



aerospace
climate control
electromechanical
filtration
fluid & gas handling
hydraulics
pneumatics
process control
sealing & shielding





Process Filtration

A guide to products and services







Parker Hannifin Ltd
Parker domnick hunter - Process Division

phone +44 (0)191 4105121

fax +44 (0)191 4105312

email: dhprocess@parker.com www.parker.com/processfiltration

Contents

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		Ge	neral
Process filtration	6-7	DEMICAP options	152-153
Quality & control	8-9	MURUS and syringe options	154
Innovation	10-11	Installation and operating guidelines	155
Technical support	12-13	Conversion tables	156-157
A scientific approach	14-15	Chemical compatibility	158-161
Dedicated product range	16-17	Glossary of terms used in filtration	162-165
e-Learning & training	18-19	Compressed air treatment	166
Filter discs	148-149	Gas generation	166
Endcap styles	150-151	Parker motion & control technologies	167

				Air / Ga	S
HIGH FLOW PREPOR GFA		 			
PEPLYN AIR BIO-X II	MEMER	 	OR II Vent Autoc	lave ZTA	38-39
HIGH FLOW BIO-X HIGH FLOW BIO-X Vent Autocl		 HIGH FLOW TETP	OR H.T	ZCHT	40-41

	Steam filte

ZCSS.ZCHS......44-47

			Liquid filte	rs
PROSPUN	PRSC_PRST_PRSA50-51	PREPOR GF	ZCGF.ZEGF.ZSGF	62-63
PROPLEAT PP	PRPP 52-53	PREPOR GP	ZCGP.ZEGP	64-65
PROSTEEL A	ZCCF.ZCMF54-55	PREPOR PES	ZCPS.ZEPS	66-67
PROSTEEL N	ZCCM.ZCPM56-57	TETPOR PLUS	ZCTP	68-69
PEPLYN NE	ZCNE.ZENE 58-59	CARBOFLOW MX		70-71
PEPLYN PLUS	ZCPP ZEPP ZSPP 60-61			

			Beverage filters
PEPLYN HD	PHD 74-75	BEVPOR PS	BPS 88-89
	PHA76-77	22 0	BPH 90-91
PREPOR GF	PFD78-79	BEVPOR PT	BPT92-93
PREPOR GP	PGP 80-81	BEVPOR PW	BPW94-95
PREPOR PP	PPP 82-83	BEVPOR MS	BMS96-97
CRYPTOCLEAR PLUS	ZCCP.ZECP84-85	BEVPOR MT	BMT98-99
CRYPTOCLEAR PES	ZCCS ZECS 86-87	BEVPOR MH	BMH 100-101

		Pn.	armaceutical filters
PROCLEAR GE	PCGE PLGE PEGE 7SGE 104-107	PROPOR HC	7CHC 7LHC 7FHC HSHC 124-127
	PCGP PLGP PEGP ZSGP 108-111		zclr zllr zelr zslr 128-131
PROCLEAR PP	PCPP PLPP PEPP ZSPP 112-115	TETPOR HP	ZCMT'HP' 132-135
PROPOR BR	ZCBR ZLBR ZEBR ZSBR 116-119	TETPOR LIQUID	ZCMT ZLMT ZEMT ZSMT 132-135
PROPOR SG	ZCSG ZLSG ZESG ZSSG 120-123		

Housings

136-139

		Integrity	test equipment
VALAIRDATA II	WVA 142-143	BEVCHECK	WBC 146-147
PORECHECK IV	WPK 144-145	BEVCHECK PLUS	

Parker domnick hunter has a continuous policy of product development and although the Company reserves the right to change specification, 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 product's suitability for specific applications. All products are sold subject to the company's Standard conditions of sale.

4

A dedicated housing range....

Process filtration

Adding value to your business





Parker domnick hunter specializes in the manufacture and supply of high quality products for the clarification, stabilization and sterilization of liquids and gases, providing full scaleability from membrane flat stock to multi element filter systems. Each filter has been specifically developed to meet industry applications and requirements.

As a company it is our goal to deliver innovative quality products on time while responding to the needs of the end user with premier customer service. We know our success is only possible through increasing our customers' productivity and profitability.

Parker domnick hunter manufacture products in the most efficient, effective and environmentally conscious way building on a culture of continuous improvement.

With nearly 50 years filtration experience in markets such as pharmaceutical, beverage and water treatment we have developed innovative and cost-effective solutions that will add value to your manufacturing process, providing reliable products and services that meet or exceed your expectations.

Our worldwide assistance extends to on-site evaluations, design, manufacture, validation, quality control and ongoing support long after the filters are installed.

In 2005 domnick hunter became part of the Parker Hannifin Corporation. Parker Hannifin is the world's leading diversified manufacturer of motion and control technologies and systems, providing precision-engineered solutions for a wide variety of commercial, mobile, industrial and aerospace markets. The company employs more than 50,000 people in more than 50 countries around the world.

- Continued investment in research & technology
- Application driven approach to new products
- Market specific experience leading to tailored solutions
- Global network providing technical, service and sales support
- Excellent reputation gained through working with some of the world's leading companies
- Highly skilled and trained employees





Quality & control

At the forefront of manufacturing excellence



Parker domnick hunter's commitment to leading quality standards in the filtration industry led to us being the first UK based filter company to achieve BS 5750 Pt 1 in 1984 and then BS EN ISO 14001 in 2001. The company is now certified to current version of ISO9001, ISO 13485 and is again leading the way through the implementation of a new application guide PS9100 in 2007.

In support of our on going commitment to quality, Parker domnick hunter has recently completed a £5 Million investment programme to upgrade and increase capacity at our Birtley, UK manufacturing facility. As well as investing in the latest clean room and custom manufacturing technologies, Parker domnick hunter has invested in key lean and six sigma initiatives.

Our focus on the selection of materials in accordance with current regulations such as FDA CFR's, cGMP guidelines and specifications from our Scientists, Engineers and validation experts, together with the use of validated manufacturing and test methodologies ensures high batch-to-batch reproducibility.

A controlled approach

- Both lot number and serial number are recorded for all products providing complete traceability back to base materials
- Products, processes and software are validated at regular intervals
- Integration of productivity, product quality and employee safety into the design and construction of facilities and equipment
- Clean room environment used for all manufacturing operations
- Extensive supplier quality assurance program in place
- Regular process audits conducted by trained auditors from across the business
- Extensive customer audits completed





Innovation

Putting your future needs at the forefront of product development



Parker domnick hunter understands the need to be innovative and deliver real solutions to customer problems. As a company we are always striving to create a culture that will achieve this goal, both through individual team creativity and measured risk taking.

Project teams with members from technical, marketing, manufacturing and procurement functions are necessary for the success of this process. Working closely with our customers has enabled us to design innovative products with value-added benefits.

People are vital to this process and Parker domnick hunter recognizes and supports the need for continuous learning to ensure that its employees have the skills to meet the demands of the changing world we live in.

Winovation

Parker Hannifin has developed an NPD system called Winovation, focusing on long term development of products that will grow our business together.

"Winovation, creates value by determining customer needs and developing products that meet those needs".

- Focus on value proposition
- Unique customer benefits
- Provide a differentiated solution
- An effective discovery stage to generate great ideas
- Accountable and empowered cross functional teams
- Dedicated resource
- Strong market and voice of the customer input
 - Products that are linked to customer goals and initiatives

A forward thinking team provide:

- Introduction of new materials
- Sustained engineering
- Rapid response team
- Engineer existing products to meet demands of new applications
- Development to meet ever changing industry regulations
- Joint engineering projects, combining expertise
- Cross fertilization of ideas with industry leaders
- Cost reduction exercises
- Increased throughputs and lifetime as your business grows • New products that can set new
- industry standards Helping to establish industry best
- practice
- Provide solutions to application driven
- Maximize value and user friendliness of products
- Joint projects with leading universities and institutions
- Access to Parker design and development global resource







Technical support

Dedicated team committed to improving the efficiency of your filtration process



Parker domnick hunter has a multi-disciplinary team of Scientists and Engineers committed to the technical support of our products around the world, providing pro-active practical support in all areas. The aim is to improve economy of filter use and to improve product yield and quality. We understand the practical needs within the process. If system performance is found to be out of specification, or showing deviation from the norm, you can count on active support on-site to identify and resolve problems.

A process audit is an excellent way of identifying and addressing the main risks that may compromize the quality of your production process. From utilities through to your aseptic filling line we can help identify improvements and advise on areas such as applicable products, system layouts, steam sterilization and integrity testing.

System design and implementation
A full operationally qualified filter system
can be implemented using sample and
used cartridge analysis from laboratory
and pilot scale investigations. This can
include the specification for a fully
automated filter system design. This
allows the filter user to have the difficult
task of commissioning a filter system
shared and facilitated through the Parker
domnick hunter team of process experts.

- Filter system audits to optimize system performance
- Contract integrity testing
- Practical laboratory scale testing for continuous process improvements
- Sample and used cartridge analysis to aid in filter system design
- Process simulation
- Chemical compatibility
- Microbial analysis
- Customer specific validation strategy and protocol
- Remote monitoring of system performance

Existing system optimization

Where a process is altered through increased operational demand, e.g. through extension of a production campaign, higher production volumes, an increased number of product changes or a more rigorous sanitization / sterilization regime, Parker domnick hunter offer support to ensure the system remains appropriate for these changed process demands.

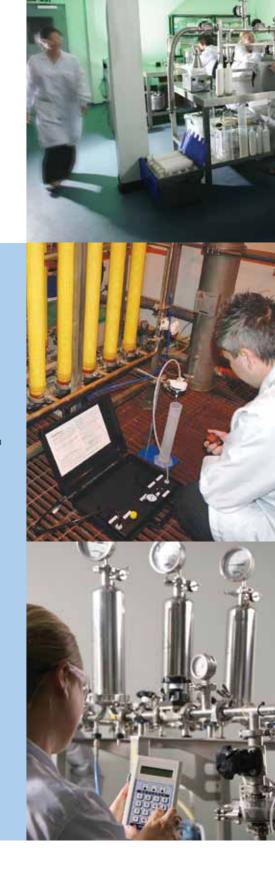
Training

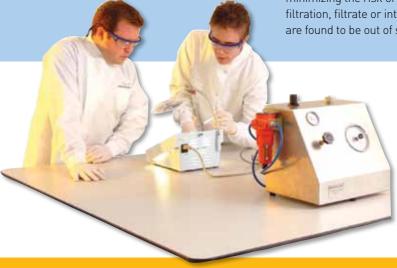
Specialists from across our business can provide training at our state-of-theart facilities or at your own site, which includes:

- Filtration theory and practice
- Integrity testing and validation
- SIP, CIP and compatibility testing

Fault diagnosis

Often filtration is a critical step or control point within a process. Therefore, when finished product quality is not achieved, the filter is often the first point of call. The Parker domnick hunter TSG group can provide a reactive service to enable rapid 'root cause' analysis and assist in minimizing the risk of recurrence where filtration, filtrate or integrity test values are found to be out of specification.





A scientific approach

Consistent performance put to the test



Parker domnick hunter employ a combination of Engineers and Scientists with advanced degrees in a wide range of fields including bioscience, biotechnology, microbiology and chemistry.

Using state-of-the-art equipment and facilities, the Parker Laboratory Services Group are equipped to become a valued partner in your validation process.

Providing step-by-step validation support to the customer by developing and executing process-specific protocols based on your application.

The Laboratory Services Group (LSG) at Parker domnick hunter provides documented evidence that gives the customer a high degree of assurance that our filters will consistently produce a level of performance that meets its predetermined specifications and quality attributes.

Quality control testing

- Water testing: TOC, endotoxin, bioburden, pH and conductivity
- Environmental monitoring, microbial assay
- Filter characteristics, visual bubble point, liquid and air flow rates, porometry analysis, water intrusion
- Quality control testing of incoming filter materials including bacterial challenge to ASTM 838-05 for sterilizing grade products
- Lot release of finished products and rinse water / effluent analysis

Customer validation

- A bespoke service offering a full validation package to support sterile filtration steps
- Includes protocol and experimental design, technical support and production of an audit reference of each filter and filtered product
- Establish integrity test parameters
- Develop customer specific validation strategies
- Examination of filter extractables
- Documented assurance

Scientific research

- Microbial assays standard and bespoke
- Protein binding analysis via SDS PAGE and gel imagery
- Process simulation and scale-up support
- New product design and optimization
- Process characterization and filtration analysis





Dedicated product range

Choice and flexibility to suit your application



Parker domnick hunter manufacture a range of microfiltration cartridges for liquid and gas applications that utilize the latest production techniques, combining the most suitable membranes and filtration media with the latest easy to use formats.

All of Parker domnick hunter's filters meet strict validation guidelines that provide a high degree of assurance that they will consistently achieve a high level of performance in a given application and meet the needs of the industry that they have been specifically designed for.

- Wide choice of filtration media and filter formats
- Technical and validation support
- Industry specific designed filters
- Fully retrofitable range of products
- Manufactured in state-of-the-art facilities

Scaleability provides flexibility

The ability to scale up from small area discs to process scale systems with minimal revalidation is paramount.

Parker domnick hunter provides a wide range of filter formats to ensure that the transition from pilot-scale through to full production is as smooth as possible.

Single use systems

Disposable systems can eliminate cleaning validation, reduce capital costs, minimize health & safety risks and lower the chance of product contamination.

Single use systems also provide a more convenient way of processing a product.

Close working relationships

Parker domnick hunter have partnered engineering companies on large-scale projects around the world that require filtration expertise and a capability to fabricate large-scale systems.







Understanding the principles of filtration

e-learning and training at your own speed



What is e-Learning?

e-Learning is an effective learning process created by interaction with digitally-delivered content, learning support and services. It uses a combination of text, voice-over and moving images to explain ideas and concepts.

Why has Parker domnick hunter developed e-Learning?
Parker operates in more than 50 countries and employs more than 50,000 people worldwide. e-Learning enables us to reach all the relevant people with a consistent and a clear message. e-Learning content has been developed in-house and we believe we have a unique and innovative package which provides world-class filtration training. We are now enabling our customers to access the same learning.

What courses are available?

We can provide access to the Certificate in Filtration Technology course. This course consists of 9 modules of e-learning. It is intended as an introductory level course which looks mainly at the management of compressed air. Two further modules cover sterile air filtration and the filtration of liquids. Taken together they provide an excellent introduction to the world of filtration.

Each module has its own test and these test results are retained by the Learning Management System for later review.

Further Parker domnick hunter Certificate courses include a Certificate in Compressed Air Quality Management which consists of three modules covering ISO 8573.1 Air Quality standards, dryers and compressed air filter solutions. How can I access e-learning?
The e-Learning is held on a LMS
(Learning Management System) at
www.dhelearning.com.

To access the e-Learning you will need a user name and password, supplied by Parker domnick hunter.

How long will the course take to complete?
Learners are able to complete the
course at their own pace and can fit the
course around the demands of a busy
working day. The time taken to complete
the course varies from person to person
but for most people the Certificate in
Filtration Technology represents 20 hours
of study.

How do I find out more?
It is possible to demonstrate the e-Learning package (and some of the other e-Learning materials) to you and your learning and development specialists. We firmly believe that in-house e-Learning represents world-class learning which is not available elsewhere.

For further information, email: FGE.training@parker.com





Air / Gas filters



Filtration of air and gas

There is an increasing demand in the food & beverage industry for sterile air / gas which can be used in applications such as line clearing, storage tanks, machines and the venting of gas from storage tanks. It is essential that whenever gases come into contact with product or process equipment, any microbiological contamination is removed to guarantee product safety, uniform quality and extended shelf life. Parker domnick hunter provide a range of class-leading products with a proven track record.

Filters include

- PTFE impregnated glass microfibre

 (PTFF / GF)
- Polypropylene (PP)
- Glass microfibre (GF)
- Polytetrafluoroethylene (PTFE)

TETPOR filters from Parker domnick hunter utilize a PTFE membrane to provide competitive performance and value in sterile air applications. Also available in high temperature formats.

HIGH FLOW BIO-X - High flow rates and high dirt holding capacity make
HIGH FLOW BIO-X the filter of choice within the fermentation and beverage industries. A combination of PTFE and glass fibre media provides a product with high voids volume and added strength giving unrivalled performance in applications such as the provision of sterile gas to filling machines.





HIGH FLOW PREPOR GFA Filter Cartridges

- air / gas filters
- · glass microfibre



HIGH FLOW PREPOR GFA is a high capacity glass fibre prefilter specifically designed for the removal of bulk particulate from compressed air and gases.

It is used extensively for prefiltration duties in dry compressed air systems and provides excellent protection for final sterile

HIGH FLOW PREPOR GFA utilizes pleated glass fibre filter media encased within an upstream and downstream expanded polypropylene mesh filter support. The pleat pack is supported by an inner stainless steel core and outer heat stabilized polypropylene cage, heat bonded to heat stabilized polypropylene end caps.

The combination of high voids volume filter media and pleated construction results in a filter cartridge with exceptional dirt holding capacity, able to operate at very low differential pressures.

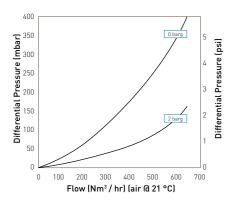
Features and Benefits

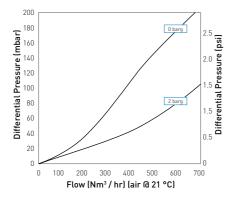
- High surface area and voids volume filter media
- Exceptionally high flow rates with low pressure drops
- Reliable efficient protection of final sterilization filters
- Heat stabilized componentry to allow operation at elevated temperatures



Note: PREPOR is a registered trademark of Parker domnick hunter

Performance Characteristics





Cartridge flow rates 10" Size (250 mm)

Cartridge flow rates 20" Size (500 mm)

HIGH FLOW PREPOR GFA Filter Cartridges

Specifications

Materials of Construction

■ Filtration Media: Glass Microfibre ■ Upstream Support: Polypropylene Downstream Support: Polypropylene ■ Inner Support Core: 316L Stainless Steel Outer Protection Cage: Polypropylene ■ End Caps: Polypropylene ■ End Cap Insert: Stainless Steel

Food and Biological Safety

■ Standard o-rings/gaskets: Silicone

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

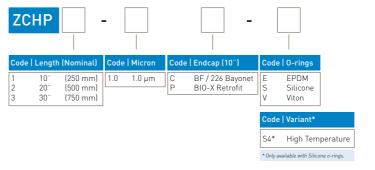
Recommended Operating Conditions

The maximum differential pressure in direction of flow (outside to in) is 3.5 barg (50.76 psig) at 20 °C (68 °F).

The maximum recommended continuous operating temperature is 70 °C (158 °F). Note: For temperatures from 70 °C (158 °F) to 100 °C (212 °F) a special product with polyester supports is available.

Effective Filtration Area (EFA)

10" (250 mm) 0.48 m² (5.16 ft²)



PEPLYN AIR Filter Cartridges

- air / gas filters
- meltblown polypropylene



PEPLYN AIR filter cartridges have been specifically designed to guarantee removal of particulate from gas streams.

They can be used to protect sterilizing grade filters in pressurized systems or in exhaust gas vent applications.

PEPLYN AIR is particularly suitable for:

- Inlet gas in the fermentation industry as protection to sterilizing grade filters where polypropylene media is preferred
- As protection to sterilizing grade filters in exhaust gas systems
- Vent applications
- Systems where high particulate loading is expected

PEPLYN AIR has the ability to be steam sterilized and has a broad range of chemical compatibility

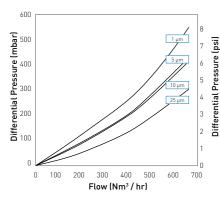
Features and Benefits

- Cost-effective prefiltration
- Absolute micron rating range from 1.0 - 25 micron
- High flow rates and long life
- Steam sterilizable
- Graded density for excellent particle retention
- No release of particles even during system pressure fluctuations



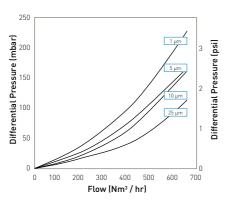
Note: PEPLYN is a registered trademark of Parker domnick hunter

Performance Characteristics



Flow rates for other sizes available upon request

Cartridge flow rates @ 0 barg 10" Size (250 mm)



Flow rates for other sizes available upon request

Cartridge flow rates @ 2 barg 10" Size (250 mm)

PEPLYN AIR Filter Cartridges

Specifications

Materials of Construction

■ Filtration Media: Polypropylene ■ Upstream Support: Polypropylene ■ Downstream Support: Polypropylene ■ Inner Support Core: 316L Stainless Steel Outer Protection Cage: Polypropylene ■ End Caps: Polypropylene

Food and Biological Safety

■ Standard o-rings/gaskets: Silicone

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions The maximum differential pressure in

direction of flow (outside to in) is 3.5 barg (50.76 psig) at 20 °C (68 °F).

The maximum recommended continuous operating temperature is 50 °C (122 °F).

Effective Filtration Area (EFA)*

10" (250 mm) 0.49 m² (5.27 ft²) *Varies with micron rating

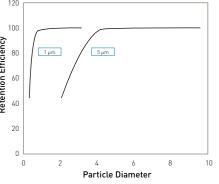
Cleaning and Sterilization

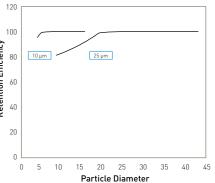
PEPLYN AIR cartridges can be repeatedly in situ steam sterilized or autoclaved up to 142 °C (287.6 °F).

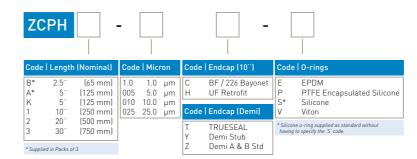
Determination of Micron Ratings

Particle removal efficiencies of PEPLYN AIR cartridges have been determined independently by challenging with a cut silica test dust, generated by BUS1701 dust injector used in conjunction with laser particle counters.

Micron Efficiency Ratings









BIO-X II air sterilization filter cartridges utilize a borosilicate microfibre media. This media has proven to be particularly effective in the removal of sub-micron particles as small as 0.01 micron, therefore ensuring the removal of all microorganisms, including bacteria and viruses.

The media is sandwiched between Nomex support materials to provide additional strength and prevent media migration. This is rigidly held between stainless steel support cylinders and finally encapsulated into stainless steel end caps. The result is a filter cartridge with the exceptional strength and efficiency necessary for absolute security in the most testing of applications.

BIO-X II filter cartridges are particularly suitable for the increasing number of high temperature applications. They also fulfil the sterile compressed air and gas requirements of the dairy, brewery and food processing industries.

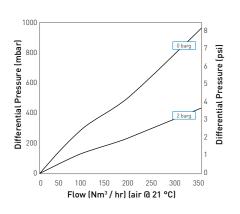
Features and Benefits

- Nomex support materials for high temperature operation
- Robust stainless steel construction
- High temperature operation 200 °C (392 °F)
- 100% integrity tested prior to despatch
- Unique serial number for full traceability
- Fully validated by aerosol bacterial challenge



Note: BIO-X is a registered trademark of Parker domnick hunter

Performance Characteristics



ME10AB7SRH Cartridge

Specifications

Materials of Construction

■ Filtration Media: Borosilicate Microfibre ■ Upstream Support: Nomex* ■ Downstream Support: Nomex* ■ Inner Support Core: Stainless Steel

Outer Protection Cage: Stainless Steel ■ End Caps: Stainless Steel ■ Encapsulant: Epoxy Resin

■ Standard o-rings / gaskets: Silicone *Nomex is a registered trademark of E.I. du Pont de Nemours and Co. Inc.

Recommended Operating Conditions

The maximum differential pressure is 700 mbar for economical element change.

Maximum Continuous Inlet Air Temperature

200 °C (392 °F) Intermittent 170 °C (388 °F) Continuous

Sterilization

BIO-X II filter elements can withstand a maximum of 100 in-line sterilization cycles with purified saturated steam. In-line sterilization 142 °C (287.6 °F), 2.8 barg (40.7 psig) for 30 minutes.

Integrity Test Data

All cartridges are integrity tested prior to despatch by the aerosol challenge test method using the Parker domnick hunter VALAIRDATA II.

Validation

The BIO-X II range of cartridges have been fully validated by bacterial challenge of aerosolized Brevundimonas diminuta.

BIO-X II Filter Cartridges

Ordering Information

Cartridges

Element Code	Cartridge Length		Endcap Location	
MER-BZ	2.5"	(65 mm)	Demi A & B Std	(Z)
MER-AZ	5"	(125 mm)	Demi A & B Std	(Z)
ME10-AB7SRH	10"	(250 mm)	BS226	(C)
ME20.AB7-SRH	20"	(500 mm)	BS226	(C)
ME30.AB7-SRH	30	(750 mm)	BS226	(C)

BIO-X II Retrofit Cartridge Part Numbers

Parker domnick hunter Cartridge	ME3/1	ME3/1.5	ME4/1.5	ME4/2.5	ME5/2.5	ME5/3	ME10/3	ME15/3	ME20/3	ME30/3	ME30/5		
Retrofit Cartridge	SRF3/1	SRF3/1.5	SRF4/1.5	SRF4/2.5	SRF5/2.5	SRF5/3	SRF10/3	SRF15/3	SRF20/3	SRF30/3	SRF30/5		
Parker domnick hunter Cartridge	MER2/10	MER3/10	MER4/20	MER5/20	MER5/25	MER7/25	MER7/30	MER10/30	MER15/30	MER20/30	MER30/30	MER30/50	
Retrofit Cartridge	SRF02/10	SRF03/10	SRF04/20	SR05/20	SRF05/25	SRF07/25	SRF07/30	SRF10/30	SRF15/30	SRF20/30	SRF30/30	SRF30/50	
Parker domnick hunter Cartridge	ME2/10	ME3/10	ME4/20	ME5/20	ME5/25	ME7/25	ME7/30	ME10/30	ME15/30	ME20/30	ME30/30	ME30/50	
Retrofit Cartridge	P-SRF02/10	P-SRF03/10	P-SRF04/20	P-SRF05/20	P-SRF05/25	P-SRF07/25	P-SRF07/30	P-SRF10/30	P-SRF15/30	P-SRF20/30	P-SRF30/30	P-SRF30/50	

- air / gas filters
- PTFE impregnated borosilicate microfibre



HIGH FLOW BIO-X combines proven depth filter technology and a pleated construction to provide retention down to 0.01 micron in gas.

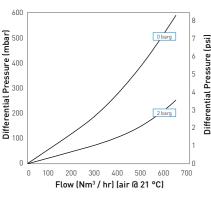
Flow rates typically 2-3 times that of membrane filters make HIGH FLOW BIO-X the filter that can dramatically reduce cartridge usage and installation size within the fermentation, food and beverage industries.

The specially developed PTFE impregnation process imparts greater strength and permanent hydrophobicity to the borosilicate microfibre media. This leads to excellent performance in applications such as the provision of sterile gas in filling machines.

Features and Benefits

- 94% voids volume PTFE impregnated microfibre
- Wide bore cartridge construction to maximize flow rate
- Stainless steel inner core
- Exceptionally high flow rates with low pressure drops
- Fully validated by aerosolized bacterial and viral challenge

Performance Characteristics



Flow rates for other sizes available upon request

10" Size (250 mm) Cartridge

10 20 30 40 50 60 70 80 90 Flow (Nm3 / hr) (air @ 21 °C)

Flow rates for other sizes available upon request

A Size (125 mm) Cartridge

Note: BIO-X is a registered trademark of Parker domnick hunter

Specifications

Materials of Construction

■ Filtration Media: PTFE Impregnated Borosilicate

Microfibre

■ Upstream Support: Polypropylene ■ Downstream Support: Polypropylene 316L Stainless Steel ■ Inner Support Core:

Outer Protection Cage: Polypropylene Fnd Caps: Polypropylene ■ End Cap Insert: 316L Stainless Steel

■ Standard o-rings/gaskets: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

The maximum differential pressure in direction of flow (outside to in) is 3.5 barg (50.76 psig) at 70 °C (158 °F).

The maximum recommended continuous operating temperature is 70 °C (158 °F).

Effective Filtration Area (EFA)

10" (250 mm) 0.38 m² (4.09 ft²)

Sterilization

HIGH FLOW BIO-X cartridges can be in situ steam sterilized or autoclaved up to 142 °C (287.6 °F) for a maximum of 150 steam cycles.

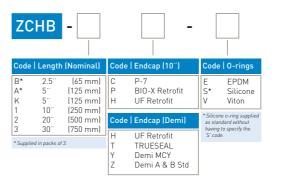
Retention Characteristics

The HIGH FLOW BIO-X range of cartridges has been fully validated by aerosol bacterial challenge levels of 10¹² Brevundimonas diminuta per 10" (250 mm) filter cartridge. Independent test work also shows full retention to MS-2 Coliphage.

Integrity Test Data

HIGH FLOW BIO-X Filter Cartridges

All cartridges are integrity tested prior to despatch by the aerosol challenge test method using the Parker domnick hunter VALAIRDATA II.



- air / gas filters
- PTFE impregnated borosilicate glass microfibre



HIGH FLOW BIO-X Vent Autoclave filter cartridges are designed for critical applications where sterile air is required to break the vacuum formed by the condensation of steam inside the autoclave chamber.

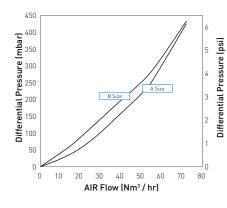
At the heart of the HIGH FLOW BIO-X Vent Autoclave filter cartridge is the latest inherently hydrophobic PTFE impregnated microfibre. With a voids volume of 94%, this media gives exceptional flow rates when compared to membranes. It will remove all particles down to 0.01 micron therefore ensuring the removal of microorganisms, including bacteria and viruses. The filter cartridges are manufactured using a heat sealed construction and no adhesives or resins are used in fabrication. The result, a product of not only exceptional quality, but also a very cost effective solution for the production of sterile air.

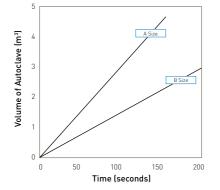
Features and Benefits

- High flow rates
- Hydrophobic filter medium
- Exceeds requirements of HTM10 and EN285
- Detachable prefilter layer
- Exceptional strength
- Repeatedly autoclavable



Performance Characteristics





Cartridge flow rates @ 0 barg

Vacuum break time against autoclave volume

HIGH FLOW BIO-X Vent Autoclave Filter Cartridges

Specifications

Materials of Construction

■ Filtration Media: PTFE Impregnated Glass Microfibre ■ Upstream Support: Polypropylene ■ Downstream Support: Polypropylene ■ Inner Support Core: Polypropylene

Outer Protection Cage: Polypropylene ■ Prefilter Sock: Polyurethane ■ End Caps: Polypropylene ■ Standard gaskets: **EPDM**

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

The maximum differential pressure in direction of flow (outside to in) is 4.5 barg (65.26 psig) at 70 °C (158 °F).

The maximum recommended continuous operating temperature is 70 °C (158 °F).

Effective Filtration Area (EFA)

5" (125 mm) 0.2 m² (2.3 ft²)

Sterilization

HIGH FLOW BIO-X Vent Autoclave filter cartridges can be repeatedly autoclaved up to 142 °C (288 °F) for a maximum of 150 cycles.

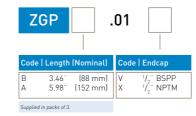
Note: Remove prefilter layer before steaming.

Retention Characteristics

The HIGH FLOW BIO-X Vent Autoclave range of cartridges has been fully validated by aerosol bacterial challenge levels of >10⁷ Brevundimonas diminuta per cm². Independent test work also shows full retention to MS-2 Coliphage.

Integrity Test Data

All cartridges are integrity tested prior to despatch by the aerosol challenge test method using Parker domnick hunter's VALAIRDATA II.



TETPOR AIR Filter Cartridges

- air / gas filters
- expanded PTFE



TETPOR AIR sterilization filter cartridges offer exceptional filtration performance while providing the highest levels of biosecurity throughout the process industry.

Operating at ambient temperature conditions, TETPOR AIR filter cartridges provide a cost-effective filtration solution. A unique polypropylene prefilter layer extends service life in heavily contaminated environments.

TETPOR AIR filter cartridges also utilize a well-proven inherently hydrophobic expanded PTFE membrane with an absolute removal rating of 0.01 micron for gas applications. This ensures the removal of all airborne bacteria, viruses and bacteriophage.

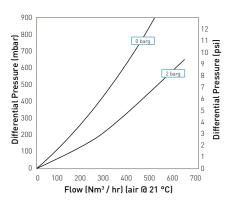
Features and Benefits

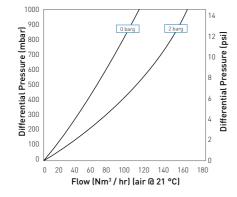
- · Assured biosecurity with absolute rated filtration
- High flow rates with low pressure drops
- High voids volume PTFE membrane
- Steam sterilizable to 142 °C (287.6 °F)
- Unique prefilter layer



Note: TETPOR is a registered trademark of Parker domnick hunter

Performance Characteristics





10" Size (250 mm) Cartridge

B Size (65 mm) Cartridge

Specifications

Materials of Construction

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

Filter Cartridges

Inner Support Core:	Potypropytene
Outer Protection Cage:	Polypropylene
■ End Caps:	Polypropylene
■ End Caps Insert:	316L Stainless Steel
■ Standard o-rings/gaskets:	Silicone

MURUS Disposable Filter Capsules

THORICO DIOPOSODIO I ILLOI	oupoutos
Core:	Polypropylene
■ Sleeve:	Polypropylene
■ Standard o-rings:	Viton
■ Capsule Body:	Polypropylene
■ Capsules Vent Seals:	Silicone

NEMICAP Filter Capsules

Core:	Potypropyter
■ Sleeve:	Polypropyler
■ End Caps:	Polypropyler
■ Capsule Body:	Polypropyler
Canculae Vant Spale:	Silicone

Filling Bell: Polycarbonate

Syringe Filters

■ Body: Polypropylene

Recommended Operating Conditions

Filter Cartridges

Up to 60 °C (140 °F) continuous operating temperature and higher short-term temperatures during CIP to the following limits:

	erature	Max. For	
°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 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):	0.77m^2	(8.28 ft ²)
K Size:	0.36 m^2	(3.87 ft ²)
A Size:	0.25m^2	(2.69 ft ²)
B Size:	0.12m^2	(1.29 ft ²)
E Size:	0.06m^2	(0.64 ft ²)
Syringe ø50 mm:	14.50 cm ²	(2.25 in ²)

Sterilization

	Aut	oclave	Steam-in-Place					
	Cycles	Temp	Cycles (30 min.)	Temp				
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)	-	-				

For detailed operational procedures and advice on cleaning and sterilization, 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

TETPOR AIR Filter Cartridges

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.

32 33

TETPOR AIR Filter Cartridges

Performance Characteristics

TOC / Conductivity

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

Endotoxins

Aqueous extracts from the 10" [250 mm] TETPOR AIR 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 <5 mg.

Pharmaceutical Validation

A full validation guide is available upon request from Laboratory Services Group [LSG].

Oxidizable Substances

TETPOR AIR 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.

our a rage		ssure	Flow	Intru Test Pr	sion	Intrusion	Flow
	(barg) (psig)	(ml / min)			(ml / 10 min)	(µl / 10 min)
E	0.8	11.6	1.5	2.5	36.3	1.3	371
В	0.8	11.6	3.0	2.5	36.3	2.6	742
A	0.8	11.6	6.0	2.5	36.3	5.3	1514
К	0.8	11.6	8.5	2.5	36.3	7.5	2142
10"	0.8	11.6	18.0	2.5	36.3	16.0	4571

Retention Characteristics

TETPOR AIR 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.

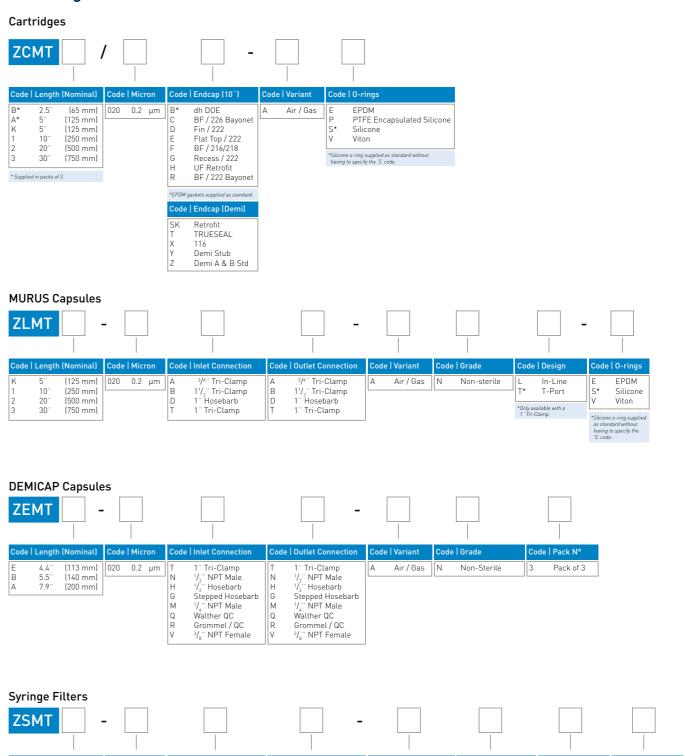
34

TETPOR AIR Filter Cartridges

Ordering Information

020 0.2 μm

Stepped Hosebarb



Stepped Hosebar

Pharmaceutical N

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.

