Evaluation of different cleaning agents used for cleaning ultra filtration membranes fouled by surface water

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Abstract

This paper reviews the published literature on potential membrane fouling components, available cleaning agents and possible interactions between cleaning agents and fouling components. It also lists the cleaning models available in the literature, and evaluates the advantages and disadvantages of these models. Based on this outcome, a new cleaning model is proposed to capture cleaning dynamics for 10 different cleaning agents, varying from acidic, alkali and oxidizing to sequestering agents and detergents that were used to clean dead-end ultra filtration membranes fouled by surface water. The model is effectively fitted to the experimental data of the different cleanings. Two criteria are subsequently introduced to quantify the overall cleaning effect of a cleaning agent in terms of cleaning rate and cleaning effectiveness. For membranes fouled by surface water with high organic content it was found that caustic-and oxidizing cleaning agents give the best overall cleaning results.

Keywords: Ultra filtration; Surface water; Chemical cleaning; Modeling

1. Introduction

Membrane ultra filtration is increasingly used as a total or intermediate surface water purification technique. In the last 15 years, production of polymeric membrane fibers as well as ceramic membranes became economically attractive. Membrane performance, however, is influenced by fouling. In addition to hydraulic cleaning, chemical cleaning in which cleaning agents are used is a common way to remove foulants. Although chemical cleaning is a useful way to restore membrane performance, membrane cleaning procedures are often based on rules of thumb and are usually conservative.

This paper presents a brief literature review on potential fouling components, cleaning agents and possible interactions between fouling components and cleaning agents. It also addresses the modeling efforts reported in the literature. From a practical viewpoint, a component-balance-based model will be used in the further evaluation of different cleaning agents. The model is fitted to the data collected for the different cleaning agents, and subsequently the instant cleaning rate and overall cleaning effectiveness can be calculated. The cleaning rate and effectiveness form good evaluation criteria for the selection of a proper cleaning agent. The proposed model and selection criteria are suitable to evaluate different cleaning agents. However, it should be noted that some cleaning procedures are more sensitive to measurement errors than others. For modeling data, the response from turbidity measurements and pH measurements is collected. In some cases, the turbidity response was found to be small, compared to the measurement noise, resulting in considerable modeling errors.

For Twente canal water (characterized as water with reasonable amount of organic content, DOC = 9 mg/l, TOC = 9.5 mg/l), caustic and oxidative cleaning procedures are most effective. Also commercial blends can be used to clean the membrane. Commercial blends often contain detergents. It was found that detergents should be used in low concentrations (around 0.05–0.1%), otherwise negative effects occur, such as TMP increase, instead of a decrease.

The main objective of this research is to evaluate the cleaning model for different cleaning agents and to develop criteria to value the overall effect of a cleaning agent.
2. Theory

2.1. Fouling components

Surface water is known to contain three potential membrane fouling categories, namely microbial content (bacteria, viruses, etc.) [7], organic content (natural organic matter) [8–13] and inorganic content (minerals) [14,16–18]. Sometimes a fourth category is added, namely: colloidal content (e.g. clay). The physicochemical mechanisms of fouling components attaching to the membrane surface are complex and poorly understood. Physical adsorption, electro-static charge and bridge-formation are fouling mechanisms frequently reported in the literature. An overview of the research conducted in the field of membrane chemical cleaning in relation to membrane fouling is presented in Table 1. The plus and minus sign in the fouling section of the table indicate if a certain type of fouling was present (+) or absent (−). The symbols in the chemical cleaning agents part of the table mean: (+) tests were performed with this type of cleaning agent, (−) no tests were performed with this cleaning agent, (0) this type of cleaning agent found to be most effective and (*) no information available. Table 1 gives an overview of what kind of foulants were found on membrane surfaces and which cleaning agents were accordingly found to be most effective removing the foulant. It shows that for surface water and waste water the feed is a complex mixture of the three basic fouling categories. Organic fouling has been reported in many publications, in addition to that, was found that caustic cleaning procedures are often most effective.

