





BEVPOR PS wine filters protect the unique characteristics of wine by removing yeast and other spoilage organisms to ensure microbial stabilization prior to packaging.

The inert and highly asymmetric PES membrane provides validated microbial retention to typical spoilage organisms whilst preserving the wine's unique properties to ensure it reaches the consumer as the wine maker intended. Combined with hydrophilic properties for easy integrity testing, BEVPOR PS filters provide assured performance throughout their service life.

BEVPOR PS filters have been designed to provide a cost-effective solution to wine microbial stabilization by providing increased process control with increased operational efficiency.

Features

Validated retention to spoilage organisms

Inert materials of construction

Easily integrity tested in-situ

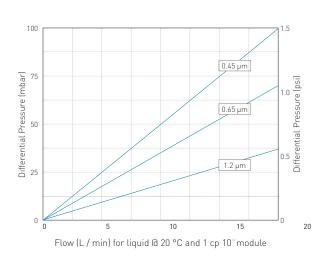
Benefits

Ensures effective microbial stabilization of wine

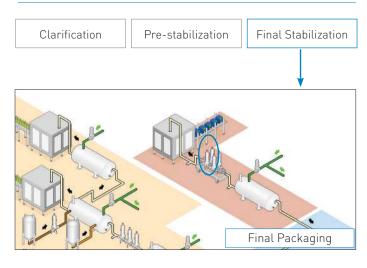
Preserves the desirable characteristics of the wine

Assured filtration performance

Performance Characteristics



Filtration Stage





Specifications

Materials of Construction

Filtration Membrane:
 Upstream Support:
 Downstream Support:
 Inner Support Core:
 Outer Protection Cage:
 End Caps:
 Tod Cap Insert:
 Polyethersulphone
 Polyester
 Polypropylene
 Polypropylene
 Nylon
 3161 Stainless Stee

End Cap Insert: 316L Stainless SteelO-rings: Silicone / EPDM

Food Contact Compliance

Materials conform to the relevant requirements of FDA 21 CFR Part 177, current EC1935 / 2004 and current USP Plastics Class VI - 121 °C.

Recommended Operating Conditions

Up to 70 °C (158 °F) continuous operating temperature and higher short-term temperatures during CIP to the following limits:

Temperatur	re	Max Fo	Max Forward dP		
°C	°F	(bar)	(psi)		
20	68	5.0	72.5		
40	104	4.0	58.0		
60	140	3.0	43.5		
80	176	2.0	29.0		
90	194	1.0	14.5		
>100 (steam)	>212 (steam)	0.3	4.0		

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.6 m² (6.45 ft²)

Cleaning and Sterilization

BEVPOR PS cartridges can be repeatedly steam sterilized in-situ or autoclaved at up to 130°C (266°F). They can be sanitized with hot water at up to 90°C (194°F) and are compatible with a wide range of chemicals. Please refer to our Clean-in-Place support guide or contact your local Parker representative for more information.

Retention Characteristics

The retention characteristics of BEVPOR PS filters have been validated by challenges performed with the following organisms.

Organism	LRV wh	LRV when challenged with a minimum of 10 ⁷ cfu per cm ²			
		0.45	0.65	1.2	
Saccharomyces & Brettanomyces & Lactobacillus bro Acetobacter oen Pseudomonas ac Serratia marceso	eruxellensis evis eruginosa	FR FR FR FR 9.1 FR	FR FR FR FR 8.9 FR	FR FR 2.0 7.6 4.8 2.4	

*FR - Fully retentive during challenge

When expressed as titre reduction "FR" equates to >10" per 10" module.

Integrity Test Data

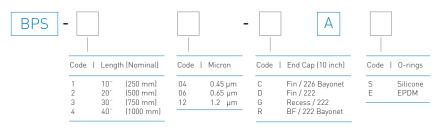
All filters are flushed with pharmaceutical grade purified water prior to despatch. They are integrity tested to the following limits:

Diffusional Flow	Micron Rating	
Test Parameters	0.45 0.65 1.2	
Test Pressure (barg) Test Pressure (psig)	1.4 1.0 0.6 20.0 15.0 9.0	
Max Diffusional Flow per 10" (ml /min)	16.0 16.0 16.0	

Manufacturing Traceability

Each filter cartridge displays the product name, product code and lot number.
Additionally, each module displays a unique serial number providing full manufacturing traceability.

Ordering information



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