Standard 025 25 per box

HIGH FLOW TETPOR II Filter Cartridges

- air / gas filters
- polytetrafluoroethylene PTFE



HIGH FLOW TETPOR II gas sterilization filters have been developed to benefit from technological advances within the manufacture of PTFE membranes. This new generation of filter sets the standard with an unrivalled combination of efficiency, flow rate and strength.

The HIGH FLOW TETPOR II is validated as a 0.2 micron sterilizing grade filter in liquids through ASTM 838-05 and 0.01 micron in gas through full retention to an aerosol challenge of MS2 phage. This ensures the filter will quarantee the sterility of your process in the worst-case scenario where the filter may be subjected to bulk liquid due to a process problem. Subtle changes to the structure of the PTFE have also resulted in the production of an extremely robust product now validated for 225 steam sterilization cycles @ 142 °C (287.6 °F). The combination of nonwoven supports upstream of the membrane and an expanded net layer downstream has significant benefits. It provides increased protection and service life while guaranteeing zero fibre shedding into the process.

HIGH FLOW TETPOR II is suitable for all sterile gas applications including fermentation inlet and off gas streams, venting, lyophilisers, autoclave vacuum breaks and blow-fill-seal equipment as well as the provision of particle free air within the electronics industry.

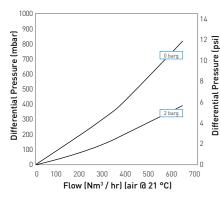
Features and Benefits

- Optimum pleat configuration
- Steam sterilizable up to 225 cycles at 142 °C (287.6 °F)
- Unrivalled flow rates combined with low pressure drops
- Fully validated to ASTM 838-05 for liquid bacterial challenge
- Fully validated to aerosol and viral challenge
- Integrity testable by all methods including water intrusion test



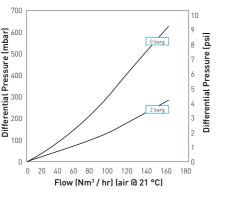
Note: TETPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



Flow rates for other sizes available upon request

10" Size (250 mm) Cartridge



Flow rates for other sizes available upon request

A Size (125 mm) Cartridge

HIGH FLOW TETPOR II Filter Cartridges

Specifications

Materials of Construction

■ Filtration Membrane: Polytetrafluoroethylene ■ Upstream Support: Polypropylene ■ Downstream Support: Polypropylene ■ Inner Support Core: 316L Stainless Steel ■ Outer Protection Cage: Polypropylene ■ End Caps: Polypropylene ■ End Cap Insert: Polysulphone

Silicone

Food and Biological Safety

Standard o-rings:

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

The maximum differential pressure in direction of flow (outside to in) is 3.5 barg (50.76 psig) at 60 °C (140 °F).

The maximum recommended continuous inlet air temperature is 60 °C (140 °F). Note: HIGH FLOW TETPOR II cartridges can be used as WFI vents in heated housings if changed on a 4-6 monthly basis.

Sterilization

HIGH FLOW TETPOR II cartridges can be in situ steam sterilized for up to 225 cycles at 142 °C (287.6 °F).

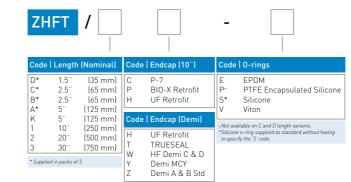
Retention Characteristics

HIGH FLOW TETPOR II cartridges have been fully validated as 0.2 micron sterilizing grade filter cartridges, for compressed air and gas applications. They exceed liquid bacterial challenge levels as recommended by ASTM+. In addition, HIGH FLOW TETPOR II is also validated by aerosol bacterial and MS-2 Coliphage challenge testing. +ASTM American Society for Testing and Materials

Integrity Test Data

All cartridges are integrity tested prior to despatch by the pressure decay and aerosol challenge test methods. Values are for cartridges wetted with 60 / 40 IPA / Water.

Cartridge	Test Pressure		Diffusional Flow	Wat Intru	sion	Water Intrusion	Water Flow
	(bai) (psi)	(ml / min)	Test Pressure (barg) (psig)		(ml / 10 min)	(µl / 10 min)
D	0.8	11.6	0.6	2.5	36.2	N/A	N/A
С	0.8	11.6	1.1	2.5	36.2	N/A	N/A
В	0.8	11.6	2.8	2.5	36.2	2.3	657
Α	0.8	11.6	5.6	2.5	36.2	4.6	1314
K	0.8	11.6	7.70	2.5	36.2	6.4	1828
10"	0.8	11.6	16.50	2.5	36.2	13.5	3857
20"	0.8	11.6	33.00	2.5	36.2	27.0	7714
30	8.0	11.6	49.50	2.5	36.2	40.5	11571



HIGH FLOW TETPOR II Vent Autoclave Filter Cartridges

- air / gas filters
- polytetrafluoroethylene PTFE



HIGH FLOW TETPOR II Vent Autoclave filter cartridges are designed for critical applications where sterile air is required to break the vacuum formed by the condensation of steam inside the autoclave chamber.

At the heart of the HIGH FLOW TETPOR II Vent Autoclave filter cartridge is the latest inherently hydrophobic PTFE membrane. This absolute rated membrane will remove all particles down to 0.01 micron, thus removing airborne bacteria, viruses and bacteriophage.

The filter cartridges are manufactured using a heat sealed construction, thus eliminating the need for adhesives or resins in fabrication. The result is a product of exceptional strength and quality.

• Exceptional strength

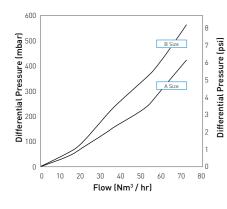
Repeatedly autoclavable

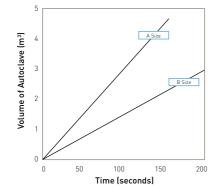
Features and Benefits

- Hydrophobic PTFE membrane
- Fully validated
- Detachable prefilter layer



Performance Characteristics





Cartridge flow rates @ 0 barg

Vacuum break time against autoclave volume

HIGH FLOW TETPOR II Vent Autoclave Filter Cartridges

Specifications

Materials of Construction

Filtration Membrane: Polytetrafluoroethylene Upstream Support: Polypropylene Downstream Support: Polypropylene Inner Support Core: Polypropylene Outer Protection Cage: Polypropylene Prefilter Sock: Polyurethane End Caps: Polypropylene Standard gaskets: EPDM

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

The maximum differential pressure in direction of flow (outside to in) is 3.5 barg (50.76 psig) at 60 °C (140 °F).

The maximum recommended continuous operating temperature is 60 °C (140 °F).

Effective Filtration Area (EFA)

5" (125 mm) 0.3 m² (3.22 ft²)

Sterilization

HIGH FLOW TETPOR II Vent Autoclave filter cartridges can be repeatedly autoclaved up to 142 °C [287.6 °F] for a maximum of 100 cycles.

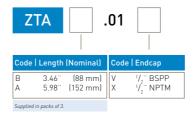
Note: Remove prefilter layer before steaming.

Retention Characteristics

The HIGH FLOW TETPOR II Vent Autoclave range of cartridges has been fully validated by aerosol bacterial challenge levels of >10⁷ Brevundimonas diminuta per cm². Independent test work also shows full retention to MS-2 Coliphage.

Integrity Test Data

All cartridges are integrity tested prior to despatch by the aerosol challenge test method using Parker domnick hunter's VALAIRDATA II.



HF TETPOR H.T. Filter Cartridges

- air / gas filters
- expanded PTFE



HIGH FLOW TETPOR H.T. gas sterilization filter cartridges provide unrivalled performance in process industry applications where continuous cartridge operation of up to 100 °C (212 °F) is a requirement.

Applications include specific biological fermentations which use high inlet air temperatures and heated vent filters on storage tanks whose contents are at elevated temperatures >80 °C (176 °F), e.g. WFI tanks.

HIGH FLOW TETPOR H.T. cartridges utilize a proven inherently hydrophobic, expanded PTFE membrane with an absolute removal rating of 0.01 micron. This ensures the removal of all airborne bacteria, viruses and bacteriophage. Nomex membrane support layers facilitate continuous operation at temperatures up to 100 °C (212 °F).

• Steam sterilizable

drops

to 142 °C (287 °F)

Exceptionally high flow

rates with low pressure

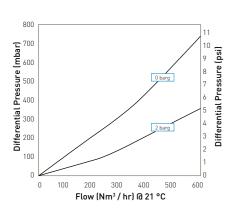
Features and Benefits

- Long service life even at elevated temperatures 100 °C (212 °F)
- Assured biosecurity with absolute rated filtration
- Stainless steel inner core



Note: TETPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



10" Size (250 mm) Cartridge

HIGH FLOW TETPOR H.T. Filter Cartridges

Specifications

Materials of Construction

Filtration Membrane: Expanded PTFEUpstream Support: Nomex*

■ Downstream Support: Nomex*

Inner Support Core: 316L Stainless SteelOuter Protection Cage: Heat Stabilized

Polypropylene

End Caps: Heat Stabilized
Polypropylene

End Cap Insert: Stainless Steel

*Nomex is a registered trademark of E.I. du Pont de Nemours

Silicone

Food and Biological Safety

■ Standard o-rings:

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

The maximum differential pressure in direction of flow (outside to in) is 3.5 barg (50.76 psig) at 100 °C (212 °F).

The maximum recommended continuous operating temperature is 100 °C (212 °F).

Effective Filtration Area (EFA)

10" (250 mm) 0.9 m² (9.8 ft²)

Sterilization

HIGH FLOW TETPOR H.T. cartridges can be in situ steam sterilized for up to 120 cycles at 142 °C (287.6 °F).

Retention Characteristics

HIGH FLOW TETPOR H.T. cartridges have been fully validated as sterilizing grade filter cartridges, for compressed air and gas applications. They exceed liquid bacterial challenge levels as recommended by ASTM+. In addition, HIGH FLOW TETPOR H.T. is further validated by aerosol bacterial challenge testing.

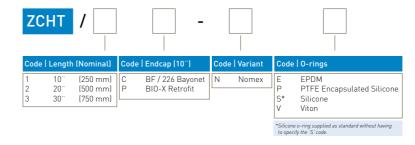
+ASTM American Society for Testing and Materials

Integrity Test Data

All cartridges are integrity tested prior to despatch by the pressure decay and aerosol challenge test methods. Values are for cartridges wetted with 60 / 40 IPA / Water.

Micron Rating		0.2	
Diffusional Flow	(barg)	0.80	
Test Pressure	(psig)	11.6	
Minimum Bubble	(barg)	1.00	
Point	(psig)	14.5	
Max. Diffusional Flo (ml / min)	ow (10")	16.0	

Ordering Information



© +44 (0)191 4105121 ≢=7 dhprocess@barker.com €3 www.barker.com/broce

Steam filters





Filtration of steam

Steam is utilized in many areas of process manufacturing both directly and indirectly coming into contact with product, process lines and equipment. The quality of this steam varies considerably depending on methods of generation, additives, condition of supply pipelines and condensate management. If not treated, poor quality steam that is used to sterilize downstream process filters will lead to premature blockage.

Steam filters from Parker domnick hunter have been specifically designed to protect process equipment and pipework from particulate contamination, extending their overall life. Pleated Steam filters from Parker domnick hunter are designed to provide a culinary grade steam coupled with exceptionally high flow rates. The 1 micron version guarantees steam to 3A.609-03 standard.

Sintered Steam filters from Parker domnick hunter are manufactured from a highly porous sintered stainless steel available in 1 and 25 micron. The 1 micron filter provides culinary grade steam that meets 3A standards. The general purpose 25 micron filter provides protection for membrane filters located downstream in the process.





43

42

STEAM Filter Cartridges

- steam filters
- 316L stainless steel



Steam is an often neglected part of a process, regarded as an add on to a customers liquid or gas filtration needs.

It has however, large specific applications in its own right and should be treated with the same level of importance as air, gas and liquid systems if long filter lifetimes and system cost effectiveness are to be achieved.

The quality of steam used within the food and dairy industries has been raised higher on the agenda in an ever increasing number of companies. Minimum acceptable standards are now being quoted on a more regular basis with particular reference to 'culinary grade' steam. Steam serves several purposes in the food & beverage industry. It is critical that this steam is of a high quality to ensure effective and continuous operation of the process.

Features and Benefits

- 316L stainless steel filter cartridges
- Exceptionally high flow
- Available in culinary grade 1 micron
- High dirt holding capacity
- 'JUMBO' filter configuration ensures maximum utilization of pipework capacity



Which Filter for Which Application?

Process Steam

- · Direct from boiler
- No direct contact with product being manufactured



Applications

- General heating
- Steam jackets
- Bio waste kill systems



Cartridges

Required if steam is used to sterilize liquid and gas cartridge filters



Sintered 25 µm

Use for relatively low flow rates

Culinary Steam (3A Standard 609-03)

- 95% retention of >2 micron particles in the liquid
- Manufactured from 300 series stainless steel
- Any additives to the boiler feed should conform to CFR Title 21, Chapter 1, Part 173, Section 173.310



Applications

- · Used in direct contact with food
- Direct contact with food processing equipment and HVAC systems



Cartridges

Selection dependant on flow parameters



Pleated 5 µm

High flow rate and dirt holding capacity

Sintered 1 µm

Use for relatively low flow rates

Pleated 1 µm

Used to maximize steam capacity of pipe

JUMBO Filters

Highest available capacity

Clean Steam (HTM 2031:1997)*

STEAM Filter Cartridges

Condensate to WFI standards



Applications

- Pharmaceutical products
- Pharmaceutical plant HVAC systems



Cartridges

For removal of magnetite particles generated from stainless steel pipes due to corrosive purity of steam



HIGH FLOW TETPOR II

PTFE membrane 100% removal of magetite particles generated from stainless steel pipes

Culinary 1µm

To conform to HTM 2031 Point of Use filter rated at <5 μm

45 44

Specifications - PLEATED

Materials of Construction

- Filtration Media: 316L Stainless Steel ■ Inner Support Core: 316L Stainless Steel Outer Support Cage: 316L Stainless Steel End Caps: 316L Stainless Steel
- Standard o-rings/gaskets: EPDM (standard)

Silicone and Viton

Recommended Operating Conditions

The maximum differential pressure in direction of flow (outside to in) is 10 barg (145.03 psig).

The maximum differential pressure in direction of flow (in to outside) is 2 barg (29.00 psig).

The maximum recommended continuous operating temperature range is -75 °C (-103 °F) to +200 °C (392 °F). Note: Temperature dependant on o-ring compound

0.15 m² (1.61 ft²)

Effective Filtration Area (EFA)

10" (250 mm)

Housing Materials of Construction

316L Stainless Steel Surface Finish Electropolished Ra 0.8 Single Internal:

Single External: Mechanical Polish (Commercial Bright) Jumbo Internal: Upstream - Beadblast

> Outlet Assembly Linished 180 grit

Beadblast Jumbo External:

Vent / Drain Single / Jumbo:

Seal Material:

1/4" BSPP Female Thread EPDM Aseptic Seal

Housing Design Pressure and Temperature

Single:

16 barg (232 psig) @ 200 °C (392 °F)

Jumbo:

7 barg (101 psig) @ 170 °C (338 °F)

1	2		Figure	Housing Code	Connection Size	Capacity Kg / hr @ 1 barg	Overall Height	Replacement Filter Code
						<100 mbar or 40 m / sec		
		هم مه	1 1	HBAHP01KY HBAHP011C	1.5" (38.1 mm) 2" (50.8 mm)	150 280	14.8" (376 mm) 20.7" (526 mm)	ZCHS-KC ZCHS-1C
			2 2 2 2	VISCE-01J-D VISCE-01J-E VISCE-03J-G VISCE-03J-H	3" (50.8 mm) 4" (101.6 mm) 6" (152.4 mm) 8" (203.2 mm)	750 1300 2300 3750	30.0" (763 mm) 35.2" (895 mm) 41.2" (1049 mm) 48.7" (1237 mm)	ZCHS-J3 ZCHS-J4 3 x ZCHS-J3 3 x ZCHS-J4

Note: For efficient steam distribution it is recommended that steam velocities are restricted to 25 m/sec⁻¹. For more information on the HBA range, please contact Parker domnick hunter.

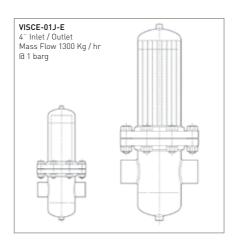
Correction Factors

To use the table above, the steam flow rates must be at 1 barg (14.50 psig). For system flows at different line pressures, divide the system flow by the correction factor below to find the equivalent flow @ 1 barg (14.50 psig).

Table showing the relative system size difference between pleated cartridges left and sintered cartridges right.

eam Pressure	0	1	2	3	4	5	6	7	8	9	10
Correction Factor					2.5			4.0	4.5	5.0	5.5





STEAM Filter Cartridges

Specifications - SINTERED

Materials of Construction

Filtration Media:

Steel (316L) End Caps: Stainless Steel (316L)

Standard o-rings/gaskets: EPDM (standard)

Silicone and Viton

Sintered Stainless

Recommended Operating Conditions

The maximum differential pressure in direction of flow (outside to in) is 10 barg (145.03 psig).

The maximum differential pressure in direction of flow (in to outside) is 5 barg (72.51 psig).

The maximum recommended continuous operating temperature range is -75 °C (-103 °F) to +200 °C (392 °F). Note: Temperature dependant on o-ring compound

Housing Materials of Construction

316L Stainless Steel

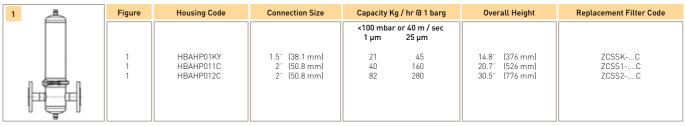
Surface Finish Internal: Electropolished Ra 0.8 External Mechanical Polish (Commercial Bright)

Vent / Drain: 1/, " BSPP Female Thread

(Supplied with Plug) EPDM Aseptic Seal Seal Material:

Housing Design Pressure and Temperature

16 barg (232 psig) @ 200 °C (392 °F)



PLEATED

20^{..}

Note: For efficient steam distribution it is recommended that steam velocities are restricted to $25\,\mathrm{m/sec^{-1}}$. For more information on the HBA range, please contact Parker domnick hunter.

Correction Factors

To use the table above, the steam flow rates must be at 1 barg (14.50 psig). For system flows at different line pressures, divide the system flow by the correction factor below to find the equivalent flow @ 1 barg (14.50 psig)

Steam Pro	essure	0	1	2	3	4	5	6	7	8	9	10
Corr	rection Factor	0.5	1.0	1.5	2.0	2.5	3.0	3.5	4.0	4.5	5.0	5.5

Ordering Information

[125 mm] | 025

(125 mm)

(250 mm

(500 mm)

(750 mm)

SINTERED

ZCSS

20^{..}



25 ft ijm



Code | Endcap (Demi)

TRUESEAL

Demi A & B Std

BF / 226 Bayonet





SINTERED Stainless Steel Retrofit Cartridge Part Numbers - 1.0 μ m & 25 μ m

Parker domnick hunter Cartridge	DS-R 3/1	DS-R 3/1.4	DS-R 4/1.5	DS-R 4/2.5	DS-R 5/2.5	DS-R 5/3	DS-R 10/3	DS-R 15/3	DS-R 20/3	DS-R 30/3	DS-R 30/5				
Retrofit Cartridge	GS3/1 SS3/1	GS3/1.5 SS3/1.5	GS4/1.5 SS4/1.5	GS4/2.5 SS4/2.5	GS5/2.5 SS5/2.5	GS5/3 SS5/3	GS10/3 SS10/3	GS15/3 SS15/3	GS20/3 SS20/3	GS30/3 SS30/3	GS30/5 SS30/5				
Parker domnick hunter Cartridge	DS-R 02/05	DS-R 02/10	DS-R 03/05	DS-R 03/10	DS-R 04/10	DS-R 04/20	DS-R 05/20	DS-R 05/25	DS-R 07/25	DS-R 07/30	DS-R 10/30	DS-R 15/30	DS-R 20/30	DS-R 30/30	DS-R 30/50
Retrofit Cartridge	GS02/05 SS02/05	GS02/10 SS02/10	GS03/05 SS03/05	GS03/10 SS03/10	GS04/10 SS04/10	GS04/20 SS04/20	GS05/20 SS05/20	GS05/25 SS05/25	GS07/25 SS07/25	GS07/30 SS07/30	GS10/30 SS10/30	GS15/30 SS15/30	GS20/30 SS20/30	GS30/30 SS30/30	GS30/50 SS30/50
Parker domnick hunter Cartridge	PDS-R 02/05	PDS-R 02/10	PDS-R 03/05	PDS-R 03/10	PDS-R 04/10	PDS-R 04/20	PDS-R 05/20	PDS-R 05/25	PDS-R 07/25	PDS-R 07/30	PDS-R 10/30	PDS-R 15/30	PDS-R 20/30	PDS-R 30/30	PDS-R 30/50
Retrofit Cartridge	P-GS02/05 P-SS02/05	P-GS02/10 P-SS02/10	P-GS03/05 P-SS03/05	P-GS03/10 P-SS03/10	P-GS04/10 P-SS04/10	P-GS04/20 P-SS04/20	P-GS05/20 P-SS05/20	P-GS05/25 P-SS05/25	P-GS07/25 P-SS07/25	P-GS07/30 P-SS07/30	P-GS10/30 P-SS10/30	P-GS15/30 P-SS15/30	P-GS20/30 P-SS20/30	P-GS30/30 P-SS30/30	P-GS30/50 P-SS30/50

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TRUESEA Demi A & B Std

Liquid filters





Covering a wide range of process applications, Parker domnick hunter manufacture a range of filters that exceed industry requirements, providir high flow rates and long life in often demanding environments. With the ability to withstand aggressive chemicals and high temperature operations, Parker domnick hunter has a liquid filter that will match your requirements.

As an industry focussed manufacturer, Parker domnick hunter understand that every process or application can be different, which is why we have a Sustaining Engineering Group whose purpose is to tailor our product range to meet your exacting needs, making our filters truly fit for purpose.

Filters include

- Polypropylene (PP)
- Glass microfibre (GF)
- Polyethersulphone (PES)
- Polytetrafluoroethylene (PTFE)

PEPLYN filters from Parker domnick hunter are used for clarification and prefiltration in a wide range of applications. The polypropylene construction makes them the ideal choice for aggressive and viscous chemicals and solvents.





48 l



PROSPUN C is the most economical solution for delivering general liquid clarification and particle retention. It can be used as a guard filter to protect the process against high variable levels of particulate.

- Economical general clarification
- High strength bonded fibre construction
- Ideal for primary stage filtration
- Nominal retention efficiency for general clarification duties

PROSPUN T offers consistent retention characteristics and a high level of security that is enhanced by the option to incorporate plug-in o-ring seal adapters on the cartridge. The service life of PROSPUN T is maximized through the use of closely controlled density and diameter fibre technology.

- High dirt holding capacity
- Range of end cap adapters and seals
- Excellent protection of downstream process
- >90% efficiency at given rating

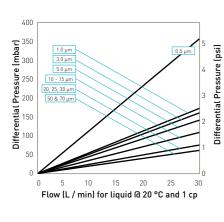
PROSPUN A - Closely controlled fibre diameter and density in a multiple layered construction serve to maximize service life of PROSPUN A whilst delivering absolute particle retention.

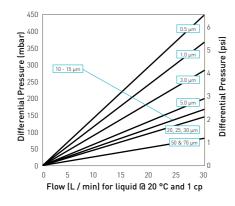
- High dirt holding capacity
- Range of end cap adapters, seals and additional support for backwash applications
- Consistent absolute retention under a wide range of operating conditions

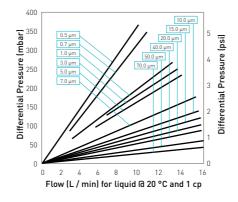


Note: PROSPUN is a registered trademark of Parker domnick hunter

Performance Characteristics







PROSPUN C

PROSPUN T

PROSPUN A

PROSPUN Filter Cartridges

Specifications

Materials of Construction

Filtration Media: Polypropylene ■ End Caps: Polypropylene Seals: As Required

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

Up to 60 °C (140 °F) continuous operating temperature and higher short-term temperatures during CIP to the following limits:

Temp °C	erature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Cleaning and Sterilization

PROSPUN cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 121 °C (249.8 °F). They 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.

Dimensions

Nominal outside diameter: 2.4" (62 mm) 1.1" (29 mm) Nominal inside diameter:

Connection Configuration				
Length	B Seal-Seal	L and 0 Seal-Seal Single Open Ended Shoulder-Shoulder		
1	9.87" (251 mm)	10" (254 mm)		
2	19.50" (498 mm)	20" (508 mm)		
3	29.37" [746 mm]	30" (762 mm)		
4	39.12" [994 mm]	40" (1016 mm)		

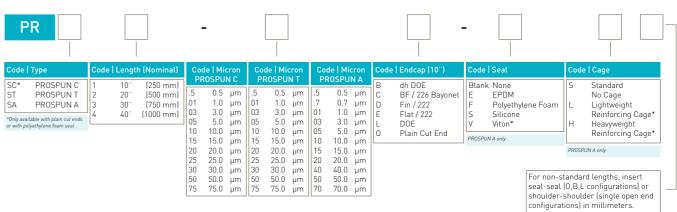
Optional reinforcing cage available for PROSPUN A, contact Parker domnick hunter for details.

Minimum Box Quantities

Cartridge Size	Quantity	
10" (254 mm)	40	
20" (508 mm)	20	
30" (762 mm)	20	
40" (1016 mm)	20	

Recommended Rinse Volume

Prior to use - 10 litres per 10" (250 mm) filter cartridge.



PROPLEAT PP Filter Cartridges

- liquid filters
- polypropylene



PROPLEAT PP cartridges have been developed to bridge the gap between meltblown depth filters and absolute rated pleated media filters.

Their continuous length and all-polypropylene construction results in a robust yet economical design that maximizes the effective filtration area and provides wide chemical compatibility, coupled with low extractable levels.

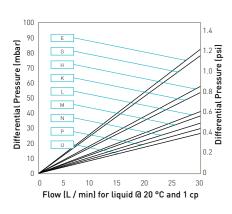
All PROPLEAT PP cartridges exhibit 99% efficiency at their given retention rating, providing consistent and economical clarification in a diverse range of applications.

Features and Benefits

- Continuous length rigid sleeve and core provide high strength during normal and reverse flow operations
- Retention ratings to suit a wide range of clarification applications
- Excellent chemical compatibility
- Elevated temperature option available for hot water sanitization and steam sterilization



Performance Characteristics



Cartridge flow rates

PROPLEAT PP Filter Cartridges

Specifications

Materials of Construction

- Filtration Media: Polypropylene
 Upstream Support: Polypropylene
 Downstream Support: Polypropylene
 Inner Support Core: Polypropylene
 Outer Protection Cage: Polypropylene
- End Cap Insert (if specified): 316L Stainless Steel*
 *Not available in B & L endcap variants

Polypropylene

■ Standard o-rings/gaskets: Silicone / EPDM

Food and Biological Safety

■ End Caps:

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

Up to 60 °C (140 °F) continuous operating temperature and higher short-term temperatures during CIP to the following limits:

Temp °C	erature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 [steam]	0.3	4.0

Effective Filtration Area (EFA)

40" (1000 mm) 2.2 m² (23.2 ft²)

Cleaning and Sterilization

PROPLEAT PP cartridges can be repeatedly in situ steam sterilized or autoclaved at up to 121 °C (250 °F). They 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.

Retention Characteristics

The retention characteristics of PROPLEAT PP have been determined by a single-pass technique using suspension of ISO 12103 Part 1 A2 Fine and A4 Coarse test dust in water.

Media Code ß ratio	99% & approximate 99% 100	ratings at lowe 95% 20	er efficiencies 90% 10
Е	0.8	0.7	0.6
G	1.0	0.9	0.7
Н	3.5	2.3	1.0
K	4.8	3.8	2.8
L	7.2	6.0	4.5
M	10.0	8.0	6.0
N	12.0	9.0	7.0
Р	18.0	13.0	10.0
U	40.0	30.0	25.0

Recommended Rinse Volume

Prior to use - 10 litres per 10" (250 mm) cartridge.

Minimum Box Quantities

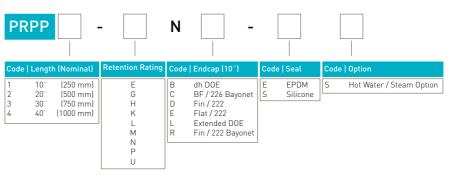
All cartridges supplied in boxes of 6.

Dimensions

- Nominal Outside Diameter: 2.8" (70 mm) C,D,E,R Style
- 2.5" (64 mm) B,L Style
- Nominal Inside Diameter: 1.1" (28 mm)

Standard Lengths (DOE seal to seal) - mm (inch)

1	9 7/8"	(250 mm)	9 7/8"	(250 mm)
2	19 1/2"	(498 mm)	20"	(508 mm)
3	29 3/8"	(746 mm)	30 1/8"	(766 mm)
4	39 1/8"	(994 mm)	40"	(1014 mm)





PROSTEEL A filters provide the ideal solution in applications where traditional polymer based filters are limited by compatibility, exposure time or a combination of high temperature and viscosity.

They are ideally suited to filtration of the solvents used in a wide range of process industries from pharmaceuticals, food &beverage and electronics through to paints and inks. The Parker domnick hunter range of stainless steel filters provides a solution to compatibility issues while maintaining absolute retention ratings down to 3.0 micron. 316L stainless steel fibres are sintered together into a graded pore structure.

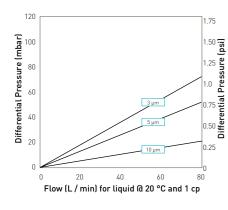
The efficiency of the media increases through the filtration bed resulting in excellent dirt holding capacity while maintaining high relative flow rates compared to alternative technology such as sintered powder tubes and metal membranes. The filters are available in two formats both using the same filtration media but one manufactured in a pleated construction and one in a cylindrical wrap. This allows a cost-effective selection depending on flow rate and dirt holding requirements.

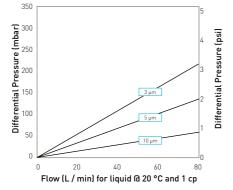
Features and Benefits

- Absolute rated stainless steel liquid filters
- Ideal for aggressive solvents, viscous and hot solutions
- Removal rating 3, 5 and 10 microns
- Compatible with most solvents
- Graded density metal fibre technology provides exceptional dirt holding capacity while retaining excellent flow rates
- Available in two formats; pleated and wrapped, for complete system optimization



Performance Characteristics





Pleated cartridge flow rates 10" Size (250 mm) Cartridge Cylindrically wrapped cartridge flow rates 10" Size (250 mm) Cartridge

PROSTEEL A Filter Cartridges

Specifications

Materials of Construction

■ Filtration Media: 316L Stainless Steel 316L Stainless Steel ■ Inner Support Core: Outer Protection Cage: 316L Stainless Steel

■ End Caps: 316L Stainless Steel ■ Standard o-rings/gaskets*:EPDM

TIG Welded

Assembly Method:

Recommended Operating Conditions

	rating erature °F		mum ard DP (psi)		imum rse DP (psi)
200	392	25	364	3	44

Note: The maximum operating temperature is dependant on o-ring selection and properties of the liquid being filtered.

Effective Filtration Area (EFA)

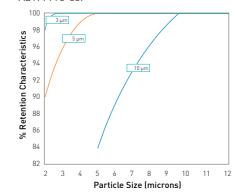
■ ZCCF Cylindrical Wrap 10" (250 mm) 0.05 m² (0.53 ft²)

ZCMF Pleated

10" (250 mm) 0.13 m² (1.39 ft²)

Retention Characteristics

The retention characteristics of the stainless steel filters are determined using ACFTD in accordance with the single pass test ASTM 795-88.



Dirt Holding Capacity

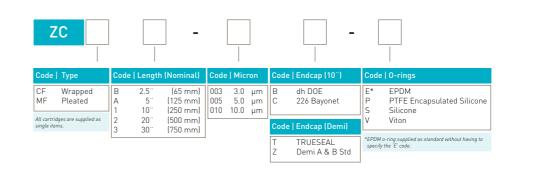
The table below gives an indication of dirt holding capacity in grams when tested in accordance with the Multipass method ISO 168892.

	M	icron Ratio	ng	
Туре	3.0	5.0	10.0	
ZCCF	3.0	3.5	4.0	
ZCMF	7.0	7.6	8.4	

Integrity Test Data

The general condition of the cartridge can be tested via the bubble point method. Typical values are detailed in the table below.

Micron Rating	9	3.0	5.0	10.0
Bubble Point	(mbarg)	125.0	76.0	37.0
in Water	(psig)	1.78	1.1	0.54





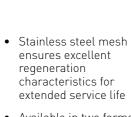
PROSTEEL N filters provide the ideal solution in applications where traditional polymer based filters are limited by compatibility, exposure time or a combination of high temperature and viscosity.

They are ideally suited to filtration of solvents used in a wide range of processes in pharmaceuticals, food & beverage and electronics through to paints and inks.

The Parker domnick hunter range of stainless steel filters provides the solution to compatibility issues while maintaining excellent flow rates for clarifying duties. The filters are available in two formats both using the same filtration media but one manufactured in a pleated construction and one in a cylindrical wrap. This allows a cost-effective selection depending on flow rate and dirt holding requirements.

Features and Benefits

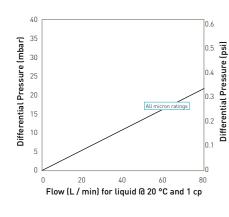
- Nominally rated stainless steel liquid filters
- Ideal for aggressive solvents, viscous and hot solutions
- Removal rating from 5 to 100 microns
- Compatible with most solvents

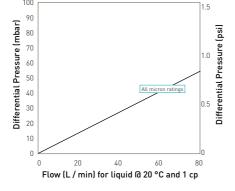


 Available in two formats; pleated and wrapped, for complete system optimisation



Performance Characteristics





Pleated cartridge flow rates 10" Size (250 mm) Cartridge Cylindrically wrapped cartridge flow rates 10" Size (250 mm) Cartridge

PROSTEEL N Filter Cartridges

Specifications

Materials of Construction

- Filtration Media: 316L Stainless Steel
 Inner Support Core: 316L Stainless Steel
- Outer Protection Cage: 316L Stainless SteelEnd Caps: 316L Stainless Steel
- Standard o-rings/gaskets*:EPDMAssembly Method: TIG Welded

*All o-rings are manufactured from FDA approved compounds.

Recommended Operating Conditions

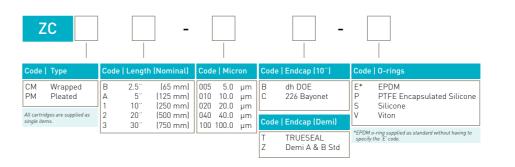
Operating Temperature		Maximum Forward DP			imum rse DP
°C	°F	(bar)	(psi)	(bar)	(psi)
200	392	25	364	3	44

Note: The maximum operating temperature is dependant on o-ring selection and properties of the liquid being filtered.

Effective Filtration Area (EFA)

- ZCCM Cylindrical Wrap
- 10" (250 mm) 0.05 m² (0.53 ft²)
- ZCPM Pleated

10" (250 mm) 0.13 m² (1.39 ft²)



PEPLYN NE Filter Cartridges

- liquid filters
- polypropylene



PEPLYN NE liquid filter cartridges are designed for use in the microelectronics industry for filtration of water, process chemicals, photochemicals, solvents and etchants.

PEPLYN NE filters resist hydrolysis in aggressive solutions which would otherwise result in the contamination of the process fluid. The filter media has graded fibre diameter and density, resulting in progressively finer retention through the depth of the media. This graded density depth mechanism, combined with optimized pleated pack configuration and high surface area, affords high flow capability and exceptional dirt holding capacity when compared with competitive pleated cartridges and meltblown depth filters. PEPLYN NE provides consistant retention and stability over a wide range of operating conditions.

