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Fouling control using critical, threshold and limiting fluxes concepts for cross-flow NF of a complex matrix: Membrane BioReactor effluent

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A B S T R A C T

The optimization of permeate flux is a particularly interesting strategy to control fouling and, as a consequence, enhance productivity for nanofiltration (NF) processes. Critical flux, threshold flux and limiting flux theories represent significant advance in this strategy. The aim of this study was to apply these concepts to achieve fouling control during NF of a real complex matrix: hospital wastewater after Membrane BioReactor treatment (MBR permeate). At low pressure (3 bar) no flux decline was observed, revealing no fouling conditions. By applying a range of transmembrane pressure and using the square-wave method, the critical flux for irreversibility ($70 \text{ L h}^{-1}\text{m}^{-2}$) and the corresponding critical pressure (3.4 bar) were then determined for the NF process in this complex matrix. Above these critical conditions, irreversible fouling started to occur. The threshold pressure and related flux (transition points between low and high fouling regions) were then searched by critical flux data conversion. Our results suggest, even if an exact value for the threshold pressure could not be determined, that it could be located in the range 3.4–10 bar. Operating in this pressure range should lead to acceptable fouling rate and flux decline.

During filtrations conducted above the critical flux in the MBR effluent, two stable fluxes behaviours were observed indicating that different fouling stages occur: pseudo stable flux was $67 \text{ L h}^{-1}\text{m}^{-2}$ at 5 bar, whereas $33 \text{ L h}^{-1}\text{m}^{-2}$ at 10–35 bar. It can be then confirmed that a limiting flux occurred in this system, the value of which $33 \text{ L h}^{-1}\text{m}^{-2}$ is rather lower than that of the critical flux. This flux behaviour was proved to be related to a severe fouling in the pressure range 10–35 bar due to a combined effect of colloidal silica and organics fouling and calcium phosphate scaling. The early fouling stage at 5 bar was expected to be solely related to colloidal silica and organics accumulation. To characterise this change in fouling behaviour, a method allowing the estimation of the permeability before scaling was proposed. The combination of the permeability before scaling and critical flux has enabled a NF working diagram to be drawn from which the fouling stage for a given transmembrane pressure and corresponding permeate flux was able to be determined.

1. Introduction

Nanofiltration (NF) membranes are widely used for liquid-based separations, such as water purification. These membranes are often applied as tertiary treatment for effluents of Conventional Activated Sludge (CAS) or Membrane BioReactors (MBRs), to remove biorefractory micropollutants, when high water quality is required. Although the efficiency for the removal of organic contaminants is satisfactory [1–3], membrane fouling is a major problem in the operating of NF. Fouling increases the cost due to the increase in energy demand, the use of chemicals for cleaning, and the frequency of membrane replacement [4].

NF membranes are normally subject to fouling by colloidal materials and soluble inorganic compounds, which are not removed by pre-

treatment. Firstly, colloidal sized substances, consisting of organic material, silica, metal oxides and microorganisms are the foulants of greatest concern for nanofiltration [4]. Their deposition on the membrane surface is, in a complex way, due to various phenomena: hydraulic drag force, electrostatic repulsion, van der Waals attraction or hydrophobic/hydrophilic interactions with different interaction lengths [5]. Secondly, membrane scaling (or inorganic fouling) is a serious problem in applications of NF processes. Scaling leads to all of the following: irreversible membrane pore blocking [4,6], a dramatic decrease of flux, and physical damage of the membrane due to the difficulty in the removal of scalant. Scaling results from the concentration of one or more inorganic substances beyond their solubility limits and their precipitation onto the membrane surface [7]. In real effluents, the complexity of the matrix composition makes it difficult

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to understand and anticipate fouling only through feed analysis.

One strategy for fouling control is to optimise the permeate flux which is a particularly interesting parameter. Cohen and Probstein [8] found that fouling may be prevented simply by operating below a threshold transmembrane flux during the filtration of a colloidal suspension in reverse osmosis systems. These authors hypothesised that foulant deposition occurs only if the hydrodynamic drag force exceeds the interaction forces. Based on this study, critical flux concept was firstly defined as “the flux below which no fouling occurs” [9,10], and as “the flux below which a decline of flux with time does not occur; above it fouling is observed” [11]. For colloidal suspensions [9,10], this concept was developed to generally describe the balance between foulants–foulants or foulants–membrane repulsive forces and permeate flux drag forces. Above the critical flux, permeate drag forces overcome repulsive forces which induce the deposition of foulants on the membrane surface. At that time fouling occurs and the permeate flux declines. The original forms of critical flux were described to be the strong form and the weak form by Field et al. [11,12]. The strong form of critical flux, J_{cs} , was developed to discriminate no fouling conditions and defined as the point at which the flux starts to deviate from the pure water flux-transmembrane pressure line. The weak form, J_{cw} , is observed in systems where a very rapid fouling (due to surface or pore adsorption that takes place independently of flux) occurs immediately at the start of the operation, leading to a linear portion of the flux-transmembrane pressure curve with a slightly lower slope than the pure water flux line. In addition to J_{cs} and J_{cw} , Bacchin et al. [13] introduced the concept of critical flux for irreversibility, J_{ci} , which has a clear link to a physical phase transition. Above the critical flux for irreversibility, multilayers of irreversible fouling appear and grow in the boundary layer whereas below J_{ci} only a concentration polarization layer exists in all cases with an additional monolayer of adsorbed species in some cases. This research group [5,14] also developed a filtration procedure, namely square-wave method, which enables the identification of the reversible and irreversible parts of fouling and the determination of the critical flux for irreversibility. Theoretically, operating below J_{ci} can minimise irreversible membrane fouling and flux decline.

In the case of real wastewater filtration, membranes system may not exhibit a critical flux as fouling immediately occurs even at very low fluxes. Moreover, in water industry the main objective is not to totally avoid fouling but to define operating conditions in which fouling is minimized to limit cleaning frequency. The concept of threshold flux was then recently introduced. Threshold flux, J_{th} , was proposed to be that flux at or below which a low and nearly constant rate of fouling occurs and above which the rate of fouling increases markedly [12]. Instead of “no fouling” condition, threshold flux focuses on the rate of fouling with no distinction between reversible and irreversible fouling. The goal of operating below the threshold flux is to obtain an acceptable fouling rate and sufficiently high flux over a long period of run.

Beside critical and threshold fluxes, limiting flux, J_L , is another important concept for fouling control. This limiting flux represents the maximum stationary permeate flux, which can be reached by increasing the transmembrane pressure with a given solution or suspension [13,15,16]. For this limit, further increments in transmembrane pressure do not increase flux. At a high flux, the drag force overcomes repulsive forces and the flux continues to decrease until the hydrodynamic drag force is lower or sufficiently close to the foulant-membrane and/or foulant deposited-foulant interaction forces [13].

Considering the fact that critical, threshold and limiting fluxes can be potentially powerful tools for understanding and controlling fouling, numerous studies attempted to develop these concepts and to investigate their existence in synthetic solutions and real wastewater. Wu et al. [17] determined the value of critical fluxes with pressure steps, composed of regular positive and negative variations. Both strong and

weak forms of critical flux were observed for two colloidal silica suspensions in this study. Manttari and Nystrom [18] evaluated the critical flux with a stepwise filtration procedure, and only a weak form of the critical flux was observed in effluents from the paper industry. In other cases, some authors noted that fouling cannot be completely inhibited during the treatment of complex fluids such as wastewater even at low flux rates [19]. In that conditions, the critical flux (J_{cs} and J_{cw}) cannot be defined strictly. Le Clech et al. [20] pointed out that a zero rate of transmembrane pressure (TMP) increase was never attained during the filtration of real and synthetic sewage. As the critical flux may not exist in such effluents, threshold flux can be suitable for the description of this system. Luo et al [21] validated and applied the concept of threshold flux in the treatment of dairy wastewater. Ochando-Pulido et al. [22,23] used the pressure-cycling method to analyse and identify the critical and threshold fluxes of an ultrafiltration membrane in several effluents. Based on the study of Field et al. [11,12], Stoller et al. [24] developed a threshold flux determination method by means of critical flux data conversion for a nanofiltration membrane. Stoller and Ochando-Pulido [25,26] then tried to merge threshold and critical flux concepts into a single concept: boundary flux. On the other hand, few studies have investigated the critical flux for irreversibility (J_{ci}), since it was defined by Bacchin et al [13]. Rayess et al. [27] measured J_{ci} during cross-flow microfiltration of wine by the square-wave method and noticed that no apparent critical flux for irreversibility can be determined in the range of tested pressures, as irreversible fouling always takes place from the beginning of the filtration, even at low pressures. A criterion ($R_{if}/R_m \leq 1$) was suggested to determine the “threshold flux” below which fouling remains acceptable. Based on these different works, the existence of a critical flux for irreversibility in real wastewater and its difference or potential relationship with threshold flux deserve to be investigated in further study.

