

TETPOR LIQUID filters are particularly suitable for sterilization and particulate removal from aggressive chemicals (including acids, bases and solvents) within a wide range of critical processing industries.

The superior performance, strength and durability of TETPOR LIQUID filters stems from the use of a single layer, high security PTFE membrane, which has a high dirt holding capacity due to its high voids volume. This results in low pressure drops and long service life.

High flow rates are achieved due to the optimized pleat pack density and the superior design construction of TETPOR LIQUID filters.

## **Features and Benefits**

- Superior chemical resistance of PTFE membrane combined with polypropylene hardware
- Integrity tested prior to despatch
- Validated to current ASTM F838 methodology
- Comprehensive range of end cap configurations for retrofitting

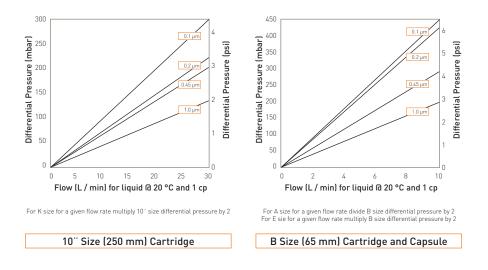
## **TETPOR LIQUID Filters**

- liquid filters
- PTFE



Note: TETPOR is a registered trademark of Parker Hannifin Corporation.

## **Performance Characteristics**



## **Specifications**

### Materials of Construction

Filtration Membrane:	PTFE
Upstream Support:	Polypropylene
Downstream Support:	Polypropylene

Polypropylene

Downstream Support:

#### Filter Cartridges

- Inner Support Core:
- Outer Protection Cage: Polypropylene Polypropylene
- End Caps:
- 316L Stainless Steel End Caps Insert:
- \*Not available in B endcap variant Standard o-rings/gaskets: Viton

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:	Silicone								

#### DEMICAP Filter Capsules

Core:	Polypropylene
Sleeve:	Polypropylene
End Caps:	Polypropylene
Capsule Body:	Polypropylene
Capsules Vent Seals:	Silicone
Filling Bell:	Polycarbonate
Svringe Filters	

Polypropylene

Body:

#### **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:

Temp		Max. Forward dP						
°C		(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 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):	0.77 m <sup>2</sup>	(8.28 ft <sup>2</sup> )
K Size:	0.36 m <sup>2</sup>	(3.87 ft <sup>2</sup> )
A Size:	0.25 m <sup>2</sup>	(2.69 ft <sup>2</sup> )
B Size:	0.12 m <sup>2</sup>	(1.29 ft <sup>2</sup> )
E Size:	0.06 m <sup>2</sup>	(0.64 ft <sup>2</sup> )
Syringe ø50 mm:	14.50 cm <sup>2</sup>	(2.25 in <sup>2</sup> )

#### Sterilization

	Aut Cycles	oclave Temp	Steam-in-Place Cycles Temp (30 min.)					
Cartridges	120	142 °C (287.6 °F)	120	142 °C [287.6 °F]				
MURUS	5	130 °C (266 °F)	-	-				
DEMICAP	100	135 °C (275 °F)	-	-				
Syringe	1	130 °C (266 °F)	-	-				

TETPOR LIQUID 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 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 and integrity tested prior to despatch. A sample of each lot is tested to demonstrate conformity to validated claims.

## **Performance Characteristics**

### TOC / Conductivity

The filtrate quality from a 10" (250 mm) TETPOR LIQUID conforms to the requirements of current USP <643> (TOC) and USP <645> (conductivity).

#### Endotoxins

Aqueous extracts from the 10" (250 mm) TETPOR LIQUID 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  $10^{\circ}$  (250 mm) cartridge are <5 mg.

#### Pharmaceutical Validation

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

#### **Oxidizable Substances**

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

#### **Integrity Test Data**

All filters are integrity testable to the following limits when wet with 60 / 40 IPA / Water and using air as the test gas.