2.2. Cleaning agents

Cleaning agents are substances that can be used to effectively remove matter that is not an integral part of the membrane surface. Chemical cleaning effectiveness depends on the cleaning flow, the concentration of the cleaning agent and/or the cleaning temperature. In principle six basic cleaning agent categories can be distinguished [14](see Table 2). In addition to the basic categories, many commercial blends are available. Commodity agents (such as sodium hydroxide or hydrochloric acid) are generally less expensive than the commercial blends, however, blends can be dosed in lower concentrations.

2.3. Interactions between fouling components and cleaning agents

Weis et al. [17] believes that a cleaning agent can affect fouling materials present on a membrane surface in three ways: (i) foulants may be removed, (ii) morphology of foulants may be changed (swelling, compaction) and/or (iii) surface chemistry of the deposit may be altered, such that the hydrophobicity or charge is modified. The cleaning agents may react chemically or physically with the foulant, to weaken the cohesion forces between the foulants themselves and the adhesion between the foulants and the membrane surface. The possible reactions between foulant and cleaning agent are: hydrolysis, peptization, saponification, solubilisation, dispersion (suspension) and chelation [14], as presented in Table 3. It is further noted that biocides (cleaning agents used to treat microbial fouling) are grouped under ‘Disinfectants’. For a more extended discussion on chemical cleaning in relation to treatment of biofouling, the reader is referred to Ref. [15].

In addition to positive effects between cleaning agent and foulant, also negative effects might occur if an inappropriate cleaning agent is chosen, membrane performance could even be adversely affected. Suppliers of cleaning agents advise in which fouling case, which cleaning agent should be used (cleaning solutions are generally applied within the range of 0.5–1.5%). In Table 4 an overview of preferred cleaning solution in relation to membrane fouling is presented.

According to Mohammadi [19], the best choice of cleaning agent selection is based on a knowledge of the feed and the foulant composition and is in most cases performed by trial and error. Other decision criteria are based on the safety, the stability, the price and the waste treatment of the cleaning agent. Especially the impact of the cleaning agent on the environment and the membrane are important factors in the selection of a cleaning agent according to Maartens et al. [5].

2.4. Modeling of membrane chemical cleaning

2.4.1. Model by Bird and Bartlett

Bird and Bartlett [1] developed a model to predict the change in flux during caustic cleaning of a flat plate cross flow micro filtration membrane fouled by whey proteins. Starting with Darcy’s equation, which relates the flux J to the total resistance R_t as:

\[ J = \frac{\Delta P}{\mu R_t} \]  

(1)

where \(\Delta P\) is the trans-membrane pressure and \(\mu\) is the viscosity. The total resistance is modeled by a resistance in series approach:

\[ R_t = R_m + R_c + R_p + R_{cp} \]  

(2)

where \(R_m\) is the membrane resistance, \(R_c\) is the hydraulic resistance as a result of fouling at the membrane surface, \(R_p\) the resistance as a result of in-pore fouling and \(R_{cp}\) is the so-called resistance as a result of concentration polarization. In Bird and Bartlett’s model the resistance as a result of concentration polarization disappears on the release of TMP and the membrane resistance is a material property. The resistance decrease during caustic cleaning at the membrane surface is modeled by a second order decrease, according to:

\[ \frac{dR_c}{dr} = -k_c R_c^2 \]  

(3)

And the in-pore resistance is correlated to the effective pore diameter by Carman–Kozeny’s equation:

\[ R_p = \frac{36\tau(1-\varepsilon)^2l}{\varepsilon^3 d_c^2} \]  

