

Clean-in-place Best Practice Guide



ENGINEERING YOUR SUCCESS.

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Parker Bioscience Filtration

Setting the standard

Parker brings extensive experience through our scientists, engineers and sales representatives to the process of offering specific filtration systems to meet the needs of your production process. Support services are available covering a wide range of activities including scale-up advice from laboratory through pilot scales to production systems, validation support on-site technical support, design and manufacturing of custom housings and filtration products.

Committed to quality

Quality is of paramount importance to Parker. As such we have been certified to ISO9001, providing a quality management system that covers the entire organization including R & D, production, warehousing, materials management and customer support. In addition, our manufacturing facilities operate to the principles of cGMP.

Our manufacturing facilities are also certified to ISO14001 Environmental Management Standard and ISO13485 Medical Device requirements.

Validation and product certification

To certify that Parker products meet the required regulatory and quality standards of the industries that we supply, all filters are supplied with a certificate of conformance. These certificates are linked to validation documents for both prefilter and sterilizing grade membrane filter products that define methodologies and data appropriate to each filter type. This information typically includes:

- Technical specifications.
- Biological safety testing information; USP <88> Class VI - 121C Plastics and equivalents.
- Extractable testing including 21CFR211.72 and 210.3(b), 6 for fibre release.
- Purified water filtration quality including TOC, bacterial endotoxins, conductivity and particle release.
- Chemical compatibility guidance information.
- Thermal stability.
- Correlation of an appropriate non-destructive integrity test to a defined bacterial challenge.
- EC Food contact safety specification.

Validation support services

Parker has extensive laboratory facilities and trained personnel capable of providing a range of validation services to support manufacturers in meeting their requirements for process validation relating to the use of filtration products.

Clean-in-place

Successful and economic use of process cartridge filters is dependent upon employing correct procedures to ensure that the filters are not damaged and their lifetime in the application is maximized. Parker cartridges are designed to withstand adverse conditions of use, but by adhering to the following guidelines, the likelihood of premature blockage, integrity test failure and process stoppages will be reduced.

This document provides guidelines that are intended for incorporation into Standard Operating Procedures. Recommendations should be considered flexible and continuous monitoring should be carried out to improve the process as on-going experience is gained.

The following 4 guidelines are key to a good CIP practice:

- Good regular cleaning can greatly increase the life of a filter. Little and often is better than an occasional strong clean.
- Improving efficiency of cleaning can result in increased filter life.
- Effective sanitization is important to maintain product quality.
- Inappropriate cleaning regimes can damage filters.

The following three-stage filtration system (Figure 1) with ancillary supplies is used as an example throughout the document. Between 'process in' and 'process out' the 3 filter stages can be seen. Over each filter system it is recommended that the differential pressure (dP) be monitored and recorded by (automated) pressure sensors. The temperature should be measured and recorded downstream of the final filter. A CIP system (upper left) can provide flushing and cleaning of the system. A steam supply can be used to steam sterilize the filter system.

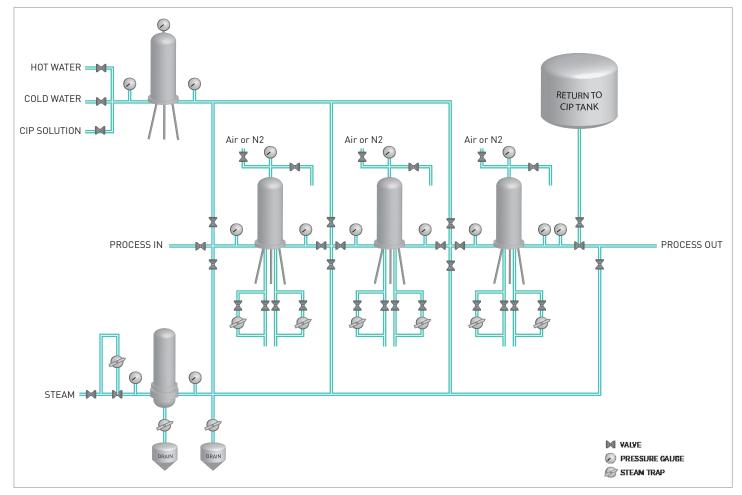
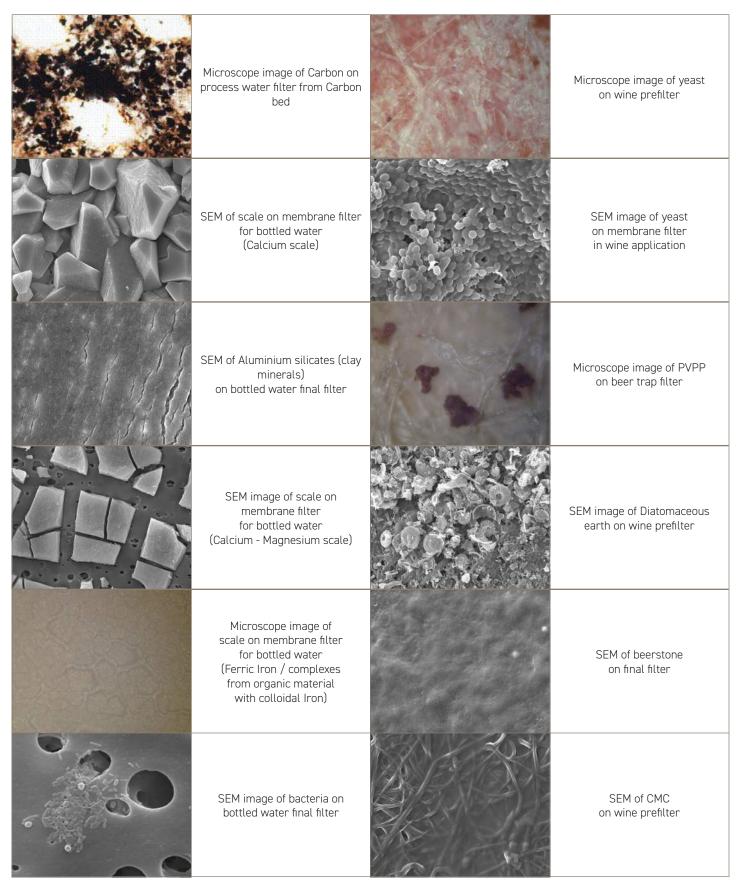


