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Preparation of multifunctional hollow fiber nanofiltration membranes by dynamic assembly of weak polyelectrolyte multilayers

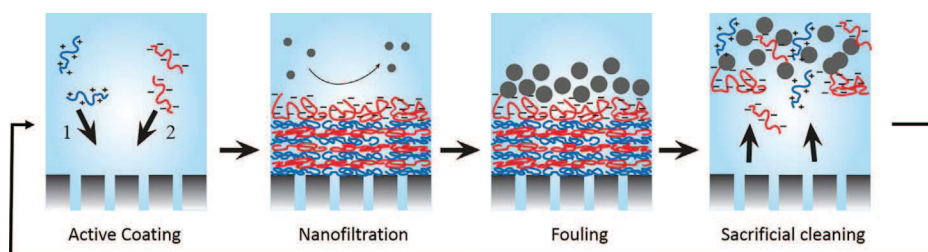
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GRAPHICAL ABSTRACT



ABSTRACT

Keywords:

Layer by layer assembly
Polyelectrolyte multilayers
Weak polyelectrolytes
Dynamic coating
Easy cleaning
Antifouling

In this work, we investigate the effect of preparation conditions for dynamic layer-by-layer (LbL) coating, to prepare multifunctional hollow fiber nanofiltration (NF) membranes. Dynamic coating was performed at constant pressure and at variable cross flow speeds. In this way, polyelectrolyte multilayers (PEMs) were formed of the weak polyelectrolytes poly(acrylic acid) (PAA) and poly(allylamine hydrochloride) (PAH), on a negatively charged polyethersulfone ultra-filtration (UF) support. The resulting membrane performance was investigated and compared to membranes prepared by different methods (dip coating and dead end filtration), and it was found to be comparable. It was shown that PAH/PAA multilayers can be fabricated reproducibly and homogeneously using optimised dynamic LbL deposition conditions on single fiber module (surface area of 6.2 cm²) as well as on a module of 15 fibers (surface area of 67 cm²). Moreover, the approach of dynamic coating could be easily up scaled to coat existing UF modules. The prepared membranes reject the solutes on the basis of size exclusion and Donnan exclusion, retaining small organic solutes and divalent ions. Membranes terminated with PAA exhibit a low fouling tendency compared to membranes terminated with PAH and compared to the UF support membrane. Moreover, if severe membrane fouling would occur after prolonged use, the PEM coating, including any attached foulants can be removed by rinsing with a solution that combines a low pH and a high salinity (pH 3, 3 M). The surface of bovine serum albumin (BSA) fouled membranes were successfully cleaned at least twice, after which a new PEM coating could be re-applied by active coating.

1. Introduction

Growing water crises around the globe and more stringent environmental regulations are increasing the interest of the water

industry to consider new, more energy efficient, treatment processes. During the last two decades, membrane separation processes in general and nanofiltration (NF) in particular have attracted enormous attention as an environmental friendly separation technology for water and

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wastewater treatment [1]. NF allows the effective removal of inorganic and organic pollutants at much lower pressures than those required for reverse osmosis (RO). NF membranes are proposed to be an alternative to RO membranes, because of their higher fluxes, their high retention of multivalent salt ions, and their low operation and maintenance costs [2].

A major challenge for all membrane processes, including NF, is the loss of membrane productivity over time, due to fouling at the membrane interface. Fouling can lead to lower fluxes and therefore higher operating costs, while frequent physical and chemical cleaning is costly and might shorten the membrane life-time [3,4]. Fouling is affected by many parameters, including the membrane surface chemistry, feed composition, operational parameters, and hydrodynamic conditions [4–6]. To deal with membrane fouling, anti-fouling materials and new cleaning methods have become the main focus of research and development within the water industry [7]. In a typical process, fouling is controlled using intermittent backwashes or backflushes, which is only effective for loosely attached foulants on the membrane surface [8,9]. However, impurities such as proteins, humics, and polysaccharides can behave as strong and irreversible foulants [10]. For these foulants, expensive chemical cleaning is required, generally consisting of harsh treatments with acids, alkalis, chlorine etc. [5,8]. Even under these cleaning conditions, complete recovery of membrane performance is not guaranteed. Additionally, there exists a high risk of damage to the membrane due to the frequency and severity of the cleaning. Other strategies to control membrane fouling are mainly focused on surface modification of the membrane to weaken the interactive forces between foulants and membrane, thus inhibiting foulant adsorption. This can be achieved by altering the membrane surface charge, increasing the hydrophilicity or decreasing the surface roughness [11–13].

In the past decade, LbL assembly of polyelectrolytes (PEs) in the form of polyelectrolyte multilayer (PEM's) onto a charged and porous support has proven to be a promising way of membrane surface modification. PEM's can provide membranes with antifouling functionalities [11,12,14–18], as well as act as the separation (skin) layer for NF membranes [19]. Most of the previous studies on PEM based NF membranes, utilized flat sheet membrane supports as reported for example by Bruening et al. [20–25] and Tieke et al. [26,27]. However, the hollow fiber configuration would offer a higher productivity [28,29]. Furthermore, the hollow fiber geometry offers the possibility of backwashing (with or without chemicals) at higher pressures for enhanced foulant removal. To our knowledge only one commercial hollow fiber NF product with the trade name HFW 1000 and a molecular weight cutoff (MWCO) of 1000 Da is available in market. So, it would be very beneficial to develop hollow fiber NF membranes at a lower MWCO, and for that the surface modification of a UF support using LbL assembly of PEMs is a very promising method. This approach was shown to successfully produce hollow fiber NF membranes with precise control over separation performance [30–34]. In these studies, LbL deposition is performed by dip coating (static coating) by immersing the hollow fibers into the solutions of PEs followed by intermittent rinsing steps. An alternative is the use of dynamic coating (or active coating), which involves running the PE solution through the lumen of the hollow fibers while controlling the pressure [35–39] and/or the flux [40]. This dynamic approach can give fast and controlled convective transport of PEs towards the membrane. Recently Menne et al. [40] studied dynamic coating of the strong PEs poly(diallyl-dimethylammonium chloride) (PDADMAC) and poly(styrene sulfonate) (PSS) in a dead end mode. He found that especially the constant flux approach was an effective way to coat the hollow fiber UF membrane support. Coating in the dead end mode quickly increases the filtration resistance due to accumulation of material, while the cross flow mode, where liquid flows over the surface of membrane, limits the accumulation of PEs, and thus provides more precise control over layer deposition.