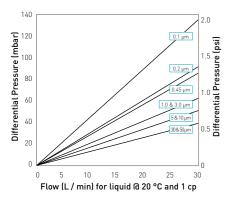
Features and Benefits

- Nominal micron ratings ranging from 0.1 to 50 micron
- · Graded density for excellent particle retention
- Pleated media for high flow rates and long life
- All polypropylene construction
- Wide range of end caps to provide retrofitting of existing systems



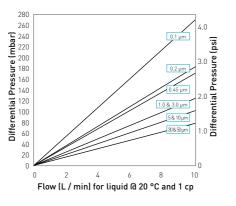
Note: PEPLYN is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" Size (250 mm) Cartridge



For A size for a given flow rate divide B size differential pressure by 2 For E size for a given flow rate multiply B size differential pressure by 2

B Size (65 mm) Cartridge

PEPLYN NE Filter Cartridges

Specifications

Materials of Construction

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

■ Inner Support Core: Polypropylene Outer Protection Cage: Polypropylene

■ End Caps: Polypropylene ■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants

■ Capsule Body: Polypropylene ■ Capsule Vent Seals: EPDM

■ Standard o-rings/gaskets: EPDM

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Ordering Information

[113 mm]

[200 mm]

.45 0.45 µm H 1.0 1.0 µm G 003 3.0 µm M 005 5.0 µm V

010 10.0 µm 050 50.0 µm

Recommended Operating Conditions

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

Temp °C	erature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules can be operated at a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.79 m² (8.50 ft²)

Cartridges **ZCNE** (65 mm) .10 (125 mm) .20 (125 mm) .45 (250 mm) 1.0 dh DOE BF / 226 Bayone EPDM PTFE Encapsulated Silicone 1.0 µm 3.0 µm Flat Top / 222 Viton Recess / 222 30" (750 mm) 005 5.0 μm H 40" (1000 mm) 010 10.0 μm J UF Retrofit SOE (no o-ring) 030 30.0 μm L 050 50.0 μm N dh DOE Internal 213 BF / 222 Bayone TRUESEAL Demi A & B Std Capsules **ZENE**

Tri-Clamp

NPT Male

... Hosebarh

Stepped Hosebark

NPT Male

3/... NPT Female

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NPT Male

" Hosebarb

Stepped Hosebarb

NPT Male

√°. NPT Female

Non Sterile 3

Pharmaceutical

PEPLYN PLUS Filter Cartridges

- liquid filters
- polypropylene



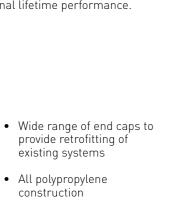
PEPLYN PLUS liquid filter cartridges are utilized for the clarification and prefiltration of a wide range of products in the pharmaceutical, beverage, ultrapure water and fine chemical industries.

The all polypropylene construction ensures a broad range of chemical compatibility making PEPLYN PLUS cartridges particularly suitable for the filtration of aggressive and viscous chemicals and solvents. They do not suffer from hydrolysis in aggressive solutions which would result in the contamination of the process fluid.

Extensive research has resulted in filter media with continuously graded fibre density giving progressively finer particulate retention through the depth of the media. This combined with optimized media pleating density gives PEPLYN PLUS cartridges exceptional lifetime performance.

Features and Benefits

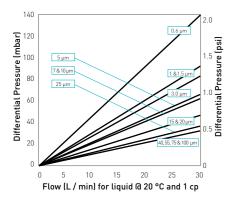
- Micron rating range from 0.6 to 100 micron
- Pleated media for high flow rates and long life
- Graded density for excellent particle retention





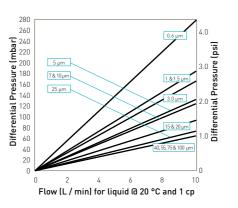
Note: PEPLYN is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" Size (250 mm) Cartridge



For A size for a given flow rate divide B size differential pressure by 2 For E size for a given flow rate multiply B size differential pressure by 2

B Size (65 mm) Cartridge

PEPLYN PLUS Filter Cartridges

Specifications

Materials of Construction

Filtration Media:	Polypropylene
Upstream Support:	Polypropylene
■ Downstream Support:	Polypropylene
■ Inner Support Core:	Polypropylene

Outer Protection Cage: ■ End Caps: Polypropylene

■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants

■ Standard o-rings/gaskets: Silicone / EPDM ■ Capsule Body: Polypropylene ■ Capsule Vent Seals: Silicone Filling Bell: Polycarbonate Syringe Filter Body: Polypropylene

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °F	Max. For (bar)	c. Forward dP (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	1.0	14.5				
>100 (steam)	>212 (steam)	0.3	4.0				

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

Up to 0.79 m² (8.50 ft²) 10" (250 mm)

Cleaning and Sterilization

PEPLYN PLUS cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 135 °C (275 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 135 °C (275 °F).

Retention Characteristics

The retention characteristics of PEPLYN PLUS have been determined by a single-pass technique using suspensions of ISO 12103 Part 1 A2 Fine and A4 Coarse test dust in water.

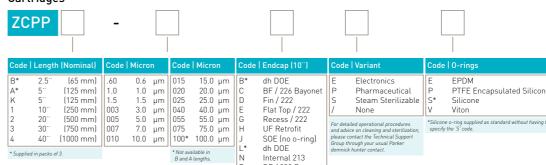
Micron Rating at Various Efficiencies											
Media	>99.99%	99.98%	99.90%	99%	90%						
Code	10000	5000	1000	100	10						
.60	0.60	0.57	0.54	0.32	0.20						
1.0	1.00	0.95	0.90	0.70	0.50						
1.5	1.50	1.40	1.10	0.80	0.60						
003	3.00	2.80	1.80	1.00	0.70						
005	5.00	4.70	4.50	3.50	1.00						
007	7.00	6.70	6.30	4.50	2.50						
010	10.00	8.00	7.00	4.80	2.80						
015	15.00	12.00	10.00	7.20	4.50						
020	20.00	16.00	14.00	10.00	6.00						
025	25.00	20.00	17.00	12.00	7.00						

Pharmaceutical Validation

A full validation guide is available upon request from Laboratory Services Group

Ordering Information

Cartridges

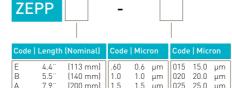


BF / 222 Bayone TRUESEAL Demi Stub Demi A & B Std

Capsules

Syringe Filters

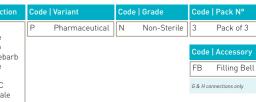
ZSPP



Length (Nominal)	Code	Mici	ron	Code	Mic	ron	Cod	e Inlet Co
4.4" (113 mm) 5.5" (140 mm) 7.9" (200 mm)	1.0	1.0	μm	020	15.0 20.0 25.0	μm	T N H	1" Tri-C 1/2" NPT 1/2" Hos
	003 005 007	3.0 5.0		040	40.0 55.0	μm	G M	Stepped 1/4" NPT Walther

Clamp T Male sebarb ed Hosebarb T Male Grommel / QC 3/, " NPT Female

" Tri-Clamp "," NPT Male epped Hosebarb "NPT Male Walther QC



Grommel / QC ³/。" NPT Female

											L					
Code	Diameter	Code	Micr	on	Code	Inlet / Outlet Connection	Code	Inlet / Outlet Connection	Code	Variant	Code	Grade	Code	Options	Code	Pack No
025 050	25 mm 50 mm	.60 1.0	0.6 1.0			Female Luer Lock Stepped Hosebarb	F G	Female Luer Lock Stepped Hosebarb	Р	Pharmaceutical	N	Non-sterile	S	Standard	025	25 per b

PREPOR GF Filter Cartridges

- liquid filters
- glass microfibre



PREPOR GF liquid filter cartridges are utilized for the clarification, stabilization and bioburden reduction of aqueous solutions, media and biologicals.

These filters have a high dirt holding capacity and exhibit exceptional flow performance compared to polypropylene filters. The hydrophilic nature of PREPOR GF filter cartridges also makes them more suitable for gravity fed systems.

PREPOR GF utilizes a glass microfibre filter medium encased within an upstream polypropylene mesh and a downstream non-woven filter support material. PREPOR GF filter cartridges are dimensionally stable with no media migration. The pleat pack is supported by an inner polypropylene core and outer polypropylene cage, heat bonded to polypropylene end caps.

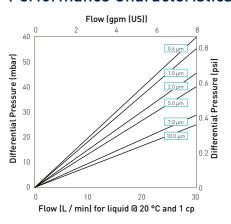
Features and Benefits

- Micron rating range from 0.6 to 10 micron
- Wide range of end caps to allow retrofitting of existing systems
- High filtration area
- High capacity filter media giving microbial retention
- · Heat bonded construction



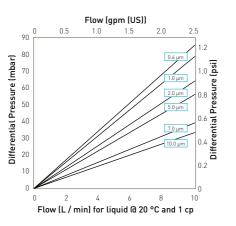
Note: PREPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" Size (250 mm) Cartridge



For A size for a given flow rate divide B size differential pressure by 2 For E size for a given flow rate multiply B size differential pressure by 2

B Size (65 mm) Cartridge

Materials of Construction

Specifications

■ Filtration Membrane: Glass Microfibre ■ Upstream Support: Polypropylene ■ Downstream Support: Polypropylene ■ Inner Support Core: Polypropylene Outer Protection Cage: Polypropylene

Polypropylene Fnd Caps: ■ End Cap Insert (if applicable): 316L Stainless Steel*

*Not available in B & L endcap variants ■ Standard o-rings/gaskets: Silicone / EPDM

■ Capsule Body: Polypropylene ■ Capsule Vent Seals: Silicone Filling Bell: Polycarbonate Syringe Filter Body: Polypropylene

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp	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	1.0	14.5			
>100 (steam)	>212 (steam)	0.3	4.0			

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.6 m² (6.3 ft²)

Cleaning and Sterilization

PREPOR GF cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 121 °C (249.8 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 121 °C (249.8 °F).

Retention Characteristics

PREPOR GF Filter Cartridges

The retention characteristics of PREPOR GF have been determined through controlled laboratory tests challenging with a standard aqueous suspension of ACFTD (AC Fine Test Dust) using on-line laser particle counters.

Micron Rating at Various Efficiencies										
fficiency : Ratio	99.99% 10000	99.98% 5000	99.90% 1000	99% 100	95% 20	90% 10				
6 & 0.8 µm	0.60	0.50	0.46	0.33	0.25	0.22				
0 & 1.5 μm	1.0	0.80	0.60	0.52	0.42	0.35				
2.0 µm	1.5	1.2	0.93	0.77	0.63	0.47				
5.0 µm	2.0	1.6	1.5	1.2	0.82	0.73				
7.0 µm	5.0	4.3	3.6	2.9	2.3	2.0				
0.0 µm	10.0	9.2	7.9	5.9	4.4	4.0				

Pharmaceutical Validation

A full validation guide is available upon request from Laboratory Services Group

Ordering Information





Code	Lengtl	h (Nominal)	Code	e Mic	ron	Code	Endcap (10")	Code	Variant	Code	0-rings
B*	2.5"	(65 mm)	.60	0.6	μm	B*	dh DOE	Р	Pharmaceutical	Е	EPDM
Α*	5"	(125 mm)	.80	0.8	μm	C	BF / 226 Bayonet	S*	Steam Sterilizable	P	PTFE Encapsulated Silicone
K	5	(125 mm)	1.0	1.0	μm	D	Fin / 222	/	None	S*	Silicone
1	10"	(250 mm)	1.5	1.5	μm	E	Flat Top / 222	_		V	Viton
2	20	(500 mm)	002	2.0	μm	G	Recess / 222		ailed operational procedures vice on cleaning and	*Silicone	o-ring supplied as standard without having to
3	30	(750 mm)	005	5.0	μm	Н	UF Retrofit	steriliza	ation, please contact the		he 'S' code.
4	40"	(1000 mm)	007	7.0	μm	J	SOE (no o-ring)		cal Support Group through your Parker domnick hunter contact.		
* Sunnlin	nd in nacks o	43	010	10.0	μm	L*	dh DOE				

Internal 213 BF / 222 Bayone

0.6 µm 0.8 µm

1.5 µm G 2.0 µm M

Retrofit TRUESEAL Demi A & B Std

Capsules

ZSGF

ZEGF	-	

(113 mm) (140 mm)

7.9" (200 mm)

NPT Male

Hosebarb

Stepped Hosebar

NPT Male

NPT Male Hosebarb

Stepped Hosebarl

" NPT Male

Walther QC Grommel / QC

3/8" NPT Female

5.0 μm Q 7.0 μm R Walther QC Grommel / QC 3/8" NPT Female

Syringe Filters

.60 0.6 µm 002 2.0 µm 6.80 0.8 µm 005 5.0 µm 1.0 1.0 µm 007 7.0 µm 1.5 1.5 µm 010 10.0 µm

Female Luer Lock

Female Luer Lock Stepped Hosebarb

Pharmaceutical N Standard

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DS_LF_07_01/11 Rev. 4A 63

25 per box

-44 (0)191 4105

PREPOR GP Filter Cartridges

- liquid filters
- · glass microfibre / polypropylene



PREPOR GP is a new prefilter that combines the strength of polypropylene with the microbial retention of glass fibre for demanding applications such as long term exposure to steam, high differential pressures or aggressive chemicals.

The combined media will also provide a significant microbial reduction that makes PREPOR GP equally suitable for bioburden reductions in pharmaceutical liquids as well as offering excellent protection to sterilizing grade membrane cartridges. By using graded density media, PREPOR GP has a higher voids volume (95%) and greater dirt holding capacity than surface filtration membranes which means that filtration costs are reduced without affecting the product quality. PREPOR GP can also provide excellent prefiltration to membrane filters in proteinaceous and high contamination applications by extending the life of the membrane cartridge and hence reducing filtration costs.

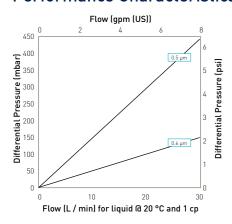
Features and Benefits

- Combined media for microbial retention and mechanical strength
- Graded density media gives increased dirt holding capacity
- Suitable for bioburden reduction and fine prefiltration
- Pleated construction with rigid core and sleeve



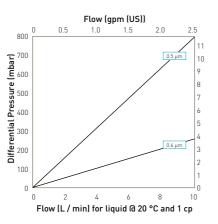
Note: PREPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" size (250 mm) filters



For A size for a given flow rate divide B size differential pressure by 2 For E size for a given flow rate multiply B size differential pressure by 2

B size (125 mm) filters

PREPOR GP Filter Cartridges

Specifications

Materials of Construction

Filtration Media: Glass Microfibre / Polypropylene ■ Upstream Support: Polypropylene ■ Downstream Support: Polypropylene ■ Inner Support Core: Polypropylene Outer Protection Cage: Polypropylene ■ End Caps: Polypropylene

■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants ■ Standard o-rings/gaskets: Silicone / EPDM

■ Capsule Body: Polypropylene ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °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	1.0	14.5				
>100 (steam)	>212 (steam)	0.3	4.0				

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.37 m² (3.9 ft²)

Cleaning and Sterilization

PREPOR GP cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 121 °C (249.8 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

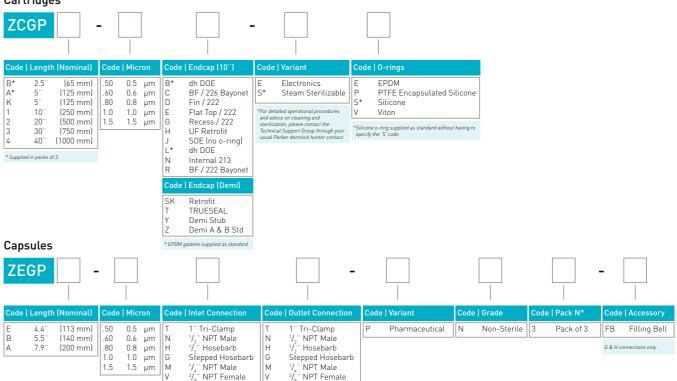
Retention Characteristics

The retention characteristics of PREPOR GP have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms. Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell	Typical Titre Reduction					
	Size (µm)*	0.5	0.6	1.0	1.5		
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0	104	10³	-	-		
Oenococcus oenos	0.5 - 0.7 x 0.7 - 1.2	104	10 ³	-	-		
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0	104	10 ³	-	-		
Saccharomyces cerevisiae	1.0 (spherical buds)	107	106	104	10³		

Ordering Information

Cartridges



PREPOR PES Filter Cartridges

- liquid filters
- · polyethersulphone



PREPOR PES is an innovative particulate grade membrane prefilter cartridge designed to work in harmony with final sterilizing filters, to guarantee the highest levels of performance and security.

PREPOR PES combines high flow rate characteristics with good microbial reduction and minimum product adsorption by using the latest hydrophilic polyethersulphone membrane technology.

PREPOR PES uses all polypropylene hardware to offer good chemical compatibility and low extractables and is suitable for use in many pharmaceutical applications including terminal and aseptic filtration, ophthalmics, biologicals, serum, SVPs, LVPs and other complex liquids.

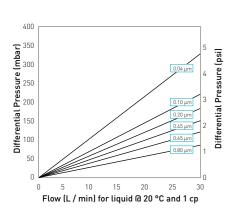
Features and Benefits

- Micron rating from 0.04 to 0.8 micron
- Versatile particulate grade membrane filter for bioburden reduction and prefiltration duties
- High filtration area with asymmetrical membrane giving long life and high flow rates
- Available in a comprehensive range of end cap configurations for retrofitting existing applications



Note: PREPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply $10^{\circ\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge

Specifications

Materials of Construction

Fnd Caps:

■ Filtration Membrane: Polyethersulphone ■ Upstream Support: Polypropylene Downstream Support: Polypropylene ■ Inner Support Core: Polypropylene Outer Protection Cage: Polypropylene

Polypropylene ■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants

■ Standard o-rings/gaskets: Silicone / EPDM ■ Capsule Body: Polypropylene Capsule Vent Seals: Silicone Filling Bell: Polycarbonate

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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	1.0	14.5				
>100 (steam)	>212 (steam)	0.3	4.0				

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.69 m² (7.42 ft²)

Cleaning and Sterilization

PREPOR PES cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

To maximize the life of the cartridge, the differential pressure across the cartridge should not exceed 0.3 barg (4.35 psig) at 130 °C (266 °F).

Retention Characteristics

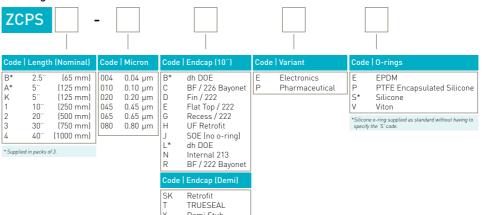
PREPOR PES Filter Cartridges

While the PREPOR PES product is rated for particulate retention, the performance of PREPOR PES products has been assessed to bacterial titre reduction using a challenge methodology based on the ASTM F838-05 methodology applied to sterilizing grade filters. Typical levels are given below:

Organism	Approx. Cell*	Typical Titre Reduction						
	Size (µm)	0.2	0.45	0.65	8.0	1.2		
Brevundimonas diminuta	0.5 - 1.0 x 1.5 - 5.0	>1010	105	10²	-	-		
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0	>1012	1010	10 ⁷	104	10²		
Oenococcus oenos	0.5 - 0.7 x 0.7 - 1.2	>1012	1012	10 ⁸	10 ⁵	10³		

Ordering Information

Cartridges



Demi A & B Std

Capsules

ZEPS		•	
Code Length	(Nominal)	Code Micron	Code Inlet Conn

010

045

(140 mm)

(200 mm) 020

0.10 μm N 0.20 μm H

0.45 µm G 0.65 µm M

" NPT Male " NPT Male " Hosebarb Hosebarb Stepped Hosebarb Walther QC

Pharmaceutical Stepped Hosebark Walther QC Grommel / QC

3/8" NPT Female

Supplied

Non-Sterile

Supplied Gamma

Grommel / QC

³/₈ NPT Female

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FB

Filling Bell

Pack of 3

Approx. values as in "Holt, J.G., Krieg, N.R., Sneath, P.H.A., Staley, J.T., Williams, S.T., 1994. Bergey's Manual of Determinative Bacteriology, Ninth Edition, Williams & Wilkins



TETPOR PLUS filters are manufactured entirely from fluoropolymers making them extremely resistant to a wide range of aggressive chemicals.

TETPOR PLUS filter cartridges have been specifically designed for the filtration of liquids and gases in the bulk pharmaceutical, chemical and biopharmaceutical industry where particulate removal, bioburden reduction and guaranteed sterility is required.

The increasing use of ozonation for the treatment of WFI systems has highlighted compatibility issues with vent filters based on standard polypropylene components. The introduction of a fully validated 0.2 micron sterilizing grade TETPOR PLUS filter cartridge provides guaranteed long term performance in these applications with the additional benefit that the filters integrity can be validated by the water intrusion test method.

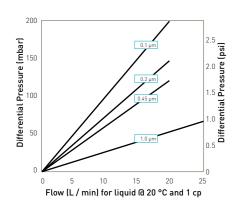
The high voids volume single layer PTFE membrane ensures an excellent combination of flow rate and retention.

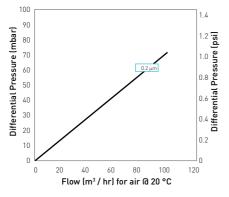
Features and Benefits

- Sterile filtration of oxygen / oxygen enriched feeds in cell culture
- Exceptional resistance to solvents and oxidative environments
- Ideal for sterile venting on ozonated water systems
- Fully validated to ASTM F838-05 for sterilizing grade filters
- PTFE membrane
- Available in a wide range of micron ratings to suit all applications



Performance Characteristics





10" Size (250 mm) Cartridge

10" Size (250 mm) Cartridge

TETPOR PLUS Filter Cartridges

Specifications

Materials of Construction

■ Filtration Membrane: Polytetrafluoroethylene ■ Upstream Support: Polytetrafluoroethylene ■ Downstream Support: Polytetrafluoroethylene ■ Inner Support Core:

Outer Protection Cage: PFA ■ End Caps:

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

Up to 125 °C (257 °F) continuous operating temperature and higher short-term temperatures during CIP to the following limits:

Temperature		Max. Forward dP		
°C	°F	(bar)	(psi)	
20	68	5.5	80.0	
75	167	3.8	55.0	
125	257	2.0	30.0	

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.63 m² (6.78 ft²) K Size (125 mm) Up to 0.32 m² (3.44 ft²)

Cleaning and Sterilization

TETPOR PLUS cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 142 °C (287.6 °F) for a maximum of 30 cycles.

Retention Characteristics

TETPOR PLUS 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) module.

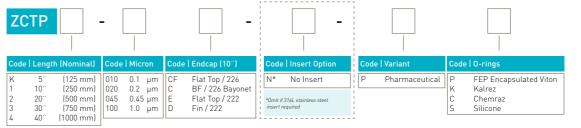
Integrity Test Data

The following is the integrity test information for the micron ratings available within the TETPOR PLUS product range. Diffusional flow and bubble point values are given for cartridges wetted in 60:40 v/v IPA:Water solution

Micron Rating		0.1	0.2	0.45	1.0	
Diffusional Flow	(barg)	1.45	1.0	0.45	3.0	٦
Test Pressure	(psig)	19.0	15.0	0.5	0.2	
Max. Diffusional Flo	w (10")	35.0	16.5	50.0	-	
(ml / min)	(K)	16.3	7.7	23.3	-	
Min. Bubble Point	(barg)	1.45	1.0	0.48	3.0	
	(psig)	19.0	15.0	0.5	0.2	
Water Intrusion	(barg)	-	2.5	-	-	
Test Pressure	(psig)	-	36.3	-	-	
Max. Water Intrusio	n (10)	-	13.5	-	-	
(ml / 10 min)	(K)	-	6.4	-	-	

Pharmaceutical Validation

A full validation guide is available upon request from Laboratory Services Group



CARBOFLOW MX Filter Cartridges

- carbon activated filters
- carbon



CARBOFLOW MX cartridges are offered in both high efficiency and general grades. They consist of bituminous coal sourced carbon, extruded together with an FDA listed thermoplastic binder, to produce an extremely porous yet rigid structure.

The result is a filter offering unsurpassed adsorptive capacity, up to 20 times that of traditional granular carbon or carbon impregnated filters, and high particle removal efficiency.

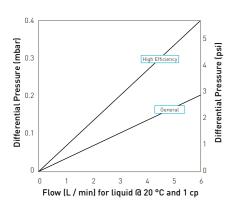
The rigid structure of CARBOFLOW MX not only minimizes any possibility of channelling, bypass or fluidizing, but also the release of carbon fines during start up and operation. Such problems are common with more traditional carbon filters. CARBOFLOW MX is available in lengths up to 40" (1016 mm) together with end fittings to suit most industry standard housings.

Features and Benefits

- Available in lengths 5" to 40"
- Ideal for chlorine and chloroform reduction
- Available in 2 grades
- FDA approved materials



Performance Characteristics



10" Size (250 mm) Cartridge

CARBOFLOW MX Filter Cartridges

Specifications

Materials of Construction

 Carbon: Bituminous Coal
 Carbon Type: Steam Activated, Acid Wash
 Carbon Weight (per 10"): 350 q

■ End Caps: Polypropylene

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Maximum Operating Temperature 60 °C (158 °F)

Maximum Differential Pressure

7 bar (101.52 psi)

Recommended Changeout Differential Pressure

2 bar (29.00 psi)

Retention Characteristics

	High Efficiency	General
Particle Removal	99.9% @ 2 mic	98% @ 10 mic
Chlorine Reduction**	76 cu.m @ 4 l / min	22.7 cu.m @ 4 l / min
Chloroform Reduction*	3 cu.m @ 2 l / min	n/a

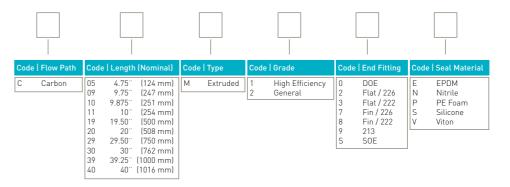
 Per 10" element, for longer lengths multiply pro-rata for details of test conditions contact Parker domnick hunter for details.

**Based on an inlet concentration of 2 ppm chlorine.

Applications

- Pre and post R.O. filtration
- Domestic drinking water
- De-chlorination
- Process water
- Product rinse waters
- Plating solutions
- De-colourization

Ordering Information



1191 4105121 🖅 dhprocessidparker.com 🖏 www.parker.com/processfi

Beverage filters





Beverage liquids

Parker domnick hunter has supplied the beverage industry with high quality filter products since 1963. During this time the company has worked hand in hand with leading beverage manufacturers to develop an industry specific range of filter products.

Experience in local markets, supported by a dedicated team of Engineers and Scientists allows Parker domnick hunter to maximize your manufacturing process and support your future development plans. PREPOR - Prefiltration liquid filters from Parker domnick hunter provide high efficiency removal of spoilage organisms and yeast removal, providing economic stabilization of your product.

BEVPOR - PES membrane range of filters from Parker domnick hunter have been specifically designed for the beverage industry to provide microbial stabilization that extends shelf-life, while maintaining colour and flavour of the final product.





PEPLYN HD Filter Cartridges

- liquid filters
- polypropylene



The two ways to increase the lifetime of a filter are to increase the amount of contamination it can handle, or to improve the effectiveness of cleaning procedures.

PEPLYN HD combines both of these capabilities in an advanced pleated construction. PEPLYN HD utilizes high depth pleated polypropylene media that balances high contaminant loading capacity with efficient cleaning.

Capture of particles is throughout the depth of the media, larger $% \left(1\right) =\left(1\right) \left(1\right) \left$ particles being retained in the outer prefiltration layers, while the inner graded density PEPLYN media provides accurately defined retention under wide extremes of operating conditions. The lifetime of PEPLYN HD is enhanced by its ability to withstand frequent backwash cleaning.

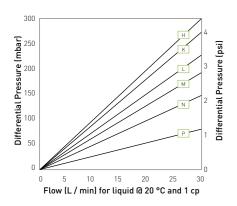
Features and Benefits

- Raw water filtration for the protection of downstream process such as RO membranes
- Trap filtration removing pre-coat and body fed particles that have been released from powder filters
- Removal of carbon and resin fines downstream from treatment processes



Note: PEPLYN is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10 $\!\!^{\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge

PEPLYN HD Filter Cartridges

Specifications

Materials of Construction

■ Filtration Media:	Polypropylene
■ Prefilter Media:	Polypropylene
Upstream Support:	Polypropylene
Downstream Support:	Polypropylene
Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
■ End Caps:	Polypropylene
■ End Cap Insert (if applicable):	316L Stainless Steel*

*Not available in B & L endcap variants

■ Standard o-rings/gaskets: Silicone / EPDM ■ Capsule Body: Polypropylene ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °F	Max. Forv (bar)	vard dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.3 m² (3.22 ft²)

Cleaning and Sterilization

PEPLYN HD cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 135 °C (275 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 135 °C (275 °F).

Retention Characteristics

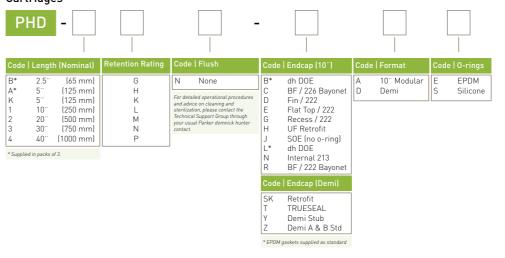
The retention characteristics of PEPLYN HD filter cartridges have been determined by a single-pass technique using suspensions of ISO 12103 Pt. 1 A2 Fine and A4 Course test dust in water.

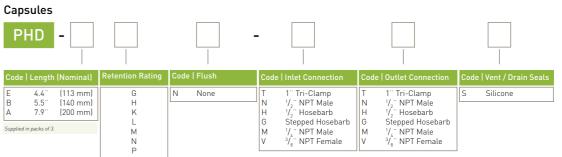
ficiency ta Ratio		99.98% 5000	99.90% 1000	99% 100	95% 20	90% 10
G	3.00	2.80	1.80	1.00	0.85	0.70
Н	4.80	4.00	3.20	2.60	1.90	1.60
K	9.00	8.20	6.90	5.00	3.70	3.40
L	12.00	10.00	7.80	5.90	4.60	4.00
М	14.00	10.00	9.20	6.90	6.10	5.00
Ν	17.00	14.00	12.00	9.00	7.00	6.00
Р	22.00	18.00	15.00	12.00	9.40	6.80

Micron Rating at Various Efficiencies

Ordering Information

Cartridges





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Two ways to increase the lifetime of a filter are to increase the amount of contamination it can handle or to improve the effectiveness of cleaning procedures. PEPLYN HA combines both of these features in its advanced pleated construction.

PEPLYN HA utilizes polypropylene filter media and support materials, which balance a high surface area and closely controlled porosity, in a configuration that maximizes the cleaning efficiency of the cartridge.

Capture of larger particles is predominantly on the surface of the media, where the rigid, open pleat structure ensures that backwash cleaning provides effective removal. Smaller particles are retained throughout the depth of the graded density PEPLYN media, providing accurately defined retention under wide extremes of operating conditions.

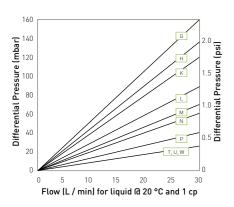
Features and Benefits

- Ideally suited for raw water filtration where the longevity of the filter can be enhanced by repetitive backwashing
- Trap filtration (also known as police or quard filtration) removing precoat and body fed particles that have been released from powder filters. for example, in a brewing process
- Removal of carbon and resin fines downstream from treatment processes
- Clarification of CIP solutions prior to their use with fine prefilter cartridges and microporous membranes



Note: PEPLYN is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10° size differential pressure by 2

10" Size (250 mm) Cartridge

Specifications

Materials of Construction

■ Filtration Media:	Polypropylene
Upstream Support:	Polypropylene
Downstream Support:	Polypropylene
Inner Support Core:	Polypropylene

Outer Protection Cage: ■ End Caps: Polypropylene

■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants ■ Standard o-rings/gaskets: Silicone / EPDM

■ Capsule Body: Polypropylene ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	oerature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.7 m² (7.53 ft²)

Cleaning and Sterilization

PEPLYN HA cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 135 °C (275 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 135 °C (275 °F).

Retention Characteristics

PEPLYN HA Filter Cartridges

The retention characteristics of PEPLYN HA filter cartridges have been determined by a single-pass technique using suspensions of ISO 12103 Pt. 1 A2 Fine and A4 Course test dust in water.

Efficiency Beta Ratio				ious Effic 99% 100	95% 20	90% 10
	4.00	0.05		0.70	0.40	0.50
D	1.00	0.95	0.90	0.70	0.60	0.50
Е	1.50	1.40	1.10	0.80	0.70	0.60
G	3.00	2.80	1.80	1.00	0.90	0.70
Н	5.00	4.70	4.50	3.50	2.30	1.00
K	10.00	8.00	7.00	4.80	3.80	2.80
L	15.00	12.00	10.00	7.20	6.00	4.50
М	20.00	16.00	14.00	10.00	8.00	6.00
N	25.00	20.00	17.00	12.00	9.00	7.00
Р	32.00	27.00	24.00	18.00	13.00	10.00
T	50.00	40.00	34.00	28.00	20.00	17.00
U	70.00	55.00	50.00	40.00	30.00	25.00
W	125.00	100.00	80.00	70.00	50.00	40.00

Ordering Information

Cartridges

Capsules



B*	2.5"	(65 mm)	D	N
A*	5"	(125 mm)	E	
K	5	(125 mm)	G	For detaile and advice
1	10"	(250 mm)	Н	sterilizatio
2	20"	(500 mm)	K	Technical S your usual
3	30	(750 mm)	L	contact.
4	40"	(1000 mm)	M	
			N	
* Supplie	ed in packs (of 3.	P	
			T	
			U	
			W*	
			* Not available in A and B lengths.	

S	C D E G H J L*	BF / 226 Bayonet Fin / 222 Flat Top / 222 Recess / 222 UF Retrofit SOE (no o-ring) dh DOE Internal 213	D	Der
	R	BF / 222 Bayonet		
	Code	Endcap (Demi)		
	SK	Retrofit		

Demi A & B Std

4.4" 5.5" 7.9" [113 mm] " Tri-Clamp Tri-Clamp NPT Male (200 mm) . Hosebarb Hosebarb Stepped Hosebarb Stepped Hosebarb NPT Male

NPT Male NPT Female

PREPOR GF Filter Cartridges

- liquid filters
- · glass microfibre



PREPOR GF filter cartridges have been specifically developed for fine clarification of water, products and ancillary liquids.

The higher efficiency grades also provide excellent bioburden reduction and protection to microporous membranes.

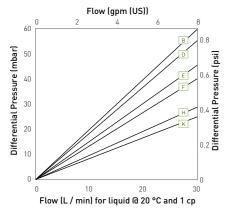
The high porosity of the microfibre filter media means that the filters have high dirt holding capacity and exhibit exceptional flow performance compared to similarly rated polypropylene filters. Coupled with the hydrophilic nature of the media, this makes them more suitable for low pressure and gravity fed systems, viscous liquids and an option for all systems where long-term elevated temperature and chemical cleaning are not required.



Features and Benefits

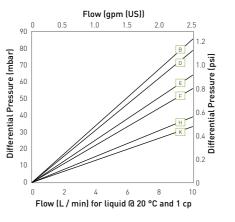
- Clarification of products for the purpose of visual aesthetics
- Fine clarification of products and ancillary liquids to extend the lifetime of microporous membrane filters
- Removal of low levels of bioburden, such as natural yeasts, from incoming liquids
- Clarification of viscous liquids such as syrups, especially where low transfer pressures are used

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" Size (250 mm) Cartridge



For A size for a given flow rate divide B size differential pressure by 2 For E size for a given flow rate multiply B size differential pressure by 2

B size (125 mm) Cartridge and Capsule

Specifications Materials of Construction

■ Filtration Membrane:	Glass Microfibre
■ Upstream Support:	Polypropylene
■ Downstream Support:	Polypropylene
■ Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
■ End Caps:	Polypropylene

■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants ■ Standard o-rings/gaskets: Silicone / EPDM

■ Capsule Body: Polypropylene ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp		Max. For	ward 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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.6 m² (6.3 ft²)

Cleaning and Sterilisation

PREPOR GF cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 121 °C (249.8 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

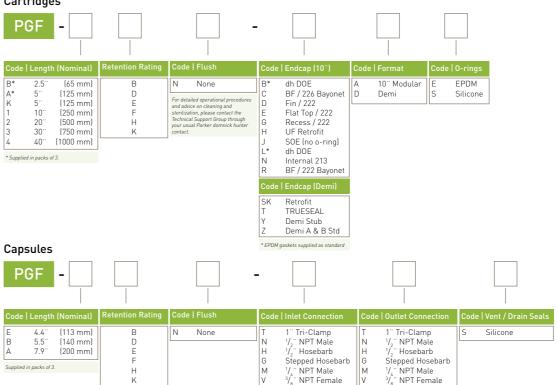
PREPOR GF Filter Cartridges

The retention characteristics of PREPOR GF have been determined through controlled laboratory tests challenging with a standard aqueous suspension of ACFTD (AC Fine Test Dust) using on-line laser particle counters.

ficiency ta Ratio			ng at Vari 99.90% 1000	ous Effic 99% 100	iencies 95% 20	90% 10
В	0.60	0.50	0.46	0.33	0.25	0.22
D	1.0	0.80	0.60	0.52	0.42	0.35
E	1.5	1.2	0.93	0.77	0.63	0.47
F	2.0	1.6	1.5	1.2	0.82	0.73
Н	5.0	4.3	3.6	2.9	2.3	2.0
K	10.0	9.2	7.9	5.9	4.4	4.0

Ordering Information

Cartridges



PREPOR GP Filter Cartridges

- liquid filters
- glass microfibre / polypropylene



PREPOR GP filter cartridges will significantly reduce numbers of yeast and spoilage organisms in beverage products to provide extremely cost-effective microbiological stabilization.

