling bell	4 = 2 sanitary valves with hose barb			
	$M = 0.28 \text{ ft}^2 (260 \text{ cm}^2)$ $K = 0.35 \text{ ft}^2 (325 \text{ cm}^2)$					SLH	0.4, 0.6	41 = 1/4" MNPT; 1/4" hose barb out	5 = 1 sanitary valve with hose barb
			SPH	0.2, 0.4	44 = 1/4" MNPT	connection, inlet side			
					71 = 3/4" TC in; 1/4' hose barb out	6 = 1 sanitary valve with hose barb, outlet side			
					72 = 3/4" TC in; 3/8' hose barb out				
					77 = 3/4" sanitary (TC) flange				



Capsule (CF Ordering Ma	-					
CF	2	STW	<b>0.1</b> ·	- 33	А	1
$\square$	$\square$				$\square$	
Sterile Option	Material Code	Filter Grade	Absolute Rating (µm)	Inlet/Outlet Connections	Filtration Area (Nominal)	Vent/Drain Ports
CF = Standard (non-sterile)	(Blank) or 1 = Polypropylene capsule shell material	e SMH	0.4, 0.6	33 = Stepped barb (1/4" - 3/8")	B = 0.70 ft² ( 650 cm²)	0 = No vent/drain port
GF = Gamma Irradiated	2= Animal component free polypropylene capsule shell material , optimized for gamma irradiation compatibility	SLH SPH	0.4, 0.6	<ul> <li>3B = Stepped barb (1/4" - 3/8") inlet; filling bell outlet</li> <li>41 = 1/4" MNPT inlet; 1/4" hose barb outlet</li> <li>44 = 1/4" MNPT</li> <li>73 = 3/4" sanitary flange in; stepped barb outlet</li> </ul>		<ul> <li>1 = 1 luer port with cap, outlet side</li> <li>2 = Standard - 2 luer ports with caps</li> <li>4 = 2 sanitary valves with hose barbs</li> <li>5 = 1 sanitary valve with hose barb connection, outlet side</li> </ul>
				77 = 3/4" sanitary flange		6 = 1 sanitary valve with hose barb connection, inlet side



Capsule (C Ordering N			-	<b>₽</b> ₽			
С	S	2	STW	0.1	- 02		2
$\square$	<b>P</b>	$\square$		$\Box$			$\square$
Sterile Option	Filtration Area (Nominal)	Material Code	Filter Grade	Absolute Rating (µm)	Inlet/Outlet Connections		/ent/Drain Ports
= Standard (non-sterile)	For SMH, SLH, SPH grade filters:	capsule shell	SMH	0.4, 0.6	00 = 1" sanitary flange	0 =	No vent/dra port
	S = 1.6 ft² (0.15 m²) L = 3.3 ft² (0.31 m²)	material 2 = Animal	STW	0.1, 0.2	02 = 1" sanitary flange inlet;	1 =	1 luer port with cap,
= Gamma Irradiated	$J = 5.2 \text{ ft} 2 (0.37 \text{ m}^2)$	component free	STS	0.2	3/8" hose barb outlet		outlet side
	For STW, STS, STC grade filters:	polypropylene capsule shell material, optimized	STC	0.1	09 = 1" sanitary flange inlet; 1/2" hose	2 =	Standard - 2 luer ports with caps
	$S = 1.2 \text{ ft}^2 (0.11 \text{ m}^2)$	for gamma irradiation	SLH	0.4, 0.6	barb outlet (flexible tubing)	4 =	2 sanitary valves with
	L = 2.6 ft² (0.24 m²) J = 5.2 ft2 (0.37 m2)	compatibility	SPH	0.2, 0.4	0C = 1" sanitary flange inlet; 1/2" hose	5 =	hose barb 1 sanitary valve with
					barb outlet 22 = 3/8" hose barb		hose barb connection, outlet side
					2B = 3/8" hose barb with filling bell	6 =	valve with
					CC = 1/2" hose barb		hose barb, inlet side
					LL = 1" sanitary flange, long neck		
					44 = 1/4" MNPT		
					55 = 3/8" FNPT		
					66 = 3/8" MNPT		
					72 = 3/4" sanitary flange inlet; 3/8" hose barb outlet		
					77 = 3/4" sanitary flange		
					88 = 3/4" hose barb		
					99 = 1/2" hose barb flexible tubing		



