

Pre-treatment of surface waters for ceramic microfiltration

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Abbreviations:

BBs – Building blocks
BPs – Biopolymers
BW – Backwash
CEB – Chemically enhanced backwash
CIP – Cleaning in place
CMF – Ceramic microfiltration
DOC – Dissolved organic carbon
HMW – High molecular weight
HSs – Humic substances
IEX – Ion exchange
ILCA – Inline coagulation
IR – Irreversible
LC-OCD – Liquid chromatography-organic carbon detection
LORIV – Lowland river (River Tamar)
LMW – Low molecular weight
LMW-N – Low molecular weight-neutrals
NOM – Natural organic matter
NTU – Nephelometric turbidity unit
PACl – poly-aluminium chloride
SIX – Suspended Ion exchange
TMP – Trans-membrane pressure
UPRES – Upland reservoir (Burrator Reservoir)
UPRIV – Upland river (River Tavy)
UVT – Filtered UV transmittance at 254nm
WTWs – Water treatment works

Abstract

The influence of pre-treatment on the suppression of irreversible (IR) fouling of ceramic membranes challenged with three UK surface waters has been studied at pilot scale. An initial scoping study compared the efficacy of suspended ion exchange (SIX) and clarification (coagulation followed by sludge blanket clarification) individually and in combination. Direct membrane filtration following in-line coagulation (ILCA) was also investigated with and without SIX. The impact on the various organic fractions, specifically high molecular weight (HMW) biopolymers (BPs) and humic substances (HSs), and lower molecular weight (LMW) building blocks (BBs) and neutrals, was studied using liquid chromatography – organic carbon detection (LC-OCD).

Results revealed SIX and coagulation to preferentially remove the LMW and HMW organic fractions respectively. Residual HMW organic matter (primarily BPs) following SIX pre-treatment were retained by the membrane which led to rapid irreversible fouling. Coagulation pre-treatment provided stable membrane operation and the residual LMW organics were not significantly retained by the membrane. Combining clarification and SIX resulted in

significantly increased removal of organics and lower membrane fouling rates. Tests performed using SIX and ILCA revealed comparably high dissolved organic carbon (DOC) removal to SIX with clarification. However, unlike the case for clarification with SIX, the addition of SIX to optimised ILCA dosing offered no additional suppression of membrane fouling compared to ILCA alone. Optimised ILCA pretreatment led to very low IR fouling rates of $<0.3\text{kPa/day}$ trans-membrane pressure, despite highly challenging operating conditions of elevated fluxes ($185\text{ L m}^{-2}\text{ h}^{-1}$) and highly variable feedwater dissolved organic carbon concentrations.

Keywords: Ceramic membrane; pretreatment; ion exchange; coagulation; organic fouling

1 Introduction

Historically, the conventional processes of coagulation, clarification and granular media filtration have been used for removing suspended material and natural organic matter (NOM) from surface water prior to disinfection (Bond *et al.*, 2011). Membrane filtration offers some key advantages over such processes, including higher removal efficiency, compactness, robustness against fluctuating feed water quality, and the provision of an absolute barrier against suspended particles and pathogens such as *Cryptosporidium* (Huang *et al.* 2009; Vreeburg *et al.* 2008). Membranes have thus been increasingly applied for water treatment (Huang *et al.*, 2012), with polymeric materials being most commonly used.

Recently there has been increased interest in ceramic membranes for potable and industrial water treatment applications due to their greater operational lifetime, solids loading capacity, sustainable flux rates (from reduced organic fouling), mechanical robustness and resistance to aggressive cleaning protocols (Hofs *et al.*, 2011; Lee & Kim, 2014). These and other technological benefits have meant that the higher capital costs associated with ceramic membranes can be significantly offset by lower operating costs over the life of the installation, making ceramic membranes potentially economically competitive on a whole life cost basis (Freeman and Shorney-Darby, 2011; Meyn *et al.* 2012; Shang *et al.* 2015).

Membrane fouling nonetheless remains a major obstacle to the application of membranes *per se*. Much research has been focussed on understanding fouling mechanisms and identifying pretreatment capable of removing highly-fouling compounds (Huang *et al.* 2009). Such research has generally revealed that the high molecular weight (HMW) biopolymer (BP) fraction of NOM is primarily responsible for irreversible (IR) membrane fouling, i.e. demanding intensive chemical cleaning in place (Fan *et al.* 2008; Tian *et al.* 2013; Kimura *et al.* 2014). Pretreatment methods which substantially remove the BP fraction have been shown to provide stable membrane operation, with coagulation being the most consistently successful method (Fabris *et al.* 2007; Huang *et al.* 2012; Humbert *et al.* 2007). Coagulation preferentially flocculates the HMW organics, including BPs. The resulting solids are then either removed by clarification (e.g. sludge blanket clarification or dissolved air flotation) or, where “in-line” coagulation with direct filtration is used, by the backwash cycle of the membrane process.

Adsorption processes (anion exchange and activated carbon) have been shown to improve treated water quality through the removal of low molecular weight (LMW) organics but, since they remove only small amounts of the HMW fraction, in most cases fouling reduction has been shown to be minimal (Humbert *et al.* 2007; Fabris *et al.* 2007; Fan *et al.* 2008; Huang *et al.* 2012). Against this, some studies have revealed LMW organics to cause or contribute to fouling through synergistic action with the higher MW fraction (Gray *et al.* 2011; Subhi *et al.*

2012; Lai *et al.* 2015). Such differences in findings highlight the complexity of interactions between the organic constituents and the membrane material and fouling layer.

Since coagulation and adsorption, and specifically ion exchange processes (IEX), have been shown to preferentially remove the high and low MW fractions of the NOM respectively, it may be surmised that their use in combination may both improve permeate water quality and suppress membrane fouling. However, previous studies of the use of combined IEX and coagulation upstream of membrane filtration have not unequivocally demonstrated membrane fouling benefits: reported fouling rates have been similar to those possible with coagulation alone (Humbert *et al.* 2007; Fan *et al.* 2008).

