

# **Certificate of Quality**

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

## AseptiCap KSO PES Membrane Capsule Filters

Catalog No. : DKOX5201EEXX101

Type : DKSO

Pore Size : 0.2 μm (0.45 μm + 0.2 μm)
Lot Number : DK7772I SI.No. 007

#### **SPECIFICATION**

Length	2"
Filter Media	Polyethersulfone Membrane
Drainage Layers	Polypropylene
Differential Pressure	< 4Kg/cm² at 30 °C
Housing	Polypropylene
Maximum Operating Temperature	80 °C @ < 2 Kg/cm²
Sterilization	25 autoclaving cycles at 125 °C of 30 minutes each

### **LOT RELEASE CRITERIA**

**100% Integrity Tested** : The capsule filter has been tested for integrity by Bubble point test using DI water.

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**  $\ge 3.5 \text{ lpm} @ 0.70 \text{ Kg/cm}^2 @ 27 °C$ 

Microbial Challenge Test : Retains ≥ 10<sup>7</sup> organisms/cm² of *B. diminuta* ATCC 19146 challenge as per

ASTM F838-05 methodology.

**VALIDATED FOR** 

**Heat Stability** : Maintains integrity after 25 autoclaving cycles at 125 °C of 30 minutes each.

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

Oxidizable matter : Passes test as per USP.

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

in USP <88>.

**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.

Our I

Head of Quality Assurance

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An ISO 9001 Company