

# **Certificate of Quality**

The Hydrophilic PVDF Membrane Capsule filters have been manufactured in a **mdi** facility in compliance with **ISO 9001** regulations using **validated production processes**.

## AseptiCap WS Hydrophilic PVDF Membrane Capsule Filters

Catalog No. : LWSX5301EEXX101

Type : LWS

Pore Size : 0.2 μm (0.45μm + 0.2μm)
Lot Number : LV4477I SI.No. 004

#### **SPECIFICATION**

Length	5"
Filter Media	Hydrophilic PVDF Membrane
Drainage Layers	Polyester
Housing	Polypropylene
Differential Pressure	< 4Kg/cm² at 30 °C
Maximum Operating Temperature	80 °C @ < 2 Kg/cm <sup>2</sup>
Sterilization	2 Autoclaving cycles at 125 °C of 30 minutes each

#### **LOT RELEASE CRITERIA**

**100% Integrity Tested** : The capsule filter has been tested for integrity by Air Diffusion Flow test and Bubble

Point test using DI water.

Diffusion flows with DI water were: ≤ 15 ml/min @ 2.60 kg/cm² Bubble point value with DI water was: ≥ 50 psi (3.44 Bar)

The capsule filter is also certified for integrity by Bubble point test using 50%

IPA/Water solution.

Bubble point with 50% IPA/Water solution is ≥ 16 psi (1.10 Bar)

**Water Flow Rate** :≥ 10 lpm @ 0.70 Kg/cm² @ 27 °C

**Microbial Challenge Test** : Retains  $\ge 10^7$  organisms/cm<sup>2</sup> of *B. diminuta* ATCC 19146 challenge as per

ASTM F838-05 methodology.

**VALIDATED FOR** 

**Heat Stability** : Maintains integrity after 2 autoclaving cycles at 125 °C of 30 minutes each.

**Extractable** : Within limits as specified in USP.

Oxidizable matter : Passes test as per USP.

**Bacterial Endotoxins** : Filtrate meets the USP requirements for Sterile WFI of ≤ 0.25 EU/mI as determined

by Limulus Amebocyte Lysate (LAL) test.

**Biosafety** : Passes Biological Reactivity Tests, *In Vivo* for Class VI plastic as described in

USP <88>.

**Cytotoxicity**: Passes Biological Reactivity Tests, In Vitro as described in USP <87>.

indirect Food Additives : Passes as per FDA 21CFR 177.1520(a)1(i).

Particle Release : Passes test as per USP <788>, "Particulate matter in Injections".

Fiber Release : Complies with FDA 21CFR 210.3(b)(6).

### **CUSTOMER SUPPORT**

**mdi** offers its unique interdisciplinary skills to provide solutions to specific problems. Please contact our factory or the local application specialist.

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Head of Quality Assurance

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