The current study aimed to evaluate suppression of irreversible fouling of ceramic membranes associated with pretreatment by suspended ion exchange (SIX), coagulation or a combination of both. The analysis proceeded through quantification of the organic fractions removed by both pretreatment and the membrane itself, and examined the resulting impact on membrane fouling. Experiments were conducted for three UK surface waters at large pilot scale over a 26 month period under conditions appropriate for potable water production. The irreversible fouling rate was assessed from trans-membrane pressure (TMP) transients generated under operating conditions (including physical and chemically enhanced backwashing) pertaining to those applied at full-scale.

2 Methods and materials

2.1 Raw waters

Three raw waters were tested, either individually or as a blend:

- An upland reservoir (UPRES) of low turbidity and low-to-moderate dissolved organic carbon (DOC) (Burrator Reservoir).
- A soft, upland river (UPRIV), prone to rapid changes in quality following rain, of low-to-high DOC and low-to-moderate turbidity (River Tavy).
- A lowland river (LORIV) prone to rapid changes in quality following rain of low-to-high DOC and turbidity (River Tamar).

2.2 Pilot plant

The 150 m³/day pilot plant (Fig. 1) comprised the SIX® and ceramic membrane filtration (CMF) (CeraMac®) processes and subsequently the SIX®, in-line coagulation (ILCA®) and CMF (PWN Technologies, Netherlands). These processes have been described elsewhere (Galjaard *et al.* 2011; Metcalfe *et al.* 2015) and feature:

- SIX®: an acrylic quaternary amine, gel-type strongly basic anion exchange resin in the chloride form was used throughout the trial (Lewatit S5128, Lanxess, Germany). The resin was generally dosed at 18 mL/L with a contact time of 30 min, with dosing conditions informed by preliminary bench-scale tests. Tests were also performed with lower or zero resin doses (i.e. with ILCA® only) as appropriate. The resin was in continuous use over the 2 years of the trial.
- A Lamella separator was used for separating the resin from the treated flow and the resin regenerated with 30 g/L NaCl. SIX-treated water samples were collected directly after resin separation.
- ILCA® using polyaluminium chloride (WAC®, Water Treatment Solutions, UK, 0.53-4.23 mg/L as Al) was used alone or following SIX pretreatment. Water was pH-corrected with NaOH or HCl (Brenntag, U.K), injected with coagulant, and mixed by a static mixer

and flocculated for 2.4-3.9 minutes prior to CMF (direct filtration). The coagulation pH was circa 6.4 for all tests.

- CMF was carried out using a vertically mounted 25 m² ceramic membrane element (Metawater, Japan, nominal pore size 0.1 mm) operating by dead end filtration.

2.3 Full scale WTW process

The water treatment works ((WTWs) Crownhill WTWs, South West Water (SWW), Plymouth, U.K) treated water using optimised coagulation with aluminium sulphate dosed at 3.39-6.36mg/L as Al (Kemira, U.K) and Magnafloc LT25 (BASF, U.K) anionic polyelectrolyte at 0.1-0.2 mg/L. Powdered activated carbon (PAC, Aquasorb BP2, Jacobi, U.K) was dosed at 2-3 mg/L prior to coagulation. The WTWs coagulant dose was optimised through jar testing and works operation. Clarified water following flash mixing, flocculation and solids-liquid separation (by sludge blanket clarification) was supplied to the pilot plant for some of the tests.

2.4 Pretreatment

The pilot plant was fed with either raw or, during the clarification or clarification + SIX campaigns, clarified water (Fig. 1).

Tested pretreatment options for CMF comprised:

- 1) Clarification only, or clarification followed by SIX: Raw water was treated by the full scale clarification process and the water piped from the clarifier outlets to the pilot plant feed tank. During clarification-only tests the SIX process was taken off-line and all resin removed. Additional tests were performed where the clarified water was further treated by SIX within the pilot plant, prior to CMF.
- 2) SIX followed by ILCA, or ILCA alone: ILCA was employed following the SIX process, or when ILCA was used alone the SIX process was taken off line and the resin removed. For most tests the coagulant dose and pH correction was manually controlled, which led to periods of sub-optimal operation due to the rapidly changing raw water sources. The pH and coagulant dosing systems were automated in March 2015.

2.5 Ceramic microfiltration (CMF)

The CMF membrane flux was varied between 109 and 250 LMH (L/(m²h)). A new membrane was installed at the start of the trials in March 2013, replaced by a new element after 2 years in operation (March 2015) prior to extended tests with automated coagulant and pH control.

Backwashing and chemically enhanced backwashing (BW and CEB) were performed following a given filtration load, measured in litres of feed solution filtered per m² membrane surface (L/m²). BW used 75 L permeate pressurised to 5 bar and passed through the membrane in a reverse flow direction. CEBs with NaOCl (100 mg/L), NaOH (pH 12) or acid + H₂O₂ (pH 2.5, 100 mg/L H₂O₂) were performed at 2 bar pressure. The membrane was soaked in the reagent for 10 minutes prior to a standard BW. Process conditions for the tests are reported in Table 1.