The cartridges will also 'condition' liquids and can be used to improve the filterability of products prior to terminal stabilization by thermal or filtrative methods.

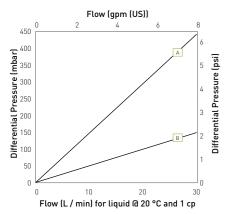
The filters utilize a unique combination of graded density glass microfibre and polypropylene media. Combined together in a pleated construction, this configuration provides a high surface area and couples the advantages of glass microfibre with the inherent strength and durability of polypropylene.



Features and Benefits

- Microbial reduction in beverage applications
- Ideally suited for yeast removal and bacterial reduction to provide shortterm microbiological stability
- Fine clarification to provide bright finished product
- Adjustment of filterability of bulk liquids after tank storage transport
- Prefiltration duty to extend the lifetime of downstream microporous membrane filters

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" size (250 mm) Cartridge

PREPOR GP Filter Cartridges

Specifications

Materials of Construction

■ Filtration Membrane:	Glass Microfibre /
	Polypropylene
■ Upstream Support:	Polypropylene
■ Downstream Support:	Polypropylene
■ Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
■ End Caps:	Polypropylene
■ End Cap Insert (if applicable):	316L Stainless Stee

*Not available in B & L endcap variants

Standard o-rings/gaskets: Silicone / EPDM

Capsule Body: PolypropyleneCapsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °F	Max. Ford (bar)	ward dP (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	1.0	14.5
100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.37 m² (3.9 ft²)

Cleaning and Sterilization

PREPOR GP cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 121 °C (249.8 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

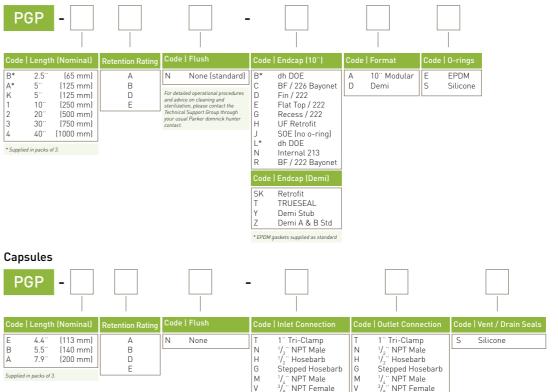
Retention Characteristics

The retention characteristics of PREPOR GP have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms. Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size (µm)*	Typica A	al Titre B	Redu D	ction E
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0	104	10³	-	-
Oenococcus oenos	0.5 - 0.7 x 0.7 - 1.2	104	10 ³	-	-
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0	104	10 ³	-	-
Saccharomyces cerevisiae	1.0 (spherical buds)	107	106	104	10³

Ordering Information

Cartridges



^{*} Approx. values as in "Holt, J.G., Krieg, N.R., Sneath, P.H.A., Staley, J.T., Williams, S.T., 1994. Bergey's Manual of Determinative Bacteriology, Ninth Edition, Williams & Wilkins".

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PREPOR PP Filter Cartridges

- liquid filters
- polypropylene



PREPOR PP filter cartridges will significantly reduce numbers of yeast and spoilage organisms from beverage products, to provide extremely cost effective microbial stabilization.

The cartridges will also 'condition' liquids and can be used to improve the filterability of products prior to terminal stabilization by thermal or filtrative methods.

The filters will withstand harsh operational conditions and repeated cleaning, making them ideal for extended use in the bulk conditioning of products prior to membrane 'sterilization' and pasteurization. Their mechanical strength and wide chemical resistance also make them suitable for long-term contact with strong cleaning agents and detergents.

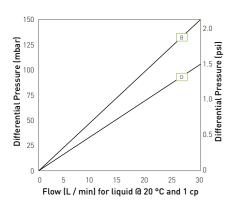
Features and Benefits

- Yeast and bacterial reduction to provide short term microbial stability
- Adjustment of filterability of bulk liquids after tank storage or transport
- Fine clarification to provide bright finished product
- Prolonged contact with hot water, steam and chemicals
- Prefiltration duty to extend the lifetime of downstream microporous filters



Note: PREPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10 $\!\!^{\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge

Specifications

Materials of Construction

■ Filtration Media:	Polypropylene
Upstream Support:	Polypropylene
Downstream Support:	Polypropylene
Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
■ End Caps:	Polypropylene

■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants ■ Standard o-rings/gaskets: Silicone / EPDM

■ Capsule Body: Polypropylene ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.5 m² (5.38 ft²)

Cleaning and Sterilization

PREPOR PP cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 135 °C (275 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 135 °C (275 °F).

Retention Characteristics

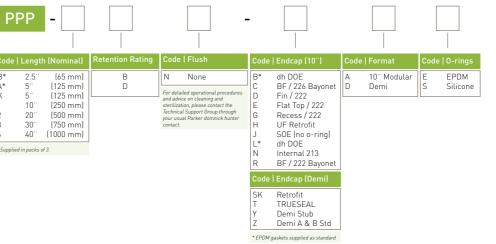
PREPOR PP Filter Cartridges

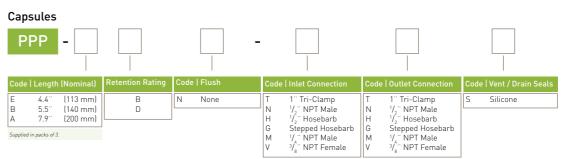
The retention characteristics of PREPOR PP have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms. Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size (µm)*	Typical Titre B	Reduction D
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0	10 ²	-
Oenococcus oenos	0.5 - 0.7 x 0.7 - 1.2	10²	-
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0	10²	-
Saccharomyces cerevisiae	1.0 (spherical buds)	104	10²

Ordering Information

Cartridges





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CRYPTOCLEAR PLUS Filter Cartridges

- liquid filters
- polypropylene



CRYPTOCLEAR PLUS pleated filter cartridges have been designed specifically for the removal of Cryptosporidium parvum and Giadia intestinalis from water in the food, beverage and healthcare industries.

Extensive research, including live oocyst challenge has resulted in a graded density filtration medium that maximizes loading capacity of the filters whilst accurately defining particle and oocyst retention under a variety of operating conditions.

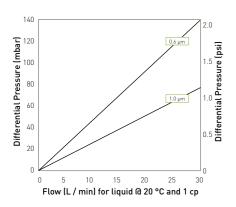
CRYPTOCLEAR PLUS cartridges can be repeatedly sanitized using hot water, steam and a wide range of chemicals.

Features and Benefits

- Specifically designed for the reduction of Cryptosporidium parvum oocysts
- 0.6 and 1.0 micron retention ratings
- All polypropylene construction
- Graded density pleated media optimized dirt capacity and oocyst retention
- Independently tested with viable *Cryptosporidium* parvum oocysts



Performance Characteristics



For K size for a given flow rate multiply 10 $\!\!^{\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge

CRYPTOCLEAR PLUS Filter Cartridges

Specifications

Materials of Construction

■ Filtration Media:	Polypropylene
Upstream Support:	Polypropylene
Downstream Support:	Polypropylene
Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
■ End Caps:	Polypropylene

■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants

■ Standard o-rings/gaskets: Silicone / EPDM ■ Capsule Body: Polypropylene ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents. CRYPTOCLEAR PLUS is listed as a WRAS Approved Product.

WRAS - Water Regulations Advisory Scheme BS6920 Test of Effect on Water Quality

Recommended Operating Conditions

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

Temp °C	erature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.57 m² (6.13 ft²)

Cleaning and Sterilization

CRYPTOCLEAR PLUS cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 142 °C (287.6 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 135 °C (275 °F).

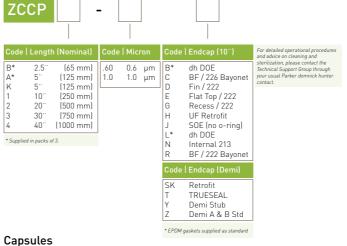
Retention Characteristics

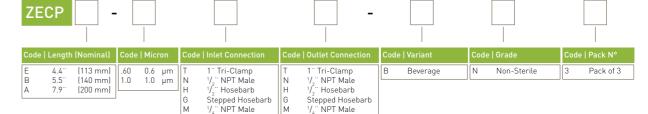
The removal efficiencies of CRYPTOCLEAR PLUS cartridges have been determined from tests conducted by Thames Water Utilities Limited on live Cryptosporidium oocysts.

Product	Micron	Retention	
CRYPTOCLEAR PLUS	0.6	>99.997%	
CRYPTOCLEAR PLUS	1.0	>99.3%	

Ordering Information

Cartridges





3/8" NPT Female

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NPT Female

CRYPTOCLEAR PES Filter Cartridges

- liquid filters
- polyethersulphone



CRYPTOCLEAR PES utilizes the unique properties of a microbially retentive polyethersulphone membrane that provides absolute retention of *Cryptosporidium parvum* oocysts to meet the specific needs of the food, beverage and potable water industries.

CRYPTOCLEAR PES membrane has an asymmetrical pore structure with a high voids volume which offers unrivalled retention capacity resulting in higher throughputs and higher flow rates than symmetrical membranes.

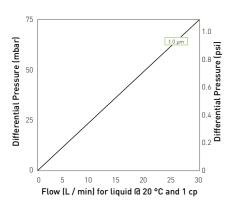
The microporous membrane is inherently hydrophilic and can be integrity tested repeatedly, providing a valuable quality assurance tool that fits well into a HACCP framework.

Features and Benefits

- Specifically developed for the removal of Cryptosporidium parvum oocysts
- 1.0 micron absolute rated polyethersulphone membrane
- High throughputs and flow rates
- Can be repeatedly steam sterilized or chemically sanitized
- Repeatedly integrity testable
- 100% retention of oocysts



Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" Size (250 mm) Cartridge

CRYPTOCLEAR PES Filter Cartridges

Specifications

Materials of Construction

Filtration Membrane:
 Prefilter Layer:
 Upstream Support:
 Downstream Support:
 Inner Support Core:
 Outer Protection Cage:
 End Caps:
 Polyethersulphone
 Polyester
 Polyester
 Polypropylene
 Nylon

■ End Cap Insert (if applicable): 316L Stainless Steel*
*Not available in B & L endcap variants

■ Standard o-rings/gaskets: Silicone / EPDM

Capsule Body: NylonCapsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents. CRYPTOCLEAR PES is listed as a WRAS Approved Product.

WRAS - Water Regulations Advisory Scheme BS6920 Test of Effect on Water Quality

Recommended Operating Conditions

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

Temp °C	erature °F	Max. Ford (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.8 m² (8.61 ft²)

Cleaning and Sterilization

CRYPTOCLEAR PES cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

The removal efficiencies of CRYPTOCLEAR PES cartridges have been determined from tests conducted by Thames Water Utilities Limited on live *Cryptosporidium* oocysts.

Product	Micron	Retention	
CRYPTOCLEAR PES	1.0	100%	

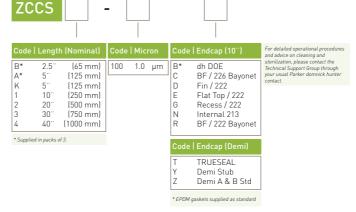
Integrity Test Data

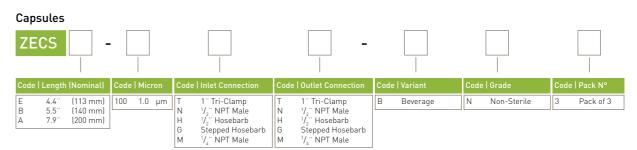
All filters are flushed with purified water prior to despatch. They are integrity testable to the following limits:

	1.0	
(barg)	0.6	
(psig)	9.0	
low (10)	21.0	
(K)	9.8	
(A)	8.0	
(B)	3.9	
(E)	1.8	
	(psig) low (10") (K) (A) (B)	(barg) 0.6 (psig) 9.0 (low (10") 21.0 (K) 9.8 (A) 8.0 (B) 3.9

Ordering Information

Cartridges





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.

r44 (0)191 4105121

BEVPOR PS Filter Cartridges

- liquid filters
- polyethersulphone



Minimizing the cost of microbiological stabilization per unit volume while maintaining quality and product characteristics is a key requirement within beverage production.

BEVPOR PS is an advanced membrane filter cartridge designed for the beverage industry to meet and surpass these criteria.

Specifically developed as a beverage grade cartridge, BEVPOR PS utilizes an advanced polyethersulphone membrane configured to provide high flow and cost-effective performance. The membrane has an asymmetric pore structure which provides graded filtration throughout its depth, resulting in increased capacity to hold contaminants. Componentry has been selected to maximize mechanical strength and chemical compatibility enabling the filter to withstand repeated chemical cleaning and sterilization.

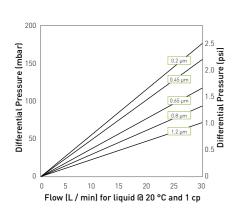
Features and Benefits

- Removal ratings from 0.2 to 1.2 micron
- Repeatedly integrity testable
- Cartridge can be regenerated and sanitized for extended service life
- Low adsorption of protein, colour and flavour components
- Asymmetrical pore structure provides high capacity contaminant loading



Note: BEVPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" Size (250 mm) Cartridge

BEVPOR PS Filter Cartridges

Specifications

Materials of Construction

- Filtration Membrane: PolyethersulphoneUpstream Support: Polyester
- Downstream Support: PolyesterInner Support Core: Polypropylene
- Outer Protection Cage: PolypropyleneEnd Caps: Nylon
- End Cap Insert (if applicable): 316L Stainless Steel*

 *Not available in B & L endcap variants
- Standard o-rings/gaskets: Silicone / EPDM
 Capsule Body: Nylon
 Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.6 m² (6.45 ft²)

Cleaning and Sterilization

BEVPOR PS cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

The retention characteristics of BEVPOR PS have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms.

Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size* (diameter x length µm)	
Brevundimonas diminuta°	0.3 x 0.6 - 0.8	
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0	
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0	
Lactobacillus brevis	0.5 - 1.2 x 1.0 - 10.0	
Saccharomyces cerevisiae	1.0 (Spherical Buds)	
Brettanomyces*	1.5 - 3.5 x 2.0 - 19.0	

			0.65		
Brevundimonas diminuta	6	-	-	-	-
Serratia marcescens	9	8	6*	-	-
Escherichia coli	>9	>9	6	2	1
Lactobacillus brevis	>9	>9	5	-	-
Saccharomyces cerevisiae	>7	>7	-	-	-
Brettanomyces	>6	>6	4	2	1

* Results based on BEVPOR

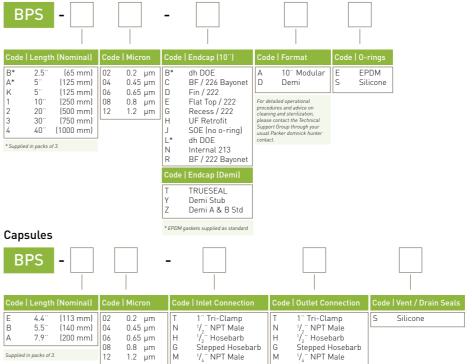
Integrity Test Data

All filters are flushed with pharmaceutical grade purified water prior to despatch. They are integrity tested to the following limits:

Micron Rating		0.2	U.45	0.65	0.8	
Diffusional Flow	(barg)	1.7	1.4	1.0	0.8	0.6
Test Pressure	(psig)	25.0	20.0	15.0	12.0	9.0
Max. Diffusional Fl	ow (10")	16.0	16.0	16.0	16.0	16.0
(ml / min)	(K)	7.5	7.5	7.5	7.5	7.5
	[A]	6.1	6.1	6.1	6.1	6.1
	(B)	3.0	3.0	3.0	3.0	3.0
	(E)	1.4	1.4	1.4	1.4	1.4

Ordering Information

Cartridges



^{*} Approx. values as in "Holt, J.G., Krieg, N.R., Sneath, P.H.A., Staley, J.T., Williams, S.T., 1994. Bergey's Manual of Determinative Bacteriology, Ninth Edition, Williams & Wilkins".

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PDA Technical Report 26, Sterilizing Filtration of Liquids

BEVPOR PH Filter Cartridges

- liquid filters
- · polyethersulphone



Minimizing the cost of microbiological stabilization per unit volume while maintaining quality and product characteristics is a key requirement within beverage production.

BEVPOR PH is an advanced membrane filter cartridge designed for the beverage industry to meet and surpass these criteria.

Specifically developed as a beverage grade cartridge, BEVPOR PH utilizes an advanced polyethersulphone membrane and an integral prefilter layer to give high flow rates, long life and improved throughput. The combination of prefilter and the asymmetrical pore structure of the membrane provides graded filtration through the depth of the media, resulting in increased capacity to hold contaminants. Componentry has been selected to withstand repeated chemical cleaning and steam sterilization.

Features and Benefits

- Removal ratings from 0.2 to 1.2 micron
- Integral prefilter layer and high surface area combine to maximize service life
- Repeatedly integrity testable
- Cartridge can be regenerated and sanitized for extended service life
- Low adsorption of protein, colour and flavour components Asymmetrical membrane pore structure provides

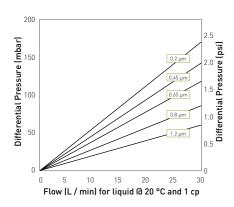
high contaminant loading

capacity



Note: BEVPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10 $\!\!^{\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge

BEVPOR PH Filter Cartridges

Specifications

Materials of Construction

- Filtration Membrane: Polyethersulphone ■ Prefilter Layer: Polyester Upstream Support: Polyester ■ Downstream Support: Polyester ■ Inner Support Core: Polypropylene Outer Protection Cage: Polypropylene ■ End Caps:
- End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants
- Standard o-rings/gaskets: Silicone / EPDM
- Capsule Body: Nylon ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.8 m² (8.61 ft²)

Cleaning and Sterilization

BEVPOR PH cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

The retention characteristics of BEVPOR PH have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms. Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size* (diameter x length μm)
Brevundimonas diminuta°	0.3 x 0.6 - 0.8
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0
Lactobacillus brevis	0.5 - 1.2 x 1.0 - 10.0
Saccharomyces cerevisiae	1.0 (Spherical Buds)
Brettanomyces*	1.5 - 3.5 x 2.0 - 19.0

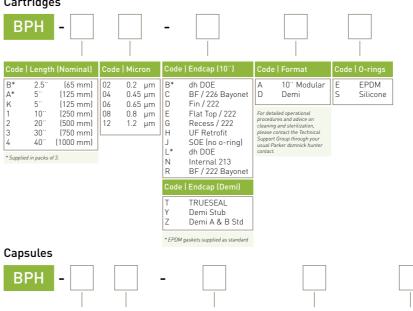
Organism			LRV		
			0.65		
Brevundimonas diminuta	6	-	-	-	-
Serratia marcescens	9	8	6*	-	-
Escherichia coli	>9	>9	6	2	1
Lactobacillus brevis	>9	>9	5	-	-
Saccharomyces cerevisiae	>7	>7	-	-	-
Brettanomyces	>6	>6	4	2	1

Integrity Test Data

All filters are flushed with pharmaceutical grade purified water prior to despatch. They are integrity tested to the following limits:

Micron Rating		0.2	0.45	0.65	8.0	1.2
Diffusional Flow	(barg)	1.7	1.4	1.0	0.8	0.6
Test Pressure	(psig)	25.0	20.0	15.0	12.0	9.0
Max. Diffusional FI	.ow [10"]	21.0	21.0	21.0	21.0	21.0
[ml / min]	(K)	9.8	9.8	9.8	9.8	9.8
	(A)	8.0	8.0	8.0	8.0	8.0
	(B)	3.9	3.9	3.9	3.9	3.9
	(E)	1.8	1.8	1.8	1.8	1.8

Ordering Information Cartridges



Supplied in packs of 3		Stepped Hosebarb G 1/, " NPT Male M	Stepped Hosebarb 1/," NPT Male
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^{*} Approx values as in "Holt, J.G., Krieg, N.R., Sneath, P.H.A., Staley, J.T., Williams, S.T., 1994. Bergey's Manual of Determinative Bacteriology, Ninth Edition, Williams & Wilkins Kuzmann, C.P., Fell, J.W., 1998 The Yeasts. A Taxonormic Study, Elsevier Science Publisher BV, Amsterdam, The Netherlands.
POI Rechnical Report AS, Sterlining Filtration of Liquids.

0.2 μm 0.45 μm

1" Tri-Clamp

Silicone

BEVPOR PT Filter Cartridges

- liquid filters
- · polyethersulphone



Minimizing the cost of microbiological stabilization per unit volume while maintaining quality and product characteristics is a key requirement within beverage production.

BEVPOR PT is an advanced membrane filter cartridge designed for the beverage industry to meet and surpass these criteria.

Specifically developed as a beverage grade cartridge, BEVPOR PT utilizes an advanced polyethersulphone membrane and an integral membrane prefilter layer to give high flow rates, long life and improved throughputs. Both prefilter and final membrane layers have an asymmetrical pore structure, providing graded filtration throughout their depth and resulting in increased capacity to hold contaminants. BEVPOR PT is especially suited to filtration of products that contain submicron colloidal species that may block unprotected sterilising-grade membranes.

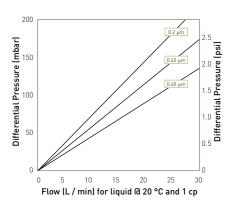
Features and Benefits

- Removal ratings from 0.2 to 0.65 micron
- Prefilter layer selected to provide removal of colloidal species providing long service life
- Repeatedly integrity testable
- Cartridge can be regenerated and sanitized for extended service life
- Low adsorption of protein, colour and flavour components
- Asymmetrical membrane pore structure provides high contaminant loading capacity



Note: BEVPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" Size (250 mm) Cartridge

BEVPOR PT Filter Cartridges

Specifications

Materials of Construction

Filtration Media: Polyethersulphone ■ Prefilter Layer: Polyethersulphone Upstream Support: Polyester ■ Downstream Support: Polyester ■ Inner Support Core: Polypropylene Outer Protection Cage:

■ End Caps: ■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants

Polypropylene

■ Standard o-rings/gaskets: Silicone / EPDM

■ Capsule Body: Nylon ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °F	Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.6 m² (6.45 ft²)

Cleaning and Sterilization

BEVPOR PT cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

The retention characteristics of BEVPOR PT have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms. Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size* (diameter x length µm)
Brevundimonas diminuta°	0.3 x 0.6 - 0.8
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0
Lactobacillus brevis	0.5 - 1.2 x 1.0 - 10.0
Saccharomyces cerevisiae	1.0 (Spherical Buds)
Brettanomyces*	1.5 - 3.5 x 2.0 - 19.0

Organism		LRV 0.45	
Brevundimonas diminuta	6	-	-
Serratia marcescens	9	8	6
Escherichia coli	>9	>9	6
Lactobacillus brevis	>9	>9	5
Saccharomyces cerevisiae	>7	>7	-
Brettanomyces	>6	>6	4

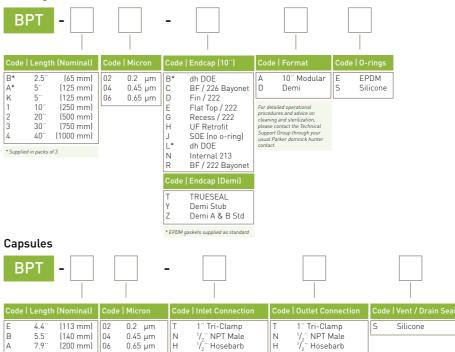
Integrity Test Data

All filters are flushed with pharmaceutical grade purified water prior to despatch. They are integrity tested to the following limits:

Micron Rating		0.2	0.45	0.65
Diffusional Flow	(barg)	1.7	1.4	1.0
Test Pressure	(psig)	25.0	20.0	15.0
Max. Diffusional Fl	ow (10")	16.0	16.0	16.0
(ml / min)	(K)	7.5	7.5	7.5
	(A)	6.1	6.1	6.1
	(B)	3.0	3.0	3.0
	(E)	1.4	1.4	1.4

Ordering Information

Cartridges



Stepped Hosebarb

Stepped Hosebarb

Approx. values as in "Holt, J.G., Krieg, N.R., Sneath, P.H.A., Staley, J.T., Williams, S.T., 1994. Bergey's Manual of Deter Kurzmann, C.P., Fell, J.W., 1998 The Yeasts. A Taxonomic Study. Elsevier Science Publisher BV, Amsterdam, The Net PDA Technical Report 24, Stentilings Filtration of Liquid.

BEVPOR PW Filter Cartridges

- liquid filters
- · polyethersulphone



Minimizing the cost of microbiological stabilization per unit volume while maintaining quality and product characteristics is a key requirement within beverage production.

BEVPOR PW is an advanced membrane filter cartridge designed to meet and surpass these criteria.

Specifically developed for the microbiological stabilization of bottled water, BEVPOR PW utilizes an advanced polyethersulphone membrane and integral prefilter layer to give high flow rates, long life and improved throughput. The combination of prefilter and the asymmetrical pore structure of the membrane provides graded filtration through the depth of the media, resulting in increased capacity to hold contaminants. Componentry has been selected to withstand repeated chemical cleaning and steam sterilization.

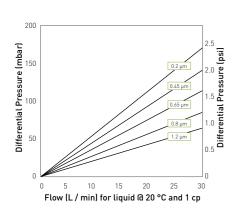
Features and Benefits

- Optimized for the microbiological stabilization of bottled water
- Removal ratings from 0.2 to 1.2 micron
- Integral prefilter layer and high surface area combine to maximize service life
- · Repeatedly integrity testable
- Cartridge can be regenerated and sanitized for extended service life
- Asymmetrical membrane pore structure provides high contaminant loading capacity



Note: BEVPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10 $\!\!^{\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge

BEVPOR PW Filter Cartridges

Specifications

Materials of Construction

■ Filtration Membrane: Polyethersulphone ■ Prefilter Layer: Polyester Upstream Support: Polyester ■ Downstream Support: Polyester ■ Inner Support Core: Polypropylene Outer Protection Cage: Polypropylene

■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants

■ Standard o-rings/gaskets: Silicone / EPDM

■ Capsule Body: Nylon ■ Capsule Vent Seals: Silicone

Food and Biological Safety

■ End Caps:

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temp °C	erature °F	Max. For	
٠٠	*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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.6 m² (6.45 ft²)

Cleaning and Sterilization

BEVPOR PW cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

The retention characteristics of BEVPOR PW have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms. Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size* (diameter x length µm)
Brevundimonas diminuta°	0.3 x 0.6 - 0.8
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0
Lactobacillus brevis	0.5 - 1.2 x 1.0 - 10.0
Saccharomyces cerevisiae	1.0 (Spherical Buds)
Brettanomyces*	1.5 - 3.5 x 2.0 - 19.0

Organism			LRV		
			0.65		
Brevundimonas diminuta	6	-	-	-	-
Serratia marcescens	9	8	6*	-	-
Escherichia coli	>9	>9	6	2	1
Lactobacillus brevis	>9	>9	5	-	-
Saccharomyces cerevisiae	>7	>7	-	-	-
Brettanomyces	>6	>6	4	2	1

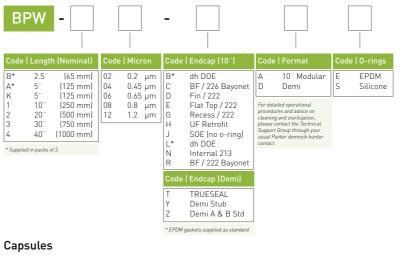
Integrity Test Data

All filters are flushed with pharmaceutical grade purified water prior to despatch. They are integrity tested to the following limits:

Micron Rating		0.2	0.45	0.65	0.8	1.2
Diffusional Flow	(barg)	1.7	1.4	1.0	0.8	0.6
Test Pressure	(psig)	25.0	20.0	15.0	12.0	9.0
Max. Diffusional F	low (10")	16.0	16.0	16.0	16.0	16.0
(ml / min)	(K)	7.5	7.5	7.5	7.5	7.5
	[A]	6.1	6.1	6.1	6.1	6.1
	(B)	3.0	3.0	3.0	3.0	3.0
	(E)	1.4	1.4	1.4	1.4	1.4

Ordering Information

Cartridges



BPW	V - [-					
Code Ler	ngth (Nor	minal)	Code	Micron	Code	Inlet Connection	Cod	e Outlet Connection	Code	Vent / Drain Seals
E 4.4 B 5.5 A 7.9	5" (14)	0 mm) 0 mm)	02 04 06	0.2 μm 0.45 μm 0.65 μm	T N H	1" Tri-Clamp 1/_" NPT Male 1/_" Hosebarb	T N H	1" Tri-Clamp 1/2" NPT Male 1/2" Hosebarb	S	Silicone
Supplied in pack	ks of 3.		08 12	0.8 μm 1.2 μm	G M	Stepped Hosebarb 1/4" NPT Male	G M	Stepped Hosebarb 1/4" NPT Male		

domnick hunter

The BEVPOR range of membrane cartridge filters is available in a selection of retention ratings to provide protection of beverages from the effects of common spoilage organisms or to enable them to meet regulatory requirements.

However, it is possible that other smaller microorganisms may be present that, while not affecting microbiological stability, may nonetheless be undesirable from a quality viewpoint. BEVPOR MS provides higher removal efficiency than BEVPOR PS, the basis of which is the recognized standard in the pharmaceutical industry for a 0.2 micron sterilizing grade membrane^[1]. Specifically developed as a beverage grade cartridge, BEVPOR MS utilizes an advanced polyethersulphone membrane configured to provide high flow and cost-effective performance. The membrane has an asymmetric pore structure which provides graded filtration throughout its depth, resulting in increased capacity to hold contaminants. Componentry has been selected to maximize mechanical strength and chemical compatibility enabling the filter to withstand repeated chemical cleaning and sterilization. [1]ASTM F838-05

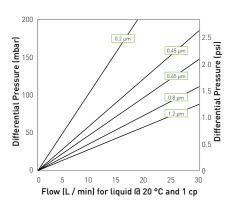
Features and Benefits

- Enhanced microbial retention based on pharmaceutical industry specifications
- Repeatedly integrity testable
- Cartridges can be regenerated and sanitized for extended service life
- · Low adsorption of protein, colour and flavour components
- Asymmetrical membrane pore structure provides high contaminant loading



Note: BEVPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply $10^{\circ\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge

BEVPOR MS Filter Cartridges

Specifications

Fnd Caps:

Materials of Construction

- Filtration Membrane: Polyethersulphone ■ Upstream Support: Polyester
- Downstream Support: Polyester Polypropylene Inner Support Core: Outer Protection Cage: Polypropylene
- End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants

Nylon

■ Standard o-rings/gaskets: Silicone / EPDM ■ Capsule Body: Nylon ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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	°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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Whilst BEVPOR MS can withstand reverse pressure, poor control of backwash procedures can result in damage to the product. Consult Parker domnick hunter before using reverse flow or pressurisation techniques.

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.6 m² (6.45 ft²)

Cleaning and Sterilization

BEVPOR MS cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

The retention characteristics of BEVPOR MS have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms. Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size* (diameter x length µm)
Brevundimonas diminuta°	0.3 x 0.6 - 0.8
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0
Lactobacillus brevis	0.5 - 1.2 x 1.0 - 10.0
Saccharomyces cerevisiae	1.0 (Spherical Buds)
Brettanomyces*	1.5 - 3.5 x 2.0 - 19.0

Ol gallisili			LRV		
	0.2	0.45	0.65	8.0	1.2
Brevundimonas diminuta	>10	6	-	-	-
Serratia marcescens	>9	9	8	6*	-
Escherichia coli	>9	>9	>9	6	2
Lactobacillus brevis	>9	>9	>9	5	-
Saccharomyces cerevisiae	>7	>7	>7	-	-
Brettanomyces	>6	>6	>6	4	2

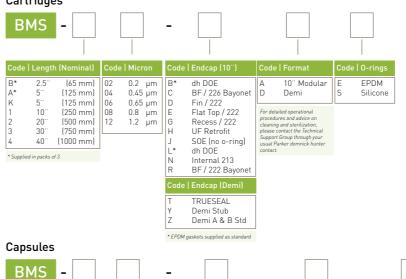
Integrity Test Data

All filters are flushed with pharmaceutical grade purified water prior to despatch. They are integrity tested to the following limits:

Micron Rating		0.2	0.45	0.65	8.0	1.2
Diffusional Flow	(barg)	2.4	1.7	1.4	1.0	0.8
Test Pressure	(psig)	35.0	25.0	20.0	15.0	12.0
Max. Diffusional F	low (10")	16.0	16.0	16.0	16.0	16.0
[ml / min]	(K)	7.5	7.5	7.5	7.5	7.5
	(A)	6.1	6.1	6.1	6.1	6.1
	(B)	3.0	3.0	3.0	3.0	3.0
	(E)	1.4	1.4	1.4	1.4	1.4

Ordering Information

Cartridges



E 4.4" [113 mm] 02 0.2 µm T 1" Tri-Clamp T 1" Tri-Clamp N 1/2" NPT Male N 1/2" NPS Male N 1/	



The BEVPOR range of membrane cartridge filters is available in a selection of retention ratings to provide protection of beverages from the effects of common spoilage organisms or to enable them to meet regulatory requirements.

However, it is possible that other smaller microorganisms may be present that, while not affecting microbiological stability, may nonetheless be undesirable from a quality viewpoint. BEVPOR MT provides higher removal efficiency than BEVPOR PT, the basis of which is the recognized standard in the pharmaceutical industry for a 0.2 micron sterilizing grade membrane^[1]. Specifically developed as a beverage grade cartridge, BEVPOR MT utilizes an advanced polyethersulphone membrane configured to provide high flow and cost-effective performance. The membrane has an asymmetric pore structure which provides graded filtration throughout its depth, resulting in increased capacity to hold contaminants. Componentry has been selected to maximize mechanical strength and chemical compatibility enabling the filter to withstand repeated chemical cleaning and sterilization. ^[1]ASTM F838-05

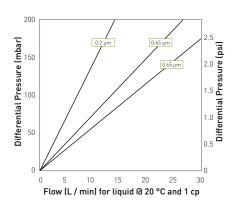
Features and Benefits

- Enhanced microbial retention based on pharmaceutical industry specifications
- Prefilter layer selected to provide removal of colloidal species providing long service life
- Repeatedly integrity testable
- Cartridge can be regenerated and sanitized for extended service life
- Low adsorption of protein, colour and flavour components
- Asymmetrical membrane pore structure provides high contaminant loading capacity



Note: BEVPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply $10^{\circ\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge

BEVPOR MT Filter Cartridges

Specifications

■ End Caps:

Materials of Construction

■ Filtration Membrane: Polyethersulphone ■ Prefilter Layer: Polyethersulphone Upstream Support: Polyester ■ Downstream Support: Polyester ■ Inner Support Core: Polypropylene Outer Protection Cage: Polypropylene

■ End Cap Insert (if applicable): 316L Stainless Steel* *Not available in B & L endcap variants

Nvlon

■ Standard o-rings/gaskets: Silicone / EPDM

■ Capsule Body: Nylon ■ Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

-			1.15
Temp	erature	Max. For	ward dP
°C		(bar)	
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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Whilst BEVPOR MT can withstand reverse pressure, poor control of backwash procedures can result in damage to the product. Consult Parker domnick hunter before using reverse flow or pressurisation techniques.