In term of limiting flux, Bacchin et al. [28] tried to link limiting and critical fluxes and found from a modelling study that the limiting flux value is probably equal to 3/2 of the critical flux value in a colloidal system. Tang and Leckie [29] investigated the existence of limiting flux during nanofiltration of a synthetic solution of purified humic acid and concluded that the limiting flux is independent of the type of membrane while has a strong dependence on the feed water composition. The value of the limiting flux can be equal to, greater than or even less than the value of the critical flux, depending on feed water chemistry, theoretically. A series of NF and RO semi-aromatic and fully-aromatic polyamine membranes was tested for the filtration of a synthetic solution of purified humic acid. These different membranes exhibit a limiting flux that is equal to or greater than the critical flux. The opposite case, i.e. a limiting flux that is lower than the critical flux, has not been observed under the experimental conditions investigated by these authors. This research group also developed a limiting flux model for predicting the fouling behaviour of nanofiltration membrane [30]. The limiting flux model is capable of anticipating the effect of solution chemistry on the degree of humic acid fouling. Nevertheless, these studies were carried out in synthetic solution. In real wastewater matrices, the correlation between the limiting flux and the fouling mechanisms, the relationship between the limiting flux and the critical flux, and even the existence of a limiting flux, are still unclear.

The objective of this study was to apply the concepts of critical flux for irreversibility, threshold flux and limiting flux for fouling control in a cross-flow NF system which is used as a tertiary treatment of Membrane BioReactor effluent. Firstly, the critical flux for irreversibility was determined by the square-wave method [5,14]. Secondly, the threshold flux was estimated by critical flux data conversion [24]. Then, the existence of a limiting flux and the associated fouling mechanisms were investigated. Based on this study of flux behaviour and fouling mechanisms, strategies for fouling control through correlation between flux behaviour and fouling stages were proposed.

Table 1
Characteristics of the NE 70 membrane.^a

| MWCO (Da) | Max. temperature (°C) | Max. pressure (bar) | pH range | Surface charge | Rejection (%) |
|-----------|-----------------------|---------------------|----------|----------------|---|
| 250 | 45 | 41.4 | 3–10 | Negative | 40–70 (NaCl) 99.5 (MgSO ₄) |

^a Provided by the manufacturer: Woongjin Chemical Co., Ltd., considering specific conditions: 2000 mg/l MgSO₄, 25 °C, recovery 15%, pressure 0.5 MPa, pH 6.5–7.0; 1500 mg/l NaCl, 25 °C, recovery 15%, pressure 1 MPa, pH 6.5–7.0.

2. Material and methods

2.1. The water matrix

The study of fluxes of the NF membrane was performed in MBR effluent. The MBR system using ultrafiltration hollow fiber membranes (polysulfone 0.2 μm, reference: UF50 IMM, Polymem), was installed at Purpan, hospital located in Toulouse, France. This system was directly fed from the hospital's sanitary collection system. MBR effluent contains salts, organic matter (total organic carbon around 20 mg L⁻¹) and around 50 pharmaceuticals from 10 different therapeutic classes [31]. Water characteristics for filtration are shown in Section 2.3.1.

2.2. Membrane and nanofiltration apparatus

2.2.1. The NF membrane

A NE 70 nanofiltration membrane was examined in this study. NE 70 is a thin film composite polyamide membrane which consists of a polysulfone sub-layer and polyamide active layers. The characteristics of this membrane are shown in Table 1.

The membrane samples were stored in a preservative solution (NaHSO₃ 0.1 g L⁻¹, 4 °C). They were rinsed and cleaned with ultrapure water before filtrations and then installed in the cross-flow cell. Membrane compaction was performed by the filtration of ultrapure water at a TMP equal to 40 bar until a constant flow rate was obtained (around 3 h).

2.2.2. Nanofiltration apparatus

Nanofiltration experiments were conducted in a cross-flow filtration unit (Fig. 1). The volume of the feed tank (1) is 10 L. The membrane was installed into a stainless cross-flow cell (Sepa CF II, Osmonics) (3) in which the effective membrane area was 1.4 × 10⁻² m². Permeate can be collected in a vessel and weighed on scale (5) connected to a computer.

In order to simulate a feed-and-bleed continuous mode at Volume Reduction Factor (VRF) of 5, the filtration experiment was applied in two steps: firstly, the concentration mode and secondly the recircula-

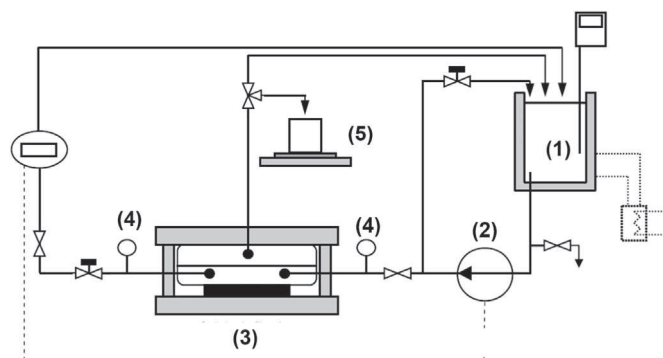


Fig. 1. Nanofiltration set-up, (1) feed tank, (2) high pressure pump, (3) cross-flow filtrate cell, (4) pressure monitor, (5) permeate tank and weighing scales.

tion mode. This VRF of 5, correlating to a 80% recovery, is within the range that would be applied in a full-scale system (30–90%) [32]. During the concentration mode, the retentate stream was recycled into feed vessel while the permeate was collected in the permeate tank. This mode corresponds to a transient state as the retentate concentration increased with filtration time. After the VRF reached 5, both retentate and permeate streams were recycled back to the feed tank, corresponding to a recirculation mode. The permeate flux was monitored during both filtration steps. The square-wave filtration method, developed by Espinasse et al. [5,14], was applied during the recirculation step. The cross-flow velocity was fixed to 0.2 m s⁻¹.

2.3. Analytical methods

2.3.1. Water matrix characterization

Concentrations of salts in MBR effluent were measured by ionic chromatography with an ICS 3000 system (Dionex, France). The injection volume was 25 μL and the column temperature was set to 30 °C. The concentrations of anions and cations were analysed with two columns (Thermo Scientific, Dionex): IonPac™ AS11 (mobile phase: 95% of 5 mM NaOH and 5% of 100 mM NaOH) and IonPac™ CS12 (mobile phase: CH₄O₃SO₃ 20 mM), respectively. Analytical errors for anions and cations ranged from 1.5% (K⁺) to 6% (SO₄²⁻). In this mobile phase, all H₃PO₄, H₂PO₄⁻, HPO₄²⁻ and PO₄³⁻ present in the solution were referred to as PO₄³⁻.

Total Organic Carbon (TOC) and Inorganic Carbon (IC) were measured with a TOC-VCSN instrument (Shimadzu). The concentration of inorganic carbon was measured after acidification and degassing. TOC was calculated from the difference between the total carbon and inorganic carbon. Chemical Oxygen Demand (COD) was determined by photometry using disposable test tubes (HI93754H-25 LR from HANNA Instruments) and a HACH DR/2400 photometer. After sample addition, test tubes were heated at 150 °C for 2 h and left to cool down at room temperature before measuring.

pH (A-32908-06, Labcor Technical Sales Inc.), conductivity (23226-523, VWR, Mississauga, Ontario) and UV₂₅₄ were measured. UV₂₅₄ was analysed using a spectrophotometer (Hewlett Packard 8452A) at a wavelength of 254 nm and using a quartz crystal cuvette with a 1 cm path length.

As previously explained, the MBR effluent from a hospital was used as the feed solution. This effluent was first concentrated 5 times in a concentration mode. Then the concentrated MBR effluent (recirculation solution) was filtrated at a constant retentate concentration (recirculation mode) to determine critical, threshold and limiting fluxes. The characteristics of the feed and the recirculation solution are shown in Table 2.

2.3.2. Membrane surface characterization

Virgin and fouled membranes after filtration were characterized after drying at 60 °C with a vacuum degree fixed at -0.8 bar (C3000 - XF020 France, Etuves).

