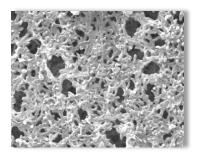


The BNM cartridge and capsule filters are made with Nylon 6,6 membrane. These long-proven filters are used for bioburden control in water, solvents and aqueous solutions. Pore sizes range from 0.10 to 0.65 μm and the filter sizes scale from laboratory to full production using identical materials to ensure consistent results.

BNM filters have broad chemical compatibility making them well suited for the filtration of solvents and other harsh chemicals. They are used to prevent bacteria contamination and remove microbials before final sterilizing filters.

The BNM filters have high retention and throughput. They are flushed to remove manufacturing debris and reduce extractables. Products are 100% integrity tested. BNM capsules are available pre-sterilized.

Critical Process provides unrivaled delivery times, technical consulting before purchasing, and very competitively priced high-performance products. Our comprehensive testing & analysis and validation services support your team whenever they need it. Your process experts partnering with our filtration experts is how we deliver your company's solution right the first time.



BNM bioburden control filters are recommended for:

- SVPs & LVPs
- Diagnostics
- Buffers
- WFI, Water Purification
- Non-protein Solutions
- Chemicals

Bioburden Control Clarification & Prefiltration



CARTRIDGES – Nominal Dimensions Length: 5 to 40 in. (12.7 to 101.6 cm) Outside Diameter: 2.75 in. (7.0 cm)



CAPSULES – Nominal Dimensions Length: 2 to 30 in. (5.1 to 76.2 cm) Outside Diameter: 3.50 in. (8.9 cm)

Maximum Operating Parameters

	CARTRIDGES	CAPSULES	
Liquid Operational Pressure	N/A	80 psi at 68 °F (5.52 bard at 20 °C)	
Gases Operational Pressure	N/A	60 psi at 68 °F (4.14 bar at 20 °C)	
Operating Temperature (water)	180 °F at 30 psid (82 °C at 2.07 bard)	110 °F at 30 psid (43 °C at 2.07 bard)	
Forward Differential Pressure	80 psid at 68 °F (5.52 bard at 20 °C)	80 psid at 68 °F (5.52 bard at 20 °C)	
Reverse Differential Pressure	50 psid at 68 °F (3.45 bard at 20 °C)	50 psid at 68 °F (3.45 bard at 20 °C)	
Recommended Changeout Pressure	35 psid (2.41 bard)	35 psid (2.41 bard)	

Sanitization & Sterilization

Filtered Hot Water*	90 °C (194 °F), 30 minutes, multiple cycles, max 3 psid forward flow	N/A	
Inline Steam*	275 °F (135 °C), 30 min, 25+ cycles	N/A	
Autoclave*	250 °F (121 °C), 30 min, 25+ cycles	250 °F (121 °C), 30 min, 25+ cycles	
Chemical Sanitization	Performed using industry standard concentrolly other selected chemicals.	Performed using industry standard concentrations of hydrogen peroxide, peracetic acid, and other selected chemicals.	

^{*}Cartridge Filters – For all elevated temperature procedures above, a stainless-steel support ring is required.

Filtration Area (Nominal)

	CAPSULES	CARTRIDGES AND CAPSULES			CARTRIDGES	
	2"	5"	10"	20"	30"	40"
Length	5.08cm	12.7cm	25.4cm	50.8cm	76.2cm	101.6cm
Area	1.2 ft ²	3.3 ft ²	7.0 ft ²	14.0 ft ²	21.0 ft ²	28.0 ft ²
	0.11m ²	0.31m ²	0.65m ²	1.30m²	1.95m²	2.60m ²

Integrity Testing

PORE SIZE	DIFFUSION TEST PRESSURE*		BUBBLE POINT MINIMUM*	
μm	PSIG	BARG	PSIG	BARG
0.10	48	3.31	**	**
0.22	35	2.41	50	3.4
0.45	20	1.38	25	1.7
0.65	15	1.03	19	1.3

DIFFUSION SPECIFICATIONS*						
Length	2"	5"	10"	20"	30"	40"
mL/min	≤ 4.3	≤ 12.9	≤ 30	≤ 60	≤ 90	≤ 120

^{*} All specifications are for water wetted membrane

^{**} Test pressure exceeds operational limits of capsule filters.
Use the Diffusion Test method.

Construction Materials

Filtration Media	Nylon 6,6 Membrane with polyester support		
Media Support	Polypropylene		
End Caps, Center Core, Outer Support Cage, Capsule Housing	Polypropylene		
Sealing Method	Thermal Bonding		
O-Rings/Gaskets Cartridges only	Buna, Viton® (or FKM), EPDM, Silicone, FEP Encapsulated Silicone, FEP Encapsulated Viton (or FKM)		

Validation

BNM filters are validated using test procedures that comply with ASTM F 838-15(ae1) protocols for the determination of bacterial retention in filters used for liquid filtration. The filters are challenged with the organisms listed below.

