

PROPOR MR filters have been specifically designed for fast, effective and economical removal of mycoplasma from cell culture media in the biopharmaceutical industry.

Incorporating a highly retentive 0.1 micron rated PES membrane, PROPOR MR is validated against the industry standard *Brevundimonas diminuta* as well as *Acholeplasma laidlawii*, a common mycoplasma species found in contaminated cell cultures.

An asymmetric integral membrane prefilter layer provides PROPOR MR with the optimal membrane configuration for maximum capacity and flow rate. Quick processing times minimize the risk of contamination while still offering maximum protection from mycoplasma.

Features and Benefits

- Fully validated and integrity testable for assurance of sterility
- LRV >10 for *Acholeplasma laidlawii* for effective mycoplasma control
- Integral prefilter layer increases throughputs for reduction of filter trains
- Exceptional flow rates for quick processing of cell culture media

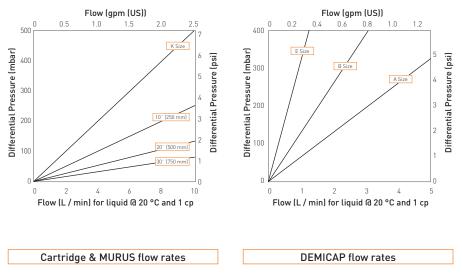
PROPOR MR Filter Cartridges

- liquid filters
- polyethersulphone



Note: PROPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



Specifications

Materials of Construction

Core:	Polypropylene
Sleeve:	Polypropylene
End Caps:	Nylon
Filtration Membrane:	Polyethersulphone

Prefilter Membrane: Polyethersulphone

Filter Cartridges

Upstream Support:	Polypropylene /
	Polyester
Downstream Support:	Polyester

Standard o-rings/gaskets: Silicone

MURUS Disposable Filter Capsules

Upstream Support:	Polypropylene /
	Polyester
Downstream Support:	Polyester
Standard o-rings/gaskets:	Silicone
Capsule Body:	Polypropylene
Capsules Vent Seals:	Silicone

DEMICAP Disposable Filter Capsules

Upstream Support:	Polyester
Downstream Support:	Polyester
Membrane Separation	
Layer:	Polyester
Capsule Body:	Nylon
Capsules Vent Seals:	Silicone
Filling Bell:	Polycarbonate

Recommended Operating Conditions Filter Cartridges

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

	erature	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	1.7	24.6					

MURUS Disposable Filter Capsules

Up to 25 °C (77 °F) @ 5.5 barg (79.7 psig) Up to 60 °C (140 °F) @ 2.8 barg (40.6 psig)

Parker Hannifin certify that this product complies with the European Council Pressure Equipment Directive (PED) 97/23/ EC Article 3, Paragraph 3 - Sound Engineering Practice (SEP). This product is intended for use with Group 1 & 2 Dangerous and Harmless Liquids and Group 2 Harmless Gases at the operating conditions stated in this document : In compliance with PED Article 3, Paragraph 3, SEP, this product does not bear the CE mark.

DEMICAP Disposable Filter Capsules

Up to 40 °C 104 °F) at line pressures up to 5.0 barg (72 psig).

Effective Filtration Area (EFA)

			-,
10" (250 mm):	0.50	m ²	(5.38 ft ²)
K Size:	0.24	m ²	(2.58 ft ²)
A Size:	0.19	m ²	(2.09 ft ²)
B Size:	0.10	m ²	(1.03 ft ²)
E Size:	0.05	m ²	(0.49 ft ²)

	Aut Cycles	oclave Temp	Steam Cycles (30 min.)	-in-Place Temp
Cartridges	10	130 °C (266 °F)	10	130 °C [266 °F]
MURUS	10	130 °C (266 °F)	-	-
DEMICAP	3	130 °C (266 °F)	-	-

Sterilization

PROPOR MR filter cartridges can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals.

For detailed operational procedures and advice on cleaning and sterilisation, please contact the Technical Support Group through your usual Parker domnick hunter contact.

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Quality Standards

Pharmaceutical grade products are manufactured in accordance with cGMP, 100% flushed with pharmaceutical purified water and integrity tested prior to despatch. A sample of each lot is tested to demonstrate conformity to validated claims.

Gamma-Irradiation

PROPOR MR MURUS disposable filters can be gamma-irradiated up to a maximum dosage of 40 kGy.

Performance Characteristics

TOC / Conductivity

The filtrate quality from a 10" (250 mm) PROPOR MR conforms to the requirements of current USP <645> (conductivity) within the first 1L flush of purified water and USP <643> (TOC) following a 10L flush.

Endotoxins

Aqueous extracts from the 10" (250 mm) PROPOR MR contain < 0.25 EU / ml when tested in accordance with the Limulus Amoebocyte Lysate test.