Apart from the mode of coating, PEs with different sizes, charge densities degrees of swelling and diffusion velocities may have

completely different deposition speeds on the membrane. Now depending upon the used PEs, these multilayers can be erasable by some external stimuli such as a shift in pH [41–43] or by increasing the ionic strength [44,45]. For a weak PE system such as PAH/PAA, where the charge density depends on solution pH, a combination of both of these stimuli (low pH and high ionic strength) can add a trigger to sacrifice (remove) the PEM coating along with attached foulants, as shown previously [34]. This sacrificial approach to cleaning is highly promising as a new approach to help alleviate fouling, but it could only really be used in practice if the coating and recoating steps are both simple, and can be applied on module level. In the current work we investigate exactly this, we study the application of PEM based NF separation layers by dynamic coating, to demonstrate that a simple method is available to coat and recoat membranes, while imbuing it with NF separation properties and a low fouling propensity. This weak polyelectrolyte system of PAH/PAA, holds great promise for membrane applications and this work intends to help to bring this system to maturity.

In this study, we have developed hollow fiber NF membranes with sacrificial layers (PAH/PAA) employing dynamic (cross flow and dead end) coating. The effects of coating parameters (deposition time, applied pressure, coating mode and cross flow speed) on membranes in terms of permeability and NF performance have been investigated in detail and compared with those prepared by a static (dip coating) LbL process. Moreover, the applied dynamic coating in cross flow mode is also shown to be applicable for upscaling, to allow coating of longer modules for industrial applications. Fouling studies of PEMs and their cleanability by releasing the PEM coating (the sacrificial layer effect) were also studied, first on model surfaces, and subsequently on membranes (hollow fiber).

2. Materials and methods

2.1. Chemicals

UF hollow fiber membranes, also known as hollow fiber silica (HFS), with a separation layer of sulfonated poly(ether sulfone) were obtained from Pentair X-Flow (The Netherlands). These membranes with a nominal molecular weight cut-off of 10,000 Da are intended for colloidal silica removal from water. The polyelectrolytes used were poly(allylamine) hydrochloride (PAH, M_w 17,500 g mol⁻¹) and poly(acrylic acid) (PAA, M_w 15,000 g mol⁻¹). The pH of the polyelectrolyte solutions was adjusted with 1 M hydrochloric acid (HCl) or 1 M sodium hydroxide (NaOH). Two model foulants, bovine serum albumin (BSA) and humic acid (HA) were used to simulate the presence of proteins and the presence of hydrophilic natural organic matter respectively. HA sodium salt was obtained from Sigma Aldrich (The Netherlands) with a reported molecular weight greater than 50,000 Da and a pH of 5.2 when dissolved in deionized water. BSA (cold alcohol precipitated) was obtained from Sigma Chemical Co. (St. Louis, MO) with a well-defined molecular weight of 67,000 Da and an isoelectric point of 4.8. At a 1 g l⁻¹ concentration, at the used pH of 7.3 the protein therefore has a net negative charge. The salts NaCl, Na₂SO₄ and MgSO₄, and organic compounds Bezafibrate and Naproxen, were used for NF performance tests and were purchased from Sigma Aldrich (The Netherlands). Deionised (DI) water used in all experiments was purified by a commercial water system (Synergy Water Purification System, Millipore) to a water resistivity of 18.2 MΩ cm or greater. All reagents were of analytical grade and used without further purification.

2.2. Membrane modification using dynamic LbL assembly

For membrane coating, PE solutions were prepared with a polymer concentration of 0.1 g l⁻¹ using a background electrolyte of 50 mM NaNO₃ at pH 6 for both PEs based on the results of our previous study [34]. Single membrane fibers were potted in 300 mm long modules

with a total surface area of 6.2 cm². Prior to coating, the membrane modules were wetted with 15 wt.% ethanol for 8 h to remove any impurities and then immersed in deionized water overnight. SEM images of the structure of membrane support used in this work were taken by JEOL JSM-6010LA and are given in Fig. S1. The coating was performed using a homemade fully automated crossflow system (schematic shown in Fig. S2) under different cross flow speeds (0.5, 2, 4 and 6 m s⁻¹) and pressures (0.1, 0.5 and 1 bar). The LbL deposition of PEs was performed from lumen to shell, and for that the membrane module was fed with a 0.1 g l⁻¹ polycation (PAH) solution. The module was then flushed with background electrolyte solution to remove excess PE, after which the switch was made to a 0.1 g l⁻¹ polyanion (PAA) solution. By repeating these steps the membrane was coated the desired number of PAH/PAA bi-layers. The coated membranes were stored in DI water until used.

Dead end coating was performed with a constant pressure setup (Amicon cell) with feed side of the system pressurized at 1 bar N₂ gas as illustrated in Fig. S2. Similar to cross-flow coating, dynamic LbL coating in dead end mode includes the following procedure: The membrane module was flushed with background electrolyte solution at 1 bar for 15 min. Then, the feed was changed to PAH solution and membrane module was flushed at 1 and 0.5 bar for 15 min followed by flushing with background electrolyte solution for 15 min as intermediate rinsing step. The flushing step, however, was performed in cross-flow mode to remove any unattached polyelectrolyte from the reservoir, and to avoid bulk polyelectrolyte complexation. Again, the feed was changed to PAA solution and module was flushed for 15 min followed by a rinsing step with background electrolyte solution to get one bilayer. Same procedure was repeated to get the desired number of bilayers. During the whole coating procedure, pure water permeability and permeability of PE solution through the UF support was measured. To check the effectiveness of dynamic coating at a much larger scale, the same cross flow modification procedure mentioned before was carried out for modules with an effective length of approximately 200 mm housing 15 fibers in one module (surface area of 67 cm²).

2.3. TOC analysis

Total organic carbon (TOC) values of PAH and PAA feed solution and UF permeate in the first filtration run were measured using TOC analyzer (TOC-V_{CSN}, SHIMADZU).

2.4. Filtration and NF performance

The pure water flux of membranes was measured for each coating step in a dead end mode. The water permeability (in l m⁻² h⁻¹ bar⁻¹) of membrane was obtained by normalizing the pure water flux measured at 20 °C by trans-membrane pressure (TMP) used (1 bar). The membrane permeability was then calculated using Eq. (1):

$$P(\text{l} \cdot \text{m}^{-2} \cdot \text{h}^{-1} \cdot \text{bar}^{-1}) = \frac{V}{A \cdot \Delta t \cdot \Delta P} \quad (1)$$

Where V is the volume (l), A is the membrane area (m²), t is the time (s), ΔP is the trans-membrane pressure (bar). The relative permeability was obtained by dividing the final permeability at a specific time with the initial value and is reported as a percentage. Each measurement was performed in duplicates and repeated only if the deviation was significant.

Retention experiments were performed using a laboratory scale crossflow filtration unit (Fig. S3) in batch mode with recirculation of the retentate in the feed tank. The inorganic salts used for salt retention measurements were NaCl, MgSO₄ and Na₂SO₄. Salt retention of the coated hollow fibers was determined by cross flow filtration at 1 bar applied pressure with a crossflow velocity of 4.5 m s⁻¹ (Reynolds number of approximately 3600). Concentration changes of the salts in the permeate and concentrate were monitored using conductivity measurements (WTW cond 3210 conductivity meter).