ATR-FTIR spectra (Attenuated Total Reflection - Fourier Transform InfraRed spectroscopy) were obtained with a Thermo-Nicolet Nexus 670 apparatus (USA). The sample was placed on diamond crystal substrates and the analytical depth was approximately 2 μm. Virgin and fouled membranes were examined using ATR-FTIR spectra to identify membrane properties and fouling composition.

Surface morphology of membranes was examined with a Scanning Electron Microscope (SEM, Hitachi Tabletop Microscope TM-1000) interfaced with an Energy Dispersive X-ray (EDX) spectroscopy system (Thermo-Fisher). Membrane samples were sputtered with a thin layer of gold before SEM analysis for better contrast. The EDX analyses were carried out at the same time. Elemental analysis began from sodium. EDX measurements were performed at different locations of the membrane surface, in order to obtain a comprehensive elemental composition. Cross-sections of samples were prepared by cryo-fracture:

Table 2

Characteristics of MBR effluent and of the recirculation solution after the concentration step.

| Item | MBR effluent (NF feed) | Recirculation solution (after concentration step VRF =5) |
|---|------------------------|--|
| pH | 7.9 ± 0.2 | 7.6 ± 0.2 |
| COD (mg L ⁻¹) | 48 ± 3 | 112 ± 3 |
| TOC (mg L ⁻¹) | 19.6 ± 0.1 | 55.1 ± 0.1 |
| HCO ₃ ⁻ | 31.3 | 72.4 |
| UV ₂₅₄ | 0.46 | 1.37 |
| Na ⁺ (mg L ⁻¹) | 62.4 ± 1.1 | 77.9 ± 1.4 |
| K ⁺ (mg L ⁻¹) | 19.6 ± 0.3 | 25.4 ± 0.4 |
| Mg ²⁺ (mg L ⁻¹) | 3.3 ± 0.1 | 4.9 ± 0.1 |
| Ca ²⁺ (mg L ⁻¹) | 29.3 ± 0.9 | 43.2 ± 0.6 |
| Cl ⁻ (mg L ⁻¹) | 44.9 ± 0.9 | 41.6 ± 0.9 |
| NO ₃ ⁻ (mg L ⁻¹) | 7.1 ± 0.1 | 5.6 ± 0.1 |
| SO ₄ ²⁻ (mg L ⁻¹) | 26.8 ± 1.6 | 26.5 ± 4.6 |
| PO ₄ ³⁻ (mg L ⁻¹) | 8.0 ± 0.2 | 18.6 ± 0.6 |

membranes were first immersed in ethanol, then fractured in liquid nitrogen.

2.4. Critical flux for irreversibility determination: the square-wave method

The square-wave method, developed by Espinasse et al. [5,14], can be applied to distinguish the reversible part of fouling, due to osmotic pressure (R_{rf}), and the irreversible fouling (R_{if}) and then to obtain the accurate value of the critical flux for irreversibility (J_{ci}). The principle of this method is to alter the operating pressure with positive and negative variations, as shown in Fig. 2. The U steps correspond to the upper pressure steps while the L steps correspond to the lower pressure steps. By comparing the fluxes between steps L_n and U_{n-1} that correspond to the same applied pressure, the decrease in flux is associated to the fouling phenomenon that occurs at step U_n . If the flux is the same in steps U_{n-1} and L_n , the fouling is considered as totally reversible and the flux is below the critical value for reversibility. If the flux is not the same in steps U_{n-1} and L_n , the fouling is partly irreversible and the flux is above the critical value.

Calculation methods of irreversible fouling are based on the relationship between the flux and the fouling resistance. The flux can be described by an osmotic pressure model, as follows:

$$J = \frac{\Delta P - \Delta \pi}{\mu_p (R_m + R_{if})} \quad (1)$$

where J is the permeate flux, ΔP is the transmembrane pressure, $\Delta \pi$ is the osmotic pressure, μ_p is the viscosity of the permeate and R_m , R_{if} are the intrinsic hydraulic membrane resistance and the resistance due to the irreversible fouling, respectively.

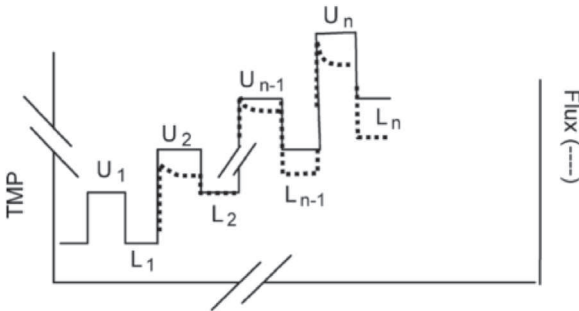


Fig. 2. Principle of the square-wave technique; pressure and flux vs time; upper and lower steps [5].

The osmotic pressure can also be incorporated into the reversible fouling resistance. In this case, the reversible resistance associated with the polarization concentration layer can be treated like a term of osmotic pressure:

$$J = \frac{\Delta P}{\mu_p (R_m + R_{if} + R_{rf})} = \frac{\Delta P}{\mu_p (R_m + R_f)} \quad (2)$$

where R_f is the total fouling resistance and R_{rf} is the resistance due to the reversible fouling, which is related to the polarization concentration.

According to the square-wave filtration method, the irreversible fouling resistance that appears at upper pressure step U_n can be reached by comparing the fouling resistance at steps L_n and U_{n-1} as follows:

$$r_{if,n} = R_{f|L_n} - R_{f|U_{n-1}} \quad (3)$$

where $r_{if,n}$ is the irreversible fouling resistance relative to the U_n step.

The value of the total R_{if} , at a given pressure step, is the sum of r_{if} at each previous pressure step.

$$R_{if} = \sum_n r_{if,n} \quad (4)$$

In the present study, the experimental procedure is shown in Fig. 3. The feed solution (MBR effluent) was firstly filtrated in concentration mode for 3 bar. Then the square-wave filtration method was applied in recirculation mode for 3, 5, 10, 15, 25, 35 bar. The operating pressure was increased or decreased when the change of flux over a 30 min period was less than 2% of the initial flux (pseudo stable flux was considered to be obtained). The irreversible fouling resistance was calculated for each pressure step.

2.5. Threshold flux estimation

The threshold flux was estimated by critical flux data conversion [24–26]. From critical flux theory, no permeability loss is observed when operating below the critical flux (J_c , Eq. (5)), while above J_c the permeability decreases according to the eq. (6) [11,24,33].

$$\frac{dL}{dt} = 0 \quad J_p(t) \leq J_c \quad (5)$$

$$\frac{dL}{dt} = B(J_p(t) - J_c) \quad J_p(t) > J_c \quad (6)$$

where L is the membrane permeability, B is a fitting parameter and $J_p(t)$ is the permeate flux at time t for the selected TMP.

Considering now the threshold flux, J_{th} , the permeability loss is unavoidable even below the threshold flux [12]. Operating below the threshold flux, the permeability loss is constant (Eq. (7)). Above J_{th} , the permeability loss follows Eq. (8) in which a and b are the fitting parameters including without distinction the contribution of reversible and irreversible fouling.

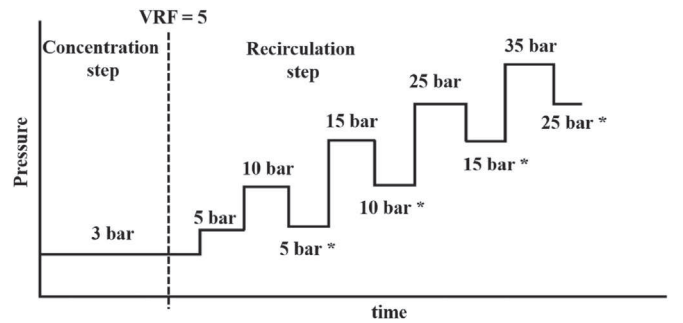


Fig. 3. Experimental procedure.

$$\frac{dL}{dt} = -a \quad J_p(t) \leq J_{th} \quad (7)$$

$$\frac{dL}{dt} = -a - b(J_p(t) - J_{th}) \quad J_p(t) > J_{th} \quad (8)$$

If a is equal to zero, the threshold flux J_{th} corresponds to the critical flux J_c . In both cases ($a = 0$ no fouling or $a \neq 0$ limited fouling), the filtration time during which $B = 0$ or $b = 0$ corresponds to long term fouling region [24].