Figure 1 - Example three-stage filtration system

Usual suspects

The following pictures show typical contaminants on filter media in water, beer and wine applications.



Guideline parameters

Water and cleaning solutions

All water and cleaning solutions should be prefiltered prior to contacting the process filters in order to avoid filter blockage occurring due to the CIP process. The protection offered to the process filters is greater if finer prefilters are used. The CIP filter rating chosen is dependent on the tightest micron rating of filters in the filter train that receives the CIP solution direct.

In the example filter train 10 micron PEPLYN HA – 1.0 micron PREPOR NG – 0.45 micron BEVPOR PH, if the CIP solution is brought first over the 10 micron filter followed by the other stages, then this filter will determine the retention rating of the CIP filter. However, if the cleaning solution is brought simultaneously over the 3 stages, then the filter for the CIP solution must be sufficiently fine to protect the finest pore size of the process filters. In the example that is the 0.45 micron BEVPOR PH filter.

The following guidelines can be used:

- When cleaning depth (media) filters, use a depth filter with a micron rating of up to two grades higher. For example, a 20 micron PEPLYN PLUS CIP prefilter for a 10 micron PEPLYN PLUS process filter.
- If back-washing a depth filter that is suitable for back flushing, use the same grade or finer for prefiltration purposes to avoid contaminating the downstream side of the filter being back-washed.
- When cleaning membrane final filters directly, prefilter using a media cartridge with an absolute rating one grade higher or a membrane filter up to two grades higher. For example for a 0.45 micron BEVPOR PH filter, use a 0.6 micron PEPLYN PLUS.

In both cases, assess performance with time in order to optimize the lifetime to blockage of the individual stages.

Cleaning solutions should be freshly prepared.

Conductivity of cleaning solutions can be monitored to ensure that the correct concentrations are used. Suppliers of cleaning solutions can provide information on conductivity values relative to concentration.

Temperature	Gradual changes, no shock
Cold	4 °C to 30 °C (i.e. ambient)
Warm	40 ℃ to 60 ℃
Hot	60 ℃ to 95 ℃
Duration	15 to 30 minutes at required temperature
Flow rate	Ideally 20% of process flow rate
Direction	Forward direction unless back flush on trap filter
Back flush	Up to 1.5 times the process flow

Gradual temperature

Maximum differential pressure (dP)

What is differential pressure?

- Differential pressure is the difference in pressure between two points in a system. In a filtration system, the differential pressure is the pressure between the upstream side of a filter and the downstream side of a filter.
- Differential pressure is directly proportional to flow rate (Where velocity does not limit flow).

The following example shows the differential pressure over a two-stage system:

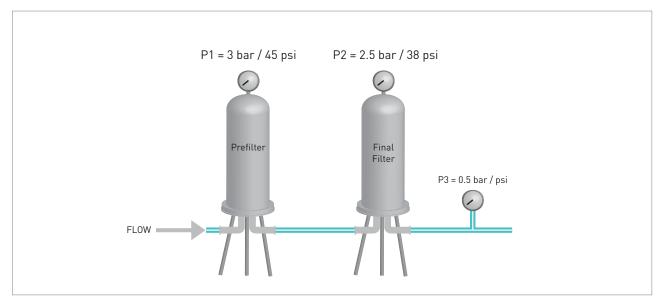


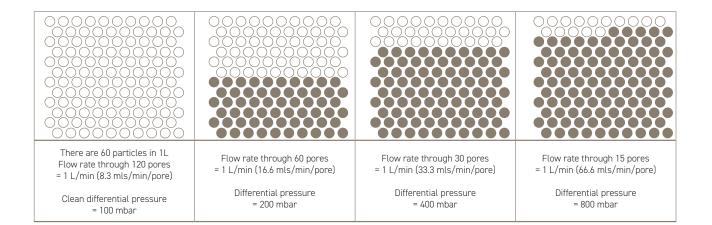
Figure 2 - Differential pressure over a two-stage system

	Prefilter	Final filter
Differential pressure (dP)	P1 - P2 = 0.5 bar	P2 - P3 = 2.0 bar

Relationship between dP and blockage:

If it is assumed that blockage is a constant process. Then when the available surface area is 50% blocked the differential pressure is increased by a factor of 2.

This is due to the fact that the same flow rate is being processed through an area half of the original filter surface area. This is shown in the following diagrams.



