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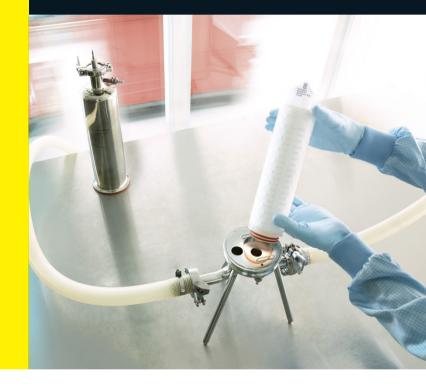
Product Datasheet

Sartopore® 2 0.45 µm

Bioburden & Particle Retentive Filter Elements

Product Information

- Extremely high flow rates and total throughputs
- Number one choice for bioburden reduction & particle removal
- Broad compatibility and easy scalability
- Outstanding filtration performance of aqueous solutions that are highly viscous & difficult to filter



No. 1 choice for bioburden reduction & particle removal

Due to the "built-in prefiltration" by a 0.8 µm membrane, Sartopore® 2 0.45 µm rated filters are ideally suited for bioburden reduction and particle removal of highly viscous and difficult to filter pharmaceutical solutions. Due to its hydrophilic Polyethersulfone membrane Sartopore® 2 0.45 µm filter elements offer outstanding total throughputs and flow rate performances. Thus ensuring highest process efficiency, minimized filtration costs and short filtration cycle times.

Broad compatibility and highest reliability

The unique Polyethersulfone membrane gives Sartopore® 2 0.45 µm broad chemical compatibility, including a pH-range from pH 1 to pH 14, and a high thermal resistance, making it suitable for a broad range of filtration liquids. To assure absolute reliability, each individual Sartopore® 2 0.45 µm filter element is integrity tested by Diffusion Point and Bubble Point Test prior to release.

Scalability

Sartopore[®] 2 0.45 µm filter elements are available in a broad range of sizes and formats to provide linear scale-up from R&D to process scale. The wide range of membrane areas from 300 cm² to 1.8 m² per filter element guarantees greatest flexibility and most economic filter sizing.

Applications

- Therapeutics
- Buffers
- Injectables
- Ophthalmics
- Large volume parenterals
- Antibiotics
- Biological Fluids
- Tissue Culture Media
- Bulk pharmaceutical products
- Acetic and basic solutions

Services

Sartorius Confidence® Validation Services is the perfect complement to Sartopore® 2 0.45 µm filters.

Our services provide

- Extractables and leachables services
- Microbiological testing
- Physicochemical testing

in compliance with regulatory requirements. Our local teams of validation experts support you with our tailored and consultative approach to determine the most costeffective solution and give you the confidence you need to succeed.

Available Sizes	Filtration Area	Max. Diffusion at 2.5 bar 36 psi [ml/min]	Min. Bubble Point [bar psi]
Cartiridges, T-Style Maxicaps®, Maxic	caps® Gamma Maxicaps®		
Size 0.5 (Only Cartridge)	0.3 m ² 3.2 ft ²	7	2.2 32
Size 1	0.6 m ² 6.5 ft ²	12	2.2 32
Size 2	1.2 m ² 12.9 ft ²	24	2.2 32
Size 3	1.8 m² 19.4 ft²	36	2.2 32
Mini Cartridges, Midicaps® Gamma	Midicaps®		
Size 7	0.05 m ² 0.5 ft ²	3	2.2 32
Size 8	0.1 m ² 1.1 ft ²	4	2.2 32
Size 9	0.2 m ² 2.2 ft ²	6	2.2 32
Size 0 (Only Midicaps®)	0.45 m ² 4.8 ft ²	12	2.2 32
Capsules Gamma Capsules			
Size 5	0.03 m ² 0.32 ft ²	2	2.2 32

Technical Specifications

Max Allowable Differential Pressure

Cartridges | Mini Cartridges

5 bar | 72.5 psi at 20°C 2 bar | 29 psi at 80°C

T-Style Maxicaps[®], Maxicaps[®] | Gamma Maxicaps[®]

5 bar | 72.5 psi at 20°C 3 bar | 43.5 psi at 50°C

Midicaps[®] | Gamma Midicaps[®] 5 bar | 72.5 psi at 20°C 2 bar | 29 psi at 80°C

Capsules | Gamma Capsules Size 5 4 bar | 58 psi at 20°C 2 bar | 29 psi at 50°C

Materials

Prefilter Membrane Polyethersulfone, asymmetric

Endfilter Membrane

Polyethersulfone, asymmetric

Support Fleece

Polypropylene (In-line steam sterilizable & autoclavable) Polyester (y-irradiatable or y-irradiatable | autoclavable)

Core Polypropylene

End Caps Polypropylene

Capsule Housing Polypropylene

O-Ring Silicone (other materials on request)