The retention values of solutes were calculated using Eq. (2):

$$R(\%) = \left(1 - \frac{C_p}{C_f}\right) \times 100 \quad (2)$$

Where c_p and c_f represent the concentration of the solute in the permeate and feed at the end of the experiment respectively. The experiment was stopped when the retention reached a constant value and the average of a minimum of three measurements was used.

The membrane performance was also investigated by performing retention experiments on the small organic pollutants Bezafibrate Mw = 361.8 Da and Naproxen Mw = 230.26 Da both with a negative charge. Dionex Ultimate 3000 U-HPLC system equipped with an RS variable wavelength detector was used to determine the concentration of micro-pollutants in the concentrate and permeate. Micro-pollutant separation was done on an Acclaim RSLC C18 2.2 μm column (Thermo Scientific) at 45 °C, while applying a gradient flow from 95 wt.% H₂O + 5 wt.% acetonitrile at pH 2–5 wt.% H₂O + 95 wt.% acetonitrile at 1 ml min⁻¹.

2.5. Fouling and subsequent cleaning by the sacrificial layer approach

2.5.1. Reflectometry

Adsorption and desorption of multilayers and model foulants (BSA and HA) was monitored on silicon wafers using optical reflectometry, a proven technique for studying adsorption onto substrates [46]. The fixed-angle optical reflectometry set-up consists of a He-Ne laser (λ = 632.8 nm) with linearly polarized light. During adsorption or desorption, the resulting change in polarization is measured by detecting the parallel (R_p) and perpendicular (R_s) components of the polarized light. The ratio of the two components is the signal, S₀ (–) and the change in the ratio, ΔS (–), is directly proportional to the amount of material adsorbed on the substrate (Eq. (3)).

$$\Gamma = Q(\Delta S/S_0) \quad (3)$$

Where Γ is the amount of mass adsorbed on the substrate (mg m⁻²) and Q is the sensitivity factor (mg m⁻²) that depends on the angle of incidence (θ), refractive indices (η), thickness of the layers (d) and the refractive index increment (dn/dc) of the adsorbate. The values used to calculate Q factor are refractive index of silicon wafer n_{Si} = 3.85, refractive index of silica layer n_{SiO₂} = 1.46, thickness of the silica layer d_{SiO₂} = 120 nm, refractive index of solutions n_s = 1.33, dn/dc = 0.185 ml g⁻¹ [41]. The corresponding Q factor was calculated to be 45 mg m⁻² for all experiments. Prior to adsorption tests, the surface of the treated silicon wafer was cleaned using O₂ plasma cleaner with 50% power (Femto Diener, Germany).

2.5.2. Membrane fouling and cleaning study

The fouling and cleaning experiments on membranes were performed in cross-flow mode by applying the minimum cross flow speed (0.5 m s⁻¹). These tests included three main steps. The coated and uncoated membranes were first fed with DI water until a stable flux was achieved. Then the feed was changed to an aqueous solution containing the model foulant at the appropriate concentration (1 g l⁻¹ BSA, pH 7). The fouling experiments were performed for 2 h at a transmembrane pressure of 2 bar, and over the time the change in flux was monitored. Similar experiments were performed for HA fouling (1 g l⁻¹ HA, pH 5.8). From the flux obtained membrane resistance was calculated using Eq. (4):

$$R = \frac{\Delta P}{\mu \times J} \quad (4)$$

Here, R is the membrane resistance (m⁻¹), μ is the dynamic viscosity of the feed (Pa·s) and J is the membrane flux (l m⁻² h⁻¹). After the fouling experiments, membrane cleaning and surface regeneration was conducted to determine the success of the sacrificial layer concept.

The fouled membranes were washed in crossflow mode with a 3 M NaNO₃ solution at pH 3 for specified time. The membranes were subsequently cleaned in black flush mode at 1 bar using the same trigger solution. After each cleaning, the membrane was rinsed with DI water and membrane resistance was measured. The extent of cleaning was characterized through comparison of the resistance before and after the cleaning. Following the cleaning steps, multilayers coating was regenerated with fresh PE solutions as outlined in Section 2.2. In order to evaluate the success of the cleaning and regeneration, the membrane resistance to pure water at each stage is compared.

3. Results and discussions

The results and discussion is divided into two main parts. In the first part we report on the preparation of hollow fiber PEM NF membranes using dynamic coating under different modes and dynamic coating conditions. The second part deals with the fouling and sacrificial cleaning studies of PEM's first on model surfaces and subsequently on dynamically coated hollow fiber NF membranes.

3.1. Membrane coating and performance

3.1.1. Effect of cross flow speed, filtration time and pressure on coating

In order to optimise the deposition of PEMs on membranes by active coating, several parameters were investigated such as adsorption time, applied pressure and cross flow speed. Typically, the adsorption time for PEs is in the order of a few seconds to a few minutes to reach a stable plateau in stagnation point flow [47]. However, this applies to a non-porous model substrate under very well defined flow conditions. Prior to coating the multilayers onto the porous UF support via dynamic assembly, the effect of filtration time and coating speed on the deposition of the first layer of PE was investigated by feeding an uncoated UF support with a PAH (+) solution at constant cross flow speed and a constant pressure of 1 bar. The same experiment was carried out at different crossflow velocities ranging from 0.5 to 6 m s⁻¹. Fig. 1(a) shows the membrane permeability during coating as a function of filtration time. The results are presented in terms of permeability of PE solution relative to the initial pure water permeability. It can be observed that initially, the flux of the UF support decreased rapidly for low cross flow speeds (0.5 and 2.0 m s⁻¹), while remaining quite high for the higher cross-flow velocities (4.0 and 6.0 m s⁻¹). For 2.0 m s⁻¹, a stable flux is reached approximately within 10–12 min of filtration, but this is not the case for lower and higher cross flow velocities (0.5, 4.0 and 6.0 m s⁻¹), where the flux keep declining even after 60 min. At high cross-flow velocities, we expect that the high shear is limiting the rate of deposition. Just as a high cross flow velocity limits the fouling in

membranes, a high cross flow applied during coating allows less material to deposit on the membrane resulting in thinner and more permeable layers. This also explain why at the lowest cross-flow velocity, a less permeable and thus thicker layer is formed. The formation of this thick layer also takes considerably more time. Overall it can be clear that the cross-flow velocity affects the rate and extend of poly-electrolyte deposition.

