



SteriPS® FILTER CARTRIDGES

PES Membrane Sterile Liquid Filters

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HYDROPHILIC PES MEMBRANE FILTERS DESIGNED FOR RELIABLE PERFORMANCE AT AN ECONOMICAL COST

SteriPS® Filter Cartridges are specially designed to provide a reliable sterilization solution at an economical cost. Hydrophilic PES membrane cartridges require no pre-wetting and are ready to use. In addition, these filters provide excellent performance for pharmaceutical applications.

| FEATURE | BENEFIT |
|--------------------------|------------------------------------------------------------------------------------------------------------------------------|
| Hydrophilic PES membrane | Requires no pre-wetting |
| | Low diffusion flow |
| | High surface area provides excellent flow rates and extended service life while maintaining high bacteria removal efficiency |
| | Low protein binding |

QUALITY STANDARDS

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|------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------|
| Quality Assurance | These products are manufactured in a facility which adheres to ISO 9001:2015 Practices. 100% Integrity tested and Lot traceable. |
| Bacterial Retention | Retention of 10 ⁷ cfu/cm² Brevundimonas diminuta (ATCC® 19146) according to ASTM F838 for ≤ 0.2µm. |
| TOC / Conductivity at 25°C | Autoclaved filter effluent meets the USP <643> for Total Organic Carbon and USP <645> for Water Conductivity per WFI requirements after a UPW flush |
| Particle Shedding | Autoclaved filter effluent meets the requirements in USP <788> for large volume parenterals/injections |
| Non-Fiber Releasing | Component materials meet the criteria for a "Non-fiber-releasing filter" as defined in 21 CFR 210.3 (b) (6) |
| Bacterial Endotoxin | Aqueous extraction of autoclaved filter contains < 0.25 EU/mL as determined by Limulus; Amebocyte Lysate (LAL), USP <85> |
| USP <87> Cytotoxicity | Meet the requirement of USP <87> In Vitro Cytotoxicity Test |
| USP <88> Biological Toxicity | Meet the criteria of the USP <88> Biological Reactivity Test for Class VI-121°C plastics. |
| Indirect Food Additive | All component materials meet the FDA Indirect Food Additive requirements cited in 21 CFR 177–182. |



Typical Applications

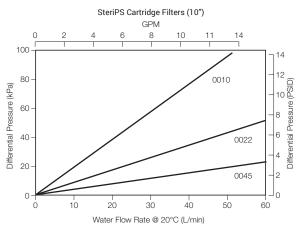
- Antibiotics
- LVP & SVP
- Large Batch Solutions
 Cleaning and Disinfectanting Liquids

SPECIFICATIONS

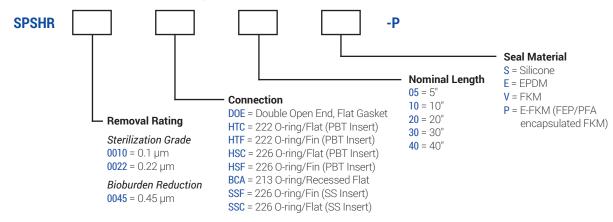
| Filter media | PES membrane (single layer) (Symmetric) |
|-----------------------------|-----------------------------------------------|
| Supports/core/cage/end caps | Polypropylene |
| Effective filtration area | 0.58m² (6.2 ft²)/ Ø69-10 inch |
| 0-rings | Silicone, EPDM, FKM, FEP/PFA encapsulated FKM |

| O-ring internal insert | PBT |
|----------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Cartridge diameter | 69 mm (2.7 in.) |
| Max. Operating pressure | 0.69 MPa (6.9 bar, 100 psi) at 25°C; 0.40 MPa (4.0 bar, 58 psi) at 60°C; 0.24 MPa (2.4 bar, 35 psi) at 80°C |
| Max. Differential pressure | Forward: 0.69 MPa (6.9 bar, 100 psi) at 25°C; 0.40 MPa (4.0 bar, 58 psi) at 60°C; 0.24 MPa (2.4 bar, 35 psi) at 80°C |
| | Reverse: 0.3 MPa (3.0 bar, 44 psi) at 25°C; 0.1 MPa (1.0 bar, 15 psi) at 80°C |
| Bubble point | 0.38 MPa (3.8 bar, 55 psi) wetted with water of 20°C, 0.1 μm 0.32 MPa (3.2 bar, 46 psi) wetted with water of 20°C, 0.22 μm 0.20 MPa (2.0 bar, 29 psi) wetted with water of 20°C, 0.45 μm |
| Steam sterilization | up to 100 cycles for 30 minutes at 135°C at Max.Differential Pressure of 0.3 bar (4.3 psi) |
| Autoclave | up to 200 cycles for 30 minutes at 130°C |
| | |

FLOW RATES



EXAMPLE: **SPSHR0022HSF10S-P** = 0.22 µm, 226/Fin, 10" filter with silicone seals



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