With this data a typical blockage curve can be drawn:

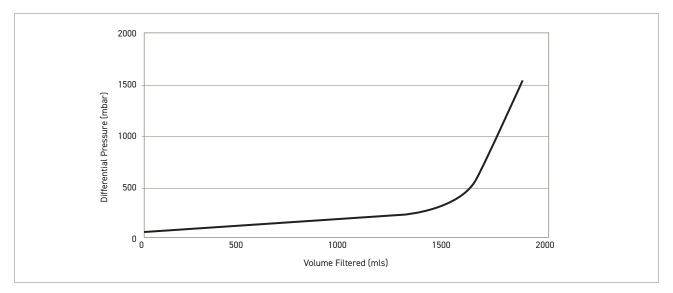


Figure 3 - Typical blockage curve

Depth (media) filters

Temperatura	Maxim	um dP (bar)	
Temperature	Forward flow	Reverse flow*	
20	5	2.0	
40	4	1.5	
60	3	1.0	
80	2	0.5	
90	1	Not recommended	
>100 (Steam)	0.3	Not recommended	

Table 1 - Maximum recommended differential pressure over depth filters

*Reverse flow is only recommended for PEPLYN range, PREPOR PP and PREPOR NG pleated polypropylene depth (media) filters. Back washing clarification filters is a generally accepted technique and some filters, such as PEPLYN HA, have been specifically designed to improve the efficiency of the back-wash operation.

Always consult individual cartridge specification before adopting backwash cleaning. Close control of the conditions is required, although these pleated polypropylene products are robust and will accept some abuse.

Membrane filters

Back washing of membrane filters is not recommended.

Temperature	Maximum dP (bar)
	Forward flow
20	5
40	4
60	3
80	2
90	1
>100 (Steam)	0.3

Table 2 - Maximum recommended differential pressure over membrane filters

When to clean ?

Cleaning frequency can be based on time or blockage. As blockage occurs exponentially, filters should be cleaned before rapid, unrecoverable blockage occurs. Typically CIP should be conducted when a rise in the dP starts to occur (ideally when the dP rises above 100 to 200 mbar over the starting dP at process flow rate with clean filters) to remove blockage particulate and prolong the lifetime of the filters.

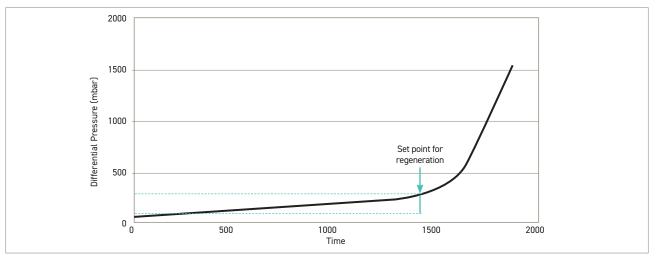


Figure 4 - Example of when to perform a CIP

CIP can also be based on time, for example weekly to control microbial levels or can be performed between batches. Good regular cleaning can greatly increase the life of a filter resulting in improved economics. Little and often (figure '4A') is better than an occasional strong clean (figure '4B').

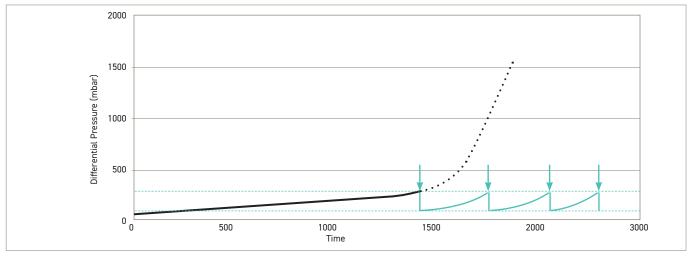


Figure 4A - Example of regular clean extending life to blockage

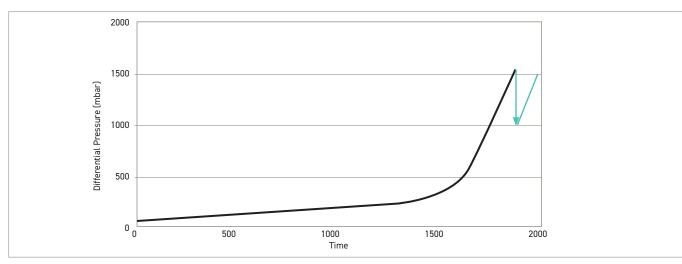


Figure 4B - Example of occasionally strong clean not reducing dP

Compatibility

Before using any chemicals, check the compatibility of the filter prior to use. If in doubt, contact Parker. As Parker exercises a policy of continual product development cartridge specifications may change, however, the general guidelines below can be applied.