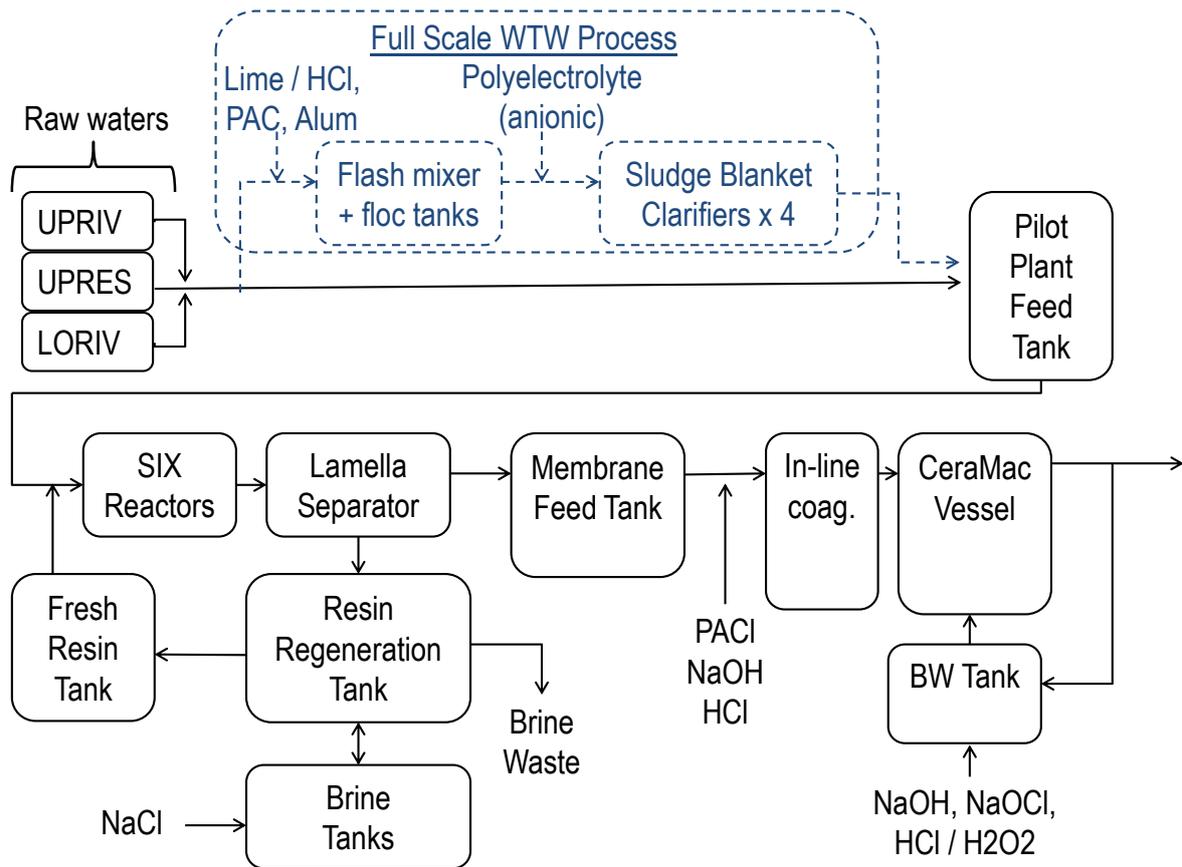


Figure 1: Process flow diagram

After each discrete test cleaning in place (CIP) was performed by circulating chemical solution through the membrane for an extended period of time (3 – 24 hours). Generally an overnight CIP using NaOH (pH 12-12.5) was performed, followed by a CIP with either NaOCl (100-500 mg/L) or HCl/H₂O₂ (pH 2.7, 100 mg/L H₂O₂) if required to increase the specific flux to >300 LMH/bar at 10°C prior to starting a new test. This low specific flux target (300 LMH/bar) did not constitute an optimised recovery of permeability but provided an attainable standard starting permeability for each test, despite the significant fouling which occurred in some of the scoping trials. Following the final extended tests, based on a virgin membrane with automated coagulant and pH control, an intensive CIP was performed which fully recovered membrane permeability (to ~1250 LMH/bar at 10°C). The conditions for the intensive CIP were 30 minutes at pH 2.7 with 100mg/l H₂O₂, followed by 1 hour at pH 12 (NaOH), followed by 2% NaOCl for 3 days.

Table 1: Test process conditions. * Backwash (BW) and chemically enhanced BW (CEB) sequences have been described as, for example, (5 BW, NaOCl CEB) x 4, 5 BW, HCl/H₂O₂ CEB – this would denote 5 BW followed by a NaOCl CEB with this sequence being repeated 4 times. Following this 5 BW and then an HCl/H₂O₂ CEB would occur and then the sequence would be repeated.

Test No.	Raw water source	Pretreatment	Flux (LMH)	Filtration load prior to BW / CEB (L/m ²)	BW number prior to CEB / CEB sequence and chemicals *	Water recovery (%)
1	UPRES	SIX	150	75 / 450	(5 BW, NaOCl CEB) x 4, 5 BW, HCl/H ₂ O ₂ CEB	95.3
2	UPRES	SIX	150	75 / 450	(5 BW, NaOCl CEB) x 4, 5 BW, HCl/H ₂ O ₂ CEB	95.3
3	UPRIV / UPRES	Clarification only	150	75 / 450	(5 BW, NaOCl CEB) x 4, 5 BW, HCl/H ₂ O ₂ CEB	95.3
4	LORIV/UPRIV/UPRES	Clarification / SIX	150	75 / 450	(5 BW, NaOCl CEB) x 4, 5 BW, HCl/H ₂ O ₂ CEB	95.3
5	UPRES	SIX + ILCA	185	100 / 1800	17BW, NaOH CEB, HCl/H ₂ O ₂ CEB	96.8
6	UPRES	SIX + ILCA	185	100 / 1800	17BW, NaOH CEB, HCl/H ₂ O ₂ CEB	96.8
6a	UPRES	SIX + ILCA	185	100 / 1800	17BW, NaOH CEB, HCl/H ₂ O ₂ CEB	96.8
7	UPRIV	SIX + ILCA	185	100 / 1800	17BW, NaOH CEB, HCl/H ₂ O ₂ CEB	96.8
8	LORIV	SIX + ILCA	185	100 / 1800	17BW, NaOH CEB, HCl/H ₂ O ₂ CEB	96.8
9	UPRES	ILCA Only	109	150 / 1800	(17BW, NaOH) x 3, 17 BW, NaOH CEB, HCl/H ₂ O ₂ CEB	97.8
10	UPRES	9ml/l SIX + ILCA	109	150 / 1800	(17BW, NaOH) x 3, 17 BW, NaOH CEB, HCl/H ₂ O ₂ CEB	97.8
11	UPRES	18ml/l SIX + ILCA	109	150 / 1800	(17BW, NaOH) x 3, 17 BW, NaOH CEB, HCl/H ₂ O ₂ CEB	97.8
12	UPRES	ILCA Only	109	150 / 1800	(17BW, NaOH) x 3, 17 BW, NaOH CEB, HCl/H ₂ O ₂ CEB	97.8
13	UPRES	18ml/l SIX Only	109	150 / 1800	(17BW, NaOH) x 3, 17 BW, NaOH CEB, HCl/H ₂ O ₂ CEB	97.8