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.6 m² (6.45 ft²)

Cleaning and Sterilization

BEVPOR MT cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

The retention characteristics of BEVPOR MT have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms. Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size* (diameter x length µm)	
Brevundimonas diminuta°	0.3 x 0.6 - 0.8	
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0	
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0	
Lactobacillus brevis	0.5 - 1.2 x 1.0 - 10.0	
Saccharomyces cerevisiae	1.0 (Spherical Buds)	
Brettanomyces*	1.5 - 3.5 x 2.0 - 19.0	

organism	0.2	0.45	0.65
Brevundimonas diminuta	>10	6	-
Serratia marcescens	>9	9	8
Escherichia coli	>9	>9	>9
Lactobacillus brevis	>9	>9	>9
Saccharomyces cerevisiae	>7	>7	>7
Brettanomyces	>6	>6	>6

Integrity Test Data

All filters are flushed with pharmaceutical grade purified water prior to despatch. They are integrity tested to the following limits:

Micron Rating		0.2	0.45	0.65
Diffusional Flow	(barg)	2.4	1.7	1.4
Test Pressure	(psig)	35.0	25.0	20.0
Max. Diffusional F	low (10")	16.0	16.0	16.0
(ml / min)	(K)	7.5	7.5	7.5
	(A)	6.1	6.1	6.1
	(B)	3.0	3.0	3.0
	(E)	1.4	1.4	1.4

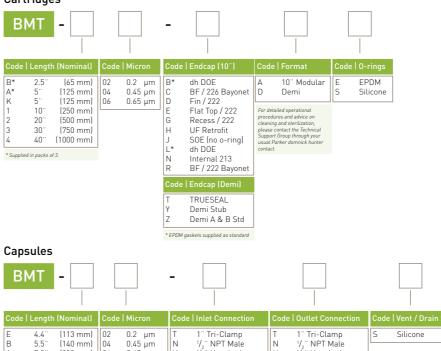
Ordering Information

(140 mm) 04

(200 mm) 06

Supplied in packs of 3.

Cartridges



" NPT Male

" Hosebarb

1/." NPT Male

0.65 um

NPT Male

· Hosebarb

1/." NPT Male



The BEVPOR range of membrane cartridge filters is available in a selection of retention ratings to provide protection of beverages from the effects of common spoilage organisms or to enable them to meet regulatory requirements.

However, it is possible that other smaller microorganisms may be present that, while not affecting microbiological stability, may nonetheless be undesirable from a quality viewpoint. BEVPOR MH provides higher removal efficiency than BEVPOR PH, the basis of which is the recognized standard in the pharmaceutical industry for a 0.2 micron sterilizing grade membrane^[1]. Specifically developed as a beverage grade cartridge, BEVPOR MH utilizes an advanced polyethersulphone membrane and integral prefilter layer to give high flow rates, long life and improved throughput. The combination of prefilter and the asymmetrical pore structure of the membrane provides graded filtration through the depth of the media, resulting in increased capacity to hold contaminants. Componentry has been selected to withstand repeated chemical cleaning and steam sterilization. MASTM F838-05

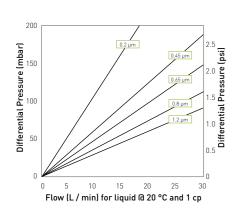
Features and Benefits

- Enhanced microbial retention based on pharmaceutical industry specifications
- Integral prefilter layer and high surface area combine to maximize service life
- Repeatedly integrity testable
- Cartridge can be regenerated and sanitized for extended service life
- Low adsorption of protein, colour and flavour components
- Asymmetrical membrane pore structure provides high contaminant loading capacity



Note: BEVPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" Size (250 mm) Cartridge

BEVPOR MH Filter Cartridges

Specifications

Materials of Construction

- Filtration Membrane: Polyethersulphone
 Prefilter Layer: Polyester
 Upstream Support: Polyester
 Downstream Support: Polyester
 Inner Support Core: Polypropylene
 Outer Protection Cage: Polypropylene
 End Caps: Nylon
- End Cap Insert (if applicable): 316L Stainless Steel*
 *Not available in B & L endcap variants
- Standard o-rings/gaskets: Silicone / EPDM
- Capsule Body: Nylon
 Capsule Vent Seals: Silicone

Food and Biological Safety

Materials conform to the relevant requirements of 21CFR Part 177, EC1935 / 2004 and current USP Plastics Class VI - 121 °C and ISO10993 equivalents.

Recommended Operating Conditions

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

Temperature °C °F		Max. For (bar)	ward dP (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	1.0	14.5
>100 (steam)	>212 (steam)	0.3	4.0

Whilst BEVPOR MH can withstand reverse pressure, poor control of backwash procedures can result in damage to the product. Consult Parker domnick hunter before using reverse flow or pressurisation techniques.

Capsules may be operated up to a temperature of 40 °C (104 °F) at line pressures up to 5.0 barg (72.51 psig) for liquids.

Effective Filtration Area (EFA)

10" (250 mm) Up to 0.8 m² (8.61 ft²)

Cleaning and Sterilization

BEVPOR MH cartridges can be repeatedly steam sterilized in situ or autoclaved at up to 130 °C (266 °F). They can be sanitized with hot water at up to 90 °C (194 °F) and are compatible with a wide range of chemicals. Capsules can be repeatedly autoclaved up to 130 °C (266 °F).

Retention Characteristics

The retention characteristics of BEVPOR MH have been determined by a combination of controlled laboratory tests and in-use monitoring for a number of organisms.

Bacterial challenge testing is carried out to methods specified in ASTM F838-05.

Organism	Approx. Cell Size* (diameter x length µm)	
Brevundimonas diminuta°	0.3 x 0.6 - 0.8	
Serratia marcescens	0.5 - 0.8 x 0.9 - 2.0	
Escherichia coli	1.1 - 1.5 x 2.0 - 6.0	
Lactobacillus brevis	0.5 - 1.2 x 1.0 - 10.0	
Saccharomyces cerevisiae	1.0 (Spherical Buds)	
Brettanomyces*	1.5 - 3.5 x 2.0 - 19.0	

Urganism			0.65		
Brevundimonas diminuta	>10	6	-	-	-
Serratia marcescens	>9	9	8	6*	-
Escherichia coli	>9	>9	>9	6	2
Lactobacillus brevis	>9	>9	>9	5	-
Saccharomyces cerevisiae	>7	>7	>7	-	-
Brettanomyces	>6	>6	>6	4	2

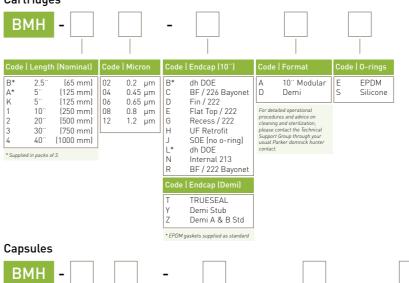
Integrity Test Data

All filters are flushed with pharmaceutical grade purified water prior to despatch. They are integrity tested to the following limits:

Micron Rating		0.2	0.45	0.65	8.0	1.2
Diffusional Flow	(barg)	2.4	1.7	1.4	1.0	0.8
Test Pressure	(psig)	35.0	25.0	20.0	15.0	12.0
Max. Diffusional FI	.ow [10"]	21.0	21.0	21.0	21.0	21.0
(ml / min)	(K)	9.8	9.8	9.8	9.8	9.8
	(A)	8.0	8.0	8.0	8.0	8.0
	(B)	3.9	3.9	3.9	3.9	3.9
	(E)	1.8	1.8	1.8	1.8	1.8

Ordering Information

Cartridges



BMH -				
Code Length (Nominal)	Code Micron	Code Inlet Connection	Code Outlet Connection	Code Vent / Drain Seals
E 4.4" [113 mm] B 5.5" [140 mm] A 7.9" [200 mm]	02 0.2 μm 04 0.45 μm 06 0.65 μm 08 0.8 μm	T 1" Tri-Clamp N 1/2" NPT Male H 1/2" Hosebarb G Stepped Hosebarb	T 1" Tri-Clamp N 1/." NPT Male H 1/.2" Hosebarb G Stepped Hosebarb	S Silicone

^{*} Approx. values as in "Holt, J.G., Krieg, N.R., Sneath, P.H.A., Staley, J.T., Williams, S.T., 1994. Bergey's Manual of Determinative Bacteriology, Ninth Edition, Williams & Wilkins'.
* Kurzmann, C.P., Fell, J.W., 1998 The Yeasts. A Taxonomic Study. Elsevier Science Publisher BV, Amsterdam, The Netherlands.

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 atterations. This publication is for general information only and customers are requested to contact on Process Efficiency Standard conditions of sale

Pharmaceutical filters



Pharmaceutical filtration

Parker domnick hunter manufacture innovative filtration solutions for critical areas of pharmaceutical production such as bulk chemicals / API, fermentation and aseptic final fill Our validated product range is fully supported by our global network of technical support Scientists and Engineers

The ability to scale up from small area discs to process scale systems with minimal revalidation is paramount.

Parker domnick hunter provides a wide range of filter formats to ensure that the transition from pilot-scale through to production is as smooth as possible.

Disposable single use systems can eliminate cleaning validation, reduce capital costs, minimize health and safety risks and lower the risk of product contamination, as well as providing a more convenient way of processing a product.

PROCLEAR filters from Parker domnick hunter represent a range of prefiltration and clarification media for particulate removal and bioburden reduction. Designed to maximize throughput in the most demanding applications.

PROPOR multi-format sterile liquid filters from Parker domnick hunter offer a PES membrane which demonstrates low preservative binding and retention of diminutive organisms, coupled with high flow and high capacity performance in critical applications.





| 103

PROCLEAR GF Filter Cartridges

- liquid filters
- · glass microfibre



PROCLEAR GF filters are designed for reliable and economical removal of particulate and microorganisms from pharmaceutical

The non-fibre releasing glass microfibre filter media gives excellent dirt holding capacity and high flow rates for long service life and efficient and cost-effective filter system design.

PROCLEAR GF filters have low extractable levels making them ideal for general clarification and prefiltration duties in pharmaceutical processing.

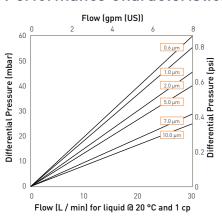


Note: PROCLEAR and DEMICAP are registered trademarks of Parker domnick hunter

Features and Benefits

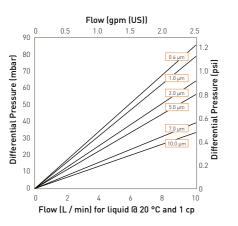
- Excellent dirt holding capacity
- Non-fibre releasing glass microfibre media
- Long service life for maximum throughput
- MURUS and DEMICAP's can be gamma-irradiated and autoclaved

Performance Characteristics



For K size for a given flow rate multiply 10 $\!\!^{\circ}$ size differential pressure by 2

10" size (250 mm) Cartridge



For A size for a given flow rate divide B size differential pressure by 2 For E size for a given flow rate multiply B size differential pressure by 2

B size (65 mm) Cartridge and Capsule

Specifications

Materials of Construction

Materials of Collstial	LIUII	Lifective i itti attoli Alea (Li A)			
■ Filtration Media:	Glass Microfibre	10" (250 mm):	0.56m^2	(6.0 ft^2)	
Upstream Support:	Polypropylene	K Size:	0.27m^2	(2.9 ft ²)	
Downstream Support:	Polypropylene	A Size:	0.20m^2	(2.2 ft ²)	
		B Size:	0.10m^2	(1.1 ft ²)	
Filter Cartridges		E Size:	0.05m^2	(0.6 ft^2)	
■ Inner Support Core:	Polypropylene	Syringe ø50 mm:	$14.50 cm^2$	(2.25 ir	
Outer Protection Cage:	Polypropylene				

MURUS Disposable Filter Capsules

*Not available in B & L endcap variants

Polypropylene

316L Stainless Steel

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:	Polypropylen
■ Sleeve:	Polypropylen
■ Capsule Body:	Polypropylen
■ Capsules Vent Seals:	Silicone
■ Filling Bell:	Polycarbonat

Syringe Filters

End Caps:

■ End Caps Insert:

■ Body: 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		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	1.5	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 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 Filter Capsules

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

Effective Filtration Area (FFA)

10" (250 mm):	0.56m^2	(6.0 ft^2)
K Size:	0.27m^2	(2.9 ft ²)
A Size:	0.20m^2	(2.2 ft ²)
B Size:	0.10m^2	(1.1 ft ²)
E Size:	0.05m^2	(0.6 ft^2)
Syringe ø50 mm:	$14.50cm^2$	(2.25 in ²)

Sterilization

	Autoclave		Steam-in-Place	
	Cycles		Cycles (30 min.)	
Cartridges	10	130 °C (266 °F)	10	121 °C [249.8 °F]
MURUS	5	130 °C (266 °F)	-	-
DEMICAP	10	130 °C (266 °F)	-	-
Syringe	1	130 °C [266 °F]	-	-

PROCLEAR GF 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.

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

PROCLEAR GF Filter Cartridges

Pharmaceutical grade products are manufactured in accordance with cGMP, 100% flushed with pharmaceutical grade purified water.

Gamma-Irradiation

PROCLEAR GF MURUS & DEMICAP disposable filters can be gamma-irradiated up to a maximum dosage of 40 kGy.

PROCLEAR GF Filter Cartridges

Performance Characteristics

TOC / Conductivity

The filtrate quality from a 10" (250 mm) PROCLEAR GF 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 GF 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" (200 mm) DEMICAP capsule are <5 mg.

Pharmaceutical Validation

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

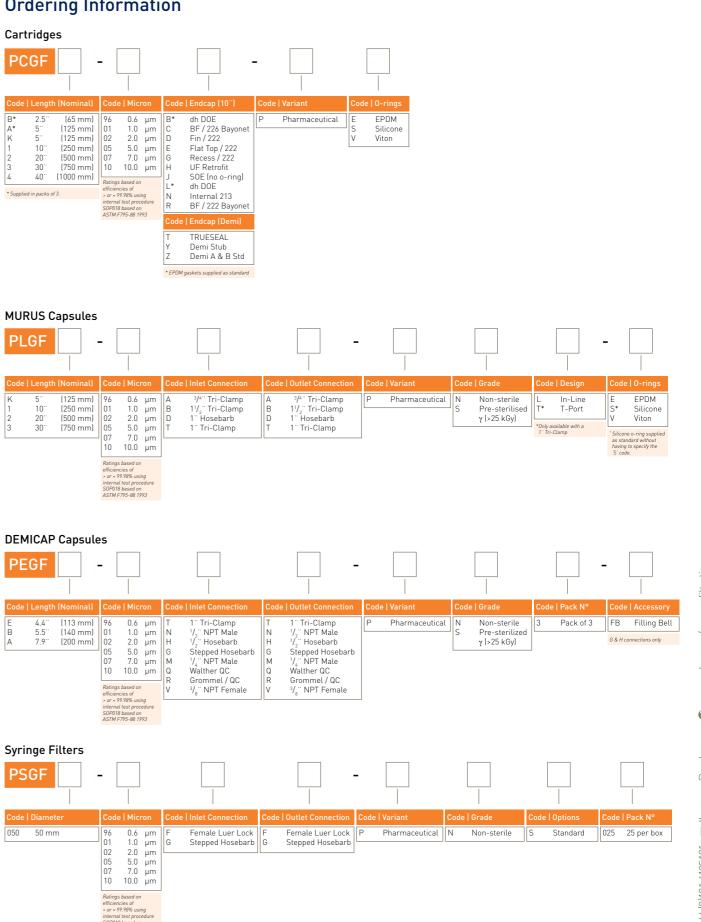
Oxidizable Substances

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

106

PROCLEAR GF Filter Cartridges

Ordering Information



DS_P_01_01/11 Rev. 4A 107

PROCLEAR GP Filter Cartridges

- liquid filters
- glass microfibre / polypropylene



PROCLEAR GP filters combine glass microfibre and polypropylene media to provide maximum protection to downstream filter membranes and equipment.

Dirt holding capacity is maximized through use of a graded density media making PROCLEAR GP cartridge filters an economical and reliable choice for prefiltration.

PROCLEAR GP filters have low extractable levels and are suitable for bioburden reduction and fine prefiltration of pharmaceutical fluids and are ideal for high contamination applications.

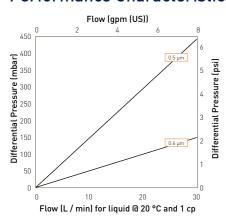
Features and Benefits

- Dual layer media or increased capacity and assurance
- Maximizes retention for protection of downstream membranes
- Ideal for difficult to filter solutions
- MURUS and DEMICAP's can be gamma-irradiated and autoclaved



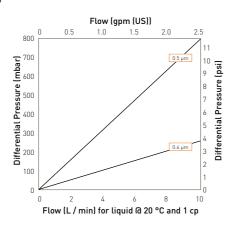
Note: PROCLEAR and DEMICAP are registered trademarks of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" size (250 mm) Cartridge



For A size for a given flow rate divide B size differential pressure by 2 For E size for a given flow rate multiply B size differential pressure by 2

B size (65 mm) Cartridge and Capsule

Specifications

Materials of Construction

Filtration Media:	Glass Microfibre
	Polypropylene
Upstream Support:	Polypropylene
Downstream Support:	Polypropylene

Filter Cartridges

Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
End Caps:	Polypropylene
End Caps Insert:	316L Stainless Steel
*Not available in B & L endcap	variants

MURUS Disposable Filter Capsules

 ortos bisposable i iller od	ipautea
Core:	Polypropylene
Sleeve:	Polypropylene
End Caps Insert:	316L Stainless Steel
Standard o-rings/gaskets:	Silicone
Capsule Body:	Polypropylene
Cansules Vent Seals:	Silicone

DEMICAP Filter Capsules

Core:	Polypropylen
■ Sleeve:	Polypropylen
■ Capsule Body:	Polypropylen
Capsules Vent Seals:	Silicone

Syringe Filters

■ Body: 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:

Temp	erature	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	1.5	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 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 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):	0.34 m^2	(3.7 ft ²)
K Size:	0.16 m^2	(1.7 ft^2)
A Size:	0.12 m^2	(1.3 ft ²)
B Size:	0.06m^2	(0.6 ft^2)
E Size:	0.03m^2	(0.3 ft^2)
Syringe ø50 mm:	14.50 cm ²	(2.25 in ²)

Sterilization

Autoclave		Steam	ı-in-Place
Cycles	Temp	Cycles (30 min.)	Temp
10	130 °C [266 °F]	10	121 °C [249.8 °F]
5	130 °C [266 °F]	-	-
10	130 °C [266 °F]	-	-
1	130 °C (266 °F)	-	-
	10 5	10 130 °C (266 °F) 5 130 °C (266 °F) 10 130 °C (266 °F)	Cycles Temp Cycles (30 min.) 10 130 °C (266 °F) 10 5 130 °C (266 °F) - 10 130 °C (266 °F) -

PROCLEAR GP 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.

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.

PROCLEAR GP Filter Cartridges

Quality Standards

Pharmaceutical grade products are manufactured in accordance with cGMP, 100% flushed with pharmaceutical grade purified water.

Gamma-Irradiation

PROCLEAR GP MURUS & DEMICAP disposable filters can be gamma-irradiated up to a maximum dosage of 40 kGy.

PROCLEAR GP Filter Cartridges

Performance Characteristics

TOC / Conductivity

The filtrate quality from a 10" (250 mm) PROCLEAR GP 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 GP 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" (200 mm) DEMICAP capsule are <5 mg.

Pharmaceutical Validation

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

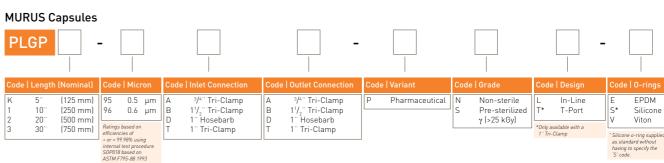
Oxidizable Substances

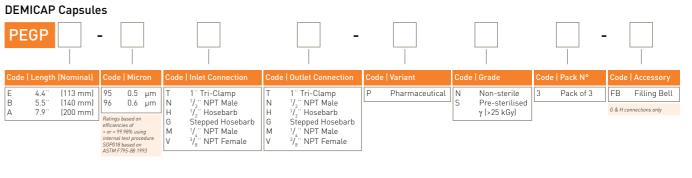
PROCLEAR GP filter cartridges meet current USP and EP quality standards for sterile purified water for oxidizable substances following a <1 litre water flush.

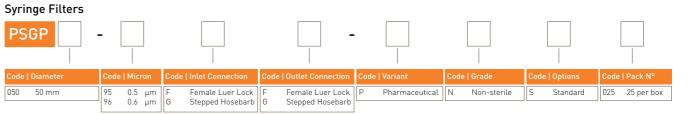
110

PROCLEAR GP Filter Cartridges

Ordering Information Cartridges 0.5 µm 0.6 µm (65 mm) (125 mm) dh DOE BF / 226 Bayor EPDM Flat Top / 222 Recess / 222 (250 mm) (500 mm) UF Retrofit SOE (no o-ring) (750 mm) internal test procedu \$000 mm ASTM F795-88 1993 dh DOE Internal 213 BF / 222 Bayon TRUESEAL Demi A & B Std







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 atterations. 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.

PROCLEAR PP Filter Cartridges

- liquid filters
- polypropylene



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.

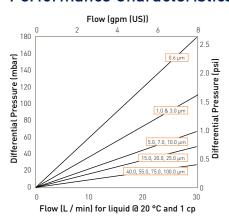
Features and Benefits

- Graded density polypropylene media for high capacity
- Extremely robust to withstand aggressive conditions
- All polypropylene construction
- MURUS and DEMICAP's can be gamma-irradiated and autoclaved



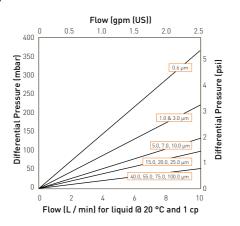
Note: PROCLEAR and DEMICAP are registered trademarks of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10" size differential pressure by 2

10" size (250 mm) Cartridge



For A size for a given flow rate divide B size differential pressure by 2 For E size for a given flow rate multiply B size differential pressure by 2

B size (65 mm) Cartridge and Capsule

Specifications

Materials of Construction

Filtration Membrane:	Polypropylen
Upstream Support:	Polypropylen
Downstream Support:	Polypropylen

Filter Cartridges

Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
End Caps:	Polypropylene
■ End Caps Insert:	316L Stainless Steel
*Not available in R & L endos	an 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:	Silicone

DEMICAP Filter Capsules

Core:	Polypropylen
■ Sleeve:	Polypropylen
■ Capsule Body:	Polypropylen
■ Capsules Vent Seals:	Silicone
■ Filling Bell:	Polycarbonat

Syringe Filters

■ Body: 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:

Temp °C	erature °F	Max. For (bar)	ward dP (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 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 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-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.

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

PROCLEAR PP Filter Cartridges

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.

PROCLEAR PP Filter Cartridges

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).

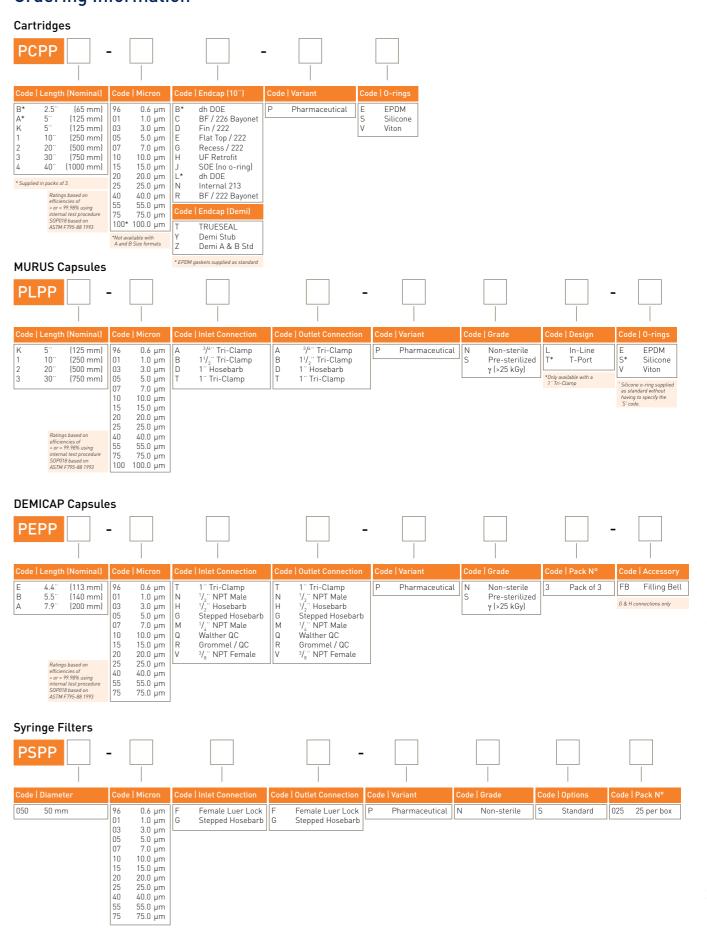
Oxidizable Substances

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

114

PROCLEAR PP Filter Cartridges

Ordering Information



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.

DS_P_03_01/11 Rev. 4A 115

PROPOR BR Filter Cartridges

- liquid filters
- polyethersulphone



PROPOR BR filters have been specifically designed for the fast and cost-effective bioburden reduction of pharmaceutical solutions.

PROPOR BR filters feature an integral meltblown prefilter layer to maximize dirt holding capacity whilst the polyethersulphone membrane guarantees a bioburden log reduction of greater than 5 giving excellent microbial protection. This makes PROPOR BR filters ideal for bioburden reduction of LVPs prior to terminal sterilization.

PROPOR BR filters are also ideally suited to prefiltration and bioburden reduction prior to sterilizing grade membrane filters. The robust construction of PROPOR BR filters guarantees consistent performance on multiple batches.

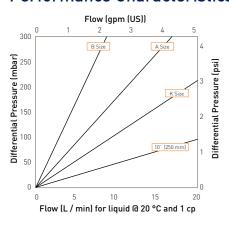
Features and Benefits

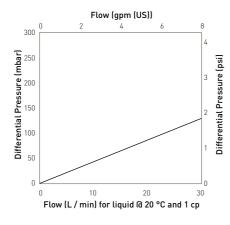
- Brevundimonas diminuta retention of LRV >5 for efficient bioburden reduction
- Additional prefilter layer gives excellent throughput to blockage
- Low binding for minimal product loss
- MURUS and DEMICAP's can be gamma-irradiated and autoclaved

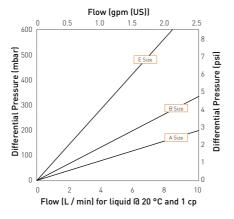


Note: PROPOR and DEMICAP are registered trademarks of Parker domnick hunter

Performance Characteristics







Cartridge flow rates MURUS flow rates (10" Size (250 mm))

DEMICAP flow rates

Specifications

Materials of Construction

■ Filtration Membrane:	Polyethersulphone
■ Prefilter Layer:	Polyester
■ Upstream Support:	Polyester
■ Downstream Support:	Polyester

Filter Cartridges

Inner Support Core:	Potypropyterie
Outer Protection Cage:	Polypropylene
■ End Caps:	Nylon
Fnd Cans Insert	316L Stainless Stee

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:	Polypropylen
■ Sleeve:	Polypropylen
■ End Caps:	Nylon
■ Capsule Body:	Nylon
■ Capsules Vent Seals:	Silicone
Filling Bell:	Polycarbonat

Syringe Filters Body:

Body: 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		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	
I				

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 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):	0.55 m^2	(5.92 ft ²)
K Size:	0.26 m^2	(2.79 ft ²)
A Size:	0.20m^2	(2.15 ft ²)
B Size:	0.10 m ²	(1.07 ft ²)
E Size:	0.05m^2	(0.53 ft ²)
Syringe ø50 mm:	14.50 cm ²	(2.25 in ²)

Sterilization

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

PROPOR BR 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.

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

PROPOR BR Filter Cartridges

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.

Gamma-Irradiation

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

PROPOR BR Filter Cartridges

Performance Characteristics

TOC / Conductivity

The filtrate quality from a 10" (250 mm) PROPOR BR 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) PROPOR BR 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" (200 mm) DEMICAP capsule are <5 mg.

Pharmaceutical Validation

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

Oxidizable Substances

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

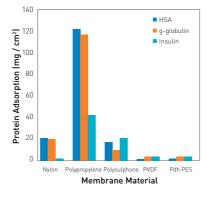
Integrity Test Data

All filters are integrity testable to the following limits when wet with water and using air as the test gas.

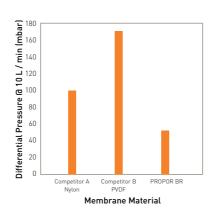
Micron Rating		0.2
Filter Cartridges /	MURUS / DEMICAP	
Min. Bubble Point	(barg)	2.5
	(psig)	36.0
Filter Cartridges /	MURUS / DEMICAP / S	yringe Filters
Diffusional Flow	(barg)	1.7
Test Pressure	(psig)	24.7
Filter Cartridges /	MURUS / DEMICAP / S	yringe Filters
Max. Diffusional FI	ow (10)	16.0
(ml/min)	[K]	7.5
	(A)	6.0
	(B)	2.9
	(E)	1.2

Retention Characteristics

PROPOR BR filter cartridges are validated to an LRV > 5 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) module.



Protein binding on membrane materials

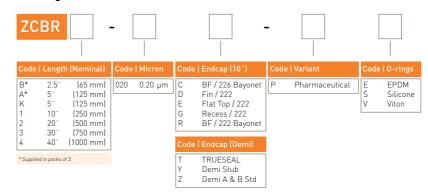


Flow rate comparison for bioburden reduction filters

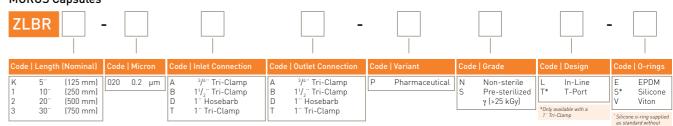
PROPOR BR Filter Cartridges

Ordering Information

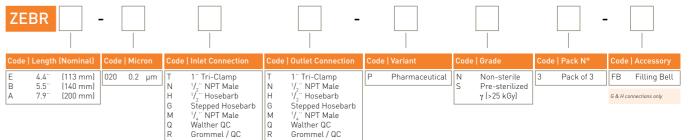
Cartridges



MURUS Capsules



DEMICAP Capsules



Syringe Filters

ZSBR	-		-				
Code Diameter	Code Micron	Code Inlet Connection	Code Outlet Connection	Code Variant	Code Grade	Code Options	Code Pack N°
050 50 mm	020 0.2 μm	F Female Luer Lock G Stepped Hosebarb			N Non-sterile	S Standard	025 25 per box

PROPOR SG Filter Cartridges

- liquid filters
- · polyethersulphone



PROPOR SG sterilizing grade filters feature a microbially retentive polyethersulphone membrane for fast, reliable and cost-effective sterile filtration of pharmaceutical fluids.

The asymmetric pore structure and high voids volume of the PROPOR SG membrane allow high throughputs and exceptionally high flow rates compared with competitive PES and alternative membranes. Low protein and preservative binding properties minimize product loss due to adsorption.

PROPOR SG filters are optimized for pharmaceutical processing. They have low extractable levels and broad chemical compatibility across the full pH range including organic solvents.

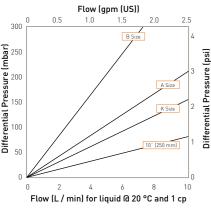
Features and Benefits

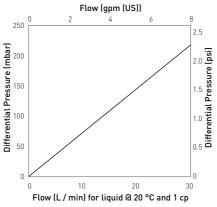
- Up to 3.5 times higher flow rates than competitive sterilizing grade filters
- Fully validated and integrity testable membrane for assurance of sterility
- Low binding for minimal product loss
- MURUS and DEMICAP's can be gamma-irradiated and autoclaved

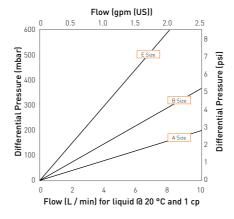


Note: PROPOR and DEMICAP are registered trademarks of Parker domnick hunter

Performance Characteristics







Cartridge flow rates MURUS flow rates (10" Size (250 mm)) 0.2 µm Cartridge 0.2 µm Capsule

0.2 µm Capsule

Specifications

Materials of Construction

Filtration Membrane:	Polyethersulphone	10" (2
■ Upstream Support:	Polyester	K Size
Downstream Support:	Polyester	A Size
		B Size

Filter Cartridges ■ Inner Support Core:

■ Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
■ End Caps:	Nylon
- Fad Casa Jasan	21/1 Ctaiplace Cta

316L Stainless Steel End Caps Insert:

MURUS Disposable Filter Capsules

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

DEMICAP Filter Capsules

Core:	Polypropylene
■ Sleeve:	Polypropylene
■ End Caps:	Nylon
■ Capsule Body:	Nylon
■ Capsules Vent Seals:	Silicone
Filling Bell:	Polycarbonate

Syringe Filters

■ Body: 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. For (bar)	ward dP (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 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):	0.55 m^2	(5.92 ft ²)
K Size:	0.26 m^2	(2.79 ft ²)
A Size:	0.20m^2	(2.15 ft ²)
B Size:	0.10m^2	(1.07 ft ²)
E Size:	0.05m^2	(0.53 ft ²)
Syringe ø50 mm:	14.50 cm ²	(2.25 in ²)

Sterilization

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

PROPOR SG 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.

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.

PROPOR SG Filter Cartridges

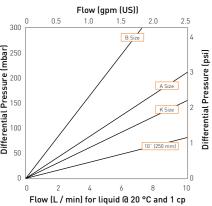
Quality Standards Pharmaceutical grade products are manufactured in accordance with cGMP, 100% flushed with pharmaceutical grade purified water and integrity tested prior to

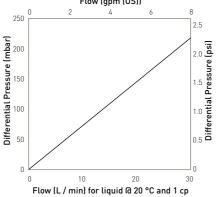
despatch. A sample of each lot is tested to

demonstrate conformity to validated claims.

Gamma-Irradiation

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





DEMICAP flow rates

PROPOR SG Filter Cartridges

Performance Characteristics

TOC / Conductivity

The filtrate quality from a 10" (250 mm) PROPOR SG 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) PROPOR SG 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" (200 mm) DEMICAP capsule are <5 mg.

Pharmaceutical Validation

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

Oxidizable Substances

PROPOR SG 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

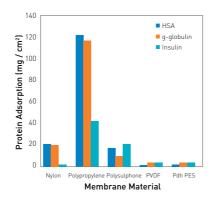
All filters are integrity testable to the following limits when wet with water and using air as the test gas.

Micron Rating		0.1	0.2	0.45
Filter Cartridges / I	MURUS / DEM	ICAP / Syringe	Filters	
Min. Bubble Point*	(barg)	2.36	3.38	2.48
	(psig)	34.2	49.0	36.0
Filter Cartridges / I	MURUS / DEM	ICAP / Syringe	Filters	
Diffusional Flow	(barg)	4.8	2.8	1.7
Test Pressure	(psig)	69.6	40.6	24.9
Filter Cartridges / I	MURUS / DEM	ICAP / Syringe	Filters	
Max. Diffusional Flo	w (10")	27.0	16.0	16.0
(ml/min)	(K)	12.6	7.5	7.5
	(A)	10.1	6.0	6.0
	(B)	4.9	2.9	2.9
	(E)	2.1	1.2	1.2

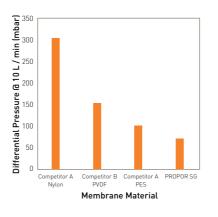
^{*}Bubble point for 0.1 µm product is in 60/40 v/v IPA/Water

Retention Characteristics

PROPOR SG 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.



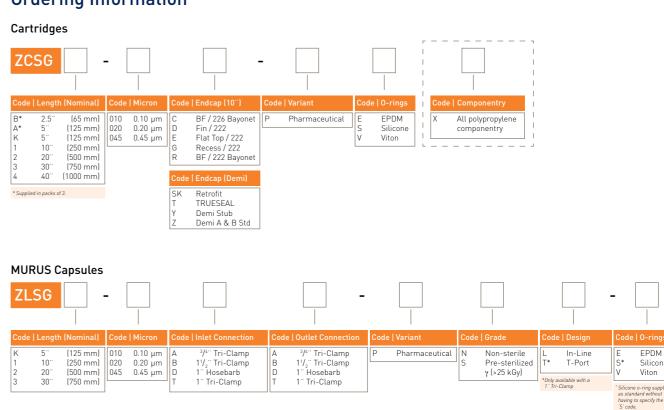
Protein binding on membrane materials

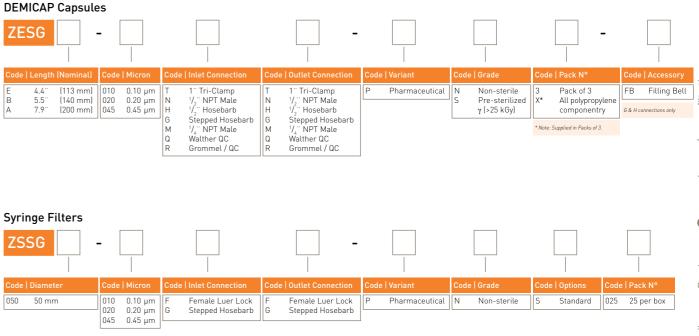


Differential pressure comparison of 10" (250 mm) sterilising grade filters

PROPOR SG Filter Cartridges

Ordering Information





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 conta 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. 122

PROPOR HC Filter Cartridges

- liquid filters
- · polyethersulphone



PROPOR HC sterilizing grade filters have been specifically designed for the effective and economical processing of difficult to filter solutions.