Active agent	Condition*	Parker Product	Guideline cumulative contact time
		BEVPOR	1000 hours ¹
Hot water	85 °C	PEPLYN, PREPOR PP, PREPOR NG	245 hours ¹
		PREPOR GF, PREPOR GP	100 hours ²
		BEVPOR	25 hours ²
Steam	121 °C	PEPLYN, PREPOR PP, PREPOR NG	100 hours ²
		PREPOR GF, PREPOR GP	10 hours ²
	0.352% NaOH (and <0.6% EDTA) @ 60°C	BEVPOR	336 hours ³
Caustic (NaOH) in a buffered sanitizer		BEVPOR, PEPLYN, PREPOR PP, PREPOR NG	>1000 hours ³
Summer	0.352% NaOH (and <0.6% EDTA)	PREPOR GF, PREPOR GP	18 hours ²
	0.54% PAA (and <0.54% ${\rm H_2O_{2'}}$ <0.54% AA)	BEVPOR	>1000 hours ³
Peracetic Acid (PAA)	0.15% PAA (and <0.15% H ₂ O ₂ , 0.15% AA) @ 50°C	BEVPOR	240 hours ⁴
	0.05% (500 ppm) PAA	PREPOR GF, PREPOR GP	168 hours ²
	0.15% H ₂ O ₂ @ 50 °C	BEVPOR	600 hours ⁴
Hydrogen Peroxide (H ₂ O ₂)	0.4% H ₂ O ₂ (and <0.13% AA, <0.1% PAA)	BEVPOR	600 hours ⁴
	0.2% $\rm H_{2}O_{2}$ (and <0.065% AA, <0.1% PAA) @ 50 $^{\circ}\rm C$	BEVPOR	240 hours ⁴
Phospheric (H ₃ PO ₄) / Nitric (HNO ₃) Acid	1% HNO ₃ and 1% H ₃ PO ₄ @ 55 °C	BEVPOR	256 hours⁵
	2% SO ₂	BEVPOR	168 hours ²
Sulphur Dioxide (SO ₂)	0.5% SO ₂	BEVPOR, PEPLYN, PREPOR GF, PREPOR GP	756 hours ²
	10ppm free Cl	BEVPOR, PEPLYN, PREPOR PP, PREPOR NG	336 hours ⁹
Chlorine Dioxide (ClO ₂) Sodium Hypochlorite	1700ppm total Cl	BEVPOR	>1000 hours ¹⁰
Chlorinated Alkaline	<500ppm Chlorine (<0.15% KOH) @ 50°C	BEVPOR	72 hours (at least) ⁸

*All conditions at ambient (25 °C) unless otherwise stated.

References: 1 = T6562, 2 = Beverage Cleaning Guidelines (2003), 3 = T9122, 4 = T9123, 5 = T8716, 7 = T9146, 8 = T8983, 9 = T7439, 10 = VSG9256/TR13341.

Notes:

The guidelines are based on tests with chemicals from commonly used industry suppliers and the concentrations are based upon the information in the TDS / MSDS.

Testing was conducted using clean, unused filters in controlled laboratory conditions and therefore can not simulate the effects caused by use in applications where the filter componentry may be weakened by prolonged exposure to aggressive process liquids and exposure to high dP through blockage or high flow.

It is recommended to regularly integrity test (for example weekly) BEVPOR filters in processes to check their performance. Integrity testing will enable the user to verify that no filter damage has occurred during storage, installation or following procedures such as chemical cleaning prior to use or during the subsequent manufacturing process.

Integrity testing

Most membrane filters can be tested to ensure that they are integral before use. This should always be carried out to ensure that the cleaning procedures have not damaged the cartridges in any way. Detailed integrity test procedures and training are available on request.

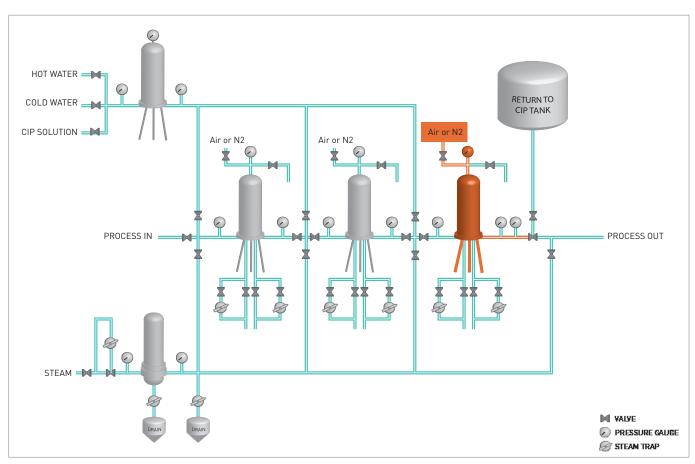


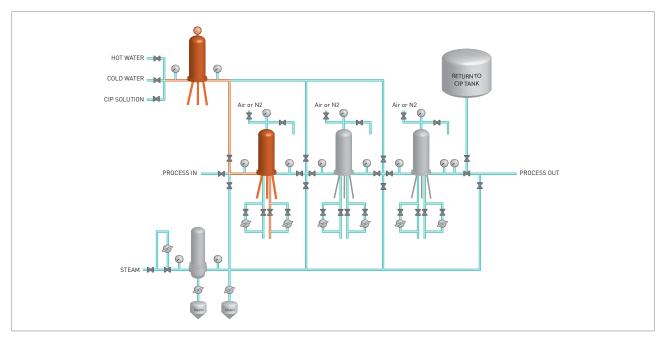
Figure 5 - Filtration system with recommended integrity test connections

Clean-in-place procedures

			Recommendations		
Procedure	When	Frequency	Temperature °C	Duration (mins)	Flow (% of process flow rate)
Cold water chase	End of batch, change of product	N/A	5-30	5	20
Cold water rinse	Change of product, regeneration / IT	Daily	5-10	10	20
Back washing	Regeneration	Daily, weekly	5-30	10	150-200 reverse
Warm or hot water rinse	Change of product, regeneration	Daily	40-80	10	20
Chemical regeneration	Change of product, regeneration	Daily, weekly	20-60	20	20 Soak after 5 mins
Chemical sanitization	Sanitization	Daily, weekly	20-60	20	20 Soak after 5 mins
Hot water sanitization	Sanitization	Daily, weekly	80-90	20	20
Steam-in-place	Sanitization	Daily, weekly	105-130	20	Very low steam flow
Autoclaving	Sanitization	Prior to storage	121	20	N/A
Short-term storage	Storage	-	20	N/A	N/A
Long-term storage	Long-term storage	-	20	N/A	N/A

Table 3 - details on CIP procedures

To avoid the transport of particles from the prefilters to the final filter, the cleaning solution over a filter should always be drained whilst that filter is being rinsed. When the cleaning solution at the drain runs clear the inlet to the next filter can be opened and the drain closed so the cleaning solution flows to the next filter. This procedure is shown in Figures 2, 3 and 4.