2.6 Coagulant dose optimisation

20 jar tests were performed on a wide range of raw or SIX treated waters from each of the 3 sources to establish a means of optimising the coagulant dose for minimising membrane fouling. The data from these jar tests revealed an excellent correlation ($R^2 = 0.96$) between the optimum coagulant dose and feed water filtered UV transmittance (UVT), the optimum coagulant dose determined as being the minimum coagulant dose in mg/L as Al providing maximum achievable UVT removal (Fig. 2):

$$C_{opt} = 8.88 - 0.0911 * UVT_{feed} \quad (1)$$

Applications of this dose resulted in negligible membrane adsorption of organics according to liquid chromatography-organic carbon detection (LC-OCD) analysis and on-site membrane

feed and permeate UVT measurements (Section 2.9). Most of the tests were conducted with daily manual adjustments to the coagulant dosing, based on the feed water UVT according to Equation 1, along with pH correction. This resulted in occasional sub-optimal coagulation conditions when rapid changes in raw water quality took place, subsequently leading to increased membrane fouling.

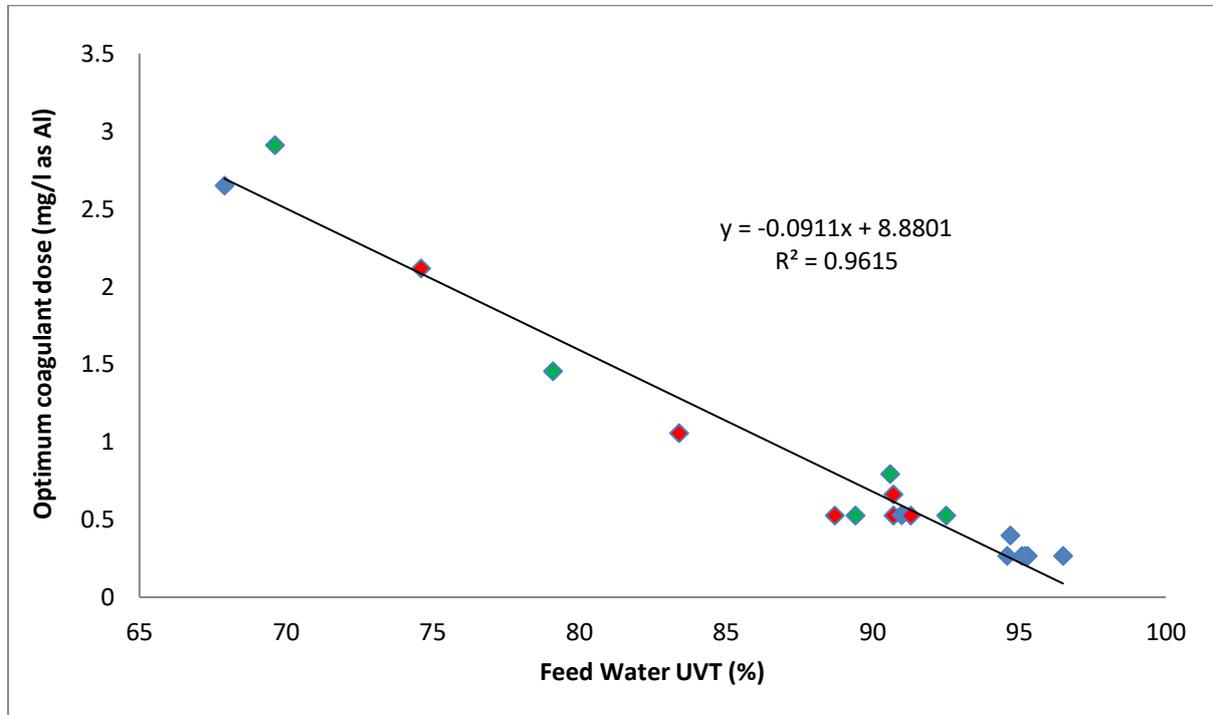


Figure 2: Feed water UVT vs. optimum coagulant dose jar tests results. Red = UPRES, Blue = UPRIV, Green = LORIV

2.7 Automated coagulant dosing and pH correction

In March 2015 automated coagulant and pH control was installed. A Spectro::lyser (S::CAN, Austria) was used to measure the solids-compensated UVT of the water in the membrane feed tank, for which measured values were similar to manual filtered UVT samples. A controller used the on-line UVT_{feed} value with Equation 1 to provide the required coagulant dose. pH control was automated to control to a set-point of 6.4 via a negative feedback loop/controller.

2.8 Membrane fouling rate

Irreversible membrane fouling (i.e. not recovered by the routine BW and CEB) was measured so as to determine the required CIP frequency during full-scale operation. Individual filtration runs were generally based on a total filtration volume of 625 m^3 , the filtration run time then varying with flux. An extended run was performed with a virgin membrane, with automated coagulant and pH control. TMP and temperature data (logged at one minute intervals) from each test was analysed. TMP data following CEBs were selected and the values corrected to 10°C . The fouling rate in kPa/day was determined by linear regression, with the corresponding R^2 value (Table 2).

2.9 Sample analysis

Turbidity was analysed using a Hach Lange 2100AN Turbidimeter (Hach Lange, Germany). 0.45 µm-filtered UVT was measured using a Hach DR6000 spectrophotometer (Hach Lange, Germany). DOC was characterised by LC-OCD analysis at Het Water Laboratorium (Netherlands), which quantified concentrations of total DOC and the discrete MW fractions of biopolymers (BP)s, humic substances (HSs), building blocks (BBs), LMW neutrals and LMW acids (Huber *et al.*, 2011). LMW acids were not detected by LC-OCD analysis since organics within this band (50 minute elution time, equating to <350 Daltons) absorbed UV and were hence classified by the analysis/software as LMW-HS. Turbidity and LC-OCD results for each test are reported in Table 2.