Since the size of PAH is approximately 50% larger than the MWCO of the virgin membrane and the PEs system is fully dissociated at pH 6, the rate of deposition is therefore expected to be primarily controlled by electrostatic interactions with the membrane support. To verify if PE chains are 100% retained by the membrane pores under pressure coating, the TOC values of the PE during the first coating step for the permeate and the feed were determined. TOC values of feed and the permeate were 47.22 mg l⁻¹ and 1.03 mg l⁻¹ respectively, suggesting that 97% of PE was retained by the membrane during deposition of the first bilayer and only 3% PE molecules may pass across the pores of the membrane support, confirming that most of the PE material is deposited inside or on top of the pores during coating. The polyelectrolyte transport that does occur, is expected to take place in the early stages of the experiment, when the pores are still fully open. As both the membrane pores and the polymers are quite polydisperse, some of the smaller chains will fit easily through the larger pores.

We then evaluated the impact of pressure on the membranes coated at a constant crossflow velocity. Fig. 1(b) shows the relative permeability for membranes coated for 15 min at varying pressures but at constant crossflow velocity. At constant cross-flow velocity of 0.5 m s⁻¹ (Reynold number ≈ 394), an increase in the pressure from 0.5 bar to 1 bar resulted in a decrease in permeability. At a higher pressure more polymers are transported to the membrane surface and are retained there, leading to a thicker less permeable layer. Unexpectedly, however, an increase from 1 bar to 2 bar resulted in an increase in the permeability of the membrane. We expect that the higher flux and the resulting higher shear leads to thinner layers at the membrane interface. On the basis of these experiments the condition of 1 bar was chosen as the optimum based on the drop in permeability observed. This will be used in the next sections.

Since preparation conditions greatly influence the efficiency of the formed membranes, the performance of these modified membranes was followed in greater detail using relative permeability measurements. Preparation under cross-flow was compared to preparation under dead-end conditions and under static conditions (dip coating). As shown in Fig. 2(a), the permeability decreases as the number of bilayers increase. The reduction in permeability with each bilayer confirms that the layers are indeed being deposited, while the small error bars confirm that all approaches give very reproducible results. It is somewhat surprising

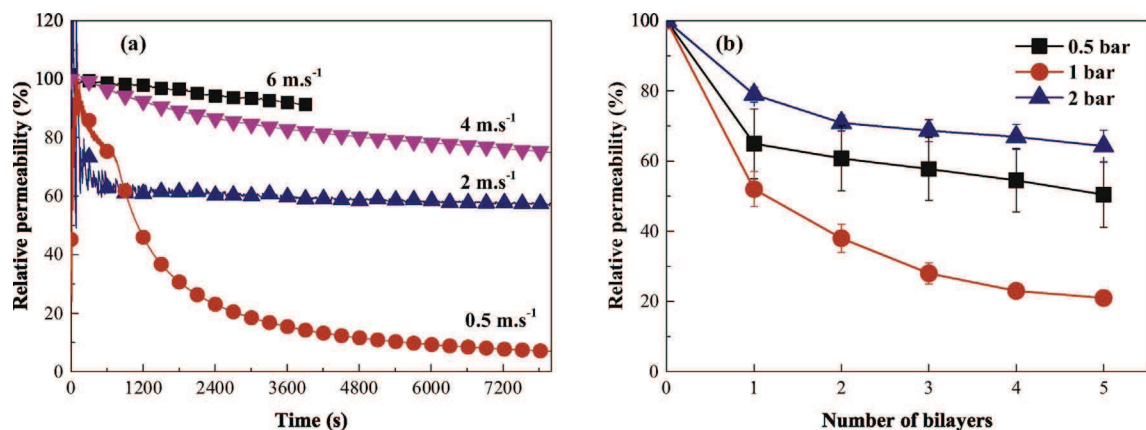


Fig. 1. (a) Time dependent variation of permeability with reference to initial water permeability during coating of the first PAH layer under different cross flow speeds; (b) Effect of coating pressure and coating steps on membrane permeability. Measurements performed in triplicates, coating solutions contained PE conc. 0.1 g l⁻¹ with 50 mM salt and pH 6. Permeability of uncoated support was 60–70 l m⁻² h⁻¹ bar⁻¹. All experiments were repeated in triplicate on separate membrane supports.

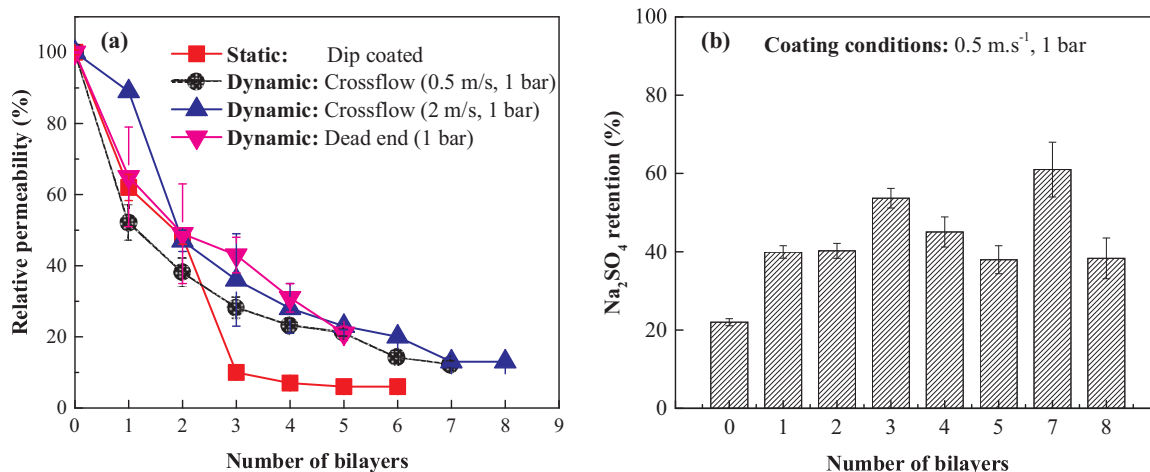


Fig. 2. Comparison of different modes of coating on membrane performance; (a) Relative permeability change with increasing bilayers coated under different coating techniques; (b) Effect of coating steps on membrane performance in terms of Na₂SO₄ rejection. All experiments repeated in triplicate.

that dead-end coating and coating under cross-flow conditions lead to such comparable permeabilities. Under dead-end coating, the polyelectrolyte concentration in the fiber would continuously increase, and one might expect thicker layers. Still, the stop mechanism for multilayer coating, even under dead-end coating, would be the build-up of an excess of charge at the membrane surface, preventing further adsorption. Moreover, while coating was performed in dead-end, the rinsing was done in cross-flow mode, possibly leading to removal of weakly attached chains.

