





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.

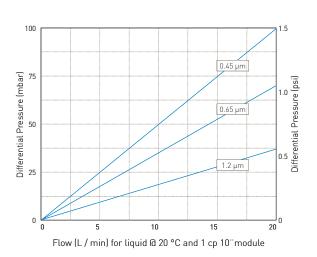
### **Features**

- I Validated retention to spoilage organisms
- I Inert materials of construction
- I Easily integrity tested in-situ

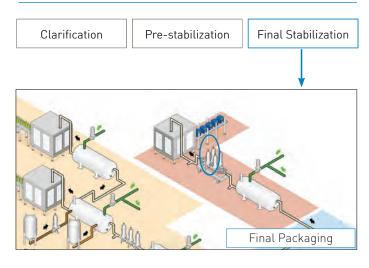
## **Benefits**

- I Ensures effective microbial stabilization of wine
- I Preserves the desirable characteristics of the wine
- I Assured filtration performance

# Performance Characteristics



# Filtration Stage





## Specifications

#### Materials of Construction

I Filtration Membrane: Polyethersulphone
Upstream Support: Polyester
Downstream Support: Polyester
Inner Support Core: Polypropylene
Outer Protection Cage: Polypropylene
I End Caps: Nylon

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:

Temperature		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 m<sup>2</sup> (6.45 ft<sup>2</sup>)

#### 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 when challenged with a minimum of 10 <sup>7</sup> cfu per cm <sup>2</sup>			
		0.45	0.65	1.2
Saccharomyces cer Brettanomyces bru. Lactobacillus brevis Acetobacter oeni Pseudomonas aeru Serratia marcescen	xellensis 5 ginosa	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)	1.4	1.0	0.6
Test Pressure (psig)	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