Adsorption of DOC and the fractions thereof onto the membrane was assessed through concentration difference between the feed and permeate. Since UVT was found to correlate reasonably well ($R^2 = 0.63-0.92$) with DOC concentration, all source water organic fractions being strongly UV absorbing, UVT data were used to derive organic carbon values in those tests where no LC-OCD data was collected.

3 Results

3.1 Scoping studies, SIX, clarification or clarification + SIX (Tests 1-4)

A number of initial tests (Tests 1-4, Tables 1-2) were performed to establish the influence of pre-treatment on irreversible fouling. These tests compared SIX treatment alone, at pilot scale, with a full scale clarification process and a combined process where the clarified water was further treated by SIX, at 150 LMH. LC-OCD analysis illustrated clear differences in the organic fractions removed by SIX, clarification and a combined process of clarification followed by SIX (Fig. 3a-d). SIX pre-treatment preferentially removed LMW organics whilst the removal of HMW fractions by SIX was low, especially for the highest MW BP fraction (25-29%). Conversely, coagulation pretreatment preferentially removed the highest MW fractions (particularly BP) whilst the LMW organics fractions were removed to a lesser extent than possible with SIX (especially lower MW HS and BB fractions) (Fig. 3c). These data are consistent with the findings of previous studies showing some LMW organics to be recalcitrant to coagulation and the removal of high MW BPs by IEX to be marginal (Fabris *et al.* 2007; Huber *et al.* 2011; Humbert *et al.* 2007; Mergen *et al.* 2009; Myat *et al.* 2012). Subsequently, clarification followed by SIX was found to yield very low residual organics concentrations (Fig. 3d), in keeping with previously reported findings (Fearing *et al.* 2004; Jarvis *et al.* 2008; Humbert *et al.* 2007; Fan *et al.* 2008; Singer and Bilyk, 2002).

In Tests 1 and 2, following SIX pre-treatment, the majority (58-80%) of the residual HMW organics were retained by the membrane (Figs. 3a and b). At the lowest DOC levels (Table 2, Test 1 and Fig. 3a), a low-to-moderate fouling rate of 3.6 kPa/day was obtained at 150 LMH. When DOC increased for the UPRES source and an organic compositional change occurred resulting in a near three-fold increase in the concentration of the HMW BP fraction, a very high membrane fouling rate (48 kPa/day) was recorded (Table 2, Test 2 and Fig. 3b). Further tests with SIX pre-treatment were performed on higher DOC reservoir and river water sources (UPRES, UPRIV and LORIV) at 100 and 150 LMH. Despite reductions in filtration volume prior to backwash and CEB, all of these tests yielded very high fouling rates (82-863 kPa/day).

Table 2: Test membrane irreversible fouling rates and water quality data.

Test No.	Irreversible (IR) fouling Rate (kPa/day)	IR fouling R ² value	Filtration time (days)	Water Source	Turbidity (NTU)	LC-OCD data							
						TOC (µg/l)	DOC (µg/l)	CDOC (µg/l)	BP (µg/l)	HS (µg/l)	BB (µg/l)	LMW-N (µg/l)	
1	3.6	0.86	5.88	Raw	0.58	1863	1751	1668	143	1003	269	254	
				SIX	0.75	727	677	565	107	243	82	134	
				CMF	0.10	552	540	454	45	214	75	120	
2	47.7	0.90	1.74 [S]	Raw	2.10	2529	2372	2372	402	1276	356	338	
				SIX	1.80	1080	1023	1020	287	409	114	210	
				CMF	<0.1	642	603	547	55	223	101	168	
3	6.2	0.97	5.33	Raw	2.00	3974	3801	3538	232	2429	458	419	
				Clarified	0.99	994	937	924	51	351	291	231	
				CMF	<0.1	973	936	826	35	342	262	187	
4	0.7	0.87	5.47	Raw	1.70	1863	1786	1677	272	855	303	248	
				Clarified	0.33	843	738	659	47	359	87	167	
				C+S	0.42	492	459	377	43	52	170	113	
				CMF	0.29	473	441	358	37	43	146	132	
5	6.1	0.99	5.91	Raw	0.57	2288	2273	2297	124	1606	284	282	
				SIX	0.60	1516	1519	1440	113	935	154	238	
				ILCA	0.86	401	303	371	21	41	144	165	
				CMF	<0.1	364	311	346	32	46	121	147	
6	5.4	0.96	3.92 *WQ	Raw	0.69	2060	1917	1976	188	1194	303	290	
				SIX	0.8	922	883	773	161	323	116	172	
				ILCA	1.4	563	425	356	30	47	161	119	
				CMF	0.1	479	466	356	41	51	147	116	
6a	571.4	1 (2 data points)	0.27 *S	Raw	NS	5188	5028	5248	365	3733	572	578	
				SIX	NS	2742	2667	2810	321	1774	312	402	
				ILCA	NS	2809	2560	2816	378	1679	297	460	
				CMF	NS	951	906	835	71	408	148	208	
7	10.6	0.86	3.94*W Q	Raw	1.20	1524	1402	1484	274	740	239	232	
				SIX	1.40	747	642	661	220	223	91	127	
				ILCA	1.30	374	334	356	52	46	137	121	
				CMF	<0.1	393	331	316	44	45	116	111	
8	3.8	0.95	5.37	Raw	5.30	4150	4011	4279	443	2683	605	548	
				SIX	5.50	1665	1555	1572	379	651	212	330	
				ILCA	5.90	1034	850	803	47	149	323	282	
				CMF	0.13	966	855	777	40	148	313	276	
9	2.1	0.83	5.6	Raw	1.10	2890	2635	2623	181	1861	292	289	
				ILCA	1.00	771	741	600	40	220	176	164	
				CMF	0.31	707	645	520	27	161	182	150	
10	1.7	0.94	5.6	Raw	0.96	2787	2630	2592	184	1828	308	272	
				SIX	1.00	2283	2117	2022	155	1370	274	223	
				ILCA	1.60	599	471	433	35	139	126	133	
				CMF	<0.1	503	485	392	31	114	124	123	
11	1.7	0.93	5.6	Raw	0.74	2334	2245	2136	132	1452	297	254	
				SIX	0.78	1812	1788	1660	121	1083	228	228	
				ILCA	1.40	523	512	441	30	156	110	143	
				CMF	0.17	471	435	359	25	114	96	123	
12	1.4	0.93	5.6	Raw	0.65	2187	2129	2040	119	1404	258	259	
				ILCA	0.72	717	688	609	29	229	163	187	
				CMF	0.11	550	512	458	26	139	147	147	
13	Very high fouling - max TMP caused shutdown			Raw	0.87		<i>2360</i>						
				SIX	1.05		<i>1100</i>						
				CMF	0.14		<i>850</i>						

