

PROCLEAR PP filters are designed for a wide range of prefiltration duties within the production of pharmaceuticals and are particularly suited to applications where chemical compatibility is an issue.

The optimum pleat configuration and graded density polypropylene media used in PROCLEAR PP filters ensure the filters have the highest possible throughput to blockage resulting in long service life.

The PROCLEAR PP range of filters are fully supported by a comprehensive validation guide to simplify its specification into new and existing processes.

Features and Benefits

Graded density . polypropylene media for high capacity

withstand aggressive

• Extremely robust to

conditions

- All polypropylene ٠ construction
- MURUS and DEMICAPs can be gamma-irradiated and autoclaved

PROCLEAR PP Filters

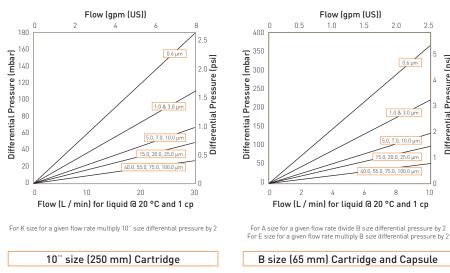
- liquid filters
- polypropylene



Note: PROCLEAR and DEMICAP are registered trademarks of Parker Hannifin Corporation.

Differential Pressure (psi)

Performance Characteristics



Specifications

Materials of Construction

Filtration Media:	Polypropylene
Upstream Support:	Polypropylene
Downstream Support:	Polypropylene

Filter Cartridges

Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene

- Outer Protection Cage:
- End Caps:
- End Caps Insert: 316L Stainless Steel
- *Not available in B & L endcap variants

MURUS Disposable Filter Capsules

- Core: Polypropylene Sleeve: Polypropylene End Caps Insert: 316L Stainless Steel
- Standard o-rings/gaskets: Silicone
- Capsule Body: Polypropylene
- Capsules Vent Seals:

DEMICAP Filter Capsules

Core:	Polypropylene
Sleeve:	Polypropylene
Capsule Body:	Polypropylene
Capsules Vent Seals:	Silicone
Filling Bell:	Polycarbonate

Syringe Filters

Body:

Polypropylene

Silicone

Polypropylene

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:

Temperature °C °F		Max. Forward dP (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	1.5	21.7

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 current European Council Pressure Equipment Directive (PED) - 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. The Pressure Equipment Directive mandates that category SEP product cannot bear the CE mark.

DEMICAP 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) up to 0.79m² (8.5 ft²)

Sterilization

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

	Autoclave		Steam	Steam-in-Place	
	Cycles		Cycles (30 min.)		
Cartridges	10	130 °C (266 °F)	30	135 °C (275 °F)	
MURUS	5	130 °C (266 °F)	-	-	
DEMICAP	10	130 °C (266 °F)	-	-	
Syringe	1	130 °C (266 °F)	-	-	

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

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 grade purified water.

Gamma-Irradiation

PROCLEAR PP MURUS & DEMICAP 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) PROCLEAR PP conforms to the requirements of current USP <643> (TOC) and USP <645> (conductivity) within the first 200 ml flush of purified water.

Endotoxins

Aqueous extracts from the 10" (250 mm) PROCLEAR PP 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 5 litre flush of purified water for a 10" (250 mm) cartridge are <10 mg.

Total NVEs extracted in the first 5 litre flush of purified water for an A size $7.9^{\circ\circ}$ (200 mm) DEMICAP capsule are <5 mg.

Pharmaceutical Validation

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

Oxidizable Substances

PROCLEAR PP filter cartridges meet current USP and EP quality standards for sterile purified water for oxidizable substances following a <1 litre water flush.

Diamete **Outlet Connection** Options 50 mm 96 0.6 µm Female Luer Lock Female Luer Lock Pharmaceutical Ν Non-sterile S Standard Stepped Hosebarb 01 1.0 µm Stepped Hosebarb G 03 3.0 µm 05 5.0 µm 7.0 μm 10.0 μm 07 10 15.0 μm 20.0 μm 15 20 25 40 55 25.0 µm 40.0 μm 55.0 μm 75 75.0 µm Parker domnick hunter has a continuous policy of product development and although the Company reserves the right to change specifications, it attempts to keep customers informed of any alterations. This publication is for general information only and customers are requested to contact our Process Filtration Sales Department for detailed information and advice on a products suitability for specific applications. All products are sold subject to the company's Standard conditions of sale.

Ordering Information

Cartridges PCPP B* dh DOE В* А* 2.5 (65 mm) 96 0.6 µm Pharmaceutical EPDM 1.0 µm RF / 226 Bayonet [125 mm] PTFE Encapsulated Silicone 5 01 C K (125 mm) 03 3.0 µm D Fin / 222 Silicone 5.0 μm 7.0 μm Flat Top / 222 Recess / 222 10 (250 mm) 05 Viton (500 mm) 07 G 20 10.0 μm 15.0 μm 3 30 (750 mm) 10 Н UF Retrofit 40 (1000 mm) SOE (no o-rina) 4 20 20.0 µm 1 * dh DOE * Supplied in packs of 3. 25.0 μm 40.0 μm Internal 213 25 Ν 40 R BF / 222 Bayonet Ratings based on efficiencies of efficiencies of or = 99.98% using nternal test procedu 55.0 μm 75.0 μm 55 75 . 100* 100.0 μm SUPU18 based on ASTM F795-88 1993 TRUESEAL Demi Stub *Not available with A and B Size formats Demi A & B Std * EPDM kets supplied as standard **MURUS** Capsules **PLPP** (Nominal) Grade Tri-Clamp EPDM Tri-Clamp Pharmaceutical 0.6 µm Ν Non-sterile In-Line 96 Δ F 11/, Tri-Clamp 11/, Tri-Clamp 10 (250 mm) 01 1.0 µm В В Pre-sterilized T-Port PTFE M** Hosebarb 20 (500 mm) 03 3.0 µm Hosebarb γ (>25 kGy) Streamline Encapsulated 5.0 µm Tri-Clamp Tri-Clamp 30 (750 mm) 05 Valves Silicone ٥7 7.0 µm Н 1/," Hosebarb н 1/." Hosebarb Silicone lable with a 10.0 µm Viton ** Not available with a 1" or 1.5" Tri-Clamp 15 . 15.0 µm 20 20.0 µm 25.0 µm Ratings based on efficiencies of > or = 99.98% using 40 40.0 µm 55.0 µm 55 75 75.0 um al test pr SOP018 based on ASTM F795-88 1993 100 100.0 µm **DEMICAP** Capsules PEPP inal) Outlet Pack N Acc 4.4 5.5 (113 mm) (140 mm) 96 0.6 μm 1.0 μm Tri-Clamp " NPT Male Tri-Clamp ¨NPT Male Pharmaceutical Ν Non-sterile Pack of 3 FB Filling Bell D 3 В 01 N Pre-sterilized Ν А 7.9 (200 mm) 03 3.0 µm Н Hosebarb н Hosebarb γ (>25 kGy) E & B-Size G & H conr 5.0 µm Stepped Hosebarb Stepped Hosebarb 07 7.0 µm M 1/4 NPT Male M NPT Male Walther QC 10.0 µm Q Q Walther QC Grommel / QC 3/8" NPT Female 15.0 µm R Grommel / QC R 20 20.0 µm 3/。" NPT Female V 25.0 µm 25 40 Ratings based or 40.0 µm 55.0 µm 75 . 75.0 µm Syringe Filters PSPP Pack N 025 25 per box 0 U

PROCLEAR PP Filters

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