0.10μm: Brevundimonas diminuta 0.22μm: Brevundimonas diminuta 0.45μm: Serratia marcescens 0.65μm: Saccharomyces cerevisiae

Endotoxins

The levels of bacterial endotoxins in aqueous extracts from BNM filters are below current USP limits as specified for water for injection.

Extractables

BNM filters typically exhibit low levels of non-volatile residues.

TOC and Conductivity

BNM filters conform with TOC standards of USP <643> and the water conductivity standards of USP <645> after an appropriate flush with purified water.

Toxicity Compliance

The materials used to construct BNM filters are non-toxic and meet the requirements for the MEM Elution Cytotoxicity Test and the requirements for Biological Reactivity Tests in the current version of the United States Pharmacopeia (USP) for Class VI - 121 °C Plastics.

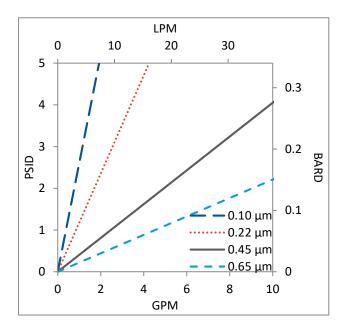
Non-Fiber Releasing

BNM filters comply with Title 21 CFR sections 210.3 (b)(6) and 211.72, for non-fiber releasing filters.

FDA Compliance

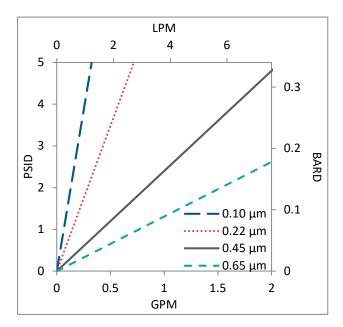
Materials meet the requirements listed by the FDA as appropriate for use in articles intended for repeated food contact as specified in Title 21 CFR sections 174.5, 177.1500, 177.1520, 177.1630, 177.2440, and 177.2600 as applicable.

Flow Rates for BNM Cartridges by Pore Size



Flow rates for Cartridge filters are per 10-inch length. The test fluid is water at ambient temperature.

Flow Rates for BNM Capsules by Pore Size

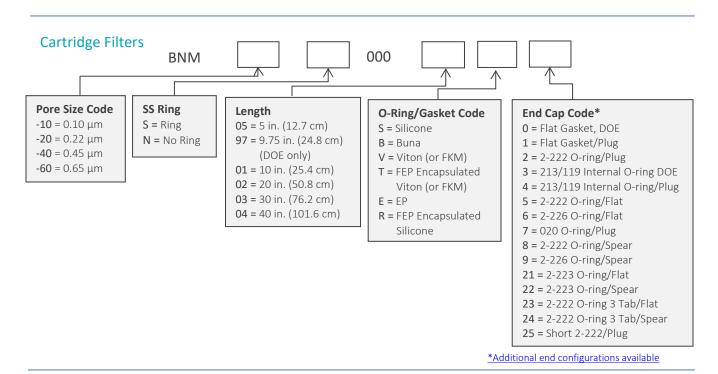


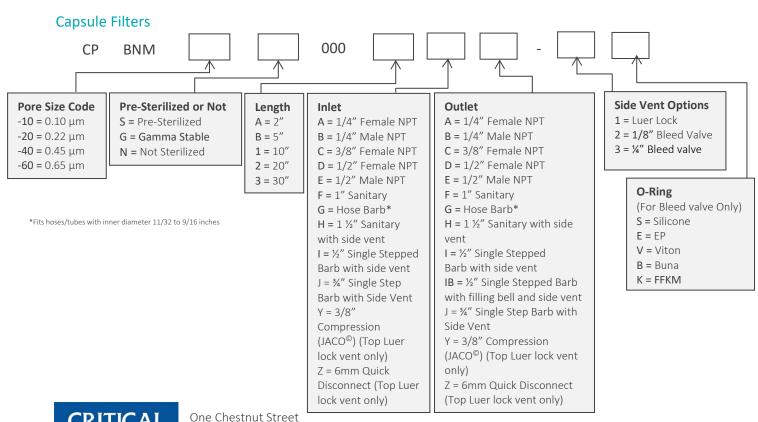
Flow rates for Capsule filters are tested using a 2" capsule filter with 1" sanitary inlet and outlet ports. The test fluid is water at ambient temperature. Flow rates for larger capsules will scale with filtration area. Rates will vary based on end configuration of the capsule.

BNM Filters Ordering Information

Fill in the corresponding codes in the boxes below to build your Part Number.

To consult with one of our technical team members, request a quote or place an order: call (603) 880-4420 or contact us here.





Nashua, NH 03060

FAX: 603.880.4536

CriticalProcess.com

603.880.4420

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