[S] run stopped due to shut-down; WQ water quality change, NS not sampled, NTU nephelometric turbidity units, TOC total organic carbon, DOC dissolved organic carbon, CDOC chromatographically detectable DOC, BP biopolymers, HS humic substances, BB building blocks, LMW-N low molecular weight neutral. Italicised DOC values (Test 13) were inferred from UVT data.

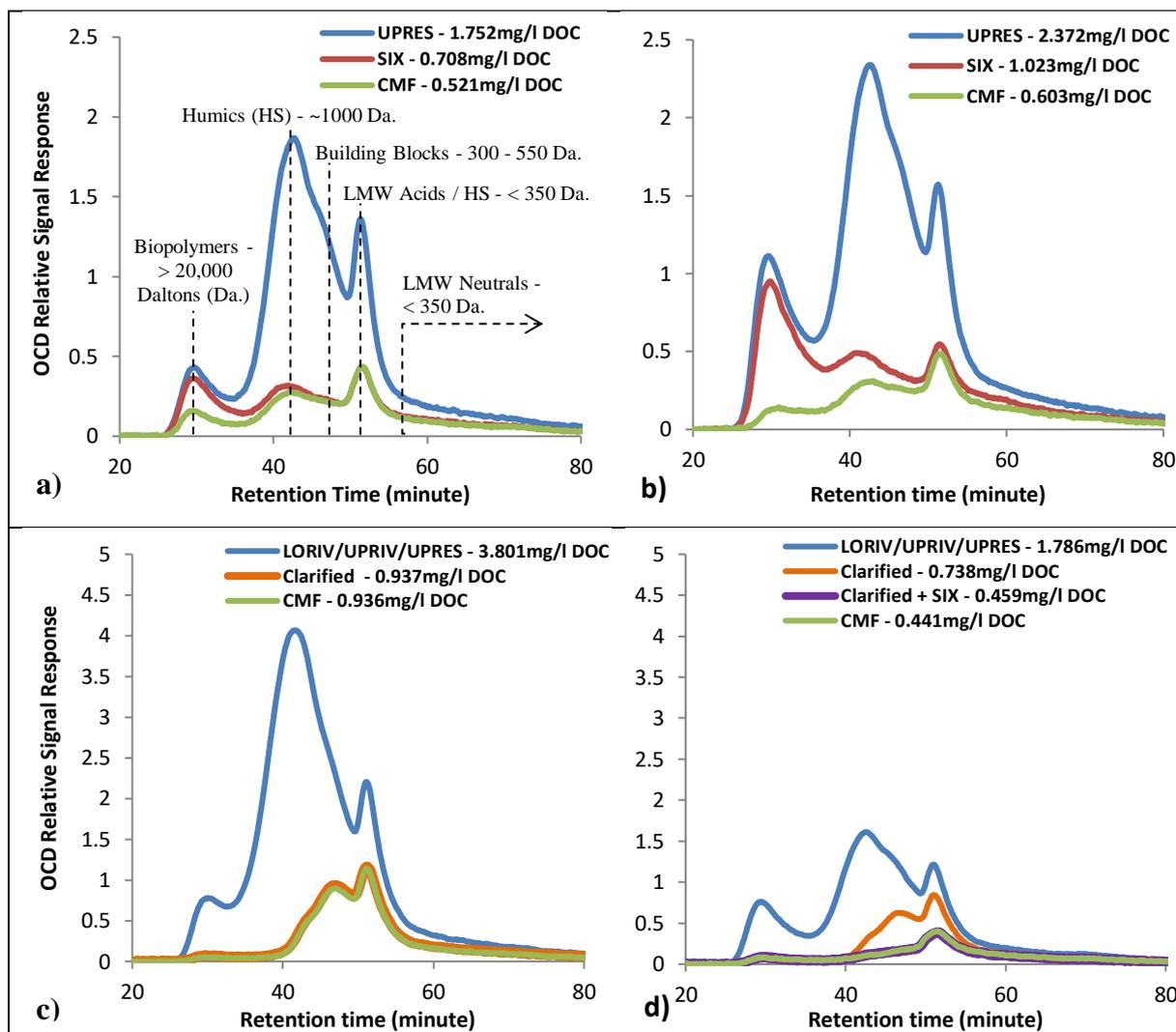


Figure 3: LC-OCD chromatograms a) Test 1: SIX only with low DOC UPRES raw water (fraction peak assignment shown for reference) b) Test 2 - SIX only with moderate DOC UPRES raw water c) Test 3 - Clarification only with moderate DOC blend of raw waters d) Test 4 - Clarification followed by SIX with low DOC blend of raw waters

For the lowest raw water organic concentrations (e.g. Table 2 – Test 1), fouling was effectively abated by the routine physical and chemical cleaning. The increased fouling recorded in Test 2 was associated with greater retention of HMW organics than for Test 1 (80% vs. 58% for BP), corroborating previous work on the deleterious impact of increasing BP mass loads on irreversible fouling (Fabris *et al.* 2007; Fan *et al.* 2008; Zheng *et al.* 2010; Huang *et al.* 2012; Myat *et al.* 2013; Tian *et al.* 2013; Shang *et al.* 2015; Kimura *et al.* 2014). Whilst ion exchange can significantly reduce overall DOC concentrations, previous research indicates that it does not lead to significant reductions in membrane fouling due to the limited removal of the HMW organics (Humbert *et al.* 2007; Fabris *et al.* 2007; Fan *et al.* 2008; Huang *et al.* 2012). Analysis of the adsorbed organics revealed that a good correlation was obtained between membrane fouling and adsorption of (a) overall DOC ($R^2 = 0.89$), (b) BPs ($R^2 = 0.95$) and (c) HSs ($R^2 = 0.98$), whereas the correlations with LMW fractions were significantly weaker ($R^2 = 0.14-0.33$).

