

# Bevigard™-M Cartridge Filters

■ For Optimal Protection of Sterilising Grade Filters or As an Excellent Final Filter for Water Applications



## Typical Applications

Bevigard-M cartridge filters are designed for beverage prefiltration applications like clarification or for final filtration stages such as sterilising grade filter protection or bioburden reduction of water.

## Efficiency

Used in last stage prefiltration applications, these versatile surface-type filters retain contaminants on the mixed cellulose esters membrane.

Used as final filter for bioburden reduction, they offer excellent microorganisms removal efficiency.

## Cost Effective

Because of their structure, Bevigard-M cartridge filters offer the best protection of final sterilising-grade filters, saving money on this critical final step.

Bevigard-M filters can also be repeatedly hot water regenerated providing highly cost effective filtration.

## Quality

Supplied with a Certificate of Quality which certifies that Bevigard-M filters meet Quality Assurance lot release criteria.

They are manufactured in a facility whose Quality Management System is approved by an accredited registering body to the ISO 9000 Quality Systems Standard.

## Specifications

### Materials

Rigid polypropylene core, end caps, and outer handling cage; cellulose ester filter material; spun bonded polyester supports; silicone O-rings.

### Cartridge Size

25cm/10", 50 cm/20", 75 cm/30", 100 cm/40" lengths; outside diameter 7.4 cm.

### Filtration Area

#### Double Layer Cartridges

0.7 m<sup>2</sup> per 25 cm/10" cartridge element.

#### Single Layer Cartridges

0.8 m<sup>2</sup> per 25 cm/10" cartridge element.

### Operating Conditions

#### Maximum Differential Pressure:

3.5 bar at 25°C.

#### Maximum Operating Temperature:

80°C.

#### Sterilisability:

Bevigard-M cartridges may be multiple steam-sterilised or hot water sanitised.

## Certificate of Quality - Example

Bevigard-M filters are designed, developed and manufactured in accordance with a Quality Management System approved by an accredited registering body to an ISO 9000 Quality System Standard.

Each Bevigard-M cartridge is shipped with a Certificate of Quality for documentation accuracy.

This certificate reports all the tests made on these cartridges to insure the consistency and the quality of the product we deliver.

<b>Bevigard-M® Cartridge Filter</b> 0,2 µm Nominal Rated Catalogue Number: CW0373SB1 Lot Number: <b>Good Manufacturing Practices</b> This product was manufactured in a Millipore facility which meets or exceeds FDA Device Good Manufacturing Practice standards. <b>ISO 9000 Quality Standard</b> This product was manufactured in a Millipore facility whose Quality Management System is approved by an accredited registering body to the appropriate ISO 9000 Quality Systems Standard. <b>Non Fiber Releasing</b> This product was manufactured with a Millipore media combination which meets the criteria for a "non-fiber releasing" filter as defined in 21 CFR 210.3 (b) (6). <b>Component Materials Toxicity</b> Component materials were tested and met the criteria for the USP Class VI Biological Test for Plastics at 50°C. <b>Indirect Food Additive</b> All component materials meet FDA Indirect Food Additive requirements cited in 21 CFR 177-182.	<b>Quality Assurance Lot Release Criteria</b> This manufacturing lot was sampled, tested and released by Quality Assurance to the following specifications: <b>Structural Integrity</b> Samples were found to be integral as measured by an aerosol challenge test. <b>Flow Rate and Pressure Drop</b> Samples met a maximum pressure drop of 6.0 psid (414 mbar) at 30 gpm (114 L/min) per 30-inch cartridge with clean water at 23°C.	<b>Quality Assurance Audit Criteria</b> This product was designed and manufactured to meet the following specifications. Performance is confirmed by testing on an audit basis: <b>Toxicity</b> This product is non-toxic per the current USP General (Mouse) Safety Test. <b>Gravimetric Extractables</b> The extractables level was equal to or less than 50 mg per 10-inch cartridge after 24 hours in ASTM Type 1 reagent-grade water at controlled-room temperature. <b>Multiple Sterilisation Cycles</b> Integrity was maintained after 10 steam-in-place cycles of 30 minutes at 121°C.
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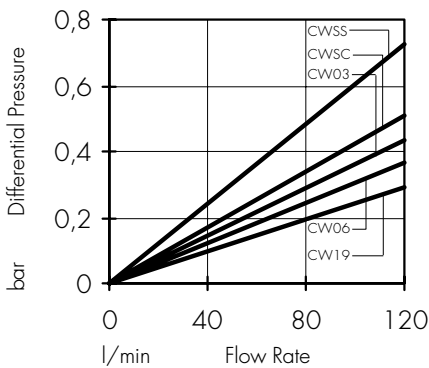
SAMPLE

  
 Nicholas Lambo  
 Vice President and General Manager,  
 BioProcess Division  
  
 John P. Tuttle  
 Manager, Worldwide Quality Systems  
 and Certification, BioProcess Division

Certificate of Quality - Example

## Water Flow Rate

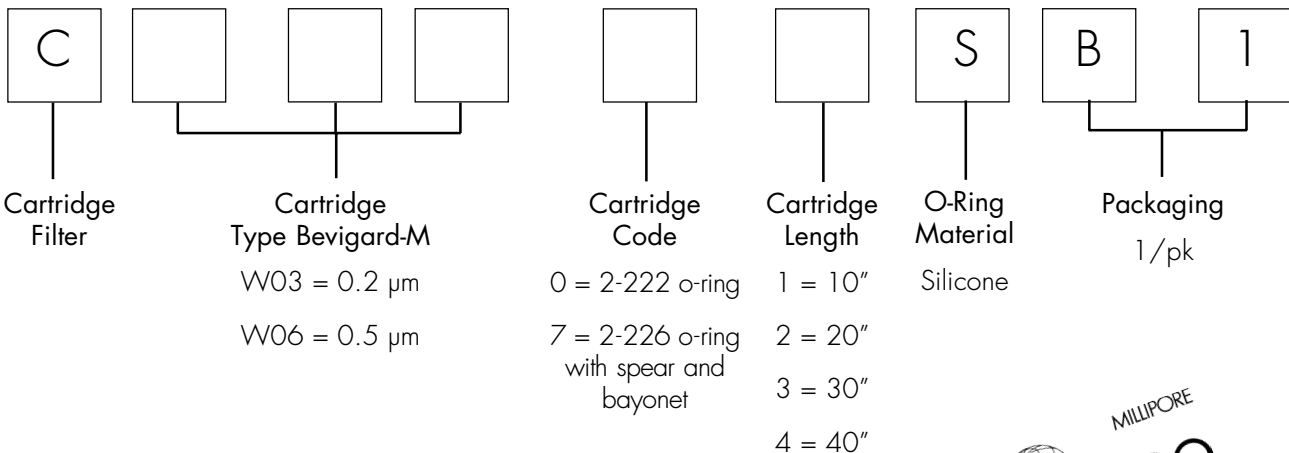
for a 75 cm/30" cartridge at 23 °C



## Construction and Pore Size

Bevigard-M	Nominal Pore Size	Mixed Cellulose Esters Membrane		
		0.2 µm	0.5 µm	1.2 µm
CW03	0.2 µm	X		
CW06	0.5 µm		X	
CW19	1.2 µm			X
CWSS	0.2 µm	X	X	
CWSC	0.5 µm		X	X

## Ordering Information



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