Fibres coated under static conditions were made via dip coating in 0.1 g l⁻¹ PE solutions with an adsorption time of 15 min per layer. As can be seen in Fig. 2 (a) the membranes prepared by dip-coating LbL assembly show sharper decrease in permeability followed by a plateau after 3 bilayers. Between 2–3 bilayers a 40% decrease in permeability of dip coated membranes occurs with the permeability being approximately 20% lower than dynamic modes. As was shown in Fig. 1a, the shear applied during active coating limits a decrease in permeability, with more shear leading to thinner layers. It is thus logical that under no shear conditions (dip coating), the thickest layers, with the lowest permeability, are formed.

Recently Menne et al. [40] demonstrated that thicker layers are produced from dynamic coatings (constant flux coating in dead end) relative to dip coating, which is in contrast to our observations. However they used a different support, a combination of strong PEs

(PDADMAC/PSS) and higher concentration of PEs (1 g l⁻¹). We expect that our combination of weak polyelectrolytes, leads to multilayers that are mechanically much weaker. Hence, the much greater role of shear in reducing the thickness of the deposited polyelectrolyte multilayers in our system. Since 0.5 m s⁻¹ proved to have the lowest permeability (better deposition of PEs) of the dynamic techniques used, this cross flow velocity was used for subsequent experiments.

3.1.2. Homogeneity of LbL coating

In order to check the homogeneity of the coating, a longer (40 cm long) membrane module was taken and divided into three segments of 10 cm active length and the permeability and retention of segmented sections were compared. A schematic of the segmented fiber is shown in Fig. S4. The segmented module was coated dynamically under 1 bar pressure, while applying 0.5 m s⁻¹ cross flow through the fiber. Fig. 3(a) shows the relative pure water permeability of each segment with a 2% deviation indicating that the sections are coated evenly. This is important to consider for longer modules where pressure drop inside the fiber can limit the proper coating, but that is not the case under the coating conditions used here. As shown in Fig. 3(b) the retention of Na₂SO₄ in each segment was comparable and no significant deviation was found. The dynamic coating process may therefore be applied to industrial systems and a uniform performance is expected from the entire module.

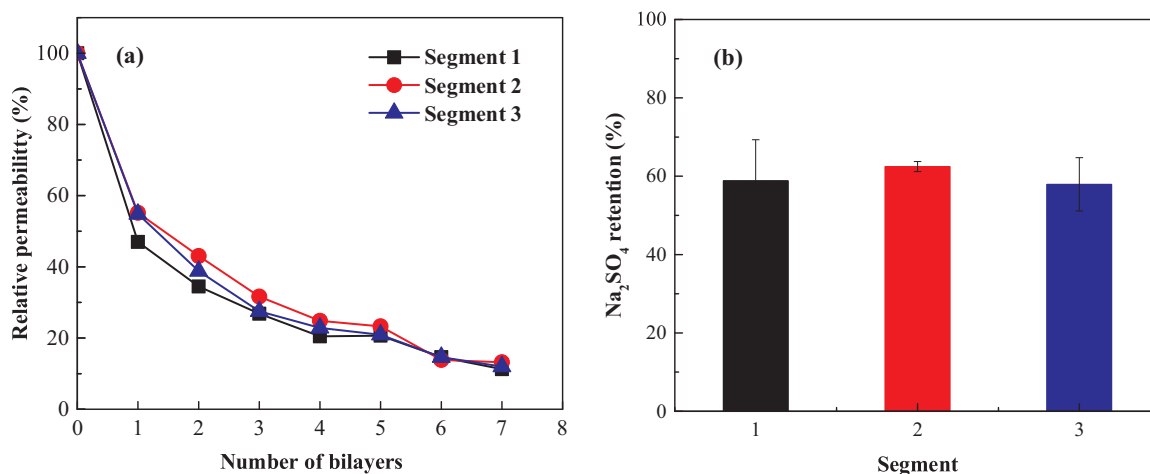


Fig. 3. Homogeneity test of dynamic coating (a) Relative permeability of each 10 cm segments; (b) Na₂SO₄ retention of each 10 cm segment, measured membrane coated with 7 bilayers (coating is performed with 0.1 g l⁻¹ PE containing 50 mM salt at pH 6, under 1 bar pressure and 0.5 m s⁻¹ cross flow speed). All experiments repeated in triplicate.

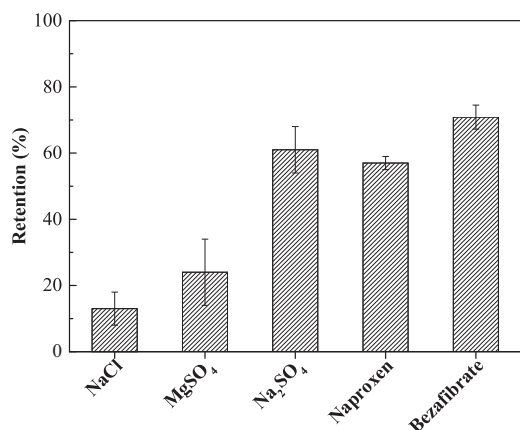


Fig. 4. NF membrane performance in terms of retention of mono and divalent salts and organic molecules. The tested membranes were coated with 7 bilayers of PAH/PAA at speed of 0.5 m s^{-1} and 1 bar pressure. Permeability of tested membrane was $6.52 \pm 0.51 \text{ m}^{-2} \text{ h}^{-1} \text{ bar}^{-1}$. All experiments repeated in triplicate.

3.1.3. Membrane performance as NF

To check if the created membranes were indeed within the NF performance regime, the retention of different salts (NaCl, Na₂SO₄ and MgSO₄) and small organic molecules (Bezafibrate and Naproxen) was investigated. Membranes coated with 7 bilayers of PAH/PAA terminating with the PAA (-) layer with 0.5 m s^{-1} cross flow and 1 bar pressure were used for NF performance test. As can be seen from Fig. 4, the prepared membranes are capable of rejecting Na₂SO₄ (60%) and organic solutes (55–70%), while monovalent ions (NaCl) and divalent cations (MgSO₄) are hardly rejected. This ion rejection is fully consistent with the results of a negatively charged nanofiltration membrane, that separates on the basis of Donnan exclusion. Here SO₄²⁻ is much more strongly rejected than Cl⁻, explaining the high Na₂SO₄ rejection compared to NaCl. On the other hand MgSO₄ is much less rejected compared to Na₂SO₄, as the divalent cation Mg²⁺ tends to accumulate in the negative membrane, drawing its counter ion along. Still the high retentions of Bezafibrate and Naproxen must have to do with their size, as they have just a single charge. Based upon the results it is expected that mainly size and charge exclusion are dominating the rejection mechanism, although other factors such as affinity between membrane, solute and solvent can also play role in separation mechanism.

