

## **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 KL-y PES Membrane Capsule Filters

Catalog No. : DKLX5702DDXX301

 $\begin{tabular}{lll} Type & : DKL-S \\ Pore Size & : 0.45 \ \mu m \end{tabular}$ 

Lot Number : DK9304K Sl.No. 234

Ster. No. : R2564

Date of Sterilization : 2024 - 11

Expiry Date : 2026 - 11

## **SPECIFICATION**

Length	8"
Filter Media	Polyethersulfone Membrane
Drainage Layers	Polyester
Housing	Polypropylene
Differential Pressure	< 4Kg/cm² at 30 °C
Maximum Operating Temperature	80 °C @ < 2Kg/cm <sup>2</sup>
Sterilization	Pre sterilized by Gamma Irradiation dose between 25 kGy to 40 kGy

**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: ≥ 30 psi (2.07 Bar)

Typical Water Flow Rate : 19 lpm @ 0.70 Kg/cm<sup>2</sup> @ 27 °C

**VALIDATED FOR** 

**Bubble point (50% IPA)** : The filter is certified/validated for integrity by Bubble point test using 50%

IPA/Water solution. Bubble point ≥ 10 psi (0.69 Bar)

Sterility : The sterilization process has been validated to assure Sterility Assurance Level

(SAL) of 10<sup>-6</sup> in accordance with ISO 11137.

**Microbial Retention**: Retains microbial challenge of *S. marcescens* (ATCC 14756).

Bacterial Endotoxin : Aqueous extracts exhibit < 0.25 EU/mL as established by Limulus Amebocyte

Lysate (LAL) test as per USP <85>.

**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).

**Total Organic Carbon**: Meets USP <643> limit of 500 ppb for total organic carbon after flushing specified

volume of water for injection.

Conductivity : Meets USP <645> limit of 1.3 µS/cm at 25 °C for water conductivity after flushing

specified volume of water for injection.

## **CUSTOMER SUPPORT**

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

T. No.: COQ/CAP/004-05

Om

Head of Quality Assurance Issue Date: 07-Nov-24 Advanced Microdevices Pvt. Ltd.

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