Test 3, using fully clarified water from the full-scale plant (Fig. 3c), was performed under the same membrane operating conditions as Tests 1 and 2 (using SIX pretreatment) but whilst

challenged with more highly fouling river water. Results indicated the removal of the HMW organics by clarification to provide a low-to-moderate fouling rate of 6.2 kPa/day (Table 2, Test 3). Whilst pre-clarification removed the LMW organics to a lesser extent than possible with SIX pretreatment (Fig. 3c), these organics were not retained by the membrane whereas the HMW organics were largely retained following SIX pretreatment. Following coagulation pretreatment no adsorption was detected. In Test 4 the clarified water was further treated by SIX (under the same operational conditions as Tests 1-3) to establish if removing additional LMW organics from the clarified water, would further suppress membrane fouling. The lower fouling rate of 0.9 kPa/day (Table 2, Test 4) was associated with negligible organics adsorption (0.018 mg/l DOC. Fig. 3d), although the apparent adsorption recorded was higher than in Test 3.

Further tests comparing clarification only with clarification + SIX using the same operational conditions as Tests 1-4 at a higher applied flux of 175 LMH, indicated a moderate fouling rate of 11 kPa/day for clarification alone and a much lower rate of 2.4 kPa/day for clarification + SIX, despite a significantly higher raw water DOC. As with Test 3 and 4, DOC adsorption was negligible (-0.022 vs. 0.037 mg/L for clarification vs. clarification + SIX). This suggests when fouling was low, the fouling was related to factors other than organic adsorption, such as floc characteristics (Jarvis *et al.*, 2008). It is also possible that the ion exchange material adsorbs any residual anionic polyelectrolyte from the clarification process, which would otherwise affect membrane fouling (Wang *et al.* 2011; Yu *et al.* 2013). Two further tests with clarification + SIX pre-treatment performed with the same operational conditions as Tests 1-4, but at higher fluxes of 200 and 250 LMH, yielded fouling rates of 5.5 and 197 kPa/day respectively, suggesting that the highest value exceeded the critical flux.

3.2 SIX/ILCA pre-treatment with enhanced CEB, manual control (Tests 5-8)

Jar tests confirmed contact times below two minutes to be sufficient for flocculating residual HMW organics following SIX treatment of the three raw waters, corroborating work performed by Meyn *et al.* (2012) on ILCA upstream of membrane filtration. Subsequent pilot trials were conducted with SIX followed by pH correction and 2-4 minutes ILCA at relatively low coagulant doses (50-90% less than that employed for the full-scale clarification process), permitted by the removal of a large amount of DOC by SIX (Huang *et al.* 2012; Humbert *et al.* 2007; Jarvis *et al.* 2008). A flux of 185 LMH was employed for these trials, with a NaOH-based CEB: supplementary trials revealed NaOH to be more effective than NaOCl for CEB.

Analysis of the organic content of the three raw water types used for these tests revealed it to be dominated by aromatic organics (high SUVA) with all organic fractions, including BPs, adsorbing UV strongly. The UVT could thus be used as a relatively accurate surrogate measure of DOC, confirmed by the strong correlation ($R^2 = 0.92$) between UVT and the LC-OCD-determined DOC. The algorithm from jar test data (Equation 1) was used to manually set the coagulant dose based on the feed water UVT. This generally led to negligible adsorption of organic matter on the membrane and corresponding low fouling rates. High observed membrane fouling, following rapid changes in raw water quality and subsequent sub-optimal coagulation conditions, was accompanied by measurable organics adsorption.

Results from Tests 5-8 (Table 2) indicated that for optimised, manually-controlled coagulant dosing and pH adjustment, membrane fouling was low to moderate (3.8–11 kPa/day) at the high flux of 185 LMH. The DOC removal by the SIX ILCA CMF process was similar to that attained in the scoping trials based on full-scale clarification followed by SIX, i.e. 74-88%

relative to the raw water. As with the scoping trials with clarification alone and clarification + SIX (Table 1-2, Tests 3 and 4), changes in fouling rate could not be related to the LC-OCD data as membrane organic adsorption was negligible when coagulation conditions were optimised. Further tests with the same operation conditions as Tests 5-8, but at a lower flux of 112 LMH, yielded low fouling rates of 1.0-1.7/day. Instances of non-optimum coagulation conditions, due to rapid changes in raw water quality, always led to rapid fouling which was reflected by measurable organics adsorption onto the membrane (Table 2, Test 6a). These results demonstrate the critical nature of optimal pre-coagulation in direct membrane filtration.

3.3 IEX dose trials in combination with ILCA (Tests 9-13)

Tests conducted at 109 LMH on standard, reduced or zero SIX pretreatment yielded low fouling rates of 1.4-2.1 kPa/day at doses of 0 to 18 mL/L SIX (Table 1-2, Tests 9-12), with optimised coagulant dosing. Coagulant demand increased with decreasing SIX dose, but under these optimised coagulation conditions there was no correlation between fouling and SIX dose with the fouling rate remaining stable at 0, 9 or 18g/l resin doses. However, when the ILCA was taken off line and SIX pretreatment alone was employed, rapid fouling was observed (Table 1-2, Test 13). This confirmed the relative importance to membrane fouling of enmeshment of HMW BP organics within coagulant flocs, compared to the additional removal of LMW fractions by SIX. In contrast to the full clarification or clarification and SIX tests (Section 3.1), the addition of SIX to ILCA pretreatment yielded no appreciable fouling suppression. This was possibly due to either the change in CEB reagent (from NaOCl to NaOH), the difference in floc morphology, or, most likely, the nature of the direct filtration process where a cake is formed on the membrane surface. Previous research has suggested that this cake layer may reduce irreversible membrane fouling by either adsorbing some unflocculated organics, and therefore preventing their adsorption to the membrane (Dong *et al.* 2007), or by rejecting fine flocs or colloids which would otherwise plug the membrane pores (Guigui *et al.* 2002).