3.1.4. Upscaling of cross flow dynamic coating approach

After optimizing the dynamic coating conditions on a single fiber module, we checked the upscaling potential of this dynamic coating approach to coat modules with an effective length of approximately

200 mm housing 15 fibers in one module yielding a total effective area of 67 cm^2 . Coating was performed at constant pressure of 1 bar under two cross flow speeds (0.5 and 2 m s^{-1}). As shown in Fig. 5 with every coating step a drop in permeability was observed and similar to single fiber module coating the drop is higher for modules coated at low cross flow speed due to accumulation of material. In contrast, the salt rejection stays relatively constant with an increasing number of bi-layers. It seems that already after 1–2 bi-layers, the PEM coating fully covers the support membrane, after which an increase in the number of layers does not lead to improved separation [32]. The results obtained on our larger module scale, is fully consistent with the results obtained at the single fiber level. We conclude that the dynamic coating approach is equally suitable for coating on module level, and could likely even be used to coat existing UF modules in various industrial applications.

3.2. Fouling and cleaning studies

In the second part of this study we test the fouling propensity of PAH/PAA multilayer system and their potential as sacrificial layers for easy membrane cleaning. Fouling and cleaning studies were performed first on model surfaces (silicon wafer) in order to optimise adsorption and release conditions, and were later replicated on LbL coated membranes.

3.2.1. Reflectometry

Adsorption, fouling and sacrificial cleaning of PEMs was monitored on silicon wafers using optical fixed angle reflectometry. The adsorption measurements on silicon wafers were started by exposing the surface to a PE solution of PAH (0.1 g l^{-1} , pH 6, 50 mM NaNO₃), then switching to a PAA (0.1 g l^{-1} , pH 6, 50 mM NaNO₃) solution. Between subsequent adsorption steps, the substrate was rinsed with the background electrolyte solution (50 mM NaNO₃) to remove excess PE and avoid the formation of bulk complexes. Continuous switching between cationic and anionic PEs leads to a stepwise growth of the multilayers. Subsequently, a foulant solution was applied on top of differently terminated multilayers (PAH or PAA as the final layer). To simulate fouling two model foulants were used in this study bovine serum albumin (BSA, 0.1 g l^{-1} , pH 7) and Humic acids (HA, 0.1 g l^{-1} , pH 5.2). Both foulants are negatively charged at the pH conditions used for this study. Similar to the spontaneous self-assembly phenomena observed in PEs, biological materials can adsorb via electrostatic interactions. Many biological foulants are negatively charged therefore negatively charged membranes generally exhibit decreased adsorption in comparison to positively charged membranes. As can be seen in Fig. 6, this phenomenon was clearly observed, when the foulants introduced to the PAA (-) terminated layer, the quantity of foulant adsorbed is significantly less at approximately 1 mg m^{-2} for BSA and HA. In contrast, for PAH (+) terminated layers the introduction of the foulant leads to a large

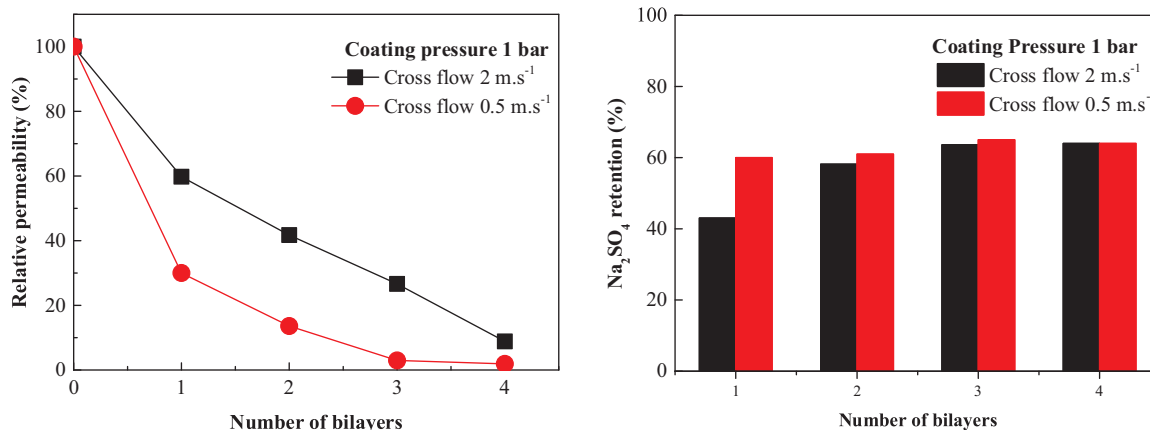


Fig. 5. (a) Variations of pure water permeability relative to initial permeability and (b) Na₂SO₄ retention of PAH/PAA coated membranes with the number of coated layers.

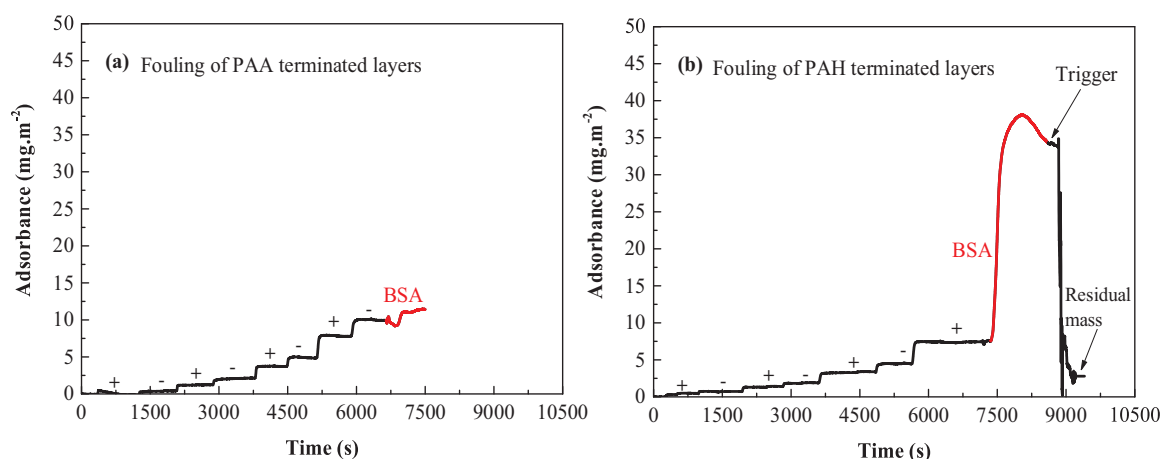


Fig. 6. Effect of terminal layer on BSA and HA adsorption and sacrificial layer effect (a) BSA fouling of multilayers terminated with PAA (–) layer; (b) BSA fouling of PAH (+) terminated layers and effect of trigger solution (pH 3, 3 M) on desorption of fouled multilayers. Concentration of BSA 0.1 g l^{-1} , pH 7. Rinsing solution used after every adsorption step is 50 mM NaNO_3 , pH 6.