The optimised PROPOR HC PES membrane configuration features a highly asymmetric membrane prefilter layer, which significantly extends throughput and prevents the problems associated with premature filter blockage with complex solutions.

PROPOR HC filters are high capacity and fast flowing. The PES membrane is inherently low binding, which minimizes product loss due to protein or preservative adsorption. The filters have low extractable levels and broad chemical compatibility.

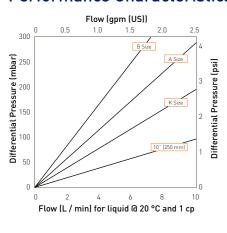
Features and Benefits

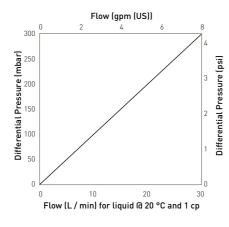
- Optimized membrane configuration allows up to ten times the throughput compared to single layer membrane products
- Integral prefilter layer can condense filter trains for greater processing economy
- Incorporates a fully validated and integrity testable 0.2 micron membrane for assurance of sterility
- Low binding for minimal product loss

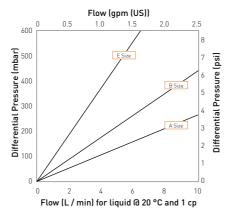


Note: PROPOR and DEMICAP are registered trademarks of Parker domnick hunter

Performance Characteristics







Cartridge flow rates

MURUS flow rates (10" Size (250 mm))

DEMICAP flow rates

Specifications

Materials of Construction

■ Filtration Membrane:	Polyethersulphone
■ Prefilter Membrane:	Polyethersulphone
Upstream Support:	Polyester
Downstream Support:	Polyester

Filter Cartridges

Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
■ End Caps:	Nylon
End Cons Incort.	2141 Ctaiplace Ctaa

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:	Polypropylen
■ Sleeve:	Polypropylen
■ End Caps:	Nylon
■ Capsule Body:	Nylon
■ Capsules Vent Seals:	Silicone
■ Filling Bell:	Polycarbonat

Syringe Filters ■ Body:

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:

erature °F	Max. For (bar)	ward dP (psi)
68	5.0	72.5
104	4.0	58.0
140	3.0	43.5
176	2.0	29.0
194	1.7	24.6
	68 104 140 176	68 5.0 104 4.0 140 3.0 176 2.0

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 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):	0.55m^2	(5.92 ft ²)
K Size:	0.26 m^2	(2.79 ft ²)
A Size:	0.20m^2	(2.15 ft ²)
B Size:	0.10m^2	(1.07 ft ²)
E Size:	0.05m^2	(0.53 ft ²)
Svringe ø50 mm:	14.50 cm ²	(2.25 in ²)

Sterilization

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

PROPOR HC 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.

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

PROPOR HC Filter Cartridges

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.

Gamma-Irradiation

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

PROPOR HC Filter Cartridges

Performance Characteristics

TOC / Conductivity

The filtrate quality from a 10" (250 mm) PROPOR HC 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] PROPOR HC 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

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

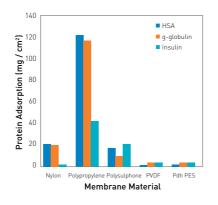
Integrity Test Data

All filters are integrity testable to the following limits when wet with water and using air as the test gas.

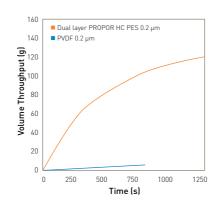
Micron Rating		0.2
Filter Cartridges /	MURUS / DEMICAP / S	yringe Filters
Min. Bubble Point	(barg)	3.4
	(psig)	49.0
Filter Cartridges /	MURUS / DEMICAP / S	yringe Filters
Diffusional Flow	(barg)	2.8
Test Pressure	(psig)	40.6
Filter Cartridges /	MURUS / DEMICAP / S	yringe Filters
Max. Diffusional Fl	ow (10)	18.0
(ml/min)	[K]	8.4
	(A)	6.7
	(B)	3.2
	(E)	1.4

Retention Characteristics

PROPOR HC 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.



Protein binding on membrane materials

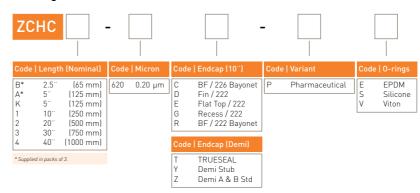


Total volume throughput (g) vs time (s) for an insulin intermediate solution

PROPOR HC Filter Cartridges

Ordering Information

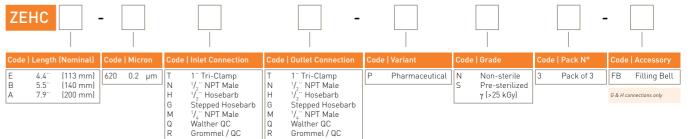
Cartridges



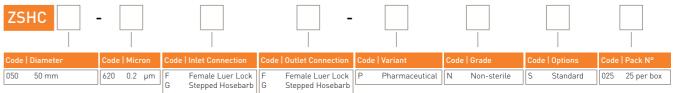
MURUS Capsules

ZL	_HC		-												-	
Code	e Length	(Nominal)	Code	Micron	Code	Inlet Connection	Code	Outlet Connection	Code	e Variant	Code	Grade	Code	e Design	Code	O-rings
K 1 2 3	5" 10" 20" 30"	(125 mm) (250 mm) (500 mm) (750 mm)	620	0.2 µm	A B D T	3/4" Tri-Clamp 1/ ₂ " Tri-Clamp 1" Hosebarb 1" Tri-Clamp	A B D T	3/4" Tri-Clamp 11/2" Tri-Clamp 1" Hosebarb 1" Tri-Clamp	Р	Pharmaceutical	N S	Non-sterile Pre-sterilized γ (>25 kGy)		In-Line T-Port wailable with a Clamp	as stan	EPDM Silicone Viton The o-ring supplied and without to specify the lee.

DEMICAP Capsules



Syringe Filters



PROPOR LR Filter Cartridges

- liquid filters
- polyethersulphone



PROPOR LR filters have been specifically designed for high flow and effective removal of *Ralstonia pickettii* and other diminutive organisms.

A number of studies have concluded that not all microorganisms are removed by 0.2 micron rated membranes under all conditions. PROPOR LR filters use a 0.1 micron rated membrane, which can remove diminutive organisms, while maintaining flow rates typical of a 0.2 micron filtration system.

Ralstonia pickettii is one organism that has frequently been shown to penetrate a 0.2 micron rated membrane and is a common contaminant in purified water systems. PROPOR LR filters have been validated directly against the removal of Ralstonia pickettii.

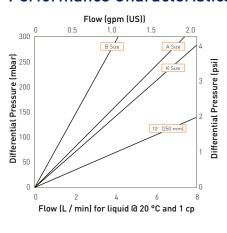
Features and Benefits

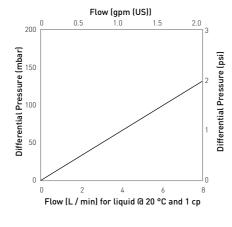
- Fully correlated against Ralstonia pickettii and integrity testable
- Increases retention efficiency whilst maintaining existing 0.2 micron rated system size
- Up to 2.5 times higher flow rate than competitive 0.1 micron rated filters
- MURUS and DEMICAP's can be gamma-irradiated and autoclaved



Note: PROPOR and DEMICAP are registered trademarks of Parker domnick hunter

Performance Characteristics





Cartridge flow rates MURUS flow rates (10" Size (250 mm))

DEMICAP flow rates

Specifications

Materials of Construction

Filtration Membrane:	Polyethersulphone	10" (
■ Upstream Support:	Polyester	K Siz
■ Downstream Support:	Polyester	A Siz
		B Siz

Polypropylene

Filter Cartridges Inner Support Core:

1.1	71 17
Outer Protection Cage:	Polypropylene
■ End Caps:	Nylon
Fnd Cans Insert	316L Stainless Ste

MURUS Disposable Filter Capsules

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

DEMICAP Filter Capsules

DEFINOR FILLOR COPCOLOG	
Core:	Polypropylene
■ Sleeve:	Polypropylene
■ End Caps:	Nylon
■ Capsule Body:	Nylon
■ Capsules Vent Seals:	Silicone
Filling Bell:	Polycarbonate

Syringe Filters

■ Body: 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:

Temp °C	erature °F	Max. For	
-0	-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 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):	0.55 m^2	(5.92 ft ²)
K Size:	0.26 m^2	(2.79 ft ²)
A Size:	0.20m^2	(2.15 ft ²)
B Size:	0.10m^2	(1.07 ft ²)
E Size:	0.05m^2	(0.53 ft ²)
Syringe ø50 mm:	14.50 cm ²	(2.25 in ²)

Sterilization

	Cycles	ociave Temp	Cycles (30 min.)	-in-Place Temp
Cartridges	10	130 °C (266 °F)	30	130 °C (266 °F)
MURUS	5	130 °C (266 °F)	-	-
DEMICAP	10	130 °C (266 °F)	-	-
Syringe	1	130 °C (266 °F)	-	-

PROPOR LR 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.

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.

PROPOR LR Filter Cartridges

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.

Gamma-Irradiation

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

PROPOR LR Filter Cartridges

Performance Characteristics

TOC / Conductivity

The filtrate quality from a 10" (250 mm) PROPOR LR 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) PROPOR LR 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" (200 mm) DEMICAP capsule are <5 mg.

Pharmaceutical Validation

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

Oxidizable Substances

PROPOR LR 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

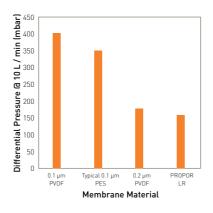
All filters are integrity testable to the following limits when wet with water (diffusional flow) and 60 / 40 : IPA / Water (bubble point) using air as the test gas.

Micron Rating		0.1
Filter Cartridges /	MURUS / DEMICAP	
Min. Bubble Point	(barg)	2.1
	(psig)	30.0
Filter Cartridges /	MURUS / DEMICAP / Sy	ringe Filters
Diffusional Flow	(barg)	4.2
Test Pressure	(psig)	61.0
Filter Cartridges /	MURUS / DEMICAP / Sy	ringe Filters
Max. Diffusional FI	ow (10")	27.0
(ml/min)	(K)	12.6
	(A)	10.1
	(B)	4.9
	(E)	2.1

(Maximum allowable diffusional flows are directly correlated to full retention of Ralstonia pickettii.)

Retention Characteristics

PROPOR LR filters are validated by bacterial challenge testing with Ralstonia pickettii and 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.

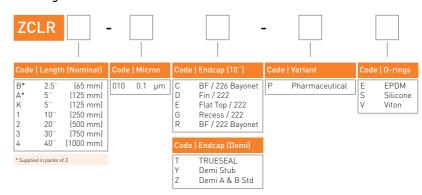


Differential pressure comparison of 10" (250 mm) sterilising grade filters

PROPOR LR Filter Cartridges

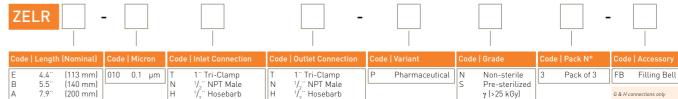
Ordering Information

Cartridges



MU	RUS C	apsules														
ZL	.LR		- [_							- [
Code	Length	n (Nominal)	Code	Micron	Cod	e Inlet Connection	Code	e Outlet Connection	Code	Variant	Code	Grade	Code	Design	Code	0-rings
K 1 2	5" 10" 20"	(125 mm) (250 mm) (500 mm)	010	0.1 µn	A B D	3/4" Tri-Clamp 1¹/₂" Tri-Clamp 1" Hosebarb	A B D	³ / ⁴ " Tri-Clamp 1¹/ ₂ " Tri-Clamp 1" Hosebarb	Р	Pharmaceutical	N S	Non-sterile Pre-sterilized γ (>25 kGy)	L T*	In-Line T-Port	E S*	EPDM Silicone Viton

DEMICAP Capsules



Stepped Hosebart Walther QC

Grommel / QC

Syringe Filters





50 mm 010 0.1 µm Female Luer Lock Female Luer Lock

Stepped Hosebarb

Grommel / QC



TETPOR HP filter cartridges have been specially designed to minimize protein and preservative binding in the sterilization of pharmaceutical and multi-dose ophthalmic solutions.

Adsorption of proteins or preservatives from a pharmaceutical preparation onto the filter membrane can complicate the manufacturing process and lead to costly product wastage. The unique hydrophilic PTFE membrane featured in the TETPOR HP exhibits lower levels of binding than other commonly used filtration membranes such as PES and PVDF which can prevent product loss during processing.

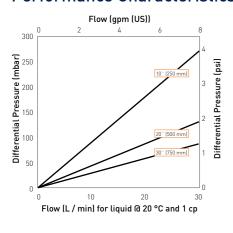
The TETPOR HP exhibits low extractable levels and the sterilizing grade membrane has comparative flow rates to PES and PVDF products. Its hydrophilicity is stable to both chemicals and heat. The product also offers an exceptionally broad range of chemical compatibility making it well suited to the processing of aggressive aqueous liquids.

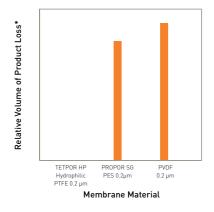
Features and Benefits

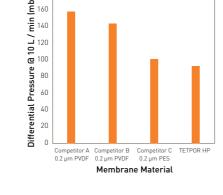
- Exceptionally low binding membrane to prevent costly product loss and process down time
- Incorporates a fully validated and integrity testable 0.2 micron membrane for assurance of sterility
- Fast flowing membrane for increased process efficiency



Performance Characteristics







Cartridge flow rates

Comparison of product loss due to preservative binding on different filter membranes for a 0.001 % solution of benzalkonium chloride (BAK)

The relative volume of product loss represents the volume at which the concentration of BAK in the filtrate recovers back to 95 % of the original concentration, which is typically the point at which the filling operation

Comparison of differential pressure of 10" (250 mm) sterilising grade cartridges filtering water

TETPOR HP Filter Cartridges

Specifications

Materials of Construction

Filtration Membrane: Hydrophilic PTFE
Upstream Support: Polypropylene
Downstream Support: Polypropylene
Inner Support Core: Polypropylene
Outer Protection Cage: Polypropylene
End Caps: Polypropylene
Standard o-rings: Silicone

Recommended Operating Conditions

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

Temp °C	erature °F	Max. Ford (bar)	ward dP (psi)
20	68	5.0	72.5
40	104	4.0	58.0
60	140	3.0	43.5
80	179	2.0	29.0
90	194	1.7	24.6

Effective Filtration Area (EFA)

10" (250 mm)	0.88 m ² (9.47 ft ²)
20" (500 mm)	1.76 m ² (18.94 ft ²)
30" (750 mm)	2.64 m ² (28.42 ft ²)

Sterilization

TETPOR HP filter cartridges are validated to withstand 10 steam-in-place cycles at 135 °C (275 °F).

TETPOR HP 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.

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.

TOC / Conductivity

The filtrate quality from a 10" (250 mm) TETPOR HP 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) TETPOR HP contain < 0.25 EU / ml when tested in accordance with the Limulus Amoebocyte Lysate test.

Non-Volatile Extractables (NVE)

The quantity of NVE's obtained from a TETPOR HP cartridge during a 24 hour static soak was undetectable compared to a control sample.

Oxidizable Substances

TETPOR HP 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 water and using air as the test gas (a minimum 20 minute purified water flush is recommended prior to integrity testing in water).

Micron Rating		0.2
Min. Bubble Point	(barg)	1.5
[60 / 40 IPA / Water (v/ v)]	(psig)	21.0
Diffusional Flow	(barg)	2.2
Test Pressure	(psig)	31.9
Max. Diffusional Flow	v*(10'')	37.0

*Note: It is also possible to integrity test the TETPOR HP in 60/40 IPA/Water (v/v). Maximum allowable diffusional flow for a 10" (250 mm) TETPOR HP in 60/40 IPA/Water is 16.8 ml/min

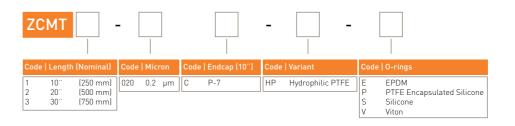
Retention Characteristics

TETPOR HP 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 1011 (250 mm) module.

Pharmaceutical Validation

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

Ordering Information



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 conta 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.

TETPOR LIQUID Filter Cartridges

- liquid filters
- PTFE



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

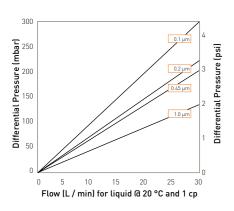
Features and Benefits

- Superior chemical resistance of PTFE membrane combined with polypropylene hardware
- Integrity tested prior to despatch
- Validated to ASTM F838-05 methodology
- Comprehensive range of end cap configurations for retrofitting



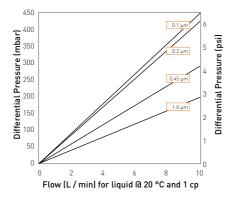
Note: TETPOR is a registered trademark of Parker domnick hunter

Performance Characteristics



For K size for a given flow rate multiply 10 $\!\!^{\circ}$ size differential pressure by 2

10" Size (250 mm) Cartridge



For A size for a given flow rate divide B size differential pressure by 2 For E sie for a given flow rate multiply B size differential pressure by 2

B Size (65 mm) Cartridge and Capsule

Specifications

Materials of Construction

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

Filter Cartridges

Inner Support Core:	Polypropylene
Outer Protection Cage:	Polypropylene
■ End Caps:	Polypropylene
■ End Caps Insert:	316L Stainless Stee
*Not available in B endcap var	iant

■ Standard o-rings/gaskets: Viton

MURUS Disposable Filter Capsules

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

DEMICAP Filter Capsules

Core:	Polypropylen
■ Sleeve:	Polypropylen
■ End Caps:	Polypropylen
■ Capsule Body:	Polypropylen
■ Capsules Vent Seals:	Silicone
Filling Bell:	Polycarbonat

Syringe Filters

■ 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 °C	erature °F	Max. For (bar)	ward dP (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 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):	0.77m^2	(8.28 ft ²)
K Size:	0.36m^2	(3.87 ft ²)
A Size:	0.25m^2	(2.69 ft ²)
B Size:	0.12m^2	(1.29 ft ²)
E Size:	0.06m^2	(0.64 ft^2)
Syringe ø50 mm:	14.50 cm ²	(2.25 in ²)

Sterilization

		oclave		n-in-Place
	Cycles		Cycles (30 min.)	
Cartridges	120	142 °C [287.6 °F]	120	142 °C [287.6 °F]
MURUS	5	130 °C (266 °F)	-	-
DEMICAP	10	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.

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.

TETPOR LIQUID Filter Cartridges

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.

TETPOR LIQUID Filter Cartridges

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" (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 / DEMI	CAP / Syringe	Filters		
Min. Bubble Point	(barg)	1.3	1.0	0.7	-
	(psig)	18.8	14.5	10.1	-
Filter Cartridges /	MURUS / DEMI	CAP / Syringe	Filters		
Diffusional Flow	(barg)	1.0	0.8	0.4	-
Test Pressure	(psig)	14.5	11.6	5.8	-
Filter Cartridges /	MURUS / DEMI	CAP / Syringe	Filters		
Max. Diffusional Flo	w (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	-

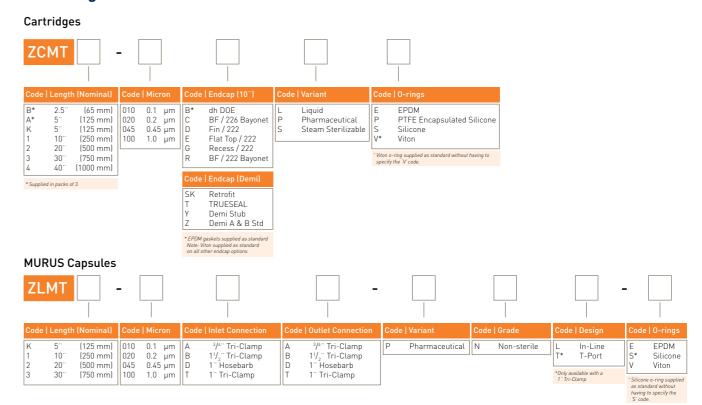
Retention Characteristics

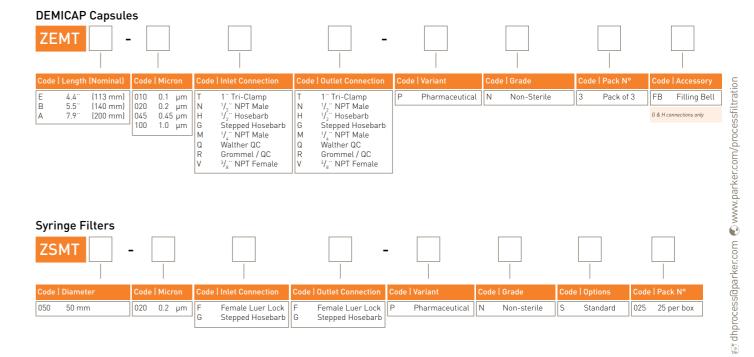
TETPOR LIQUID filter cartridges are validated by bacterial challenge testing with *Brevundimonas diminuta* to current ASTM F838-05 methodology (10⁷ organisms / cm² EFA minimum) with typical in-house challenge levels being 10¹¹ organisms per 10" (250 mm) filter cartridge.

136

TETPOR LIQUID Filter Cartridges

Ordering Information





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A dedicated housing range

That can be customized to meet the demands of your application



Parker domnick hunter manufacture stainless and carbon steel pressure vessels that are designed to meet International industry standards as well as specific customer application requirements.

A combination of highly skilled employees, dedicated manufacturing facility and nearly 50 years experience of supplying process industries around the world, Parker domnick hunter provide solutions that match your requirements for performance, quality and value.

Our fabrication facility manufactures a standard range of stainless steel housings to support our range of filters, which can be modified and adapted to meet any process requirements. Our strength is in providing a range of products that meet industry requirements and a flexibility to meet your own requirements.

Manufacturing best practice

- ISO9001
- ISO13485
- ISO14001

Vessels built to industry standards

- PED (CE)
- EN/B445
- EN / 286
- EN / 1210
- ATEXPD5500
- ASMEU
- ASME BPE

Stamp of approval

- Certificate of Authorization (U stamp)
- National Board Certificate of Authorization
- American Society of Mechanical Engineers

- Air, gas and liquid housings
- Single and multi rounds
- Multi housing skid systems
- Dedicated industry specific range
- Custom options to meet application needs
- Silicone rubber heating jackets
- Single cartridge polypropylene / nylon housings







A dedicated housing range

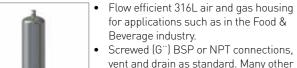


Flow efficient higher specification 316L sanitary air and gas housing for applications such as in the Pharmaceutical industry.

Sanitary tri-clamp connections, gauge port, drain and body closure as standard. Many other options in the PLUS range.

Sanitary 'C' style or demi TRUESEAL filter cartridge locations.

HSA



options available in the PLUS range, including higher specification sanitary design features.

Sanitary 'C' style or demi TRUESEAL filter cartridge locations.

HBA



Flow efficient 316L sanitary 'L' Port (side and base) air housing typically for tank venting applications.

Sanitary tri-clamp connections and body closure as standard. Other options available in the PLUS range. including electro-polished finish and integrity test sockets.

Sanitary 'C' style or demi TRUESEAL filter cartridge locations.

HSVLP



 Flow efficient 316L sanitary in-line housing for air, gas and liquid for applications in the Food & Beverage and Pharmaceutical industry.

Sanitary tri-clamp connections, gauge port and body closure as standard. Many other options in the PLUS range.

Sanitary 'C' style or demi TRUESEAL filter cartridge locations.

HSI



HIF

• Liquid Demi 316L housing for small scale applications.

Screwed (G") BSP or NPT connections. vent and drain as standard. Many other options available in the PLUS range.

'Z' style single internal o-ring filter cartridge locations.



Flow efficient sanitary 316L demi air and gas housing for smaller scale applications in the Food & Beverage and Pharmaceutical industry.

Sanitary tri-clamp connections, gauge port, drain and body closure as standard. Many other options in the PLUS range. 'Z' style single internal o-ring filter

HSA PLUS 'Z' STYLE



Flow efficient 316L sanitary open air housing typically for tank venting applications.

cartridge locations.

Sanitary tri-clamp connections and body closure as standard. Many other options, including higher specification surface finishes in the PLUS range.

Sanitary 'C' style or demi TRUESEAL filter cartridge locations.

HSV



Flow efficient 316L sanitary liquid housing for applications in the Food & Beverage and Pharmaceutical industry.

Sanitary tri-clamp connections, gauge port and body closure as standard. Many other options in the PLUS range.

Sanitary 'C' style or demi TRUESEAL filter cartridge locations.

HSL



Liquid 316L housing for prefiltration and industrial applications.

Screwed (G") BSP or NPT connections, vent and drain as standard. Many other options available in the PLUS range.

B (DOE) or D (222) style filter cartridge locations.

HII



- Liquid polypropylene, polycarbonate or nylon housing.
- Screwed (G") BSP connections, vent and drain as standard.
- B (DOE) and N or J style single internal o-ring filter cartridge locations.

ZVP (PLASTIC)



HCA

Enhanced PLUS flow efficient Alloy 22 (none wetted parts 316) air and gas housing for aggressive applications such as chemical synthesis in the Pharmaceutical industry.

Flanged connections with screwed (G), BSP or NPT vent and drain and tri-clamp body closure as standard. Other options available. Sanitary 'C' style filter cartridge



• Enhanced PLUS higher pressure 316L air and gas housing for applications such as in the Food & Beverage industry. 25 barg (363 psig) and 40 barg (580 psig) variants available.

 ANSI, ISO, [G"]BSP or NPT connections, vent and drain as standard. Other options available. Sanitary 'C' style filter cartridge

locations.



Single and 3 round flow efficient 316L steam housing for applications such as in the Industrial Biotech and Food & Beverage industry.

 Weld end or flanged connection, screwed (G") BSP or NPT, vent and drain as standard. Many other options available.

Sanitary J (Jumbo) style filter cartridge locations.

VISCE



Multi round flow efficient 316L sanitary liquid housing for applications in the Pharmaceutical industry.

 Sanitary tri-clamp connections, gauge port and body closure as standard. Sanitary screwed integrity test / sample

and drain connections. Many other options available. Sanitary 'C' style filter cartridge



SKIDS

VSLCE (MULTI)

 Custom design - Parker domnick hunter offers a specialist design and fabrication service allowing individual customer system specifications to be met.



(none wetted parts 316) liquid housing for aggressive applications such as chemical synthesis in the Pharmaceutical industry. Flanged connections with screwed (G), BSP or NPT vent and drain and tri-clamp body closure as standard.

Enhanced PLUS flow efficient Alloy 22

Other options available. Sanitary 'C' style filter cartridge

HCI



• Multi-round flow efficient 316L air and gas housing for applications such as in the Industrial Biotech and Food & Beverage industry.

Weld end or flanged connection, screwed (G") BSP or NPT, vent and drain as standard. Many other options available.

Sanitary 'C' style filter cartridge locations.

ZVACE (MULTI)



Multi-round flow efficient 316L sanitary liquid housing for applications in the Food & Beverage industry.

Many connection options as standard. Sanitary tri-clamp gauge port and drain connections (or no drains). Many other options available.

Sanitary 'C' style filter cartridge Incations

VSHCE (MULTI)



• Multi liquid 316L housing for prefiltration and industrial applications.

Screwed (G") BSP or NPT connections, vent and drain as standard. Many other options available in the PLUS range.

• B (DOE) or D (222) style filter cartridge locations.

VII CF (MULTI)



vent applications.

inter-connection wiring.

Accurate temperature control using PT100 temperature sensor with additional thermal cut out at 150 °C (302 °F).

Silicone or PTFE Glass Silk heating

jacket with integrated control unit and

HEATING **EQUIPMENT**

For more information on Parker domnick hunter's complete housing range. please contact your local Parker domnick hunter representative for a copy of the latest technical literature.

Integrity testing equipment



Whatever your industry, integrity testing plays a vital role in ensuring the performance and sterility of your process filters. The ability to integrity test a filter provides a valuable tool to gauge, not only performance of your process, but also the quality and safety of your final product. A properly conducted integrity test provides assurances that the filter will fulfil the role it was designed for ensuring your production process runs to its maximum potential.

Integrity testing of sterile grade filters is a fundamental requirement of critical process applications. FDA guidelines require integrity testing of filters used in the processing of sterile solutions. It is vital producers ensure the quality and biological safety of the product that reaches the customer. Increased shelf-life, reputation and customer well being are of paramount importance

Parker domnick hunter, have a range of instruments that have been specifically designed to meet the demands of your industry. All instrumentation is supported by our global team of dedicated instrument service Engineers on hand to provide validation, installation and performance guarantees.



Aerosol challenge

This methodology uses a high concentration of aerosol in the most penetrating particle size (MPPS) of 0.2 - 0.3 μ m. The MPPS is a function of the particle challenge for air filters.

During the test the filter system is challenged with 10° aerosol particles. The latest in laser particle detection technology measures the percentage penetration through the test system. The test is directly correlated to aersol challenges with live *Brevundimonas diminuta* and *E-coli* phage. A positive result shows that the test filter is providing bacterial and viral removal when used in gas. The integrity test method of VALAIRDATA II is unique to Parker domnick hunter and is the only integrity test method for gas filters to simulate actual filter use.

Bubble point testing

The bubble point test measures the pressure that is required to expel a wetting fluid from the largest pore in a wetted membrane. Historically this was a visual assessment indicated by bubbling on the downstream side of

the membrane, hence the term 'bubble point'. The test is typically applied to smaller filters and to remove subjectivity is now conducted using automated integrity testers.

Water intrusion

Water intrusion testing is based on the measure of the intrusion or flow of water into the pore structure of a hydrophobic filter membrane, under an applied test pressure. The flow is measured, with the test result / limit being directly correlated to the ASTM standard for a sterilizing grade filter.

Diffusional flow

The diffusional flow test measures the volume of a diffusive gas flow across a wetted membrane, under an applied test pressure. This method can be utilized to test both hydrophilic and hydrophobic membrane filters.

Diffusional flow test results are directly correlated to live bacterial challenges using industry standard organisms. For a 0.2 micron sterilizing grade filter this challenge procedure is defined in ASTM F838-05.

RESULTS:

Test pressure:
1695 mbar

Pressure drop:
12 mbar

Diffusion:
11.0 ml/min

Bubble Point:
3111 mbar

TEST PASSED

Programmer:
PR

2480 mbar





VALAIRDATA II

The most efficient test for sterile gas filters



Since 1990 and the launch of the unique VALAIRDATA aerosol integrity test system, the aerosol test method has become widely accepted in a variety of applications and industries as a routine method for integrity testing air filtration systems. The VALAIRDATA II integrity test instrument is a second generation design offering further practicality in air filter testing.

The VALAIRDATA II combines the sound principles of aerosol testing, as recommended in the 'PDA's Sterilizing Filtration of Air - Technical Report #40', with a compact, portable and ergonomic design reducing test times and improving multi cartridge system sensitivity.

The VALAIRDATA II aerosol test is correlated to an aerosolised *Brevundimonas diminuta* and bacteriophage (such as *Enterobacteria phage MS2*) challenge.

Aerosol methods are rapid, can identify filter non-integrity on very large systems, allow immediate use of filter systems after testing as drying is not required and provides direct measurement of filter performance for gas filters.

- 30 second test time for a single 10" (250 mm) cartridge challenge
- Results correlated to aerosol bacterial and viral challenge
- Increased sensitivity compared to liquid based tests especially on multi-cartridge systems
- Built-in test instrument system integrity check
- Well established with over 200 current VALAIRDATA II users

- Fully validated secure option design to GAMP 4 Guidelines and meets the FDA's 21CFR11 requirements
- Stores up to 200 test results and supported with software for PC download
- PDA recommended for use where filtered gas not in direct contact with exposed sterile product or surfaces



Physical Parameters

Instrument Material
Instrument Size
Weight
Ingress Protection Class
Power Supply
Keyboard
Inlet Pressure Required
Operating Temperature
Pneumatic Connectors
Ambient Humidity
Languages
Programmed Tests
Storable Test Programmes

Moulded Robust Polyurethane Case & Non-Slip Feet 363 mm x 155 mm x 308 mm : 14.3" x 6.1" x 12.1" 8 Kg : 18 lb

IP45

Re-chargeable Battery (12V / 3.8 Ah) & Mains (90 - 230 VAC : 50 / 60 Hz)

16 Tactile Keys with Alphanumeric Input 3.5 - 7.0 barg (50 - 100 psig) (60 Al / min)

5 - 37 °C (40 - 95.6 °F)

Rectus 21 KA Series

- 95% RH (non-condensing)

English, French, German, Spanish, Italian, Danish, Portugese & Swedish

Up to 100

Instrument Options

	Standard	Secure Environment	Electronic Signature
PC Manager Software	ST - Standard	SE - Secure Environment	ES - Electronic Signature
PC Operating Platforms	Microsoft Windows 7, 98, 2000,	Microsoft Windows 7, 98, 2000,	Microsoft Windows XP
	NT & XP	NT & XP	
Design Environment Approvals	Hardware & Software	GAMP Hardware & Software	GAMP Hardware & Software
	Development to GAMP Guidelines	Development	Development
		21CFR11 Compliant	21CFR11 Compliant
		(PC data is users responsibility)	
Operator (max. 40)	Open Access	Access Password & PIN	Access Password & PIN
Access ADMINISTRATOR	Open Access	Access Password & PIN	Access Password & PIN
Record Output	RS232 Transfer	RS232 Transfer	RS232 Transfer
Audit Trail Record	No	Yes	Yes

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PORECHECK IV

The perfect choice for the pharmaceutical industry



Parker domnick hunter, in conjunction with the pharmaceutical industry has reviewed the limitations and benefits of current integrity test equipment. This review has led to the development of the PORECHECK IV integrity test system which has been specifically designed with the needs of routine production users in mind.

The PORECHECK IV is configured for water intrusion testing, pressure decay and bubble point testing.

The PORECHECK IV comes in two versions:

'P' Pharmaceutical (CFR)

allows traceability and audit tracking capability

'C' Certified

- comes with password level protection

This market leading system incorporates a range of design features unique to the PORECHECK IV bringing true portability, enhanced ease of use, flexibility and reliability in challenging environments. All this within an instrument fully compliant with 21 CFR Part 11.