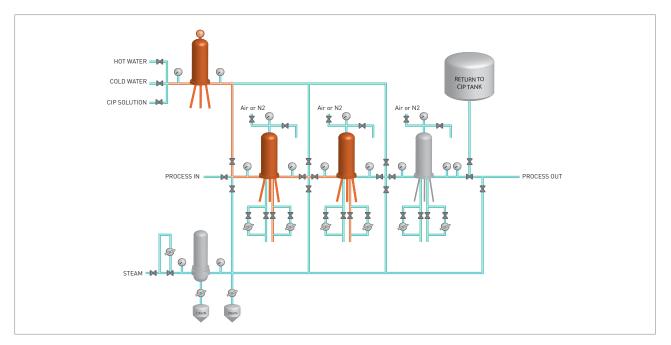


Figure 7 - Forward rinse/clean of the second filter stage

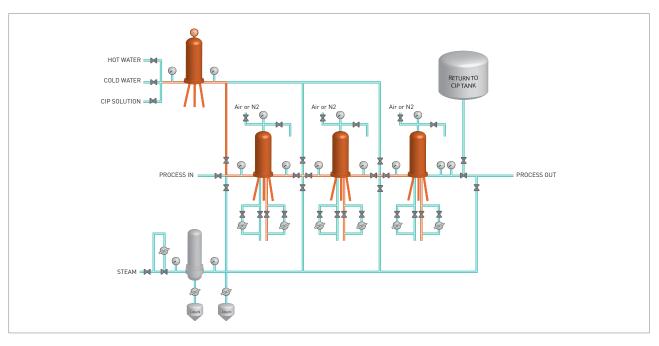


Figure 8 - Forward rinse/clean of the third filter stage

Alternatively during any rinsing or cleaning period, filter stages can be cleaned simultaneously. The advantage of rinsing simultaneously is that the overall cleaning time can be reduced, however, if this is carried out the prefilter for the water (or CIP solution) must be sufficiently fine to protect the finest pore size of process filter. This is shown in the following figure (figure 5).

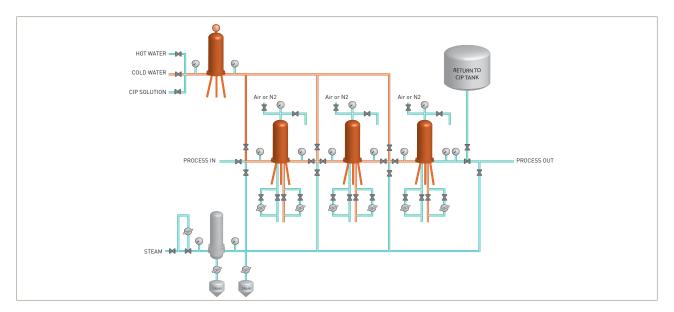


Figure 9 - Simultaneous rinse/clean of the three filter stages

Cold water chase

This is a cold water flush carried out to push product out of the line. It is often controlled manually by observing the change from product to water through a sight glass, or by on line conductivity measurement. It is carried out to maximize the volume of product transferred, but also to ensure that water-insoluble species are pushed through the media in the process liquid, thereby preventing them from binding to the filter media and causing premature blockage.

Conditions		
Temperature 5 °C to 30 °C		
Duration	Maintain for 5 minutes	
Flow rate 20% of process flow rate		
Procedure		
Cold water flush to package or transfer tank		

Stop when clean water is detected in downstream sight glass or by change in conductivity

Cold water rinse

This is a general rinse to clear the filters of product and water soluble blocking materials, or cleaning agents or before first use to ensure that no debris is in the filter after installation. This step is also used to wet out filters for integrity testing.

Conditions	
Temperature	5 °C to 30 °C
Duration	Maintain for 5 minutes
Flow rate	20% of process flow rate

Procedure
Isolate housings
Flush to drain until clear in first housing
Open the inlet to the next housing and close the drain
Repeat for all housings in series
Rinse all housings simultaneously for the specified time
Drain the system or proceed to next operation

Back washing

Back washing can be carried out as part of any of the rinsing or cleaning procedures. Back wash is usually only effective for prefiltration stages. Backwash should not be performed on membrane filters as the chance of damage is high, and blockage will be unaffected.

The flow rate used during backwashing should be as high as possible without exceeding the recommended maximum differential pressure of the cartridge. In small systems this is often possible. In larger systems, pump capacity may limit the flow. As a guideline, a flow rate of 1.5 to 2 times the process flow rate is recommended.