A linear relationship between permeability loss (ΔL) and filtration duration can be derived from the integration of Eq. (7) between time point t_1 and t_2 upon operation below the threshold flux.

$$L(t_2) - L(t_1) = \Delta L = -a(t_2 - t_1) \quad (9)$$

Where the permeability at time t can be obtained by Eq. (10), where $TMP(t)$ is the transmembrane pressure at time t . In the present study, TMP is constant at time t_1 and t_2 .

$$L(t) = \frac{J_p(t)}{TMP(t)} \quad (10)$$

The estimation of threshold flux is based on the relationship between ΔL and Δt : as long as Eq. (9) applies, the permeability loss rate ($\Delta L/\Delta t$) is constant and equal to “ a ”, indicating that the operating conditions are below the threshold flux; as soon as $\Delta L/\Delta t$ value deviates from the “ a ” value, Eq. (9) is no longer valid and Eq. (8) should be applied; the operating conditions are then above the threshold flux. The permeability loss rate value at the boundary of the two conditions allows the determination of the threshold flux, the corresponding transmembrane pressure being the threshold pressure.

2.6. Accumulation of salts at each pressure step

During the recirculation step, both the permeate and the retentate were recycled into the feed tank. The accumulated mass of salts on the membrane surface and/or structure was calculated from the measurement of the concentration in the tank before and after circulation according to a mass balance as follows:

$$Q_{ads} = C_{be}V - C_{end}V \quad (11)$$

where C_{be} and C_{end} are the salt concentrations before and after recirculation, and V is the volume in the retentate loop, assumed to be constant. The concentration of salts was measured, when a steady state was reached at each pressure step.

3. Results

3.1. The determination of the critical and limiting fluxes

The critical flux for irreversibility was determined by the square-wave method subsequent to the experimental procedure shown in Fig. 3. The evolution of fluxes and transmembrane pressure during the filtration of MBR effluents has been shown in Fig. 4.

During the concentration step (at 3 bar), the flux slightly increased at the beginning and then quasi stabilized at value $62 \text{ L h}^{-1} \text{ m}^{-2}$ when $VRF = 5$. After the recirculation step, the applied pressures ranged from 3 to 35 bar. Taking into account the experimental error in flux measurements of 8%, when the applied pressure was maintained at 3 bar the flux value was considered as stable, meaning that no significant fouling occurred at this pressure. Consequently, the flux at the beginning of the operation at 3 bar was considered to be the initial permeability for the filtration of MBR effluent in further calculations of fouling resistances. As expected, when the pressure was increased to 5 bar, the flux immediately increased. After that, a decline in flux during the recirculation time was observed until it reached a stable value again at $67 \text{ L h}^{-1} \text{ m}^{-2}$. The similar increase of flux with pressure increase was observed with an applied pressure of 10 bar. However, the

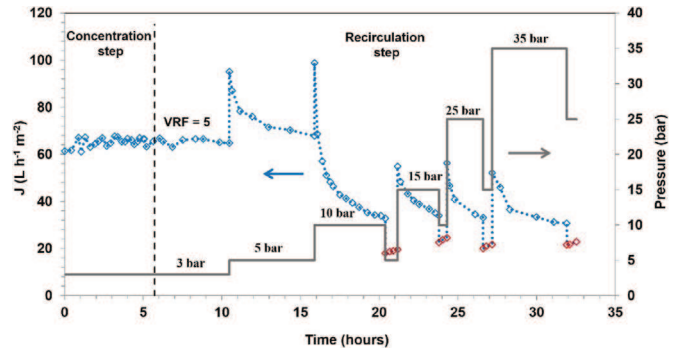


Fig. 4. Flux evolution following the experimental procedure over time: – the continuous line represents the applied pressure; the dotted line denotes the permeate flux.

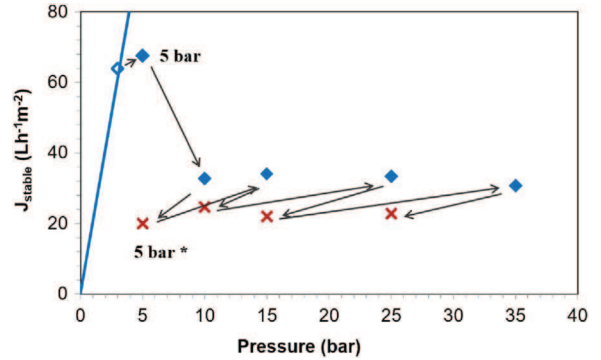


Fig. 5. Pseudo stable flux vs applied pressure following the procedure shown on Fig. 3; initial permeability —; first pseudo stable flux reached at a certain pressure ◆; hollow one is without decrease of flux over time and the bold points are in presence of flux decrease during filtration; stable flux reached after negative variation of pressure ✕ (5 bar*, 10 bar*, 15 bar*, 25 bar* in Fig. 3); Arrows represent the positive or negative variations of pressure.

pseudo stable flux reached a new value of $33 \text{ L h}^{-1} \text{ m}^{-2}$. The procedure continued with an increase in applied pressure step by step until 35 bar. With the applied pressures of 15, 25 and 35 bar, the fluxes also stabilized at about $33 \text{ L h}^{-1} \text{ m}^{-2}$, after an increase of flux at the beginning of a positive variation of pressure.

Fig. 5 shows the relationship between the pseudo-stable flux and the transmembrane pressure. The continuous line represents the Darcy law according to the initial permeability, determined at 3 bar. The difference between the pseudo stable flux in MBR effluent reached at 3 bar ($62 \text{ L h}^{-1} \text{ m}^{-2}$) and the pure water flux measured on the virgin membrane at 3 bar ($60 \text{ L h}^{-1} \text{ m}^{-2}$) is within experimental error of flux measurements (8%); which confirms that no significant fouling (even by organic adsorption) occurs at this pressure which is then below the critical one. The arrows represent the positive or negative variations of pressure. The crosses represent the stable fluxes reached after a negative variation in pressure (namely 5 bar*, 10 bar*, 15 bar*, 25 bar* in Fig. 3).

When the applied pressure increased from 3 to 5 bar, it can be suggested that fouling occurred, since a flux decline with time was observed at this pressure (Fig. 4). One can expect a strong form of critical flux has been exceeded at 5 bar, because the pseudo stable flux started to deviate from pure water permeability straight line solely from this pressure (no deviation was observed at 3 bar) [11]. By comparing the value of fluxes of two steps having the same transmembrane pressure before and after 10 bar (respectively 5 bar and 5 bar*), one can observe that the stable flux decreased drastically. This means that fouling was partially irreversible at 10 bar. As a consequence, the initial flux under 10 bar was greater than the critical flux for irreversibility (J_{ci}). This result was confirmed by the evolution of total fouling resistances at each pressure step (calculated from Eq. (2)) shown in

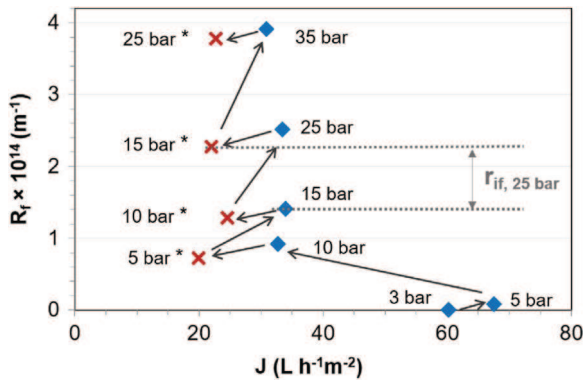


Fig. 6. Evolution of the total fouling resistance (R_f) v.s. pseudo stable flux obtained at each pressure according to the square-wave method procedure. R_f after a positive variation of pressure \blacklozenge ; R_f after a negative variation of pressure \blacktimes . Arrows represent the variations of pressure during the procedure.

Fig. 6. First, the total fouling resistance at 5 bar* (as L_n step in Fig. 2) was greater than the total fouling resistance at 5 bar (as U_{n-1} step in Fig. 2). This means that after an increase in pressure (10 bar as U_n step in Fig. 2) the resistance cannot recover its value at the lower pressure, this remaining resistance (the difference of fouling resistance at 5 bar and 5 bar*) being the irreversible part. At the same time, the total resistance decreased from 10 bar (as U_n step in Fig. 2) to 5 bar* (as L_n step in Fig. 2), which refers to the reversible fouling at 10 bar. Similarly, the total fouling resistance formed at 15 bar, 25 bar, 35 bar includes reversible and irreversible components.