- Designed to 21 CFR Part II and Annex II compliant environments
- Automatic compensation when used on housings located 10 metres above instrument
- Maintains resolution and accuracy regardless of filter system size 0.1 to 150 litres
- Highly portable and mains independent

- Configurable to automatically flush and drain filters
- Robust waterproof stainless steel casing
- Direct attachment to test disposable capsules
- 100 storable test programs defined in blocks



Physical Parameters

÷	
	Instrument Material
	Instrument Size
	Weight
	Ingress Protection Class
	Power Supply
	Keyboard
	Inlet Pressure Required
	Test Pressure Range
	Pneumatic Connectors
	Storage Temperature
	Ambient Humidity
	Display
	Printer
	Languages
	Software Protection
	Storable Test Programs

Stainless Steel 1.4301 (AISI 304)

200 mm x 300 mm x 155 mm : 7.9" x 11.8" x 6.1"

8.6 Kg : 20 lb
IP54

Re-chargeable Battery (12V / 3.8 Ah) & Mains (90 - 230 VAC : 50 / 60 Hz)
Remote Infrared - Alpha Numeric & Instrument Keypad - Numeric
6.5 - 8.0 barg (94 - 116 psig)
350 mbar to 6 barg (87 psig)
Stäubli RBE 0.3 Style : Stainless Steel 1.4404 (AISI 316L)
2 - 50 °C (35.5 - 122 °F)
1 - 80% RH
LCD - 20 Character x 4 Lines - Back Lit
Internally Housed Impact Dot Matrix , 24 Characters per Line
English, French, German, Spanish, Italian & Danish
Stored in Flash - EPROM
Up to 100 (in Flash - EPROM) Stored in 10 Blocks of 10 Programs

Test Accuracy

	Standard		High Pressure
Water Intrusion Measurement Range (ul / t)		100 - 99999	
Resolution (µl)		5	
Accuracy (for a 10" cartridge @ 4000 µl / min)	3%		6%
Test Pressure (mbar)		350 - 4000	
Stabilisation Time		60 - 999 secs	
Test Time (t)		30 - 999 secs	
Hardware Volume (ml)		1 - 32000	
Diffusional Flow Measurement (ml / min)		1 - 999	
Resolution (ml / min)		0.1	
Accuracy (for a 10" cartridge @ 16 ml / min)	3%		6%
Test Pressure (mbar)	350 - 4000		350 - 7000
Stabilisation Time		60 - 999 secs	
Test Time (t)		30 - 999 secs	
Upstream System Volume (ml)		1 - 32000	
Bubble Point Measurement Range (mbar)	450 - 3900	(min. 100 mbar above DF test pressure)	450 - 7900
Resolution (mbar)	1		2
Accuracy		1& FS	

Instrument Options

Storable Test Records
USER Accounts
Access USER
Access PROGRAMMER
Access ADMINISTRATOR
Record Output
Audit Trail Record

'P' Pharmaceutical	'C' Certified	Documentation
40	No	Installation, Operating & Maintenance Manual
25	Unlimited	Checklist of Supplied Components
Access Password & PIN	Open Access	Calibration & Pressure Vessel Certification
Access Password & PIN	Access Password	CE Declaration of Conformity
Access Password & PIN	Access Password	Operational Qualification Support Documentation
Printed Records & RS232 Transfer	Printed Record Test Result Only	Laboratory Qualification Results
256 Event Audit Trail	No	Suggested OQ Test Protocol

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BEVCHECK & BEVCHECK PLUS

Monitoring performance and product quality



BEVCHECK

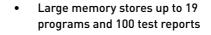
Simple routine integrity testing for the beverage industry BEVCHECK is an easy to use, portable unit that allows you to test the integrity of your membrane filters using the pressure decay method. Test data can be reported as pressure decay or diffusional flow.

BEVCHECK is a small hand held unit, or is light enough to be mounted directly on to a connection on the filter housing. Software included with the unit enables it to be connected to a pc for enhanced programming and data handling flexibility.

BEVCHECK PLUS

Provides an automated method for testing membrane filter cartridges used in beverage applications. Using the pressure decay method, the unit controls the whole test from increase of pressure, through stabilization and pressure decay measurement, to release of pressure.

Test data can be reported as pressure decay or diffusional flow and is provided in a printed summary. The unit is small enough to be portable around the production facility, or can be positioned centrally for remote connection to the filter housings.



- Flexible suitable for use with compressed air or nitrogen
- Accommodates a wide range of filter retention ratings and housing sizes
- Clear liquid crystal display and wipe clean keypad
- Self test function automatically checks the function of the unit

- PC interface and software provides additional programming and data handling flexibility
- IP53 protection class
- Hand held portability with rechargeable battery operation
- Convenient built-in printer provides printed test report (PLUS)





Physical Parameters

	nstrument Size
	Veight ngress Protection Class
	Power Supply
•	ower Supply
E	Battery Life (From Full Charge)
k	Keyboard
l	nlet Pressure Required
)peration Temperature
F	Pneumatic Connectors
c	Storage Temperature
	Ambient Humidity
	Display
	Printer
Ĺ	anquage
	Storable Test Programs
	Storable Test Records
7	est Pressure Control
T	est Pressure Range
ŀ	lousing Volume Range
	Diffusional Flow Range
2	Stabilisation Time Range
	est Time Range
h	nterfaces

ABS (WxDxH) 105 mm x 210 mm x 45 mm (4" x 8.25" x 1.75") 0.5 Kg (1.1 lbs) IP53

Re-Chargeable HiMH Battery (4.8 V / 1.5 Ah) & External Charger (100- 230V AC / 47 - 63 Hz / 7.5V 1.33A) 7 hours Typ.

BEVCHECK

16 Key - Polycarbonate Keypad 0 - 4000 mbar 3 - 33 °C (37.4 - 91.4 °F)

Compressed Air / Filter : Rectus 21 Male

3 - 35 °C (37.4 - 95 °F) 5 - 95% Rel. LCD - 16 Character x 2 Lines

English, German, Italian, French, Spanish & Portugese

100 Manual (Additional Accessory Kit Required) 0 - 4000 mbar

10 - 999999 ml

1 - 99.9 ml / min 1 - 1800 secs

1 - 1800 secs PC Data / Remote Operation :

RS232 4-Pole Jack
CE Declaration of Conformity
Calibration Certificate

Winfilter PC Software
Power Supply /
Charger with Courty Specific Mains Adaptor

Charger with Country Specific Mains Adaptor
PC Comms Cable (RS232 - 4 Pole Jack to 9 Pin Male)
Installation, Operation & Maintenance Instructions (IOMI)
Foam Lined Carry Case

BEVCHECK PLUS

(WxDxH) 315 mm x 280 mm x 150 mm (12.5"x 11" x 6") 3.9 Kg (8.6 lbs)

HiMH Battery (4.8 V / 1.5 Ah) & External Charger / Mains

[230V AC:18V DC, 1.7A / 230V AC:15V AC, 15VA] 2 hours Typ. 16 Key - Polycarbonate Keypad

0 - 4500 mbar 3 - 30 °C (37.4 - 95 °F) Compressed Air / Filter : Festo 4 mm

Stäubi RBE03 Male Vent : Festo 4 mm 3 - 35 °C (37.4 - 86 °F) 5 - 95% Rel.

LCD - 20 Character x 4 Lines

Built in Thermal Printer - 57 mm Printer

English, German, Italian, French, Spanish & Portugese
19

Fully Automatic 0 - 3900 mbar 10 - 999999 ml

1 - 999.9 ml / min 1 - 1800 secs 1 - 1800 secs

D-Sub 25 Pole PC Data / Remote Operation RS232 9-Digit Male

CE Declaration of Conformity

Calibration Certificate

Calibration Certificate Winfilter PC Software Power Supply /

Charger with Country Specific Mains Adaptor PC Comms Cable (RS232 - 4 Pole Jack to 9 Pin Male) Installation, Operation & Maintenance Instructions (IOMI) Foam Lined Carry Case

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Filter Discs

- liquid filters
- various membrane / media



Process and analytical filter discs from Parker domnick hunter are available in a range of pore size and a choice of five materials.

Membrane discs:

- Cellulose mixed esters
- Polyethersulphone
- Nylon

Fibrous media discs:

- Glass microfibre
- Polypropylene

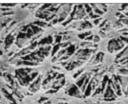
The discs are supplied interleaved between two protecting layers with the feed surface oriented upwards in the box.

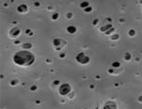
Reduced filtration time

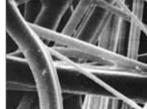
Low protein binding

Features and Benefits

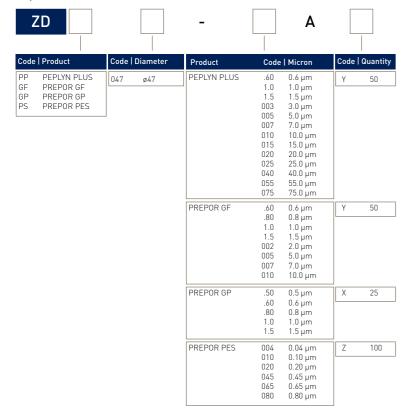
- High throughput rates
- Superior flow characteristics
- Easy to handle

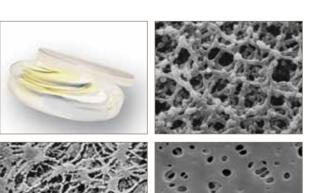


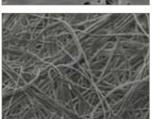




Liquid Filters

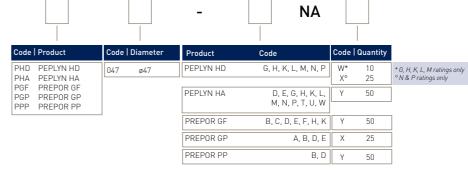






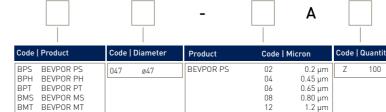
Ordering Information

Beverage Prefilters



Beverage Final Filters

BMH BEVPOR MH



	08 12	0.80 µm 1.2 µm		
BEVPOR PH	02	0.2 µm	Υ	50
	04	0.45 µm		
	06	0.65 µm		
	08	0.80 µm		
	12	1.2 µm		
BEVPOR PT	02	0.2 µm	Z	100
	04	0.45 µm		
	06	0.65 µm		
BEVPOR MS	02	0.2 µm	Z	100
	04	0.45 µm		
	06	0.65 µm		
	08	0.80 µm		
	12	1.2 µm		
BEVPOR MT	02	0.2 µm	Z	100
	04	0.45 µm		
	06	0.65 µm		
BEVPOR MH	02	0.2 µm	Υ	50
	04	0.45 µm		- "
	06	0.65 µm		
	08	0.80 µm		

12

1.2 µm

Pharmaceutical Filters

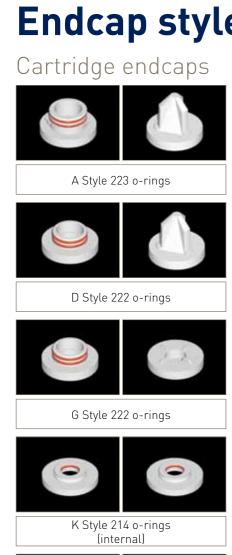
ZD -							
Code Product	Code	Diameter	Product	Code	Micron	Code	Quantity
BR PROPOR B	04/	ø47	PROPOR BR	020	0.20 µm	Y	50
HC PROPOR H LR PROPOR LI	C R		PROPOR SG	010 020 045	0.10 µm 0.20 µm 0.45 µm	Z	100
			PROPOR HC	620	0.20 µm	Υ	50
			PROPOR LR	010	0.1 µm	Z	100
			TETPOR LIQUID	010 020	0.1 µm	Z	100
				045 100	0.2 μm 0.45 μm 1.0 μm		

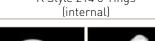
Standard diameters 047 mm.

Diameters 025mm, 090 mm & 142 mm are also available.

For full ordering information, variants, quantities and availability, please contact Parker domnick hunter.

Endcap styles







R Style 222 o-rings



T Style 126 o-rings (demi only)



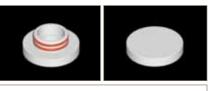
X Style 116 o-rings (demi only)



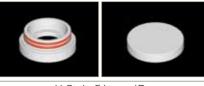
Demi H Style 217 o-rings (demi only)



B, L Style Flat Gaskets



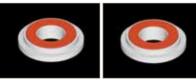
E Style 222 o-rings



H Style 54 mm ID x 4 mm o-rings



N Style 213 o-rings (internal)



S Style Flat Gaskets



U Style 222 o-rings



Y Style 116 o-rings (internal) (demi only)



C Style 226 o-rings



F Style 216 / 218 o-rings



J Style S.O.E.





P Style 227 o-rings





SK Style (demi only)



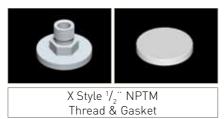


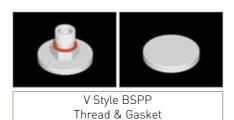
W Style 111 o-rings (demi only)

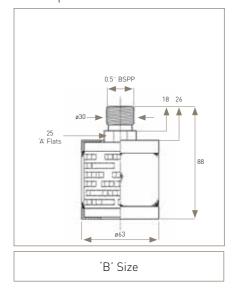


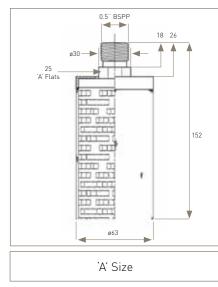
Z Style 116 o-rings (internal) (demi only)

Vent autoclave filter endcaps and dimensions





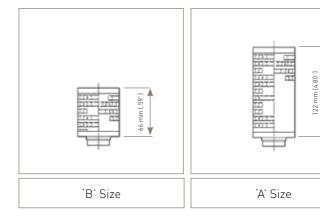


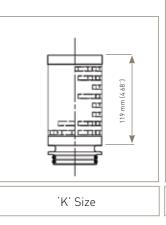


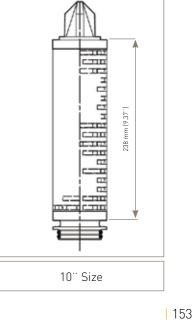
Endcap cross reference chart

Parker domnick hunter	PA	MI	SA
В	MCY 10"	F	23
C (10" Size)	7	7	25
C (K Size)	2		
D	8	5	26
E/G	E = 3 / G = 25	0	27
F	MYS	8	24
L	MCY 20" and above	F	23
R			28
X			
Υ	MCY2230		
Z	MCY2230 / 4463		

Cartridge dimensions







DEMICAP styles



Stepped Hosebarb (Code G)



 $^{1}/_{2}^{\cdot \cdot \cdot}$ Hosebarb (Code H)



1/4" NPT (Code N)



¹/₂" NPT (Code M)



Walther Male (Code Q)

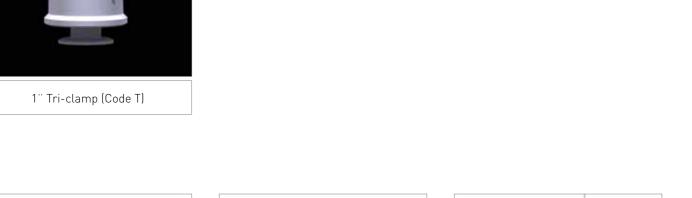


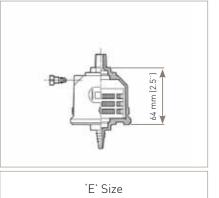
Gromelle (Code R)

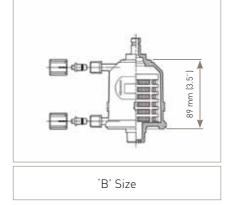


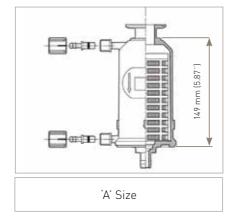


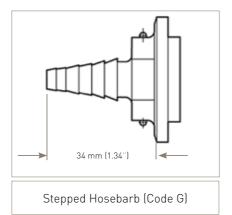


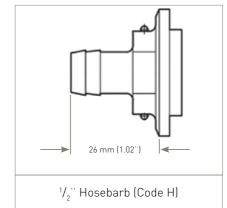


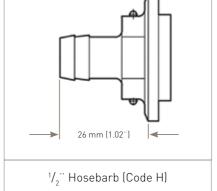


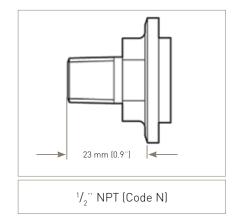


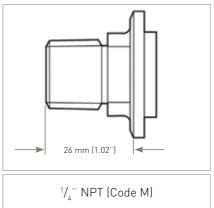


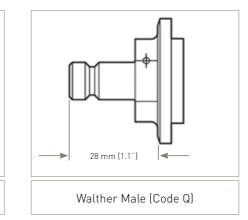


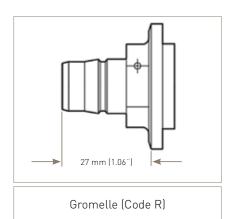


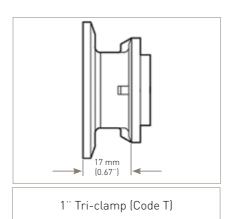












MURUS and syringe styles

Large scale disposable inlet / outlet connection styles







1" Hosebarb

3/, "Tri-clamp

1" Tri-clamp





В

11/3" Tri-clamp

(1" Tri-Clamp only)

'A' 13.07" 332 mm Cartridge Type 10.30" 262 mm 579 mm

Syringe filters



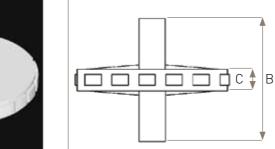
Stepped Hosebarb

uitable for tubing with 6 mm (' 12 mm ('/¸'') internal diamete



Luer Slip Male





Luer Loc Female

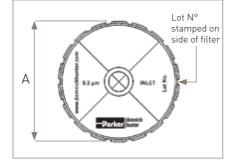
0.98" 25 mm 1.12" 28.5 mm 0.31" 8.0 mm 1.96" 50 mm 2.12" 54.0 mm 0.31" 8.0 mm







5/16 Hosebarb 1/。" BSPM Thread



Example of Syringe filter marking

Installation and operating guidelines

For liquid and gas filter cartridges

These guidelines give the correct methods for using liquid and gas filter cartridges manufactured by Parker domnick hunter. If you have any queries, our process filtration specialists will be pleased to discuss your particular filtration requirements or answer any questions you may have. We may also be contacted at any of the addresses given on the reverse of this document or through our worldwide network of subsidiary companies and distributors.

1. Storage

- 1.1 Store cartridges in a clean and dry environment and avoid placing heavy objects on the top of the cartridge tube or packaging. The cartridges should not be exposed to temperatures below 5 °C (41 °F) or above 40 °C (104 °F) or to direct sunlight
- 1.2 Keep the cartridge in it's sealed polyethylene bag until it is time to install it.
- 1.3 The shelf-life for cartridge filters is as follows:-

ASYPOR membrane variants - 2 years Liquid membrane cartridges - 3 years Liquid depth cartridges - 5 years

TETPOR membrane variants - 5 years Gas membrane cartridges

- 5 years Gas depth cartridges - 5 years Gamma irradiated cartridges - Consult Certificate

of Conformance

2. Installation

The various cartridge formats and end caps are shown on the end of this sheet, please refer to this if you are unsure which cartridge format you have.

- 2.1 New housings should be flushed out with clean water / air (dependant on the application) prior to installation of the cartridge to remove any debris. Ensure tie-rods / support plates are removed prior to flushing as vibration (especially in air) can cause components to loosen.
- 2.2 Before changing or installing a liquid or gas cartridge filter ensure that the filter vessel is depressurized and any liquid has been drained off [Most vent filter cartridges are open to atmosphere but if the filter is connected to a pressurized line then ensure that the filter vessel is depressurized before removing the filter
- 2.3 Remove the filter bowl. For plastic housings the bowl is unscrewed and for stainless steel housings the bowl is held in place using a band clamp or a bolted flange
- 2.4 Cut open the polyethylene bag at the cartridge open end and check that the o-ring seals or gaskets are clean, intact, correctly located in their grooves and not damaged.
- 2.5 Lubricate o-ring seals with a lubricant that is compatible with the process fluid (e.g. clean water) or use process liquid itself. Note: No lubricant should be used for oxygen applications.
- 2.6 Using the bag as protection and holding the cartridge as near as possible to the open end as opposed to the main body of the cartridge or the top end cap, press the

cartridge firmly into or onto the housing locations. Keep the cartridge vertical to prevent damage to the o-rings.

- a) If the vessel has a bayonet type cartridge location (A,C & R), slightly turn the cartridge clock-wise to locate the retaining lugs.
- b) For double open ended cartridges (B), take care to ensure that the cartridge gaskets on both the housing and cartridge are centred over the housing knife edge seals at both ends before closing the vessel
- c) Cartridges with a threaded end cap (V) should be screwed in until the gasket is compressed.
- d) Threaded vent filters should be screwed into position until the flat gasket is compressed (BSPP) or the thread locks (NPT).
- 2.7 Remove the polyethylene bag from the cartridge(s) before the vessel is closed.
- 2.8 Some filter housings take more than one cartridge (multi-round) and they will have a support plate that locates on top of the cartridges and prevents movement and damage. Refer to the vessel instructions for the way that this plate is secured and ensure that it is always installed before the vessel bowl is located.

3. Operation (liquid cartridges)

Filter cartridges should not be subjected to excessive hydraulic shock and should never be reverse pressurized from the downstream to the upstream side (inside to out).

- 3.1 Slowly open the upstream valve and allow liquid into the filter vessel
- 3.2 The vent valve located at the top of the vessel should be cracked open to allow air to escape and to ensure that the filter vessel is full of liquid. The vent valve should be closed when liquid starts to exit the valve.

N.B. If hazardous liquids are being filtered, please ensure that vent and drain valves are connected to a suitable drain line.

3.3 Slowly open the downstream valve and allow the filtered liquid to flow. It is recommended that newly installed cartridges are briefly flushed to drain and remove an debris that may have been inadvertently generated during cartridge installation or to remove trace levels of surfactant that may be present in some filter media. Liquid cartridges are shown to be blocked when the differential pressure across the filter has significantly increased and / or the flow of liquid through them is reduced to an unacceptable level. If you do not have pressure gauges that indicate the differential pressure then please contact Parker domnick hunter or their representative.

4. Operation (gas / vent cartridges)

Vent / Gas filter cartridges are hydrophobic and they will not operate effectively if they are covered in water or steam condensate. This can lead to tank collapse or cartridge deformation so please ensure that if vent

filters do come into contact with water they are

Gas cartridges are blocked when the differential pressure across the filter is high and/or the flow of gas through them is significantly reduced. In normal operation they should be changed at least annually.

5. Integrity testing

Some liquid and gas cartridges may be integrity tested by a number of manual or automatic methods. Please contact Parker domnick hunter or it's representative for further information on which method is most suitable for your application or refer to the appropriate product datasheet.

6. Hot water sanitization (Liquid hydrophilic cartridges)

Recirculate prefiltered water through the filter for 1 hour at 80 °C (176 °F), the maximum differential pressure across the filter should be no more than 0.3 bar (5 psi). Open all system outlet valves to sanitize the system thoroughly.

7. Steam sterilization

Please refer to the datasheets to find out if your cartridge filter and housing can be autoclaved or steamed in place (SIP) and the allowed maximum temperature. To minimize the risk of contamination to a sterile system the filter should be autoclaved or SIP'd immediately prior to use.

N.B. Plastic housings cannot be steam sterilized or autoclaved.

Steam-in-place (SIP)

It is important that both liquid and gas filter cartridges do not have bulk steam flowed through them during SIP because excessive differential presure can cause damage to the cartridge at high temperatures. It is also usual to filter the steam so that any dirt it carries does not block or damage the filter.

Vacuum autoclave sterilization

The cartridge should be installed in the housing, the vent / drain valves left open and the housing bowl left slightly open. Do not allow the cartridge to support the vessel base or allow the bowl to rest on the cartridge during autoclaving. The assembly should be autoclaved on a cycle with a slow exhaust. Where possible liquid cartridges should be flushed with clean water prior to autoclaving.

Parker domnick hunter has detailed guidelines for the sanitization and steam sterilization of liquid and gas filters so if you are unsure of the procedures please contact Parker domnick hunter or it's representative.

Disposal

All cartridge filters should be disposed of in a safe manner and in line with Health & Safety Guidelines.

Conversion tables

Volume rate of flow

CONVERT					Multiplying	Factors				
FROM TO →	litre / sec	litre / hr	m³/sec	m³/hr	ft ³ / min	ft³/hr	UK gal / min	UK gal / hr	US gal / min	US gal/hr
↓ litre / sec	1.	3600.	0.001	3.6	2.118882	127.133	13.19814	791.8884	15.85032	951.019
litre / hr	0.000278	1.	0.00000028	0.001	0.000588	0.035315	0.003666	0.219969	0.004403	0.264172
m³/sec	1000.	3 600 000.	1.	3600.	2118.88	127 133.	13 198.1	791 889.	15 850.3	951 019.
m³ / hr	0.27778	1000.	0.000278	1.	0.588578	35.3415	3.66615	219.969	4.402863	264.1718
ft ³ / min	0.471947	1699.017	0.000472	1.699017	1.	60.	6.228833	373.730	7.480517	448.8310
ft³ / hr	0.007866	28.3168	-	0.028317	0.01667	1.	0.103814	6.228833	0.124675	7.480517
UK gal / min	0.0757	272.766	0.0000758	0.272766	0.160544	9.63262	1.	60.	1.20095	72.05700
UK gal / hr	0.001263	4.54609	-	0.004546	0.002676	0.160544	0.016667	1.	0.020016	1.20095
US gal /min	0.063090	226.8	0.0000631	0.227125	7.4805	448.8	0.832674	49.96045	1.	60.
US gal / hr	0.001052	3.785411	-	0.003785	0.133681	0.133681	0.013878	0.832674	0.016667	1.

Pressure (liquid column, atmospheric, etc.)

CONVERT					Multiplying	Factors				
FROM TO→	lb/in²	InH ₂ 0	ftH ₂ O	inHg	atmos.	mmHg	mbar	kgf / cm²	N/m^2	N / mm²
↓ lb / in²	1.	27.6799	2.30667	2.03602	0.068046	51.7149	68.9476	0.070307	6894.76	0.0068948
InH ₂ 0	0.036127	1.	0.083333	0.073556	0.0024583	1.86832	2.49089	0.002540	249.089	0.0002491
ftH ₂ O	0.433528	12.	1.	0.882671	0.029500	22.4198	29.8907	0.03048	2989.07	0.0029891
inHg	0.491154	13.5951	1.13292	1.	0.033421	25.4	33.8639	0.034532	3386.39	0.003386
atmos.	14.6959	406.781	33.8984	29.9213	1.	760.000	1013.25	1.03323	101 235.	0.101325
mmHg	0.019337	0.535240	0.044603	0.03937	0.0013158	1.	1.33322	0.0013591	133.322	0.0001333
mbar	0.014504	0.401463	0.033455	0.029530	0.0009869	0.750062	1.	0.0010197	100.	0.0001
kgf / cm²	14.2233	393.700	32.8084	28.959	0.967841	735.559	980.655	1.	98 066.5	0.98066
N/m^2	0.000145	0.004015	0.0003345	0.0002953	0.000099	0.007501	0.01	0.0000102	1.	0.000001
N/mm²	145.038	4014.63	334.553	295.300	9.86923	7500.62	10 000.	10.1972	1 000 000.	1.

Mass

CONVERT				Multiplying Fa	ctors		
FROM TO →	grain	metric carat	gram	dram	drachm (apoth)	OZ	oz tr or oz apoth
↓ grain	1.	0.323995	0.064799	0.36571	0.016667	0.002286	0.002083
metric carat	3.08647	1.	0.2	0.112877	0.51441	0.007055	0.006430
gram	15.4324	5.	1.	0.564383	0.257206	0.035274	0.032151
dram	27.34375	8.85923	1.77185	1.	0.455729	0.0625	0.056966
drachm (apoth)	60.	19.4397	3.88793	2.19429	1.	0.137143	0.125
OZ	437.5	141.748	28.3495	16.	7.29167	1.	0.911458
oz tr or oz path	480.	155.517	31.1035	17.5543	8.	1.09714	1.

Conversion tables

Mass

CONVERT				Multiplyin	g Factors			
FROM TO →	lb	kg	slug	US cwt	UK cwt	oz / US ton	tonne	UK ton
↓ lb	1.	0.453592	0.031081	0.01	0.008929	0.0005	0.000454	0.000446
kg	2.20462	1.	0.068522	0.022046	0.019684	0.001102	0.001	0.000984
slug	32.1740	14.5939	1.	0.32174	0.287268	0.016087	0.014594	0.014363
US cwt	100.	45.3592	3.10810	1.	0.892857	0.05	0.045359	0.044643
UK cwt	112.	50.8023	3.481072	1.12	1.	0.056	0.050802	0.05
oz / US ton	2000.	907.185	62.1620	20.	17.8571	1.	0.907185	0.892857
tonne	2204.62	1000.	68.5218	22.0462	19.6841	1.10231	1.	0.984207
UK ton	2240.	1016.05	69.62143	22.4	20.	1.12	1.01605	1.

Volume and capacity

CONVERT					Multiplying	Factors				
FROM TO →	cm ³	in³	ft ³	yd³	m³	litre	UK pint	UK gallon	US pint	US gallon
↓ cm³	1.	0.061024	0.0000353	-	0.000001	0.001	0.001760	0.000220	0.002113	0.000264
in ³	16.3871	1.	0.0005787	0.0000214	0.0000164	0.016387	0.028837	0.003605	0.034632	0.004329
ft³	28 316.8	1728.	1.	0.037037	0.028317	28.3168	49.8307	6.22883	59.8442	7.48052
yd³	764 555.	46 656	27.	1.	0.764555	764.555	1345.429	168.1784	1615.793	201.9740
m ³	1 000 000.	61 023.7	35.3145	1.30795	1.	1000.	1759.75	219.969	2113.38	264.172
litre	1000.	61.0237	0.035315	0.001308	0.001	1.	1.75975	0.219969	2.11338	0.264172
UK pint	568.261	34.6774	0.020068	0.000743	0.0005683	0.568261	1.	0.125	1.20095	0.150119
UK gallon	4 546.09	277.420	0.160544	0.005946	0.0045461	4.54609	8.	1.	9.60760	1.20095
US pint	473.176	28.875	0.016710	0.000619	0.0004732	0.473176	0.832674	0.104084	1.	0.125
US gallon	3 785.41	231.	0.133681	0.004951	0.0037854	3.785411	6.661392	0.832674	8.	1.

Volume and capacity

CONVERT				Mult	iplying Factor	rs			
FROM TO→	UK minim	US minim	cm ³	UK fl drachm	US fl drachm	UK fl ounce	US fl ounce	litre	in³
↓ UK minim	1.	0.960760	0.059194	0.016667	0.016013	0.002083	0.002002	0.0000592	0.0036122
US minim	1.04084	1.	0.061611	0.17348	0.01667	0.002168	0.002084	0.0000616	0.0037597
cm ³	16.8936	16.2307	1.	0.281561	0.270519	0.035195	0.033814	0.001	0.061024
UK fl drachm	60.	57.64560	3.55163	1.	0.960760	0.125	0.120095	0.003552	0.216734
US fl drachm	62.45040	60.	3.696678	1.04084	1.	0.130105	0.125	0.003697	0.225585
UK fl ounce	480.	461.1648	28.4131	8.	7.68608	1.	0.960760	0.028413	1.73387
US fl ounce	499.604	480.	29.5735	8.32674	8.	1.04084	1.	0.029573	1.80469
litre	16 893.6	16 230.7	1000.	281.561	270.5125	35.1951	33.8140	1.	61.0237
in³	276.837	265.9739	16.3871	4.61395	4.432899	0.576744	0.554113	0.016387	1.

158 |

Chemical compatibility

NC = Not Compatible LC = Limited Compatibility C = Compatible - = No Data	Acetic acid 3.5N	Acetic acid 8.75N	Acetic acid conc. 17.5N	Acetone	Acetonitrile	Acidbrite 4 (Diversey) 3.0% v/v	Ammonium Hydroxide 8N	Ammonium Oxalate 0.07N	Amyl Acetate	Aqueous Ammonia 15.5N	Benzyl Alcohol	Benzyalkonium Chloride 0.1%	Boric acid, saturated	Butan-1-ol	Butan-2-ol	Carbon Tetrachloride	Chloroform
BEVPOR MH / MS / MT / PH / PS / PT	С		-	NC	-	-	LC	-	LC	LC	-	-	-	С	С	-	NC
BIO-X II	С	С	С	С	С	-	С	С	С	С	С	С	С	С	С	С	С
CRYPTOCLEAR PES	С	-	-	NC	-	-	LC	-	LC	LC	-	-	-	С	С	-	NC
CRYTOCLEAR PLUS	С	С	С	С	С	С	С	С	С	С	NC	С	С	С	С	NC	NC
HIGH FLOW BIO-X	С	С	С	С	С	-	С	С	С	С	С	С	С	С	С	С	С
HIGH FLOW BIO-X VENT AUTOCLAVE	С		-	-	-	-	С	С	С	С	С	С	С	С	С	С	С
HIGH FLOW PREPOR GFA	С	С	С	С	LC	С	С	С	LC	LC	NC	С	С	С	С	NC	NC
HIGH FLOW TETPOR II	С	С	С	С	С	-	С	С	С	С	С	С	С	LC	С	С	С
HIGH FLOW TETPOR H.T.	С	С	С	С	С	-	С	С	С	LC	С	С	С	LC	С	С	С
HIGH FLOW TETPOR VENT AUTOCLAVE	С	С	С	С	С	-	С	С	С	С	С	С	С	LC	С	С	С
PEPLYN AIR/NE/PLUS/HA/HD/PREPOR PP	С	С	С	С	С	С	С	С	С	С	NC	С	С	С	С	NC	NC
PREPOR GF / GP	-	С	С	С	LC	С	С	С	LC	LC	NC	С	С	С	С	NC	NC
PREPOR PES	С	-	-	NC	-	-	С	-	LC	С	-	-	-	С	С	-	NC
PROCLEAR PP	С	С	С	С	С	С	С	С	С	С	NC	С	С	С	С	NC	NC
PROCLEAR GF	С	С	С	С	LC	С	С	С	LC	LC	NC	С	С	С	С	NC	NC
PROPLEAT	С	С	С	С	С	С	С	С	С	С	NC	С	С	С	С	NC	NC
PROPOR BR / HC / LR	С	-	-	NC	-	-	LC	-	LC	LC	-	-	-	С	С	-	NC
PROPOR SG	С	-	-	NC	-	-	С	-	LC	С	-	-	-	С	С	-	NC
PROSPUN	С	С	С	С	С	С	С	С	С	С	NC	С	С	С	С	NC	NC
PROSTEEL A / N	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С
STEAM FILTERS	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С
TETPOR AIR / LIQUID	С	С	С	С	С	-	С	С	С	С	С	С	С	NC	С	NC	NC
TETPOR PLUS	С	С	С	С	С	-	С	С	С	С	С	С	С	NC	С	NC	NC
EPDM	С	LC	LC	С	NC	С	С	С	NC	С	С	С	С	С	LC	NC	NC
VITON	С	LC	NC	NC	NC	С	С	С	NC	С	С	С	С	С	С	С	LC
SILICONE	С	NC	NC	NC	NC	С	С	С	LC	С	С	С	С	С	С	NC	NC

Cyclohexane	1,4 - Dioxane	Diverflow (Diversey) 3% v/v	Diversey 2126 0.6% v/v	Divosan Forte 0.5% v/v	Divosan XT 1% v/v	Ethanol	Ethanol 45%	Ethyl Acetate	Formaldehyde 0.3%	Formaldehyde 37%	Formic acid conc.	Glycerol	Hexane	Hydrochloric acid 1N	Hydrochloric acid 10%	Hydrochloric acid conc.	Hydrochloric acid conc. 13%	Hydrogen Peroxide	Hydrogen Peroxide 10 Vol	Hydrogen Peroxide 100 Vol	Methanol	Methyl-Iso-Butylketone	Methylene Chloride @ 40 °C (104 °F)	Nitric Acid 2N 14.4%
-	-	С	-	С	-	С	С	NC	С	-	-	С	-	С	-	-	-	-	С	-	С	NC	-	С
С	С	-	-	-	-	С	-	LC	С	С	С	С	С	-		-	С	С	-	-	С	С	-	С
-	-	С	-	С	-	С	С	NC	С	-	-	С	-	С	-	-	-	-	С	-	С	NC	-	С
NC	С	С	С	С	С	С	С	LC	С	С	С	С	NC	С	-	С	-	С	С	С	С	С	LC	С
С	С	-	-	-	-	С	-	LC	С	С	С	С	С	-		-	С	С	-	-	С	С	-	С
С	С	-	-	-	-	С	-	LC	С	С	С	С	С	-		-	С	С	-	-	С	С	-	С
NC	LC	NC	С	С	С	С	С	LC	С	С	NC	С	-	С		NC	-	-	С	С	С	С	LC	С
-	С	-	-	-	-	С	-	LC	С	С	С	С	С	-		-	С	-	-	С	С	С	-	С
-	С	-	-	-	-	-	-	LC	С	С	С	С	С	-		-	С	-	-	С	С	С	-	С
-	С	-	-	-	-	С	-	LC	С	С	С	С	С	-		-	С	-	-	С	С	С	-	С
NC	С	С	С	С	С	С	С	LC	С	С	С	С	NC	С	-	С	-	С	С	С	С	С	LC	С
NC	LC	NC	С	С	С	С	С	LC	С	С	NC	С	-	С	-	NC	-	-	С	С	С	С	LC	NC
-	-	С	-	С	-	С	С	NC	С	-	-	С	-	С	-	-	-	-	С	-	С	NC	-	С
NC	С	С	С	С	С	С	С	LC	С	С	С	С	NC	С		С	-	С	С	С	С	С	LC	С
NC	LC	NC	С	С	С	С	С	LC	С	С	NC	С	-	С		NC	-	-	С	С	С	С	LC	С
NC	С	С	С	С	С	С	С	LC	С	С	С	С	NC	С	-	С	-	С	С	С	С	С	LC	С
-	-	С	-	С	-	С	С	NC	С	-	-	С	-	С		-	-	-	С	-	С	NC	-	С
-	-	С	-	С	-	С	С	NC	С	-	-	С	-	С		-	-	-	С	-	С	NC	-	С
NC	С	С	С	С	С	С	С	LC	С	С	С	С	NC	С	-	С	-	С	С	С	С	С	LC	С
С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С
С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С
LC	С	-	-	-	-	С	С	LC	С	С	С	С	-	С		С	-	-	С	С	С	С	-	С
LC	С	-	-	-	-	С	С	LC	С	С	С	С	С	С		С	С	С	С	С	С	С	С	С
NC	NC	С	С	С	С	С	С	С	С	С	С	С	NC	С		NC	NC	С	С	С	С	NC	-	LC
NC	NC	С	С	С	С	С	С	NC	С	С	NC	С	NC	С		NC	NC	С	С	С	NC	NC	-	С
NC	NC	LC	С	С	С	LC	С	LC	С	С	NC	С	NC	С		NC	NC	С	С	С	С	LC	-	С

Chemical compatibility

NC = Not Compatible LC = Limited Compatibility C = Compatible - = No Data	Nitric acid 15.8N	Ozone	Paraffin yellow	Pentane	Peracetic acid 0.5% (10 week test)	Peracetic acid 4%	Perchloroethylene	Petroleum spirits	Phenol (aq) (0.5N)	Phenol 5%	Phenol 0.25%	Polyethylene Glycol 600	Polyglycol 2000-E	Potassium Dichromate 0.1N	Potassium lodine 0.6N	Potassium Hydroxide 10N	Potassium Permanganate 0.1N
BEVPOR MH / MS / MT / PH / PS / PT	-	NC	-	-	-	С	NC	-	-	-	-	NC	-	-	-	LC	С
BIO-X II	С	-	LC	С	-	С	-	-	С	-	-	LC	-	С	С	С	С
CRYPTOCLEAR PES	-	NC	-	-	-	С	NC	-	-	-	-	NC	-	-	-	LC	С
CRYPTOCLEAR PLUS	С	-	С	LC		С		NC	-	С	С	С	-	С	С	С	С
HIGH FLOW BIO-X	С	-	LC	С	=	С	-	-	С	=	-	LC	-	С	С	С	С
HIGH FLOW BIO-X VENT AUTOCLAVE	С	-	LC	С	-	С	-	-	С	-	-	LC	-	С	С	С	С
HIGH FLOW PREPOR GFA	NC	-	LC	LC	-	С	-	NC	-	С	С	NC	-	С	С	NC	NC
HIGH FLOW TETPOR II	С	-	С	-	С	С		С	NC		-	С	-	С	С	С	С
HIGH FLOW TETPOR H.T.	NC	-	С	-	С	С		С	-		-	С	-	С	С	С	LC
HIGH FLOW TETPOR VENT AUTOCLAVE	С	-	С	-	С	С		С	NC		-	С	-	С	С	С	С
PEPLYN AIR/NE/PLUS/HA/HD/PREPOR PP	С	-	С	LC	-	С	-	NC	-	С	С	С	-	С	С	С	С
PREPOR GF / GP	-	NC	=	-	-	С	NC	-	-	-	-	NC	-	-	-	С	С
PREPOR PES	С	-	-	NC	-	-	С	-	LC	С	-	-	-	С	С	-	NC
PROCLEAR PP	С	-	С	LC	-	С	-	NC	-	С	С	LC	-	С	С	С	С
PROCLEAR GF	NC	-	LC	LC	-	С	-	NC	-	С	С	NC	-	С	С	NC	NC
PROPLEAT	С	-	С	LC	-	С	-	NC	-	С	С	С	-	С	С	С	С
PROPOR BR / HC / LR	-	NC	-	-	-	-	NC	-	-	-	-	NC	-	-	-	LC	С
PROPOR SG	-	NC	-	-	-	С	NC	-	-	-	-	NC	-	-	-	С	С
PROSPUN	С	-	С	LC	-	С	-	NC	-	С	С	С		С	С	С	С
PROSTEEL A / N	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С
STEAM FILTERS	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С
TETPOR AIR / LIQUID	С	-	С	LC	-	С	-	LC	-	С	С	-	-	С	С	С	С
TETPOR PLUS	С	С	С	LC	-	С	-	LC	-	С	С	-	-	С	С	С	С
EPDM	NC	-	NC	NC	С	С	-	NC	-	С	С	-	С	С	С	С	С
VITON	NC	-	С	С	С	С	-	С	-	С	С	-	С	С	С	С	С
SILICONE	NC	-	NC	NC	С	С	-	NC	-	С	С	-	С	С	С	С	С
The chemicals are arranged in alphabatical order using their				15-5-1	0	Lillia (LO	l at ambi			T O.		- 72 hour					

The chemicals are arranged in alphabetical order using their most common or trade names. If the chemical in question does not appear to be listed, it may be found elsewhere in the table under a pseudonym, in particular its IUPAC¹¹ name.

