

# 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**.

## **BioPro KS PES Membrane Capsule Filters**

Catalog No. : LBKS5601EEXX101  
 Type : LBKS  
 Pore Size : 0.2 µm  
 Lot Number : LK8961B SI.No. 015

### SPECIFICATION

<b>Length</b>	30"
<b>Filter Media</b>	Polyethersulfone Membrane
<b>Drainage Layers</b>	Polyester
<b>Housing</b>	Polypropylene
<b>Differential Pressure</b>	< 4Kg/cm <sup>2</sup> at 30 °C
<b>Maximum Operating Temperature</b>	80 °C @ < 2Kg/cm <sup>2</sup>

### LOT RELEASE CRITERIA

**100% Integrity Tested** : The Capsule filters have been tested for integrity by Air Diffusion test and Bubble point test using DI water.  
 Diffusion flows were: ≤ 100 ml/min @ 1.54 kg/cm<sup>2</sup>  
 Bubble point was: ≥ 30 psi (2.07 Bar)

**Typical Water Flow Rate** : 105 lpm @ 0.70 Kg/cm<sup>2</sup> @ 27 °C

### VALIDATED FOR

**Microbial Challenge Test** : Retains ≥ 10<sup>5</sup> organisms/cm<sup>2</sup> of *B. diminuta* ATCC 19146 challenge.

**Bacterial Endotoxin** : Aqueous extracts exhibit < 0.25 EU/mL as established by Limulus Amebocyte Lysate (LAL) test as per USP <85>.

**Sterilization** : 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.



Head of Quality Assurance  
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**An ISO 9001 Company**