Micron Rating		0.1	0.2	0.45	1.0							
Filter Cartridges / MURUS / DEMICAP / Syringe Filters												
Min. Bubble Point	(barg)	1.3	1.0	0.7	-							
	(psig)	18.8	14.5	10.1	-							
Filter Cartridges /	MURUS / DI	EMICAP / Syringe	Filters									
Diffusional Flow	(barg)	1.0	0.8	0.4	-							
Test Pressure	(psig)	14.5	11.6	5.8	-							
Filter Cartridges /	MURUS / DI	EMICAP / Syringe	Filters									
Max. Diffusional Fl	ow (10)	27.0	18.0	18.0	-							
(ml / min)	[K]	12.7	8.5	8.5	-							
	[A]	9.0	6.0	6.0	-							
	(B)	4.5	3.0	3.0	-							
	(E)	2.3	1.5	1.5	-							

### **Retention Characteristics**

TETPOR LIQUID filter cartridges are validated by bacterial challenge testing with *Brevundimonas diminuta* to current ASTM F838 methodology (10<sup>7</sup> organisms / cm<sup>2</sup> EFA minimum) with typical in-house challenge levels being 10<sup>11</sup> organisms per 10<sup>°°</sup> (250 mm) filter cartridge.

# **Ordering Information**

Cartridges

ZC	MT	-	- [													
Code B* A* K 1 2 3	2.5" 5" 5" 10" 20"		Code 010 020 045 100	Micron 0.1 μm 0.2 μm 0.45 μm 1.0 μm	Code B* C D E G R	h DOE BF / 226 Bayonet Fin / 222 Flat Top / 222 Recess / 222 BF / 222 Bayonet	Code L P S	Variant Liquid Pharmaceutical Steam Sterilizable	E P S V*	O-rings EPDM PTFE Encapsulated Silicone Viton						
4 * Supplied	40 <sup>°°</sup> (	1000 mm) t			SK T Y Z * EPDM Note: V	Endcap (Demi) Retrofit TRUESEAL Demi Stub Demi A & B Std gaskets suppleid as standard fiton suppleid as standard			specity	the V code.						
ZL	MT	apsules	- [			ther endcap options.			-						-	
Code K 1 2 3	5" 10" 20"	(250 mm)	Code 010 020 045 100	Micron 0.1 μm 0.2 μm 0.45 μm 1.0 μm	A B D T	Inlet Connection <sup>3</sup> / <sup>4</sup> " Tri-Clamp 1 <sup>1</sup> / <sub>2</sub> " Tri-Clamp 1" Hosebarb 1" Tri-Clamp	Cod A B D T	e Outlet Connectio <sup>3/4</sup> " Tri-Clamp 1 <sup>1</sup> / <sub>2</sub> " Tri-Clamp 1" Hosebarb 1" Tri-Clamp	n Co	Ide Variant	Code Gr	ade n-sterile	L T*	Design In-Line T-Port vailable with a Clamp	E S* V	e O-rings EPDM Silicone Viton hdard without to specify the le

#### DEMICAP Capsules

ZE	MT			-												
Code	Length	n (Nominal)	Code	Micron	Code	Inlet Connection	Code	Outlet Connection	Code	Variant	Code	Grade	Code	Pack N°	Code	Accessory
E	4.4"	(113 mm)	010	0.1 µm	Т	1" Tri-Clamp	Т	1" Tri-Clamp	Ρ	Pharmaceutical	Ν	Non-Sterile	3	Pack of 3	FB	Filling Bell
В	5.5"	(140 mm)	020	0.2 µm	N	1/2" NPT Male	N	1/2" NPT Male								
A	7.9"	(200 mm)	045	0.45 µm	Н	1/2" Hosebarb	Н	1/2" Hosebarb							G & H co	onnections only
			100	1.0 µm	G	Stepped Hosebarb	G	Stepped Hosebarb								
					M	1/, NPT Male	M	1/, NPT Male								
					Q	Walther QC	Q	Walther QC								
					R	Grommel / QC	R	Grommel / QC								
					V	3/。" NPT Female	V	³/。" NPT Female								

#### Syringe Filters ZSMT Female Luer Lock Stepped Hosebarb 050 50 mm 020 0.2 µm F F Female Luer Lock Pharmaceutical N Non-sterile S Standard 025 25 per box G G Stepped Hosebarb

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.