Thanks to the application of the square-wave filtration method (Section 2.4), the value of the critical flux for irreversibility J_{ci} was then estimated. The total irreversible fouling resistance at a given pressure was calculated from data in Fig. 6 and Eqs. (3) and (4). For example, the irreversible fouling resistance at 25 bar step ($r_{if, 25 \text{ bar}}$) was shown in Fig. 6 when the total irreversible fouling resistance R_{if} at 25 bar was deduced from the sum of the r_{if} at each previous step.

The irreversible fouling resistances from 10 bar to 35 bar were calculated and plotted in Fig. 7(a). Due to the fact that no fouling was observed at 3 bar, the irreversible fouling resistance is expected to be zero at this pressure. We can define a critical pressure (between 3 and 5 bar) below which irreversible fouling is not observed. In contrast, when the pressure exceeds this critical value, irreversible fouling occurs. The linear relationship between irreversible fouling resistance and transmembrane pressure, allowed us to estimate this critical value to be 3.4 bar.

The critical flux for irreversibility was evaluated at around $70 \text{ L h}^{-1} \text{ m}^{-2}$ from the critical pressure and initial permeability, as shown in Fig. 7(b). Moreover, it can be observed that the stable flux went below the expected value computed from the initial permeability, when the initial flux exceeded the critical flux. Initial flux at 5 bar was around $97 \text{ L h}^{-1} \text{ m}^{-2}$ (consistent with the pure water flux of the virgin membrane at 5 bar of $100 \text{ L h}^{-1} \text{ m}^{-2}$) then exceeding the critical value of $70 \text{ L h}^{-1} \text{ m}^{-2}$ (see Fig. 4). The pseudo stable flux at 5 bar was marginally less than the critical flux value. A further increase in pressure (10–35 bar) does not lead to an increase in the pseudo stable flux. Therefore, this behaviour may indicate that the limiting flux was reached. The value of the limiting flux around $33 \text{ L h}^{-1} \text{ m}^{-2}$ was surprisingly less than that of the critical flux one.

3.2. Estimation of threshold flux

In the present study, no fouling took place at 3 bar, which was confirmed by a permeate flux after 10 h of cross-flow filtration almost equals to pure water flux measured on the virgin membrane at 3 bar. At this pressure, permeability loss rate was then 0, thus the fitting parameter “a” is 0. In that case we can consider that our system

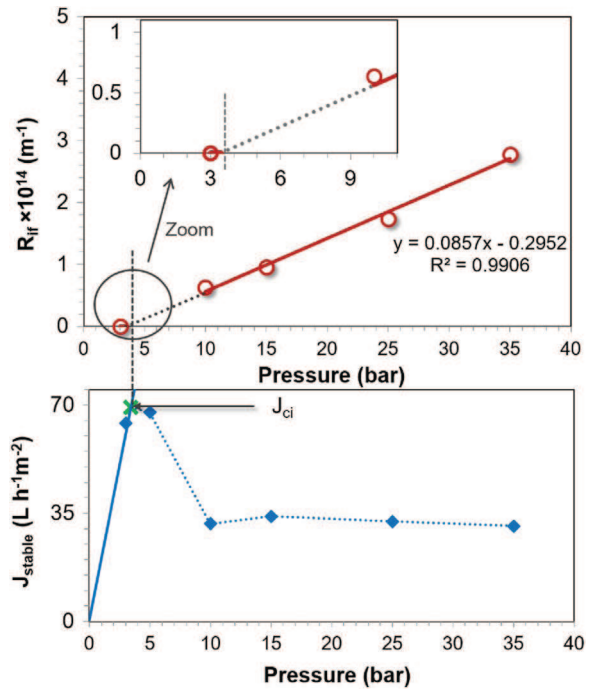


Fig. 7. Evolution of (a) irreversible fouling resistance, (b) pseudo stable flux as function of pressure and the location of the critical flux for irreversibility \blacktimes .

exhibits a critical flux, according to the boundary flux conception [25,26]. As some researchers have observed both critical flux and threshold flux on the same system [21], we decided to go further in the analysis of the results in order to determine if above this critical flux it was possible to characterize a low fouling region and a high fouling region, the transition between them corresponding to the threshold flux (Section 2.5). Fig. 8 presents for each tested transmembrane pressure the permeability loss rate ($\Delta L/\Delta t$). From 5–35 bar, the permeability loss rate was calculated from Eqs. (9) and (10) using the flux data at the last 2 h when the flux reduction was linear.

The permeability loss rate at 3 bar is equal to zero as previously discussed; whereas its value increases dramatically upon operation at 5 bar. In the pressure range from 15 to 35 bar, the calculated permeability loss rate decreases again to achieve almost zero at 35 bar. However, in this pressure range the fouling is so drastic and the permeate flux so low (Fig. 4) that this rate of permeability loss doesn't correspond to operating conditions that we could advise to industrialist. As a consequence, according to the results reported in Fig. 8, we were unable to determine a region with a constant and limited permeability loss rate that could correspond to long term fouling region before a severe fouling occurs. However, if our system exhibits a threshold pressure, this one could be expected in the range 3.4 (critical pressure) – 10 bar and its determination would have

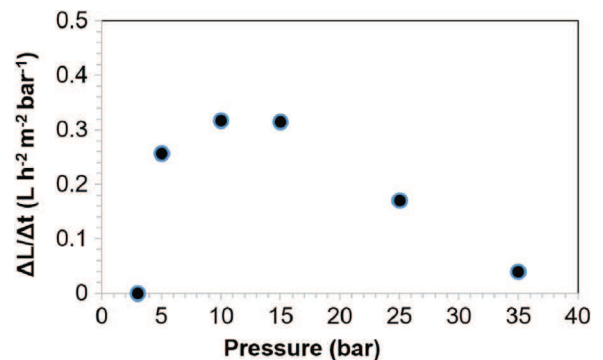


Fig. 8. Permeability loss rate obtained at different pressure values.

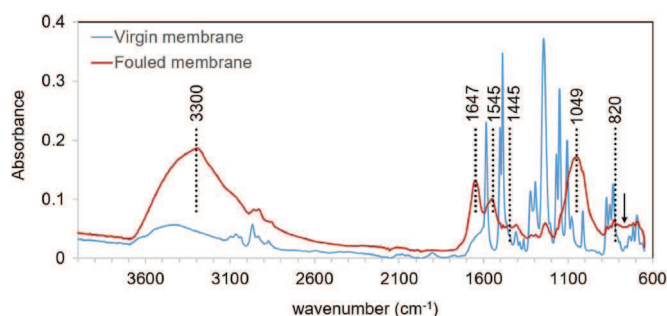


Fig. 9. ATR-FTIR spectra of fouled and clean membrane.

required more filtrations in this pressure range.

3.3. Fouling layer characterization

In order to explain fluxes behaviour reported in Fig. 7, the chemical composition of the foulant layer will be identified in this section. First, two analytical methods were applied to identify the fouling layer components: ATR-FTIR and SEM-EDX. Then, the accumulation of salts was quantified from a mass balance calculation.

3.3.1. ATR-FTIR

ATR-FTIR spectra of virgin and fouled membranes for wave lengths ranging from 4000 to 750 cm^{-1} are shown in Fig. 9. The relative penetration depth during analysis was 2 μm . The spectra of the virgin NE70 membrane is characteristic of an active polyamide top-layer, with a polysulfone sub-layer, in addition to a coating layer (top of the active polyamide layer) [34]. Concerning the polyamide top-layer, the peak at 1631 cm^{-1} , corresponding to the amide band was observed. Polysulfone groups were detected at 1587, 1504, 1488, 1365, 1350–1280, 1245, 1180–1145, and 830 cm^{-1} . Peaks observed in the range of high wave numbers (3700–2700 cm^{-1}) were associated with

the coating layer which probably consists of O–H groups [34].