	JltraCap <sup>®</sup> (T-Style & Inline) Drdering Matrix							ÂA U	₽₿			
CU	STW	0.1	-	2	т		00			2		
	$\Box$	$\square$	-	$\square$	<u> </u>		$\square$					
Sterile Option	Filter Grade	Absolute Rating (µm)	-	Cartridge Length	Body Style		Inlet/Outle Connection			Vent/Drain Po T-Style	orts	
CU = Standard (non-sterile)	SMH	0.4, 0.6		1 = 10"	T = T-style		1" sanitary 1	-		No vent or drain		
GU = Gamma Irradiated	STW	0.1, 0.2		2 = 20"	N = Inline		3/4" sanitar 1" sanitary 1		1 =	No vent; 1/4" san drain plug	itary	
inadiated	STS	0.2		3 = 30"		02 –		let; 3/8" hose		Sanitary vent; 1/4" sanitary drain plug		
	STC	0.1				= 00	1" sanitary f inlet; 1/2" h barb outlet		3 =	Sanitary vent; 3/4 flange gauge port	; 1/4"	
	SLH	0.4, 0.6				09 =	1" sanitary 1	flange	4 =	sanitary drain plug Sanitary vent; no		
	SPH	0.2, 0.4					inlet; 9/16" barb outlet			Sanitary vent; 3/4 flange gauge port	" sanitary	
						08 =	1" sanitary f inlet; 3/4" h barb outlet		6 =	No vent or drain; sanitary flange ga	3/4"	
							3/8" hose b			Vent/Drain Po Inline	orts	
						CC =	1/2" hose b	arb				
						99 =	9/16" hose	barb	0 =	No vent or drain		
			88 = 3/4" hose barb $2 =$ Two sanit		Two sanitary vent	/drain valves						
							1/2" Flarete 3/4" Flarete		4 =	One sanitary vent valve, outlet side	or drain	

### **MEISSNER**

	traCap <sup>®</sup> H.D. (T-Style & Inline) dering Matrix						e eff			
CR	CR 2 STW 0.1 – 2		- 2	т		00		2		
<u> </u>	2		<u> </u>	- 🖵	$\square$		$\square$	$\square$		
Sterile Option	Material Code	Filter Grade	Absolute Rating (µm)	Cartridge Length	Body Style		Inlet/Outlet Connections			ain Ports tyle
CR = Standard (non-sterile)	2= Animal component	SMH	0.4, 0.6	1 = 10"	T = T-style		1" sanitary flange			
GR = Gamma Irradiated	free polypropylene capsule shell	STW	0.1, 0.2	2 = 20"	N = Inline	77 =	3/4" sanitary flange	1 =	No vent sanitary	; 1/4" drain plug
	material, optimized	STS	0.2	3 = 30"		02 =	1" sanitary flange inlet; 3/8" hose	2 =		/ vent; 1/4" drain plug
	irradiation compatibility	compatibility $0C = 1"$ sanitary flang inlet; 1/2" hose barb outletSLH $0.4, 0.6$ $5 = 50"$ $0C = 1"$ sanitary flang inlet; 1/2" hose barb outletSPH $0.2, 0.4$ $09 = 1"$ sanitary flang inlet; 9/16" hose	0.1	4 = 40"	" 0C = 1" sanitary flange	3 =	sanitary			
					gauge port; 1/4" sanitary drain plug					
			1" sanitary flange inlet; 9/16" hose barb outlet	4 =	Sanitary drain	/ vent; no				
						08 =	1" sanitary flange inlet; 3/4" hose barb outlet	5 =	sanitary	y vent; 3/4" y flange port; no drair
						0D =	1" sanitary flange inlet; 1" hose barb outlet	6 =		t or drain; hitary flange port
						22 =	3/8" hose barb	A =	No vent drain va	; sanitary alve
						CC =	1/2" hose barb	B =	Sanitary	
						99 =	9/16" hose barb	C =		sanitary drain valve Sanitary vent;
						88 =	3/4" hose barb	•	sanitary drain; 3/4" sanitary flange	drain; 3/4"
						DD =	1" hose barb		gauge p	
						AA =	1/2" Flaretek®			ain Ports ine
						BB =	3/4" Flaretek®	0 =	No vent	
								2 =	Two sar drain va	nitary vent/ alves
								4 =		nitary vent or alve, outlet

Viton® and Teflon® are registered trademarks of E. I. du Pont de Nemours and Company.

Flaretek<sup>®</sup> is a registered trademark of Entegris, Inc.

EverLUX<sup>®</sup>, UltraCap<sup>®</sup> and UltraCap H.D.<sup>®</sup> are registered trademarks of Meissner Filtration Products, Inc. © 2025, 2024, 2023, 2019, 2013, 2007 Meissner Filtration Products, Inc. GD002-7.0-A