Conditions	
Temperature	5 °C to 30 °C
Duration	Maintain for 10 minutes
Flow rate	150 - 200% of process flow rate, reverse flow

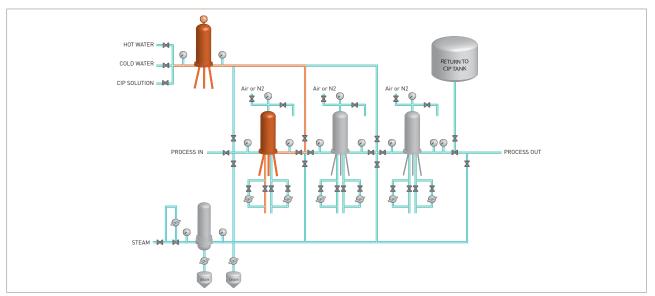


Figure 10 - Simultaneous rinse/clean of the three filter stages

Warm / hot water rinse

Again this is a general rinse to further clear the filters of product and water soluble blocking materials. This procedure can also serve to gradually increase the temperature of the filter in preparation for hot water sanitization or steam sterilization.

Conditions			
Temperature 40 °C to 80 °C			
Duration	Maintain for 10 minutes		
Flow rate	20% of process flow rate		
	Procedure		
Isolate housings			
Introduce warm or hot water to the cold water contained in the first filter housing			
Flu	Flush to drain until clear in first housing		
Open inlet to next housing and close the drain			
Repeat for all housings in series			
Rinse for specified time			
Drain the system or proceed to next operation			

Chemical regeneration

Chemical cleaning is performed to remove blocking materials that are not removed by water rinses, e.g. colloidal iron complexed with organic material, scales, trapped proteins, sugars, CMC. This can provide regeneration of the filters. The CIP could effectively increase the life to blockage of the filter by removing some of the contamination from the filter media. This can be enhanced by introducing a specific regeneration procedure.

The procedure can be simplified if the filter is compatible with chemicals already in use for other clean-in-place purposes. However, incorporating the filter CIP with a full system CIP may be false economy, as debris washed from the inner surfaces of lines and fittings may block the filter. Therefore it is recommended to prepare fresh cleaning chemicals for cleaning filters.

Conditions		
Temperature	20 °C to 60 °C	
Duration	Maintain for 20 minutes	
Flow rate	20% of process flow rate 5 mins Soak for 15 mins	

Procedure		
Cold water rinse		
Warm or hot water rinse		
Make up the required CIP solution		
Isolate housings		
Introduce CIP solution to the first filter housing		
Flush to drain until clear in first housing		
Open inlet to next housing and close the drain		
Repeat for all housings in series		
Circulate or stand for required duration		
Drain the system or proceed to next operation		

The advantage of carrying out the filter CIP in series through all of the filters is that the CIP solution is filtered by the existing process prefilters. In some circumstances it may be advisable to CIP each filtration stage separately, for example, if colour is not a good indicator of CIP solution cleanliness then apparently clean solution from the first filter may contain enough partially dissolved species to block the next filter. If the stages are cleaned separately then the prefilter for the CIP solution must be chosen so as to offer full protection to the tightest filter in the system.

Chemical sanitization

Particularly when steam is not available elevated temperature sanitization may be supported by chemical sanitizers. The following sanitizers can be used:

Solution	Concentration	Comments
Alcohol	75 % ethanol in water	Also has some regnerative effect.
Hypochlorite	150 ppm (initial) 50 ppm (long term)	May lead to flavour taints as difficult to rinse.
Sulphur dioxide	2 % in water	Only usable in sealed vessels due to liberation of gas into the atmosphere.
Peroxyacetic acid	500 ppm (initial) 100 ppm (long term)	Smells like vinegar but easily rinsed and small quantities quickly dissipate, usually leaving no flavour or odour taints.
Proprietary disinfectants	As manufacturers' instructions	Examples - Divosan Active, P3- Oxona Active.

Conditions		
Duration	Maintain for 20 minutes	
Flow rate	20% of process flow rate 5 mins Soak for 15 mins	

Procedure		
Cold water rinse		
Warm or hot water rinse		
Make up the required CIP solution		
Isolate housings		
Introduce CIP solution to the first filter housing		
Flush to drain until clear in first housing		
Open inlet to next housing and close the drain		
Repeat for all housings in series		
Circulate or stand for required duration		
Proceed to next operation		

Hot water sanitization

Hot water sanitization is no more than a continuation of the hot water rinse, although the necessity for flow is only to ensure that all parts of the system reach specified temperature for the required duration. The effectiveness of the procedure is enhanced by leaving the filter housing full of hot water and allowing it to cool naturally.

The following conditions are usually satisfactory to ensure adequate system and filter sanitization on a daily basis:

Conditions			
Temperature	Temperature 80 °C to 90 °C (85 °C recommended)		
Duration	Maintain for 15 to 30 minutes (20 minutes at 85 °C recommended) then close off housing and allow to cool down		
20% of process flow rateFlow rategentle flow to drain or circulation to stabilizetemperature throughout the system			
Procedure			
	Isolate housings		
Slowly introduce hot water to the cold water remaining in the housing from the previous step. This will ensure that the temperature increase is gradual, reducing thermal shock of the filters.			
Flus	Flush to drain until clear in first housing		
Open the i	Open the inlet to the next housing and close the drain		
Repeat for all housings in series			
Rinse all housings simultaneously for the specified time ensuring that the housings are at the specified temperature for the duration of the rinse (use temperature probes, temperature stickers on housings)			
Isolate housings and allow to cool naturally or proceed to next operation			

Steam-in-place (SIP)

Steam will effect a more efficient sanitization than hot water. By adopting the recommended temperature and duration, a properly designed system will be sterilized to standards accepted within manufacture of pharmaceuticals. The drawbacks are the availability of steam in some facilities, the additional management of the process and the corrosive and erosive effects on the filter, which may lead to shortened operational life. Where full sterility is required, or where hot water sanitization alone is known to be insufficient:

Conditions		
Temperature105 °C to 130 °C (121 °C recommended)		
DurationMaintain for 15 to 30 minutes(20 minutes at 121 °C recommended)		
Flow rate Minimal - controlled by 'cracked' valve, orifice plate or steam trap		
Procedure		
Cold water rinse		
Warm or hot water rinse		
Steam through filters in series or steam individual stages separately or steam final filter stage only		
Allow pressure and temperature to rise and fall slowly		
Close downstream valves as pressure approaches 0 barg (100 °C)		
Allow temperature to drop to 50 °C to 60 °C		

Minimal steam flow should be used, controlled through 'cracked' valves, steam traps or an orifice plate directed to drain. It is good policy to keep all external valves slightly 'cracked'. This helps prevent air traps, condensate build-up and helps to sterilize the valve itself. Good steam quality and condensate management are essential to protect the filter. Use of chemical CIP between the cold and hot water rinses will aid the SIP process by washing away contamination that may otherwise be 'baked' onto the filter by the steam. Cooling after steaming may be accelerated by flowing compressed gas across the upstream side of the cartridges (take care not to exceed maximum differential pressure).

See Application Support Document - Steam-In-Place for more information.

Autoclaving

Filter cartridges may be removed from their housings and steam sterilized in an autoclave. This would not normally be carried out as a regular procedure unless the process is prone to regular microbial contamination and on-line steam is not available. If this is the case, then when the filters are removed, the remaining system may be hot sanitized with over-strength chemicals which would otherwise damage the filter.

Procedure
Cold water rinse
Warm or hot water rinse
Chemical regeneration if needed
Cold water flush
Remove the cartridges from the housings with care
Place cartridges in an autoclave bag if possible
Place the cartridges on their side in the autoclave
Autoclave the cartridges on a 121 °C liquid cycle
When autoclave can be opened, immediately seal the autoclave bags if the cartridges are to be stored
Check o-rings prior to re-installation

Short-term storage

If the filters are not to be used straight away for filtering the next batch of product but will be used in the next 1 to 4 days or within a month, then it is recommended to store the filters in an appropriate medium to prevent microbial growth in the housing.

Before short-term storage the filters need to be sanitized (see hot water sanitization, chemical sanitization and SIP for more information).

After the sanitization of the filters, the filters can be stored in the following solutions.

Solution	Concentration	Comments
Peroxyacetic acid	100 ppm	Up to 24 hours
Peroxyacetic acid	500 - 2000 ppm	1-4 days
Hydrogen peroxide	500 - 2000 ppm	4 days - 1 month Refresh solutions every 7 days
Citric acid / $K_2S_2O_5$	1000 - 2000 ppm	Up to 30 days
Nitrogen blanket	Pure - pressure 1 - 1.5 barg	Up to 30 days Check pressure regularly

Long-term storage

For long-term storage (longer than 1 month) it is recommended to remove the cartridges from their housings. Care should be taken to ensure that they are placed in an area where they can not be damaged. If possible, they should be autoclaved or dried and placed in their original packaging for protection. Alternatively, the cartridges may be left in their housings or in a soak bath with suitable chemical sanitizer.

Procedure	
Cold water rinse	
Warm or hot water rinse	
Chemical regeneration if needed	
Chemical sanitization, hot water sanitization or SIP	
Cold water flush or allow filters to cool to below 40 $^\circ\mathrm{C}$	
Remove the cartridges from the housings with care	

After these procedures have been carried out, either autoclave the cartridges, dry them for a few hours at a temperature of 50 °C to 60 °C in a convection oven, or place them in a soak bath. If the latter is chosen, the chemical solution should be changed regularly according to the manufacturer's instructions.

If drying can not be achieved, storage in one of the following solutions:

- 70% ethanol.
- 0.1 % citric acid / K2S205, refresh the solution every 20 days.
- 50 ppm hypochlorite, refresh solution weekly or when below 40 ppm (test kits available from cleaning solution manufacturer).
- Specified storage solutions by the cleaning solution manufacturer. The chemical solution should be changed regularly according to the manufacturer's instructions. Please contact Parker for compatibility information.

Solution	Concentration	Comments
Peroxyacetic acid	100 ppm	Up to 24 hours
Peroxyacetic acid	500 - 2000 ppm	1-4 days
Hydrogen peroxide	500 - 2000 ppm	4 days - 1 month Refresh solutions every 7 days
Citric acid / $K_2S_2O_5$	1000 - 2000 ppm	Up to 30 days
Nitrogen blanket	Pure - pressure 1 - 1.5 barg	Up to 30 days Check pressure regularly

Chemical regeneration at high dP (blocked cartridges)

When dP is above 1 bar over the filter stage

Some filters could be regenerated by a strong chemical clean. This can only be used on Polypropylene type filters (PEPLYN range) and the PES type filters (BEVPOR P/M range).

Cleaning frequency should be based on time or blockage. As blockage occurs exponentially, filters should be cleaned before rapid, unrecoverable blockage occurs, therefore, a strong chemical cleaning procedure (when the dP over the filter stage at process flow rate above 1 bar) might not work to regenerate the filters.

• CIP procedure using regeneration solution. Use gentle flow of solution in forward direction to minimize the dP over the filter stage during the cleaning procedure.