For the fouled membrane after filtration, almost all of the characteristic peaks of the polysulfone sub-layer and the polyamide layer cannot be observed, and are replaced by the foulants peaks. Even the peaks related to the coating layer (top of the active polyamide layer) are weak and difficult to recognize. These observations proved that the thickness analysed by the equipment (2 μm) allowed us to characterize the fouling layer, the membrane material itself being unreachable on the fouled membrane. The absorbance at 3300, 1545 and 1647 cm^{-1} indicates the presence of organics with amide bonds, such as proteins, on the fouling layer. N–H peaks around 3300 and 1545 cm^{-1} , together with a peak at 1647 cm^{-1} associated with the stretching vibration of C=O, indicates the presence of colloidal reflected functional groups of primary and secondary amides [35,36]. The absorbance band around 1445 cm^{-1} , that corresponds to calcium carbonate, is very weak. This suggests that carbonate was not the main foulant [37]. OH stretching and the Si–OH group vibration are observed at 830–1110 cm^{-1} and 3700–3200 cm^{-1} [38]. The vibration of the Si–O bond appears at 791 and 754 cm^{-1} [39]. In particular, the absorbance around 820 cm^{-1} is associated with the symmetric bond stretching vibration of the Si–O–Si network [40]. These signals imply the presence of colloidal silica fouling. The wide and intensive peak at 1049 cm^{-1} is suggested to be characteristic of phosphate (including PO_4^{3-} , HPO_4^{2-} , H_2PO_4^-) and crystallized calcium phosphate [41–43], which probably indicates the presence of calcium phosphate scaling on the fouled membrane surface.

3.3.2. SEM-EDX analysis

SEM analysis was performed on membrane surfaces and cross-sections before and after filtration, to observe the fouling layer and to characterize the layer's structure and morphology (Fig. 10). Comparing the micrographs of virgin and fouled membranes, a variety of foulants evenly distributed over the entire membrane surface, was observed. As shown on images (b) and (d), homogenous "bright" points are superimposed on the dark foulant layer. According to the FTIR analysis and

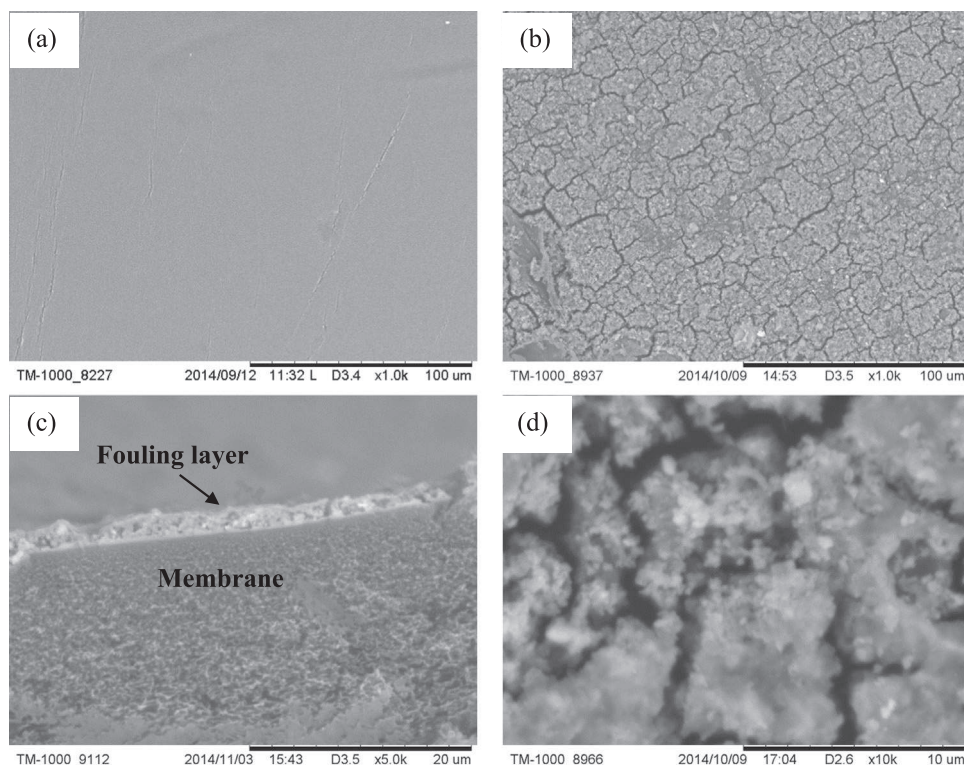


Fig. 10. SEM images of virgin and fouled membrane: (a) Clean membrane surface 1000 \times ; (b) Fouled membrane surface 1000 \times ; (c) Section of fouled membrane 5000 \times ; (d) Fouled membrane surface 10000 \times .

the morphology of the foulant, the “bright” points could be surface crystallization and the dark layer is possibly the colloidal silica and organic foulant. The (c) image shows the cross-section of a fouled membrane. The bottom and porous part is the polysulfone sub-layer. Above it, the thin, bright layer (around 0.2 μm) is the active polyamine layer. The fouling layer, which is around 2 μm , is on the top. Its thickness is relatively uniform.

Combined with SEM, EDX was used to analyse the elemental compositions on the surface of virgin membranes and fouling layer. EDX sampled different locations on the virgin and fouled membrane surfaces. The virgin membrane surface is composed of 100% sulphur, while the elements detected on the fouled membrane surface are various. The weight percentages of the main elements are shown in Table 3. These analyses allow us to verify that the main elements responsible for fouling are calcium and phosphate.

The presence of calcium is usually referred to as inverse-solubility of salts (CaCO_3 , $\text{CaSO}_4 \cdot x\text{H}_2\text{O}$, calcium phosphate etc.) which can precipitate and form a hard fouling layer on the membrane often referred to as scaling [4]. For phosphorus, it has been found in previous studies [44] that the presence of phosphate can cause the formation of a viscous gel with silica, resulting in a hard cement-like layer irreversibly bonded to the membrane surface. In addition, phosphates can precipitate with calcium and induce the common calcium phosphate scaling, as some autopsies on membrane elements have shown [4,45]. Considering the analysis of FTIR, the absorbance peak of calcium carbonate is relatively weak, while the band associated with calcium phosphate is strong. We can confirm that calcium phosphate scaling is the main fouling type, together with colloidal silica and organic fouling. From the EDX analysis, the distribution of calcium and phosphate is quite uniform, which indicates that the scaling is homogeneous on the membrane surface and structure.

Based on FTIR and SEM analyses, the composition of the fouling layer indicates the combined presence of colloidal silica and organic fouling and calcium phosphate inorganic fouling, after filtration of MBR effluent subsequent to the experimental procedure shown in Fig. 4.

3.3.3. Accumulation of ions

As discussed in Section 2.6, we assume that the difference in ions concentration in the tank before and after circulation corresponds to ions accumulated on the membrane surface and structure during the recirculation step. The accumulated mass can be calculated after each pressure step.

In order to verify the role of calcium and phosphate in irreversible fouling resistance, Fig. 11 shows the evolution of both the irreversible fouling resistance and the accumulated mass of Ca^{2+} and PO_4^{3-} on the membrane surface versus the applied pressure.

Ca^{2+} , which had slightly accumulated during the concentration step (results not shown) was partially released from the membrane surface and structure to the retentate, resulting in a negative accumulated mass at 3 bar. Desorption of Ca^{2+} probably occurred between the concentration step and the recirculation step at 3 bar when the filtration was stopped for the night. Considering that no flux decline was observed

Table 3
EDX elements' analysis at different locations on fouled membrane surfaces.

| Samples | 1 | 2 | 3 | 4 | Average |
|--------------|-----------------------------------|------|------|------|---------|
| Main element | Weight % from different locations | | | | |
| Silicon | 0 | 2.4 | 0 | 0 | 0.6 |
| Phosphorus | 24.6 | 26.6 | 23.2 | 28.8 | 25.8 |
| Calcium | 50.3 | 53.7 | 57.4 | 54.6 | 54 |
| Chromium | 4.8 | 5.0 | 0 | 7.6 | 4.3 |
| Iron | 4.7 | 4.9 | 0 | 0 | 2.4 |
| Sulphur | 0 | 0 | 19.3 | 9.8 | 14.6 |

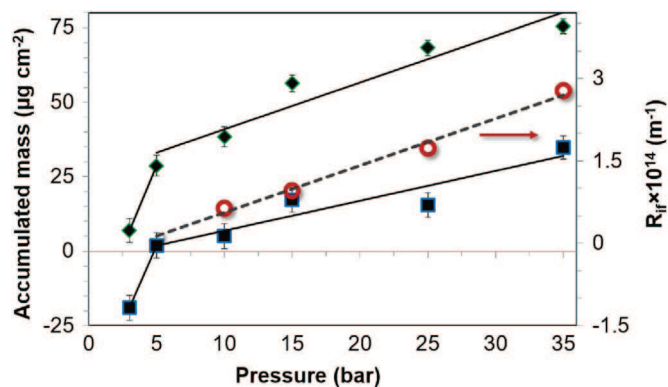


Fig. 11. Evolution of irreversible fouling resistance \circ and ions accumulated mass: Ca^{2+} \blacksquare , PO_4^{3-} \blacklozenge on the membrane with increasing applied pressure during recirculation step.

during the filtration at this pressure, we can say that the quantity of salt adsorbed in these conditions was negligible and didn't act as an additional resistance. During the recirculation operation from 5 to 35 bar, the accumulation of Ca^{2+} and PO_4^{3-} exhibited good linear relationships with applied pressure. For this range of pressures, while Ca^{2+} and PO_4^{3-} accumulation increases, irreversible fouling resistance is enhanced. Thus, the accumulation of Ca^{2+} and PO_4^{3-} on the membrane surface and structure is strongly correlated to the occurrence of irreversible fouling. This observation is in accordance with the analysis by FTIR and SEM-EDX.