Non-Volatile Extractables (NVE)

Total NVEs extracted in the first 1 litre flush of purified water for a 10" (250 mm) cartridge / MURUS capsule are <15 mg.

Pharmaceutical Validation

A full validation guide is available upon request from Laboratory Services Group (LSG).

Oxidisable Substances

PROPOR MR filter cartridges meet current USP and EP quality standards for sterile purified water for oxidisable substances following a <1 litre water flush.

Integrity Test Data

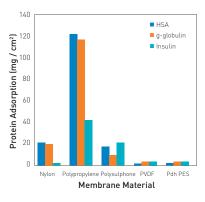
All filters are integrity testable to the following limits using air as the test gas. During diffusional flow tests filters are wet with water. Bubble points are determined in 60 / 40 v/v IPA / Water.

Micron Rating		0.1									
Filter Cartridges / MURUS / DEMICAP											
Min. Bubble Point	(barg)	2.36									
	(psig)	34.2									
Filter Cartridges / MURUS	/ DEMICAP										
Diffusional Flow	(barg)	4.80									
Test Pressure	(psig)	69.6									
Filter Cartridges / MURUS	/ DEMICAP										
Max. Diffusional Flow (10")	(ml / min)	24.2									
(K)		11.5									
(A)		9.3									
(B)		4.6									
(E)		2.2									

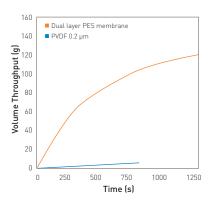
Retention Characteristics

PROPOR MR filter cartridges are validated by bacterial challenge testing with *Brevundimonas diminuta* to current ASTM F838-05 methodology (10⁷ organisms / cm² EFA minimum) with typical in-house challenge levels being 10¹¹ organisms per 10^{°°} (250 mm) filter cartridge.

To demonstrate the mycoplasma retention capabilities of the PROPOR MR, bacterial challenge testing was conducted on a number of cartridges using *Acholeplasma laidlawii* as the challenge organism with typical LRVs greater than 10.



Protein binding on membrane materials



Total volume throughput (g) vs time (s) for an insulin intermediate solution

PROPOR MR Filter Cartridges

Ordering Information

Cartridges

ZC	MR		- [-			
Code	Lengt	h (Nominal)	Code	Mic	ron	Code	Endcap (10")	Code	Variant	Code	O-rings ¹
K 1 2	5" 10" 20"	(125 mm) (250 mm) (500 mm)	610	0.1	μm	B C G	dh DOE BF / 226 Bayonet Recess / 222	P	Pharmaceutical	E S V	EPDM ² Silicone Viton
3	30" 40"	(750 mm) (1000 mm)				R	BF / 222 Bayonet			as stand having t 'S' code ² EPDM - Propylei	o-ring supplied dard without lo specify the Ethylene ne Diene er Rubber

MURUS Capsules

ZI	MR		-						-				-				
Cod	e Length	(Nominal)	Code	Mic	ron	Code	Inlet Connection	Code	e Outlet Connection	Cod	le Variant	Code	Grade	Code	 Design	Code	0-rings ¹
K 1 2	5" 10" 20"	(250 mm) (500 mm)	610	0.1	μm	A B D	³ / ⁴ " Tri-Clamp 1 ¹ / ₂ " Tri-Clamp 1" Hosebarb	A B D	^{3/4} " Tri-Clamp 1 ¹ / ₂ " Tri-Clamp 1" Hosebarb	P	Pharmaceutical	N S	Non-sterile Pre-sterilized γ (>25 kGy)	L T*	In-Line T-Port	E S V	EPDM ² Silicone Viton
3	30	(750 mm)					1" Tri-Clamp		1" Tri-Clamp					1" Tri-C		as stand	o-ring supplied lard without

DEMICAP Capsules

ZEN	MR		-					-							-	
Code L	Length (Nominal)	Code	Micron	Code	Inlet Connection	Code	Outlet Connection	Code	Variant	Code	Grade	Code	Pack N°	Code	Accessory
	4.4" 5.5"	(113 mm) (140 mm)	610	0.1 µm	T N	1" Tri-Clamp 1/," NPT Male	T N	1" Tri-Clamp 1/2" NPT Male	Р	Pharmaceutical	N S	Non-sterile Pre-sterilized	3	Pack of 3	FB	Filling Bell
A	7.9"	(200 mm)			H G	^{1/2} Hosebarb Stepped Hosebarb	H G	¹ / ₂ ² Hosebarb Stepped Hosebarb				γ (>25 kGy)			G & H co	onnections only

Syringe Filters							
ZSMR	-		-	-			
Code Diameter	Code Micron	Code Inlet Connection	Code Outlet Connection	Code Variant	Code Grade	Code Options	Code Pack N°
050 50 mm	610 0.1 µm				N Non-sterile	S Standard	025 25 per box
		G Stepped Hosebarb	G Stepped Hosebarb				