With regard to compatibility:

Propan-1-ol	Propan-2-ol	Propan-2-ol, 60:40 H ₂ 0	Pyridine	Sodium Chloride 0.5N	Saline Lactose Broth	Sodium Hydroxide 1N,4%	Sodium Hydroxide 7N,28%	Sodium Hypochlorite	Sodium Hypochlorite (14% Free CL ₂)	Sodium salts	Sodium thiosulphate 0.1N	Sulphuric acid 1N	Sulphuric acid conc.	Sulphurous acid	Toluene	1,1,1 Trichloroethane	1,1,2 Trichloroethane	Trichloroacetic Acid 80%	Trichloroacetic Acid 5N	Toluene	Xylene
С	С	С	NC	С	С	С	NC	С	С	С	-	С	NC	NC	NC	-	NC	-	-		LC
С	С	С	С	С	С	NC	NC		С		С	С	NC	-	NC	-	С	-	С	-	LC
С	С	С	NC	С	С	С	NC	С	С	С	-	С	NC	NC	NC	-	NC	-	-	-	LC
С	С	С	С	С	С	С	С	-	С	-	С	С	LC	-	NC	-	LC	С	-	-	NC
С	С	С	С	С	С	NC	NC		С		С	С	NC	-	NC	-	С	-	С	-	LC
С	С	С	С	С	С	NC	NC		С		С	С	NC	-	NC	-	С	-	С	-	LC
NC	NC	NC	NC	С	С	С	NC		С		С	LC	LC	-	-	-	LC	LC	-	NC	NC
С	С	С	С	С	С	С	С		С		С	С	LC	-	NC	-	С	-	С		LC
С	С	С	С	С	С	С	С		С		С	С	NC	-	NC	-	LC	-	С	-	LC
С	С	С	С	С	С	С	С		С		С	С	LC	-	NC	-	С	-	С	-	LC
С	С	С	С	С	С	С	С	-	С	-	С	С	LC	-	NC	-	LC	С	-	-	NC
С	С	С	NC	С	С	С	NC	С	С	С	-	С	NC	NC	NC	-	NC	-	-	-	LC
-	-	С	-	С	-	С	С	NC	С	-	-	С	-	С	-	-	-	-	-	С	-
С	С	С	С	С	С	С	С		С		С	С	LC	-	NC	-	LC	С	-	-	NC
NC	NC	NC	NC	С	С	С	NC		С		С	LC	LC	-	-	-	LC	LC	-	NC	NC
С	С	С	С	С	С	С	С	-	С	-	С	С	LC	-	NC	-	LC	С	-	-	NC
С	С	С	NC	С	С	С	NC		С		-	С	NC	NC	NC	-	NC	-	-	-	LC
С	С	С	NC	С	С	С	NC		С		-	С	NC	NC	NC	-	NC	-	-	-	LC
С	С	С	С	С	С	С	С	-	С	-	С	С	LC	-	NC	-	LC	С	-		NC
С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С
С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С	С
С	С	С	С	С	С	С	С		С		С	-	LC	-	-	-	LC	С	-	-	NC
С	С	С	С	С	С	С	С		С		С	-	LC	-	С	-	LC	С	-	-	NC
С	С	С	С	С	С	С	С		С		С	С	-	-	NC	-	NC	NC	-	NC	С
С	С	С	NC	С	С	С	С		С		С	С	-	-	LC	-	LC	LC	-	LC	LC
LC	LC	С	С	С	С	С	LC		С		С	С	-	=	NC	-	LC	NC	-	NC	NC

Any product that has Limited Compatibility (LC) at ambient temperatures should not be used at a higher temperature.
 The list of compatibilities does not take into account any synergistic effects of more than one chemical present in the solution to be filtered.

⁻ Test Conditions - 72 hours at ambient temperature and pressure, unless otherwise stated.
- Contact Parker domnick hunter for confirmation of compatibility with specific operating conditions.

Glossary of terms used in filtration



Absolute pressure

Associated with gas systems. The absolute pressure is the total pressure exerted on a system equal to atmospheric pressure plus gauge pressure, for example 2 barg = 3 bar absolute.

Absolute rating

A definitive value given to a filter that represents the smallest particle size capable of being captured by the filter. Typically it refers to 100% retention at a particular micron rating. The assigning of micron ratings is however dependant on the test methodology used. e.g.: a sterile grade absolute rated liquid filter is assigned a 0.2 micron rating if it retains all microorganisms of a predetermined size it does not mean that the filter has 0.2 micron pores. When selecting a filter for a particular application always refer to the methods and assumptions made for assigning the micron rating.

Air flow

A measure of the amount of air that flows through a filter at a certain system pressure and pressure drop. This is typically expressed in normalized units i.e.: the relative flow rate at atmospheric pressure and is quoted for a clean unused filter. Always quote system pressures when sizing gas filters.

Aerosol integrity testing

A method specifically designed for sterile gas filters whereby aerosol in the most penetrating particle size (MPPS: 0.2–0.3 micron) is used as a non-destructive challenge to the filter to determine whether it is providing sterile gas. The test can be performed using an automated test instrument such as the Parker domnick hunter VALAIRDATA II.

Autoclave

A closed pressure vessel into which steam is introduced (typically at a temperature of 121 - 134 °C (250 - 273 °F)) to sterilise the contents.

В

Backwash

A reverse flow of liquid through a filter in order to flush out trapped solids.

Bacterial challenge

This refers to a live bacterial challenge of a filter in either the liquid or gas phase. The type of organism used for the test depends on the assigned micron rating of the filter. For example a 0.2 micron sterile grade liquid filter is challenged with the organism *Brevundimonas diminuta* (test method ASTM 838-05) while a 0.45 micron absolute rated liquid filter is challenged with a suspension of *Serratia marcesens*. In some cases for critical performance validation requirements it will be necessary to challenge the filter with bacteria in the actual process fluid being filtered.

Beta rating

A measure of a filter's efficiency based on the number of particles present in the influent (upstream) to those in the effluent (downstream). Efficiency is expressed as a BETA ratio and is calculated as follows:

Beta Ratio =

Number of particles in the influent
Number of particles in the effluent

Generally a Beta Ratio at 5000 is accepted by the industry as being an 'absolute' rating for media prefilters.

C

Cartridge or filter cartridge

A filtration or separation device usually supplied in a cylindrical format which locates easily and quickly into a filter housing.

Chemical compatibility

When selecting filter materials attention needs to be given to their compatibility to the fluid which is to be filtered. A filter (depending on application) needs to be assessed for reduction in performance in terms of material degradation, integrity, etc. as well as quantifying any extractables levels. It should be noted that the compatibility of a filter is dependent on the process conditions. General material compatibility databases assume limited temperature and exposure time. They also refer to just one chemical. In an actual process there could be a combination of chemicals, high differential pressure and high temperature which all could influence filter performance. General guidance on filter performance can be given from experience and in-house data but normally it is recommended that filter compatibility is tested in the process conditions.

Clarification

This is the selective removal of particulate from a process fluid usually achieved through depth filtration. The degree of clarification is dependent on customer specification.

Colony forming unit (CFU)

The minimum number of cells on an agar plate which will give rise to a visible colony. This term is most commonly seen in the validation of sterile filters to a live bacterial challenge where the challenge and the number of organisms recovered is stated in CFU.

Coalescing

When small droplets of aerosolized liquid merge together to form larger droplets. This normally occurs in a depth filter as the process gas carrying the entrained liquid droplets passes through the filtration media. A coalescing filter such as the Parker domnick hunter OIL-X also flows from the inside of the cartridge to the outside so any coalesced liquid drains to the base of the filter and subsequently into the bottom of the filter housing

Colloid

Colloids are molecules that have not coagulated together to form a precipitate but remain in liquid suspension. These molecules are very small in size and have a molecular charge that affects their affinity for other molecules and materials. The choice of filter type and design is of paramount importance for a colloidal system if premature blockage is to be avoided.

Compaction

This can occur to a filtration medium when it is subjected to high differential pressures. The high forces on the filtration media (especially depth type) can lead to compression of the structure and subsequent changes in filtration characteristics.

Concentrate

The retained non filtered stream from a crossflow filter system.

Cross flow filtration

A filter characterized by the feed stream travelling parallel to instead of directly through the filtration medium. This has the advantage of minimizing the blockage of the membrane as the system is to some extent 'self cleaning'.

D

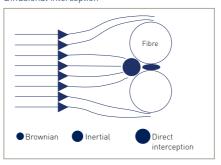
Dead leg

An area of pipework where there is potentially no flow and therefore stagnant conditions exists. It is extremely important to eliminate these if contamination issues are to be minimized.

Depth filte

A depth filter is characterised by the thickness of the filtration media as well as its structure. A depth filter is normally fibrous in nature and contaminant is retained through the depth of the filtration media rather than just the surface.

Diffusional interception



This is the dominant removal mechanism for the smallest particles captured by a filter in the gas phase. Particles as small as 0.01 µm exhibit great diffusional movement (Brownian Motion) which has the effect of increasing its nominal mean diameter to the filter. The efficiency of this capture mechanism decreases as the particle size increases.

Diffusional flow

A non-destructive integrity test method for membrane based filters. It involves wetting out every pore in the membrane structure with water or the process fluid or a low surface tension liquid in case of hydrophobic membrane. Compressed air is applied to the upstream side of the filter and gas diffuses through the wetted pores. This flow rate is either measured directly by mass flow meters or indirectly via measuring the drop in pressure on the upstream side of the filter.

Differential pressure

Differential pressure (dP) is the difference in the pressure measured upstream (influent) and downstream (effluent) of a filter. Particularly in liquid applications differential pressure will increase to a point where either filter damage or insufficient flow will result. The higher the differential pressure the higher the energy cost so it is important to balance the pressure drop requirements with the installation size and required lifetime to blockage. Units of measurement are bar and psi as opposed to barg and psig.

Glossary of terms used in filtration

Ε

Effective filtration area (EFA)

This is the area of filtration material available for filtration.

Effluent

The fluid which has passed through a filter.

Extractables

When a filter is in contact with the process fluid, chemical components may leach from the materials of construction and deposited in the filtrate. The levels of non-volatile extractables for a limited number of fluids are quoted in the filter validation guide. The level of extractables is dependent on the process conditions. Filtration of solvents, high temperature fluids and steam sterilization are three areas where extractables can increase.

F

Filter (noun) / filter cartridge / cartridge An apparatus which performs filtration.

ter (verb)

To pass a fluid or gas through a porous medium in order to remove solid particles.

Filter efficiency

Filter efficiency is a measure of the percentage of particles that are removed from the fluid by the filter. Typically these are given in terms of the % removal for a certain size of particle. A filter efficiency may also be given across a range of particle sizes . For a number of gas applications the efficiency of a filter may be quoted in relation to the filters ability to remove particles at the most penetrating particle size (MPPS) of 0.2-0.3 micron. Always ensure filter efficiency is matched to the requirements of the process.

Filterability indices (FI) and Vmax

This is an indication of a filters capacity to process certain fluids. It generally gives a measure of the rate of blockage of a filter as well as the theoretical maximum throughput. The time required to flow two consecutive 200 ml fluid samples is recorded and the filterability indices are calculated from the results. The two formulae used are as follows:

$$(Vmax) = \frac{400 + 400T}{(T_2 - 2T_1)}$$
FI =
$$(T_2 - 2T_1)$$

 T_1 = Time to filter first 200 ml

 T_2 = Time to filter second 200 ml

It should be noted that these methods give a general indication of performance and are often more useful in comparative performance measurement between different filter types.

Filtrate

Another name for effluent.

Filter sterilization

Sterilization is the act of making an organism barren or infertile (unable to reproduce). The sterilization of a filter can be achieved by a number of methods including dry heat, steam, ethylene oxide, hydrogen peroxide or irradiation. The method chosen depends on the process and

the materials of construction of the filter but by far the most widely used is that of steam, either in an autoclave or via steam-in-place (SIP).

Flux

The rate of fluid flow (gas or liquid) when expressed in terms of flow per unit area of the filter that removes the contaminants from the fluid stream. It can apply to both depth and membrane media.

G

Gauge pressure

The pressure of a system measured by a gauge, which excludes atmospheric pressure, for example 1 bar atmosphere (or 1 bar absolute) = 0 barg.

Н

Housing

An enclosure for a filter element, typically rated for pressure, that directs the fluid through the filter.

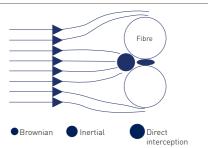
Hydrophilic

Hydrophilicity is the ability of a filtration media to 'wet out', that is, for the porous structure to be completed filled with the liquid being filtered. This is an important characteristic as incomplete wetting of the structure can lead to a reduction in flow capacity and problems with integrity testing. All liquid filters are 'hydrophilic' apart from those that may have been selected for use with aggressive solvents. These filters are typically based on a fluoropolymer and their structure needs to be wetted with a low surface tension liquid such as isopropyl alcohol. Once the structure has been wet, the filter will process aqueous solutions without a problem.

ı

Inertial impaction

This is a removal mechanism for particles captured by a filter in the gas phase. The particles follow the streamlines of gas between the filter fibres and membrane pores. Due to their mass the inertia of the particle will cause it to move out of the streamline and attach itself to a fibre or pore wall. The effect of this capture mechanism increases with particle size / mass.



Influent

The fluid entering the filter system.

In vitro

In an unnatural position e.g. outside the body "In vitro" is Latin for "in glass" an experiment performed without the involvement of a whole, living organism.

In viv

The testing of a substance or experimentation in or using a living, whole organism.

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Log reduction value (LRV)

This is a measurement of a filters removal efficiency for a specific contaminant. It is normally associated with the bacterial retention of a filter.

The I RV is:

Log₁₀ Number of bacteria in the influent
Number of bacteria in the effluent

e.g.
$$\left(Log_{10} \frac{1 \times 10^{10}}{1} \right) = LRV \text{ of } > 10$$

It is always expressed as > (greater than) as 1 has to be used for the effluent even if there are no organisms present. This can also be expressed as a 10 log reduction or a titre reduction of 10¹⁰.

M

Medium (Media)

This is the component of the filter that removes the contaminants from the fluid stream. Also commonly referring to depth - type materials, in its more generic sense a filter medium / media can refer to either depth or membrane filter materials.

Microfiltration

Microfiltration is the process of removing particles from a liquid or gas by passing it through a porous medium. It generally involves removing particles between the sizes of 10 and 0.04 micron in liquids, and down to 0.01 micron in gases.

Micron (micrometer)

Designated by the Greek letter μ a micron is 10° mm (millimeters) or 10^{4} (Angstroms) or 0.00003937 inch. For a perspective on this size a human hair is approximately 70 microns thick and the limit of resolution of the naked eye is around 40 microns.

Membrane

A membrane is a thin, porous film typically between 30 and 150 micron in thickness. It has of tens of millions of pores / cm² through which the process fluid runs. The nature of the pore structure is determined by the manufacturing method.

Solvent cast membranes such as Polyethersulphone (PES) and Mixed Esters of Cellulose (MEC) have a defined pore structure which can be asymmetric whilst membrane such as Polytetrafluoroethylene (PTFE) which is manufactured by stretching have a fibrous appearance and a less defined pore structure.

N

Nanofiltration

Filtration that removes both particles and small dissolved molecules and ions. Finer than ultrafiltration, not as fine as Reverse Osmosis.

Nanometer

A nanometer is 10⁻⁹ meters

Nominal filter rating

This rating is often quoted within the filtration industry but great care should be taken in ensuring the efficiency and test methodologies are completely understood. A 5 micron nominal filter could be 99% retentive at 5 micron, another could be 80%. It can be very misleading to compare the performance of filters on nominal ratings. When selecting a filter the duty required should be

Glossary of terms used in filtration

compared to the individual performance characteristics of filter. Parker domnick hunter has the experience to help select the most appropriate filter for the application.



Oleophobic

Oleophobic membranes and depth media have the capability to repel fluids such as oil and lubricants. This phenomena is used in some of the new generation oil coalescing filters.

Oxidation

This refers to the degradation of materials in the presence of oxygen and high temperature. It is normally associated with high temperature gas systems where the combination of steam sterilization can lead to the onset of oxidation of polypropylene filtration components in as little as 3 months. For applications where continuous (1 year and above) exposure to high temperature is required the use of a special product with oxidation resistant filtration support materials such as the HIGH FLOW TETPOR H.T. is recommended.

Oxidation can also occur on filters used in ozonated water systems. In these instances careful selection of filter components is required.



Pleating

Filtration media can be pleated or corregated to maximize the filtration area. By pleating filtration media it is possible to fit a large EFA in a relatively small cartridge volume.

Voids volume (porosity)

This is a measurement of the free space in a filtration media. The more free space the less the resistance to flow. Typical values for a membrane are in the region of 50 – 80% and for depth type media between 60 - 95%.

Pressure decay

A non-destructive integrity test method for membrane based filters. It involves wetting out every pore in the membrane structure with water or the process fluid or a low surface tension liquid in case of hydrophobic membrane. Compressed air is applied to the upstream side of the filter and gas diffuses through the wetted pores. This causes a pressure drop in the upstream side of the filter known as the pressure decay. The maximum allowable pressure decay for a filter is dependant on the upstream volume and therefore must be known.

Pressure Decay (mbar /min) =

Diffusional Flow (ml / min)
Upstream Vol (I)

Pyrogenicity

Pyrogenicity is the tendency of a substance to raise body temperature when injected into the body. Filtration materials that come in contact with injectable liquids must meet pyrogenicity standards and be classified as non-pyrogenic. Pyrogenicity can be determined by such standard tests as the Limulus Amoebocyte Lysate (LAL) test.

Permeate

Synonymous with filtrate

R

Regeneration

When a filter becomes blocked with protein based material it may be possible to regenerate, or clean the filter, so improving overall lifetime.

Reverse jetting

The application of high pressure compressed gas to the inside of a filter to release powder collected on its surface.

Reverse osmosis

Forcing a liquid through a non-porous membrane, removing particles, along with dissolved molecules and ions. Reverse osmosis is the finest form of membrane separation and is used to desalinate water for drinking, and in the preparation of ultrapure water for various industries.

S

Sanitization

Reduction not elimination of a microbial population to render a fluid/system free from spoilage organisms and increase shelf-life of products.

Sedimentation

The process by which suspended solid particles in a liquid phase gravitate downwards. Eventually they will settle on the bottom of the holding tank, pipework etc. The rate of sedimentation is governed by particle mass and fluid velocity.

Separation

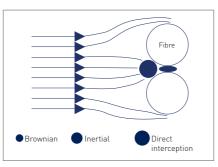
Separation is the process of dividing a fluid stream (either liquid or gas) into separate components. This can include separation of two phases (liquid from gas), separation of soluble impurities (known as purification) or solids from a fluid (filtration). The products of a separation can themselves be separated further in many cases.

Silt density index (SDI)

This is another measure of the rate of blockage and is typically used when the system is relatively clean and the difference between T_{400} and T_{200} [see Filterability Indices] is so small that large inaccuracies can occur. The SDI uses the time taken for two 500 ml samples of fluid to pass though a 47 mm diameter 0.45 μ m disc. There is typically a 15 minute gap between the two samples being taken.

Size exclusion

This is a removal mechanism for particles captured by a filter in either the liquid or gas phase. It applies to particles that are physically too large to pass through the filter structure. The mechanism is not affected by flow rate unless pressure drops cause deformation of the particle.



Solute

A solid which is dissolved in a solvent. For instance, the salt in salt water is a solute.

Solvent

A liquid substance capable of dissolving other substances. The solvent does not change its state in forming a solution.

Stabilization

This is the reduction in microbial loading in a fluid system and is generally associated with the beverage industry where partial rather than complete removal of spoilage organisms may be required to extend shelf-life.

Sterilization

In terms of filtration this means the elimination of all living microorganisms from the influent stream.

Surfactant

Acronym for a surface active agent. In filtration it is also sometimes called a wetting agent. If a filter is being used to filter aqueous solutions and incomplete wetting of the membrane pore structure is encountered a 'wetting agent' may be added to the membrane surface by flowing a quantity of surfactant through the filter. However, the use of a wetting agent is not desirable, especially in a pharmaceutical environment, as there is also the possibility of the surfactant leaching from the filter into the filtrate during processing or steam sterilization, etc.

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Thermal stability

This is most important during sterilization of the filter. The majority of cartridge and disposable type filters are manufactured from polymers such as polypropylene and nylon. During sterilization the components of the filter expand and contract putting great strain on the device. The filter performance with respect to steam sterilization should be matched closely to the requirements of the process. It should be noted that some filter configurations cannot be in-situ steam sterilized but can only be autoclaved.

Titre reduction

See I RV.

Turbidit

This is a measurement of the amount of suspended particles in a fluid and is effectively a clarity index. It is measured in NTU (Nephelometric Turbidity Units).

Glossary of terms used in filtration



Unloading

The release of contaminants which had initially been captured by a filter. This is most likely to occur in filtration systems with are subjected to high pressure pulses such as high capacity filling lines.

Ultrafiltration

Filtration of a liquid that separates suspended or dissolved substances based on their molecular weight or size. Ultrafiltration generally refers to separating everything larger than a large molecule. Compare to microfiltration, nanofiltration, reverse osmosis.



Viscosity

Viscosity is a measurement of the resistance to flow of a fluid. The more viscous the fluid, the greater the time required to filter. Viscosity will in general reduce with an increase in temperature. This is why very viscous solutions such as glucose are heated prior to filtration.

Vma

See Filterability Indices.



Water flow

Measure of the amount of water that flows through a filter. Related to the degree of contamination, differential pressure, total porosity, and filter area (ASTM:F317-72). Expressed in the membrane industry in units of millilitres / minute / square centimetre.

Water Intrusion

A non-destructive integrity test method specifically designed for hydrophobic filters. It involves filling the upstream volume of a filter housing with water and applying a pressure, typically in the order 2.5 barg. As the membrane is hydrophobic the bulk water will not pass through. However, due to the difference in pressure between the upstream and downstream side of the filter there is a net loss of water from the upstream side due to evaporation and the slight penetration of water into the pore structure. This loss of water results in a pressure drop which is displayed as either a water intrusion value or a water flow value. The water intrusion is the measure of the increase in compressible gas volume expressed at atmospheric pressure and the water flow equates to the volume of water lost from

Water flow = Water Intrusion / Absolute test pressure.

Industrial products

Parker domnick hunter, Industrial Division, is a well established global business capable of meeting the compressed air treatment product needs of all industries. Our commitment to customer satisfaction goes beyond initial supply and installation. Comprehensive aftersales support includes servicing, spare parts, quality testing and technical advice.

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- Low pressure loss / low operational cost



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generators from Parker domnick

hunter includes UHP hydrogen,

nitrogen and zero air and enables

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continuous supply of premium quality gas from a compact,

Increases safety with the

Cost-effective due to low

life-cycle ownership

UHP hydrogen generators

· Convenient, on-demand

gas supply

facilitate optimized analysis

elimination of high pressure gas

storage or cylinder handling

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- · A more efficient operation
- Improved shelf-life



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- consumption is kept to a minimum
- Totally stops corrosion / damage
- Low installation costs • Energy efficient

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way to dispose of oil contaminated

compressor condensate, ES2000

series oil water separators are a

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- Rapid payback over conventional



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- · Regional transports

Key Products

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- Fluid conveyance systems
- Fluid metering delivery & atomization devices
- Fuel systems & components
- Hydraulic systems & components
- Inert nitrogen generating systems Pneumatic systems & components Wheels & brakes



CLIMATE CONTROL Key Markets

- Agriculture
- - Food, beverage & dairy · Life sciences & medical
 - Precision cooling
 - Processing Transportation

Key Products

- CO² controls Electronic controllers
- Filter driersHand shut-off valves
- Hose & fittings
- Pressure regulating valves
- Refrigerant distributors · Safety relief valves
- Solenoid valves
- Thermostatic expansion valves



- Air conditioning
- - Paper machinery Plastics machinery & converting
 - Primary metalsSemiconductor & electronics

· Wire & cable

- **Key Products** AC / DC drives & systems
- Electric actuators, gantry robots & slides
- · Electrohydrostatic actuation
- Flectromechanical actuation
- Human machines interface Linear motors
- Stepper motors, servo motors, drives & controls





FLUID & GAS HANDLING

Key Markets

- Aerospace
- Agriculture
- Bulk chemical handling Construction machiner
- Fuel & gas deliven Industrial machinery

Welding

- Mobile · Oil & gas

Key Products

- Brass fittings & valves
- Diagnostic equipment Fluid conveyance systems
- Industrial hose . PTFE & PFA hose, tubing & plastic
- Rubber & thermoplastic hose &
- couplings

 Tube fittings & adapters



- Diagnostic equipment
- Hydraulic systems



- Oil & gas
- Hydraulic cylinders
- Rubber & thermoplastic hose &
- Quick disconnects



HYDRAULICS

- AerospaceAerial lift
- Construction machinery
- Truck hydraulics
- **Key Products**
- Hydraulic válves & controls
- couplings

 Tube fittings & adapters



- **Key Markets**
- Agriculture
- Forestry Industrial machinery
- Power generation & energy

- Power take-offs



- Aerospace
- Mining

- & accumulators Hydraulic motors & pumps



PNEUMATICS

- **Key Markets**
- Conveyor & material handling Factory automation · Life science & medical
- Machine tools · Packaging machinery · Transportation & automotive

Air preparation

Quick disconnects

Rotary actuators

- **Kev Products**
- · Brass fittings & valves Manifolds Pneumatic accessories Pneumatic actuators & grippers Pneumatic valves & controls
- Rubber & thermoplastic hose & couplings · Structural extrusions

Vacuum generators, cups &

Thermoplastic tubing & fittings



Key Markets

Chemical & refining

· Food, beverage & dairy

Analytical sample conditioning

Fluoropolymer chemical delivery

High purity gas delivery fittings

Instrumentation fittings, valves &

Medium pressure fittings & valves

nrnducts & systems

valves & regulators

Process control manifolds

fittings, valves & pumps

Medical & dental

Microelectronics

Power generation

Key Products

Oil & gas

PROCESS CONTROL **SEALING & SHIELDING**

- **Key Markets**
- . Energy, oil & gas Fluid power

Military Semiconductor

- **Key Products** Analytical sample conditioning
- Elastomeric o-rings EMI shielding
- · Homogeneous & inserted elastomeric shapes High temperature metal seals
- Thermal managemen





- Industrial machinery Life sciences Marine
 - Mobile equipment Oil & gas
 - Power generation Process • Transportation

FILTRATION

Key Markets

Food & beverage

- **Key Products**
- Analytical gas generatorsCompressed air & gas filters Condition monitor Engine, air, fuel & oil filtration &
- Process, chemical, water 8 Nitrogen, hydrogen & zero air



- Chemical processing
- General industrial Information technology Life sciences
- Telecommunications Transportation
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Parker Worldwide

Europe, Middle East, Africa

AE - United Arab Emirates, Dubai

Tel: +971 4 8127100 parker.me@parker.com

AT – Austria, Wiener Neustadt Tel: +43 (0)2622 23501-0 parker.austria@parker.com

AT – Eastern Europe, Wiener Neustadt

Tel: +43 (0)2622 23501 900 parker.easteurope@parker.com

AZ - Azerbaijan, Baku Tel: +994 50 2233 458 parker.azerbaijan@parker.com

BE/LU – Belgium, Nivelles Tel: +32 (0)67 280 900 parker.belgium@parker.com

BY - Belarus, Minsk Tel: +375 17 209 9399 parker.belarus@parker.com

CH – Switzerland, Etoy Tel: +41 (0)21 821 87 00 parker.switzerland@parker.com

CZ - Czech Republic, Klecany Tel: +420 284 083 111 parker.czechrepublic@parker.com

DE - Germany, Kaarst Tel: +49 (0)2131 4016 0 parker.germany@parker.com

DK - Denmark, Ballerup Tel: +45 43 56 04 00 parker.denmark@parker.com

ES - Spain, Madrid Tel: +34 902 330 001 parker.spain@parker.com

FI - Finland, Vantaa Tel: +358 (0)20 753 2500 parker.finland@parker.com

FR - France, Contamine s/Arve Tel: +33 (0)4 50 25 80 25 parker.france@parker.com

GR - Greece, Athens Tel: +30 210 933 6450 parker.greece@parker.com

HU - Hungary, Budapest Tel: +36 1 220 4155 parker.hungary@parker.com IE - Ireland, Dublin Tel: +353 (0)1 466 6370 parker.ireland@parker.com

IT – Italy, Corsico (MI) Tel: +39 02 45 19 21 parker.italy@parker.com

KZ - Kazakhstan, Almaty Tel: +7 7272 505 800 parker.easteurope@parker.com

NL - The Netherlands, Oldenzaal Tel: +31 (0)541 585 000 parker.nl@parker.com

NO - Norway, Asker Tel: +47 66 75 34 00 parker.norway@parker.com

PL - Poland, Warsaw Tel: +48 (0)22 573 24 00 parker.poland@parker.com

PT - Portugal, Leca da Palmeira Tel: +351 22 999 7360 parker.portugal@parker.com

RO – Romania, Bucharest Tel: +40 21 252 1382 parker.romania@parker.com

RU - Russia, Moscow Tel: +7 495 645-2156 parker.russia@parker.com

SE – Sweden, Spånga Tel: +46 (0)8 59 79 50 00 parker.sweden@parker.com

SK - Slovakia, Banská Bystrica Tel: +421 484 162 252 parker.slovakia@parker.com

SL – Slovenia, Novo Mesto Tel: +386 7 337 6650 parker.slovenia@parker.com

TR - Turkey, Istanbul Tel: +90 216 4997081 parker.turkey@parker.com

UA - Ukraine, Kiev Tel +380 44 494 2731 parker.ukraine@parker.com

UK - United Kingdom, Warwick Tel: +44 (0)1926 317 878 parker.uk@parker.com

ZA – South Africa, Kempton Park Tel: +27 (0)11 961 0700 parker.southafrica@parker.com

North America

CA – Canada, Milton, Ontario Tel: +1 905 693 3000

US – USA, Cleveland Tel: +1 216 896 3000

Asia Pacific

AU – Australia, Castle Hill Tel: +61 (0)2-9634 7777

CN - China, Shanghai Tel: +86 21 2899 5000

HK – Hong Kong Tel: +852 2428 8008

IN - India, Mumbai Tel: +91 22 6513 7081-85

JP – Japan, Tokyo Tel: +81 (0)3 6408 3901

KR - South Korea, Seoul Tel: +82 2 559 0400

MY - Malaysia, Shah Alam Tel: +60 3 7849 0800

NZ - New Zealand, Mt Wellington

Tel: +64 9 574 1744

SG – Singapore Tel: +65 6887 6300

TH - Thailand, Bangkok Tel: +662 717 8140

TW – Taiwan, Taipei Tel: +886 2 2298 8987

South America

AR – Argentina, Buenos Aires Tel: +54 3327 44 4129

BR - Brazil, Sao Jose dos Campos Tel: +55 12 4009 3500

CL - Chile, Santiago Tel: +56 2 623 1216

MX - Mexico, Apodaca Tel: +52 81 8156 6000

VE – Venezuela, Caracas Tel: +58 212 238 5422 3, 2010-09-29

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Europe & the rest of the world
Parker Hannifin Ltd
Parker domnick hunter
Process Division
Durham Road
Birtley, Co. Durham
DH3 2SF, England
phone +44 (0)191 4105121
fax +44 (0)191 4105312
email: dhprocess@parker.com
www.parker.com/processfiltration

North & South America
Parker Hannifin Corporation
Parker Process Advanced Filtration
2340 Eastman Avenue
Oxnard, California, USA 93030
toll free: +1 877 784 2234
phone: +1 805 604 3400
fax: +1 805 604 3401
email: PAFsales@parker.com
www.parker.com/processfiltration