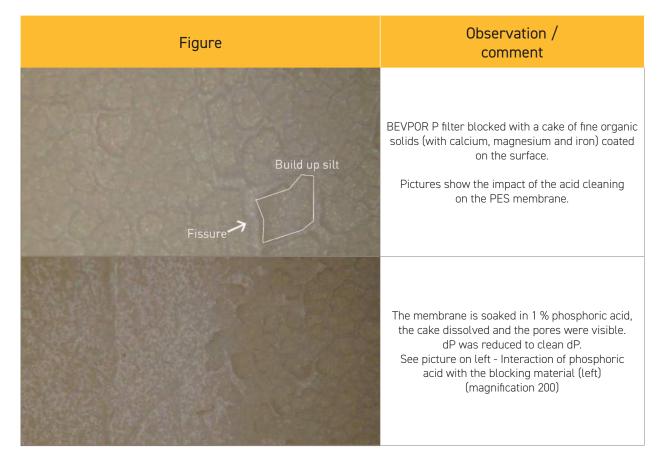
OR

• Remove cartridges from their housings, remove o-rings and place the cartridges in a bath of regeneration solution and agitate regularly.

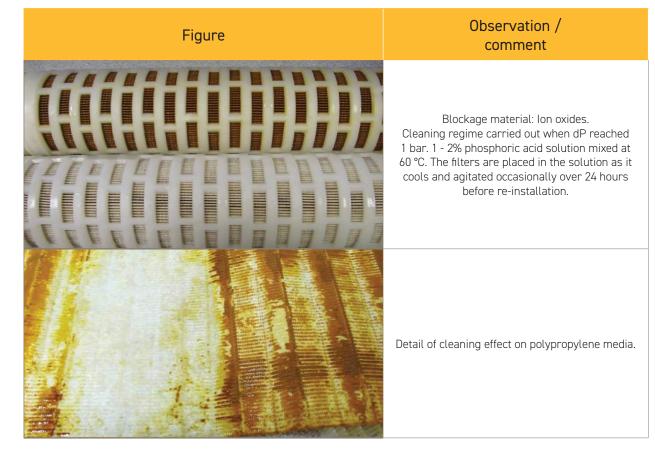
Solution	Concentration	Comments
Phosphoric acid	1% at 55 °C	Example 1 Effective in removing all alkaline hardness (calcium carbonate)
Nitric acid	1 % at 55 °C	Example 2 Also used in cleaning filters with iron oxide deposits and colloidal iron
Sodium hydroxide	0.5 M (2 %) at 60 °C	Example 3 Effective in removing soluble components
Proprietary cleaners	As manufacturers' instructions	Check with Parker domnick hunter prior to use

Regeneration examples

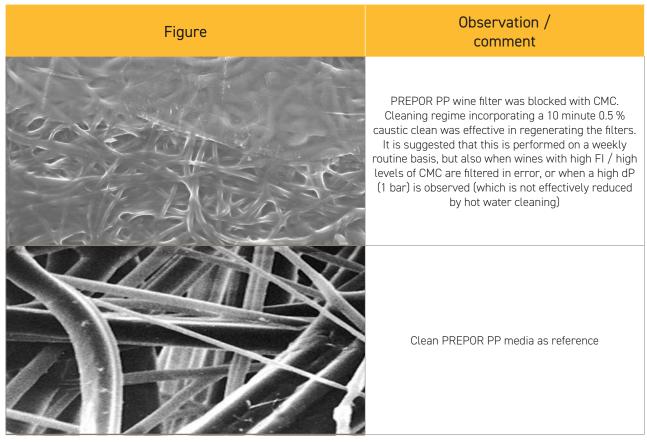
Example 1 - BEVPOR PH - water filtration



Example 2 - PEPLYN HA - water filtration



Example 3 - PREPOR PP- wine filtration



CIP approach

This section describes the effectiveness of a chemical type to a certain contaminant for performing CIP. The information is summarized in the table below.

Contaminant	CIP technique						
	Backflush water	Hot water	Alkali	Acid	Disinfectant / oxidant	Enzymes	
Debris (sediment)							
Debris (organic)							
Diatomaceous Earth							
Finings (PVPP)							
Micro-organisms							
Scale (Ca, Mg-carbonates, phosphates)							
Colloidal complexes							
Fats							
Proteins							
Polysacharides							
Acids							
CMC							

Process simulations

The following process simulations have been carried out for BEVPOR P cartridges. The table below shows the simulated process CIP, details on chemical, concentration and temperature, filter type and recommended maximum number of cycles.

Active agent	Condition	Parker product	Guideline cumulative contact time	
PAA hot water steam	1500ppm Peracetic acid (flush for 30 minutes) 80 °C hot water (flush for 30 minutes) steam @ 130°C	BEVPOR	30 cycles ¹	
Hot water NaOH/KOH/H ₂ O ₂	Hot water 80 °C for 30 minutes 1.25 % caustic for 10 minutes 0.08% Hydrogen Peroxide for 10 minutes (0.02 % AA and 0.02 % PAA)	BEVPOR	> 60 cycles ²	

References: 1 = T9050 / T9051

² = T10462

Technical Support Group activities

Parker have a trained team of scientists and engineers available to answer questions regarding the technical capabilities of our products, to assist in the selection and design of appropriate filtration systems and to provide user training programs. The following services can be delivered both on-site and in-house:

- Filterability testing to optimize filter system design
- · Advice on the development of integrity testing, steam sterilization and clean-in-place procedures
- Development of validation protocols
- Troubleshooting
- Facility audits to ensure continued optimization of filter use
- · Operator training including filtration theory, validation, filter system design and management

For more information on any of the above support services please contact your local Parker representative.

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