3.4 SIX + ILCA pretreatment with automated coagulant and pH control

A virgin membrane and an automated pH and coagulant control system was installed prior to an extended fouling rate test (21 days) at a high flux of 185 LMH on the UPRIV source water (Fig. 4). Low fouling rates were sustained despite rapid changes in raw water quality and DOC concentration associated with heavy rainfall events. Cessation of coagulation during this trial (Day 9) led to a rapid increase in the TMP, which was ameliorated on reinstating the coagulant dose, albeit at a slightly higher baseline TMP. Taking the SIX dosing off line on Day 13 did not lead to an increased fouling rate, a stable TMP being maintained. Overall a very low fouling rate of 0.3 kPa/day was sustained despite coagulant dose upset, regardless of the resin dose, whilst operating at high flux on variable quality raw water. The results further corroborate coagulation as being the most important pre-treatment for suppressing fouling and confirmed that very low fouling rates were possible at high fluxes when operating with automated coagulant dosing and pH control. The high permeability of the virgin membrane was sustained throughout the trial. Further tests of around 2 months' duration yielded an overall fouling rate 0.24 kPa/day, with a subsequent CIP returning the membrane permeability back to that of the virgin material.

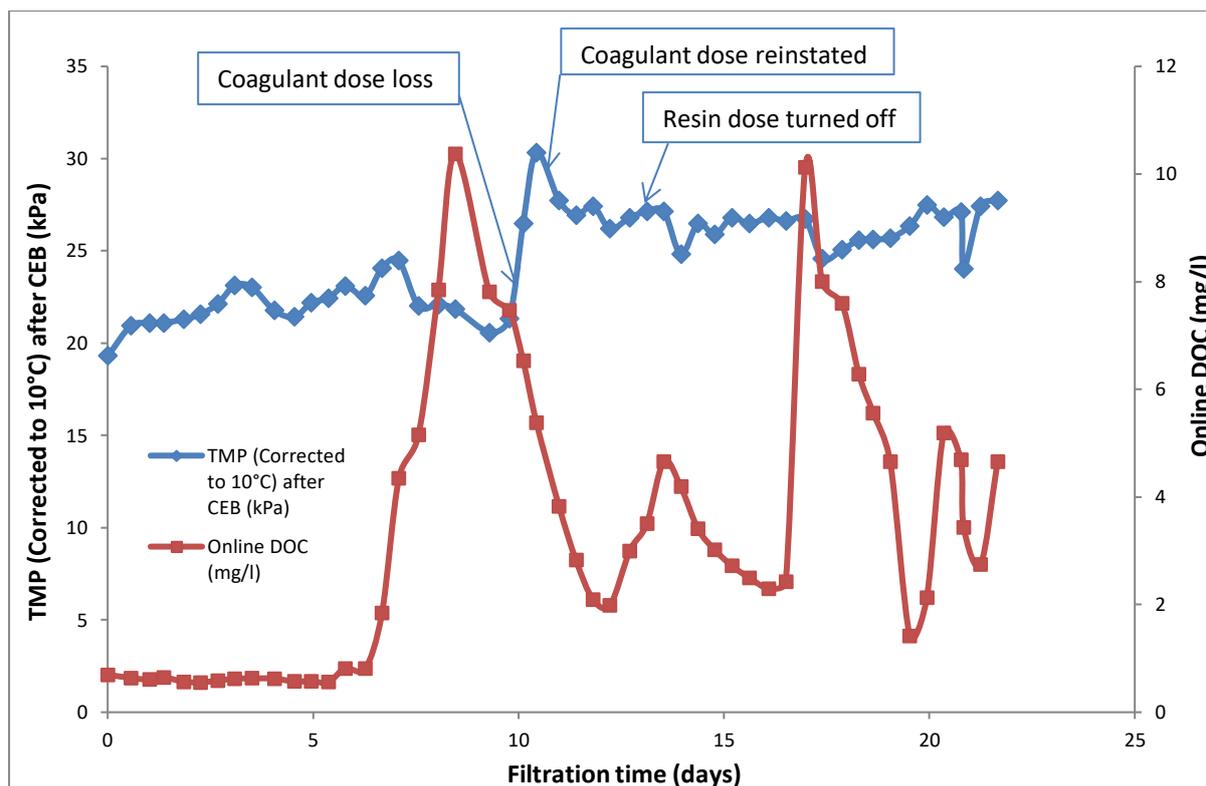


Figure 4: Extended operation during spate conditions (UPRIV) with automated coagulant and pH control

4 Conclusions

A pilot-scale study of the efficacy of pretreatment for ceramic membrane filtration of surface waters as applied to potable water supply has revealed:

- Suspended ion exchange (SIX) resin removed predominantly low molecular weight (LMW) organic matter whereas coagulation removed the high molecular weight (HMW) fractions. The combination of full clarification (coagulation with floc blanket) and SIX led to substantial removal of all organic fractions to leave a low residual dissolved organic carbon (DOC) concentration.
- The HMW organics (present following SIX), including biopolymers, were retained by the membrane and caused rapid fouling, whereas LMW organics (present following coagulation alone) were not retained when coagulation was optimised.
- When using optimised in-line coagulation (ILCA) with direct membrane filtration, as opposed to full clarification pretreatment, additional pretreatment with SIX provided no measurable benefit with regards to membrane fouling suppression.
- Sub-optimal coagulation conditions, such as under-dosing of coagulant or inappropriate pH adjustment, resulted in rapid fouling.
- SIX ILCA pretreatment provided similar DOC removal to clarification followed by SIX pretreatment whilst providing a more compact, efficient and flexible process than clarification and SIX pretreatment.
- A fully optimised system with automated pH and coagulant control allowed operation at an elevated flux of 185 LMH on a water source of highly variable organic concentration with overall membrane fouling rates below 0.3 kPa/day over a two-month test period.

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