

## Viresolve® Pro PD Kit

Catalogue Number: VPMCPDKNB9  
Lot Number: C9DN97785

### Good Manufacturing Practices

This product was manufactured in a Millipore facility which adheres to Good Manufacturing Practices.

### ISO® 9001 Quality Standard

This product was manufactured in a Millipore facility whose Quality Management System is approved by an accredited registering body to the appropriate ISO 9001 Quality Systems Standard.

### Animal Origin Statement

All Component materials used in the manufacture of this device are either animal free or in compliance with EMEA/410/01.

### Quality Assurance Lot Release Criteria

This manufacturing lot was sampled, tested and released by Quality Assurance to the following specifications:

#### USP Bacterial Endotoxin

A sample aqueous extraction contains less than 0.25EU/ml as determined using the Limulus Amebocyte Lysate (LAL) test.

#### Membrane Bacteriophage Retention Test

Samples of the Viresolve Pro membrane exhibited equal to or greater than 4 LRV retention of ØX174 bacteriophage in the presence of a model protein at a minimum challenge level  $10^7$  pfu/cm<sup>2</sup>.

#### Device Bacteriophage Retention Test

Device samples exhibited equal to or greater than 4 LRV retention of ØX174 bacteriophage in the presence of a model protein at a minimum challenge level  $10^7$  pfu/cm<sup>2</sup>.

### Quality Performance Criteria

This product was designed and manufactured to meet the following specifications.

#### Component Materials Toxicity

Component materials were tested and meet the Criteria for the USP <88> Biological Reactivity Tests for Class VI Plastics.

#### TOC and Conductivity

Affluent after 50 L/m<sup>2</sup> water flush exhibited less than 500 ppb TOC and less than 1.3 µS/cm conductivity.

#### Non-Fiber Releasing

This product was manufactured with products that meet or exceed the criteria for "Non-Fiber releasing filter" as defined in 21 CFR 210.3(b)(6).



Peter Eichert  
Quality Manager

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