increase in the quantity of material adsorbed. The factor to which the mass increased for BSA and HA was 5 and 2 respectively. Results of HA fouling of PAH and PAA terminated multilayers are given in Fig. S5. Although both foulants adsorbed onto the positive (PAH) layer, they adsorbed to different degrees with BSA being adsorbed more. While this behaviour is significant, for this particular study, the foulant-polyelectrolyte electrostatic interaction is not the primary focus here. The overall decrease in fouling propensity observed with PAA (–) terminated layers demonstrates that by careful selection of the final layer of a PEM, a low fouling layer can be achieved. But apart from low fouling surfaces, these multilayers can also be used as sacrificial layers to allow for easy cleaning. When a trigger solution was applied on the fouled PEMs, the adsorbed multilayers and the fouling agents are washed away from the silicon wafer (Fig. 6), again indicating their potential as sacrificial coatings. When the pH is low, PAA becomes considerably uncharged and an increase in the ionic strength, then further weakens the ionic interactions resulting in disintegration of multilayers, and removal of all attached foulants.

3.2.2. Membrane fouling and cleaning

In this section, we demonstrate that the promising properties of multilayers as observed via reflectometry can also be applied to a membrane system. The interactions between the foulants and the membranes are investigated through cross flow permeation tests using two model foulants (BSA and HA). Fouling experiments were conducted

under very low cross flow speed (0.5 m s^{-1}) for about 2 h with 1 g l^{-1} foulant. Initially the pure water permeability of the Virgin UF support was tested. The fouling propensity was measured as the ratio of the permeability at a point in time to the initial permeability. The UF support used contains negative charge and is well developed for retention of silica particles. Hence, the membrane is expected to already have a very low fouling propensity. Additionally, the membrane is known to be hydrophilic such that repulsive hydration forces are expected to repel the hydrophobic foulants. When exposed to the foulants, the support membranes showed an average 8% decline in permeability (Fig. 7). This decline in flux is primarily due to the deposition of the foulants on the membrane surface by adsorption and aggregation. When coated membranes (with 5 and 5.5 bilayers of PAH/PAA) were exposed to foulants, it can be observed that BSA exhibited a greater affinity for the membrane surface than HA (Fig. 7 a & b), which is in line with the results of reflectometry (Fig. 6, S5). Here we can also see the impact of differently terminated layers (PAA and PAH) on foulant adsorption. A rapid flux decline was observed for PAH terminated membrane during crossflow filtration of 1 g l^{-1} BSA (pH 7). This behaviour may indeed be explained by the electrostatic interaction between negatively charged BSA molecules and positively charged PAH membrane. The carboxylic functional groups of the PAA terminated membranes are negative and thus a strong electrostatic repulsion is expected between the negatively charged BSA molecules and the negative membrane surface. While the strong electrostatic repulsion is the most

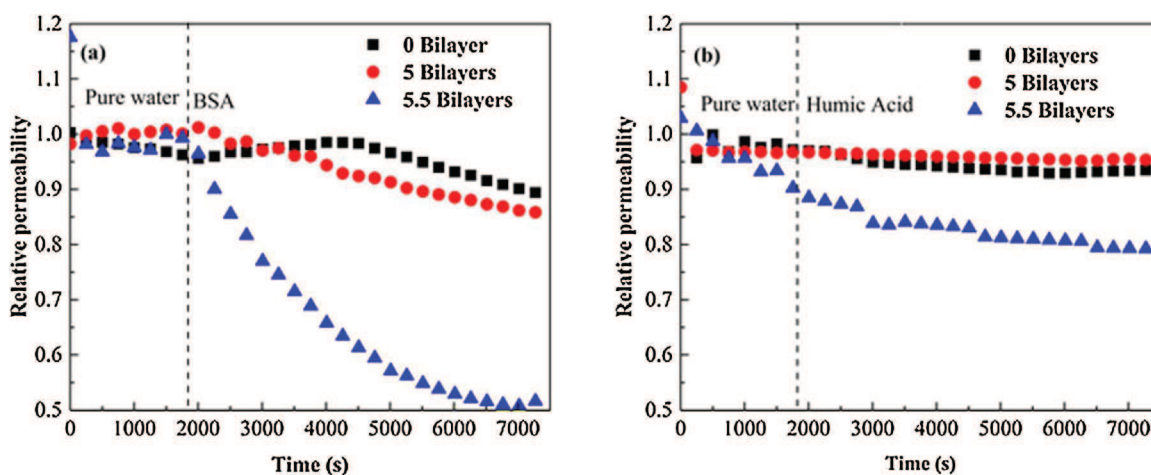


Fig. 7. Permeability versus time of un-coated and coated membranes with PAH/PAA with + and – terminal layers membranes during the filtration of BSA and HA. Fouling experiments were conducted at $25 \text{ }^\circ\text{C}$ and 1 bar operating pressure. All experiments performed in triplicate.

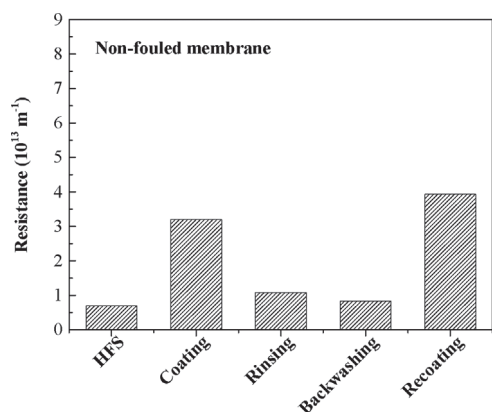


Fig. 8. Measured membrane resistance to pure water permeation at different stages of coating and sacrificial cleaning of multilayers. The sequence followed is dynamically coating the HFS membrane with 5.5 bilayers of PAH/PAA, rinsing of multilayers from inside to out for 1 h with trigger solution (pH 3, 3 M), backwashing with trigger solution from out to inside of membrane under pressure of 2 bar for 1 h and re-coating of clean membrane with 5.5 bilayers of PAH/PAA. All experiments performed in duplicate.

important reason for improved fouling prevention, the ionised carboxylic groups present on the membrane surface also interact with water molecules through Van der Waals forces and hydrogen bonds thereby preventing the deposition of the protein. The overall decrease in fouling of PAA terminated layers relative to PAH terminated layers and the uncoated membrane clearly highlights a viable approach of surface modification while retaining a very low fouling propensity. Antifouling results of PAA terminated layer are comparable to a UF membrane support optimised for fouling control.

