

# **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 KS-y PES Membrane Capsule Filters

Catalog No. : DKS15736DDXX301

Type : DKS-S

Pore Size : 0.1 μm (0.2 μm + 0.1 μm)

Lot Number : DK9979G SI.No. 005

Ster. No. : R008 Expiry Date : 2019 - 07

## **SPECIFICATION**

Length	8"
Filter Media	Polyethersulfone Membrane
Drainage Layers	Polyester
Differential Pressure	< 4Kg/cm² at 30 °C
Housing	Polypropylene
Maximum Operating Temperature	80 °C @ < 2Kg/cm <sup>2</sup>
Sterilization	Pre sterilized by Gamma Irradiation

#### **LOT RELEASE CRITERIA**

**100% Integrity Tested** : The Capsule filters have been tested for integrity by Air Diffusion Flow Test using

purified water and Bubble Point Test using 50% IPA/Water solution.

Diffusion flows were: ≤ 13.5 ml/min @ 3.52 kg/cm<sup>2</sup>

Bubble point was: ≥ 31 psi (2.14 Bars).

**Water Flow Rate** : ≥ 5.5 lpm @ 0.70 Kg/cm² @ 27 °C

**Sterility** : Samples passed the sterility test in accordance with U.S. pharmacopoeia.

**VALIDATED FOR** 

**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>.

**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.

Head of Quality Assurance

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