4. Discussion

During the filtration of the MBR effluent in a cross-flow NF system, no measurable fouling took place for the transmembrane pressure 3 bar (Fig. 4), indicating that the concept of critical flux is applicable in the present study. In a first step, the critical flux for irreversibility ($70 \text{ L h}^{-1} \text{ m}^{-2}$) and the corresponding critical pressure (3.4 bar, Fig. 7) were determined. Above these critical conditions, operating points corresponding to low fouling rate and sufficiently high flux during a long term operation were then considered. The threshold pressure has been searched but not determined due to a lack of experimental data. However, if it exists, it would have been located in the range 3.4–10 bar. Operating in this pressure range should lead to acceptable fouling rate and flux decline. For example, at 5 bar, a high initial flux ($97 \text{ L h}^{-1} \text{ m}^{-2}$) was followed by a relatively acceptable permeability loss around $0.26 \text{ L h}^{-2} \text{ m}^{-2} \text{ bar}^{-1}$ until a steady state was reached at $67 \text{ L h}^{-1} \text{ m}^{-2}$.

During the filtrations conducted in the pressure range 10–35 bar, a limiting flux was observed ($33 \text{ L h}^{-1} \text{ m}^{-2}$, see Fig. 7) which is much less than the critical flux. Through the characterization of the fouling layer after filtration (Section 3.3), we found that this flux behaviour is the result of a combination of colloidal silica and organics fouling and calcium phosphate scaling.

These results seem to show that there are two fouling stages leading to various flux behaviours in the same water matrix. The early fouling stage at 5 bar led to a pseudo stable flux, the value of which is close to J_{ci} . The permeability loss rate was acceptable in this fouling stage as we discussed above. These observations are in accordance with the results of Tang and Leckie [29]. In their work, the membrane was fouled by humic acid in synthetic solution using the same membrane as in the present study (the NE 70). Fouling at 5 bar could then be linked to colloidal silica and organics accumulation.

On the other hand, severe fouling stage at higher pressures (10–35 bar) resulted in a pseudo table flux (limiting flux) that is rather lower than J_{ci} . According to the results obtained during the fouling layer characterization and considering that calcium phosphate scaling can lead to severe flux decline during nanofiltration [7,46], the sharp

decline of the pseudo stable flux from 67 to 33 L h⁻¹m⁻² (at 5–10 bar) can be attributed to the occurrence of scaling.

A method which can correlate the flux behaviour and fouling stages is discussed. The increase in operating transmembrane pressure plays an essential role in the development of scaling. After a positive variation of pressure, flux temporarily increases, expectedly causing a more serious polarization concentration and greater ionic concentration on the membrane surface. In case the ionic concentration on the membrane surface is beyond the solubility limit of scaling salts (calcium and phosphate in the present study), scaling occurs. Moreover, in the present study we assumed that the early state fouling layer on the membrane surface could play an important role in inducing scaling. This early state fouling layer possibly serves as a “nucleation site” for scaling [47,48], resulting from an enhancement of salt concentration polarization on the membrane surface, by hindering the back-diffusion of salt ions [49]. Consequently, one assumption is that there is a transition fouling layer which can be characterized by a transition fouling resistance above which scaling occurs and fouling turns from a limited to a severe state. Corresponding to this transition resistance, we propose in this section the estimation of the permeability before scaling L_s . Once the observed permeability is below L_s , scaling starts to form.

There are two scaling pathways in membrane systems: bulk (homogeneous) and surface (heterogeneous) crystallizations [50–52]. In the present study, the bulk crystallization pathway was not observed (scaling did not occur at pressures of 3–5 bar). Moreover, the 0.2 m s⁻¹ experimental velocity is in the range of a low velocity, which is in favour of surface crystallization, according to the study of Lee and Lee [53]. We therefore consider that in our system, the main scaling mechanism is surface crystallization. Scaling due to surface crystallization can be described by the surface blockage fouling mechanism. The membrane surface may be blocked by the lateral growth of crystals meaning that only a permeate flux can pass through the membrane surface that is not occupied by crystals [53,54]. Fig. 12 shows the foulants on the membrane surface with and without scaling.

The permeability just reached before scaling occurrence, defined as permeability before scaling L_s , corresponds to a flux J_s only affected by the colloidal silica and organics fouling. J_s can be described as:

$$J_s = \frac{L_s}{\mu_p} \Delta P = \frac{L_s}{\mu_p (R_m + R_s + R_{rf})} \quad (12)$$

where μ_p and R_m are the viscosity of the permeate (Pa s) and the membrane resistance (m⁻¹) respectively, R_s is the irreversible transition fouling resistance, which is related to colloidal silica and organics fouling; R_{rf} is the reversible fouling resistance that corresponds to the concentration polarization.

When a scaling layer is formed according to a surface blockage mechanism, assuming that the area occupied by crystals is completely

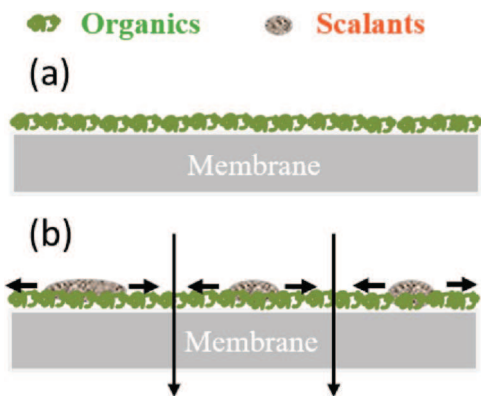


Fig. 12. Foulants on the membrane surface (a) without and (b) in presence of scaling.

impermeable, the flux could be expressed as follows:

$$J_a = \frac{\Delta P}{\mu_p R_m} \frac{A_{free}}{A_m} = \frac{\Delta P}{\mu_p R_m} \frac{A_m - A_{oc}}{A_m} \quad (13)$$

where J_a is the permeate flux estimated from the surface blockage model, A_m is the membrane active area, A_{free} and A_{oc} are the membrane areas unoccupied and occupied by surface crystals, respectively. With the assumption that the thickness of the crystal formed on the membrane surface is constant [55], the A_{oc} could be defined as follows:

$$A_{oc} = \beta m_s \quad (14)$$

where β (m² mg⁻¹) is the area occupied per mass unit and m_s (mg) is the weight of the scaling layer formed directly on the membrane surface.

When scaling occurs (at a pressure of 10–35 bar), the flux declines, due to the colloidal fouling layer, the relative concentration polarization and surface crystals hindering. This flux can be represented by combining Eqs. (12) and (13):

$$J_b = \frac{L_b}{\mu_p} \Delta P = \frac{\Delta P}{\mu_p (R_m + R_s + R_{rf})} \frac{A_m - A_{oc}}{A_m} \quad (15)$$

where J_b is the flux combining two fouling mechanisms (colloidal silica and organics in addition to scaling).

According to Eqs. (12) and (15), the reduction of permeability by surface crystallization superimposed on the colloidal silica and organics layer can be expressed as follows:

$$\frac{L_b}{L_s} = \frac{J_b}{J_s} = \frac{A_m - A_{oc}}{A_m} \quad (16)$$

where L_b is the permeability combining two mechanisms (colloidal silica and organics fouling and scaling); L_s is the permeability before scaling (corresponding to R_s). When the permeability is lower than L_s , scaling occurs.

Combining Eqs. (14) and (16), L_b can be written as:

$$L_b = -\frac{L_s}{A_m} \beta m_s + L_s \quad (17)$$

This function provides the relationship between the permeability before scaling L_s , the permeability combining two fouling mechanisms L_b , and the weight of the scaling layer directly formed on the membrane surface. m_s can be estimated from the mass balance during the filtration. L_b can be obtained from the filtration with square-wave filtration method. Thus, from this relationship, the permeability before scaling L_s can be evaluated.