Filling Bell Polycarbonate

Max Allowable Back Pressure

2 bar | 29 psi at 20°C (for all elements)

Pore Size Combination

0.8 µm + 0.45 µm

Regulatory Compliance

- Each individual element is tested for integrity by bubble point and diffusion test
- Designed, developed and manufactured in accordance with an ISO 9001 certified Quality Management System
- Meet or exceed the requirements for WFI quality standards set by the current USP
- Non pyrogenic according to USP Bacterial Endotoxins
- USP Plastic Class VI Test
- Non fiber releasing according to 21 CFR



Sterilization

Cartridges, Mini Cartridges

In-Line Steam Sterilization 134 °C, 0.5 bar, 20 min. Min. 25 Sterilization Cycles

or

Autoclaving 134°C, 2 bar, 30 min Min. 25 Sterilization Cycles

Maxicaps[®], Midicaps[®] & Capsules

Autoclaving 134°C, 2 bar, 30 min Min. 25 Sterilization Cycles (Midicaps® & Maxicaps®)

Gamma Maxicaps®, Gamma Midicaps® & Gamma Capsules Gamma Irradiation ≤ 50 kGy 1 Sterilization Cycles

T-Style Maxicaps®

Autoclaving 134°C, 2 bar, 30 min Min. 5 Sterilization Cycles

or

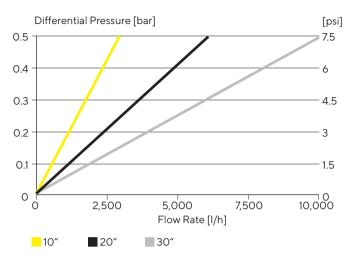
Gamma Irradiation ≤ 50 kGy 1 Sterilization Cycle

Technical References

Validation Guide	2665239
Extractables Guide	2396283

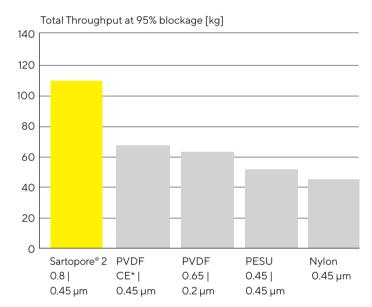
Performance

Water Flow Rate for Maxicaps® and Standard Cartridges



Standardized at 20°C

Total Throughput Comparison



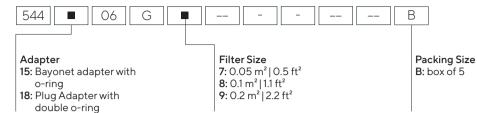
10″ Cartridges format

* Cellulose Ester prefilter

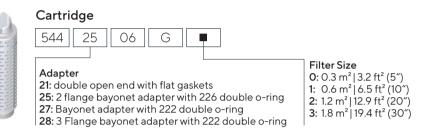
Ordering Information



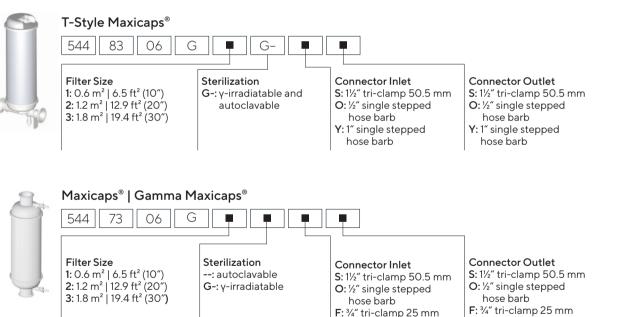
Mini Cartridge



(Standard with silicone o-ring optional with EPDM or Fluoroelastomer o-ring).



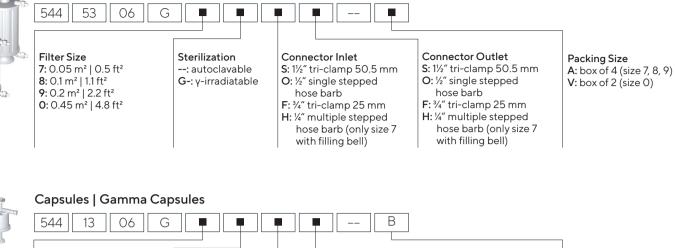
(Standard with silicone o-ring optional with EPDM or Fluoroelastomer o-ring).



(Optional with vent valve design for connection of integrity tester. Example: 5447307H1--SSIT)



Midicaps[®] | Gamma Midicaps[®]



Filter Size Sterilization 5: 0.03 m² 0.32 ft² : autoclavable G-: γ-irradiatable	Connector Inlet S: ¾" tri-clamp 25 mm O: ¼" multiple stepped hose barb	Connector Outlet S: ¾″ tri-clamp 25 mm O: ¼″ multiple stepped hose barb	Packing Size B: box of 5
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Germany

USA

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