3.2.3. Sacrificial membrane cleaning

In this section, we investigate the use of dynamically coated PEMs as sacrificial layers. The approach will be used for cleaning of fouled multilayered NF membranes along with the possibility of regeneration of the PEM membrane. Initially, the use of dynamically assembled PEMs as sacrificial layers was evaluated without fouling. For this 5 bilayers of PAH/PAA on a UF support were first rinsed with the trigger solution (pH 3, 3 M) in cross-flow mode at 2 bar and subsequently back-flushed with the trigger solution at 2 bar for an hour. It can be seen (Fig. 8) that the coating is fully removable with the applied trigger solution bringing the membrane resistance back to the level of uncoated

membrane support (HFS).

Subsequently the potential of this sacrificial coating approach to remove BSA foulant from the membrane is investigated. For sacrificial cleaning experiments we chose PAH terminated membrane (coated with 5.5 bilayers of PAH/PAA) fouled with BSA because this membrane and foulant showed the highest fouling propensity (Fig. 7a). In Fig. 9 we show the membrane resistance against pure water permeation for several stages, dynamic coating (coating at 0.5 m s^{-1} cross flow and 1 bar pressure), fouling with BSA, sacrificial cleaning and subsequent re-generation. Here the same cleaning protocol was applied for PEM removal as used for un fouled membrane, at similar conditions and number of bilayers as the un fouled membrane, the resistance of fouled membrane after cleaning went back close to the resistance of coated membrane suggesting the trigger does not effectively remove the PEM layer. This proposes that the nature of the foulant can impact the success of the layer removal. Unlike the hard spherical nature of silica particles [34], it is believed that the BSA molecules form a thick gel layer on the membrane surface that holds the layer in place. Consequently, after the cleaning step, the membrane is effectively unchanged and did not return to its pristine state as expected.

In light of these results, the order of cleaning was adjusted to include backflushing as a first step followed by rinsing, so that the polyelectrolyte on the surface can be lifted off and re-suspended in a tangential flow. This approach proved to be much more effective than the previous combination as it resulted in an 86% reduction in the resistance (Fig. S6). Rinsing, therefore had no capacity to remove the BSA molecules while backflushing did. This is believed to be due to the fact that the trigger solution now had direct contact with the PEMs through the membrane's structure resulting in immediate removal of the layers and the attached foulants. Complete recovery of the pristine membrane resistance was obtained when modules were backwashed at a bit higher pressure of 4 bar followed by rinsing for an hour. Following up with rinsing resulted in further removal of loosely bound material. The used commercial support can bear upto 5 bar backwash pressure and a pH as low as 1 so we do not expect any change in the membrane performance after cleaning procedure. This same support was already used by de Grooth et al. [31] successfully in combination with sodium hypochlorite and a backwash pressure of 5 bar without any reported damage to membrane. Two successive sets of BSA fouling, cleaning and re-generation experiments were performed on the same module to assess the longevity of this regeneration process. The pure water permeability and resistance were measured again and compared with the initial values. As shown in Fig. 9, the membrane maintained a relatively low

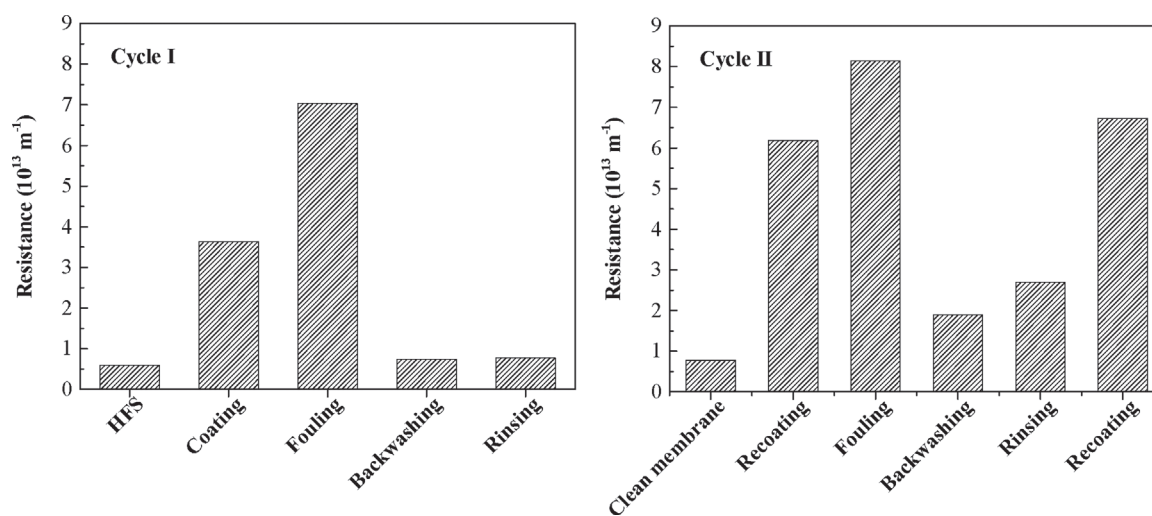


Fig. 9. Measured membrane resistance to pure water permeation at different stages of coating, fouling and sacrificial cleaning of fouled multilayers. The sequence followed is dynamically coating the HFS membrane with 5.5 bilayers of PAH/PAA, fouling with 1 g l^{-1} BSA, backwashing with trigger solution from out to inside of membrane under pressure of 4 bar for 8 h rinsing of multilayers from inside to out for 1 h with trigger solution (pH 3, 3 M), and re-coating of clean membrane with 5.5 bilayers of PAH/PAA. All experiments performed in duplicate.

resistance after a series of runs. After the first regeneration cycle, the resistance returned close to pristine membrane resistance. The second cycle resulted in a further 20% decrease in the membrane support permeability. It can therefore be concluded that the membranes are less prone to fouling if terminated with PAA layer and can be regenerated multiple times with the applied cleaning agent and cleaning protocol. However, the cleaning is not complete after more than 1 cycle, and or continuous regeneration some additional cleaning approaches need to be taken into account.

4. Conclusions

In this study sacrificial PEMs based hollow fiber NF membranes were prepared using dynamic LbL coating under constant pressure and in cross flow mode. The effect of different cross flow speeds, pressures and mode of coating was investigated extensively and compared with dip coating method. It is shown that dynamic deposition is a simple way to modify the surface (inside) of existing hollow fiber UF support to make versatile NF membranes with low fouling propensity (if terminated with PAA) and which are relatively easy to clean by sacrificial layer approach. Still, even with the sacrificial layer approach, the foulant removal after 2 cycles is not complete. This indicates that the sacrificial layer approach is beneficial, but should still be combined with other cleaning approaches. Rejection performance of solutes (salts and organic micro-pollutants) by dynamically coated membranes fall in typical NF performance range (50–70% retention). Moreover this coating method with its reproducibility and homogeneity features also shown to have upscaling potential to coat existing UF membrane modules.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.colsurfa.2017.09.003>.

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