Considering that a fouling combining organics deposition and scaling occurs from 10 to 35 bar, the observed permeability during the filtrations at 10, 15, 25 and 35 bar corresponds to L_b . This permeability can be estimated from the pseudo stable flux at a certain pressure and the stable flux after the negative variation of applied pressure according to the experimental procedure (Fig. 13). For example, L_b at 10 bar can be evaluated from the slope of the straight line joining the pseudo stable fluxes at 10 bar and 5 bar*. Moreover, due to this procedure, the determined value of L_b is not impacted by the concentration polarization and reversible fouling. Consequently, L_s which is calculated from L_b doesn't include reversible fouling. The relationship between the permeability before scaling (L_s) and the fouling transition resistance (R_s) can be then written as follows:

$$L_s = \frac{1}{\mu_p (R_m + R_s)} \quad (18)$$

As shown in Fig. 13, the first permeability of 3.42 L h⁻¹ m⁻² bar⁻¹ (the left-hand dashed slope) is L_b after reaching the pseudo stable flux at 10 bar. The following values are the permeability reached at 15, 25 and 35 bar.

In sections 3.2.3, we showed that the increase in irreversible fouling resistance (from 10 to 35 bar) is directly linked to the accumulation of

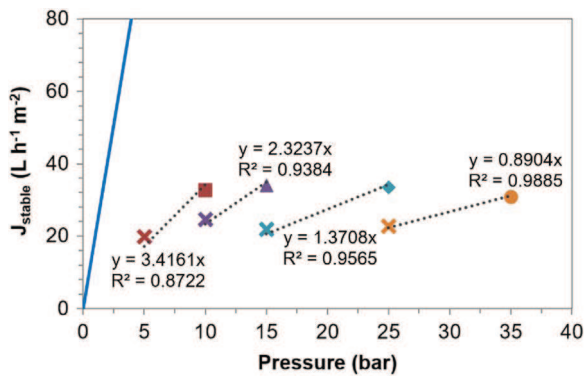


Fig. 13. Experimental permeability measurement, the continuous line - is initial permeability of membrane; bold points are the first pseudo stable fluxes reached at a certain pressure; the crosses are the stable fluxes reached after negative variation of pressure (5 bar*, 10 bar*, 15 bar*, 25 bar* in Fig. 4).

calcium and phosphate on the membrane surface. As a consequence, to estimate the weight of the scaling layer on the membrane surface (m_s), the accumulated masses of Ca^{2+} and PO_4^{3-} were calculated for the 10–35 bar pressure range and shown in Table 4.

Fig. 14 illustrates a linear relationship between the accumulated mass of Ca^{2+} and PO_4^{3-} and L_b permeability combining two fouling mechanisms. From this linear relationship, the permeability before scaling L_s is calculated as $3.85 \text{ L h}^{-1} \text{ m}^{-2} \text{ bar}^{-1}$.

With the initial permeability and the estimated value of permeability before scaling (L_s), a working diagram showing the flux behaviour versus the operating pressure was proposed. This diagram defines three separated regions: a non-fouling region A, a region of fouling by organics B and a scaling region C. Fig. 15 shows these three regions and the experimental value of fluxes as a function of transmembrane pressure. The non-fouling region A is enclosed on the left by the average initial permeability. The region B in between pure water permeability and permeability before scaling (L_s) corresponds to conditions for which colloidal silica and organics fouling was observed. Finally, fluxes in the region C on the right-hand side of L_s (region of scaling) indicates that scaling occurs. The location of L_s is in accordance with the experimental flux behaviour. Scaling occurs between 5 and 10 bar pressure in this study.

The estimation of L_s and of the critical flux then enabled a NF working diagram to be drawn which can be a new tool to optimise operating conditions. During the continuous nanofiltration process, filtration at an initial flux lower than the critical flux for irreversibility (region A) can be a strategy to totally prevent the membrane from irreversible fouling. When the flux is observed in region B colloidal fouling can be expected. We assume, even if we were unable to determine it, that the threshold flux should be located in this region corresponding to a relatively acceptable fouling rate. If the flux approaches the scaling region, some actions such as adjusting feed pH, adding anti-scaling substances and even cleaning the membrane could be considered to prevent the development of scaling, which induces low limiting flux. It should be noted that the permeability before scaling is obtained for a specific feed water composition, effects of water matrix and the verification of the model will be discussed in subsequent publications.

Table 4
Experimental permeability and accumulated mass of ions on the membrane.

| Pressure (bar) | L_b ($\text{L h}^{-1} \text{ m}^{-2} \text{ bar}^{-1}$) | $m_s = m_s(\text{Ca}^{2+}) + m_s(\text{PO}_4^{3-})$ (mg) |
|----------------|---|--|
| 10 | 3.42 | 1.79 |
| 15 | 2.33 | 6.01 |
| 25 | 1.37 | 7.43 |
| 35 | 0.89 | 11.16 |

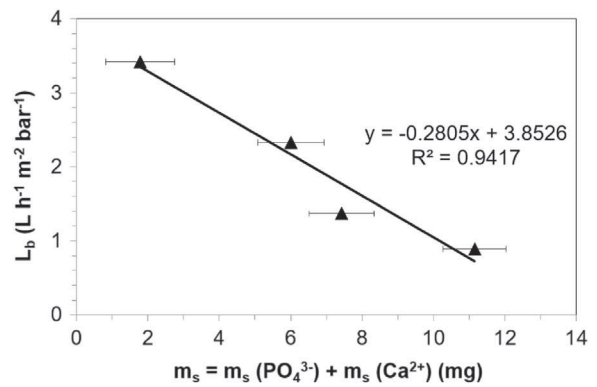


Fig. 14. Relationship between Ca^{2+} and PO_4^{3-} accumulated mass on membrane and L_b permeability combining two fouling mechanisms.

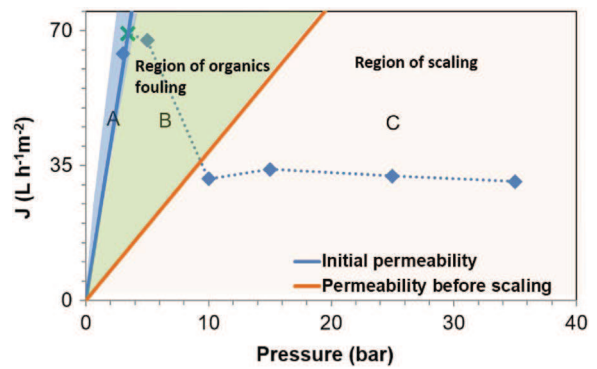


Fig. 15. Experimental pseudo stable flux as a function of transmembrane pressure and the location of the critical flux for irreversibility * in working diagram of NF considering the initial permeability and the permeability before scaling.

5. Conclusions

During the filtration in a cross-flow NF system of hospital wastewater after MBR (MBR permeate), no fouling was detected at low transmembrane pressure (3 bar), indicating that the concept of critical flux is applicable in the present study. The critical flux for irreversibility ($70 \text{ L h}^{-1} \text{ m}^{-2}$) and the corresponding critical pressure (3.4 bar) were then determined using the square-wave filtration method. Above these critical conditions, operating points corresponding to low fouling rate and sufficiently high flux during a long term operation were considered. The threshold pressure and related flux were searched by critical flux data conversion. Our results suggest, even if a value for the threshold pressure could not be determined, that it would be located in the range 3.4–10 bar. Operating in this pressure range should lead to acceptable fouling rate and flux decline. More pressure stages such as 1 bar intervals 1 between 3 and 5 bar could be interesting to obtain the exact value of threshold pressure in further work.

When the initial fluxes exceeded the critical flux for irreversibility, two pseudo stable flux behaviours were observed, versus the applied pressure, due to different fouling mechanisms in the complex water matrix used in this study. On one hand, a limiting flux value which is lower than that of the critical flux was obtained for pressures equal to, or higher than 10 bar. The low pseudo stable flux in this pressure range was linked to a severe fouling stage, combining the effect of colloidal silica and organics fouling and calcium phosphate scaling. On the other hand, in low pressure range of 5–10 bar, before reaching the limiting flux, the pseudo stable flux tends to the value of the critical flux. In these conditions limited fouling occurs, involving solely colloidal silica and organics accumulation. Based on these results, a method which can correlate flux behaviours and fouling mechanisms was discussed and permeability before scaling (L_s) was defined and calculated. The estimation of critical flux and permeability before scaling allows the

fouling stages of the membrane to be defined and quantified.

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