(4)

where \(\tau\) is the tortuosity, \(\varepsilon\) the porosity and \(l\) is the length of the pore. The effective pore diameter \(d_c\) is based on the diameter of the particle that is blocking the pore. During caustic cleaning the
<table>
<thead>
<tr>
<th>Researcher</th>
<th>Application</th>
<th>Membrane type</th>
<th>Organic fouling</th>
<th>Inorganic fouling</th>
<th>Microbial fouling</th>
<th>Acidic cleaning</th>
<th>Caustic cleaning</th>
<th>Detergent cleaning</th>
<th>Sequestering cleaning</th>
<th>Oxidative cleaning</th>
<th>Enzymatic cleaning</th>
<th>Blend cleaning</th>
</tr>
</thead>
<tbody>
<tr>
<td>Li et al. [2]</td>
<td>Gluth. Ferm. Broth</td>
<td>UF/PS</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>+</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>–</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Madaeni and Mansourpanah [3]</td>
<td>Boiler water</td>
<td>RO/PA</td>
<td>–</td>
<td>+</td>
<td>–</td>
<td>+</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Maartens et al. [5]</td>
<td>Paper effluent</td>
<td>UF/PES</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>+</td>
<td>0</td>
<td>0</td>
<td>–</td>
<td>+</td>
<td>0</td>
</tr>
<tr>
<td>Mohammadi et al. [6]</td>
<td>Waste water</td>
<td>RO/PA</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Lee et al. [19]</td>
<td>Surface water</td>
<td>UF/PES</td>
<td>+</td>
<td>–</td>
<td>+</td>
<td>0</td>
<td>+</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Liikanen et al. [20]</td>
<td>Surface water</td>
<td>NF/PPA</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>0</td>
<td>+</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Mohammadi [21]</td>
<td>Waste water</td>
<td>UF/PS</td>
<td>+</td>
<td>–</td>
<td>+</td>
<td>0</td>
<td>+</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Madaeni et al. [22]</td>
<td>WPC</td>
<td>RO/PA</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>0</td>
<td>–</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Zhu and Nystrom [23]</td>
<td>BSA/LYS</td>
<td>UF/PES</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Munoz et al. [24]</td>
<td>BSA/WPC</td>
<td>UF/PES</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>0</td>
<td>–</td>
</tr>
<tr>
<td>Bartlett [25]</td>
<td>WPC</td>
<td>MF/CER</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>+</td>
</tr>
<tr>
<td>Pavlova [26]</td>
<td>Waste water</td>
<td>UF/PAN</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>0</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Sungpet et al. [27]</td>
<td>Textile effluent</td>
<td>NF/*</td>
<td>–</td>
<td>+</td>
<td>–</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Mo and Huang [28]</td>
<td>Raw water</td>
<td>MF/PE</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>0</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Veza and Rodriguez-Gonzalez [29]</td>
<td>Waste water</td>
<td>RO/*</td>
<td>–</td>
<td>+</td>
<td>–</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Chen et al. [30]</td>
<td>Waste water</td>
<td>UF/PES</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>+</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>0</td>
</tr>
<tr>
<td>Gwon et al. [31]</td>
<td>Ground water</td>
<td>UF/PA</td>
<td>+</td>
<td>+</td>
<td>–</td>
<td>0</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Wei [32]</td>
<td>Sulphite liquor</td>
<td>UF/PES</td>
<td>+</td>
<td>+</td>
<td>–</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>+</td>
</tr>
<tr>
<td>Lim and Bai [33]</td>
<td>Waste water</td>
<td>MF/PVDF</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>0</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
</tbody>
</table>

(+): present/tested; (−): not present/tested; (0): best tested; (*): no information available.
Table 2
Basic cleaning agent categories and examples

<table>
<thead>
<tr>
<th>Caustic</th>
<th>Acidic</th>
<th>Sequestering/complexing</th>
<th>Detergent/surfactant</th>
<th>Enzymatic</th>
<th>Oxidizing/disinfec- tant</th>
<th>Blend</th>
</tr>
</thead>
<tbody>
<tr>
<td>NaOH</td>
<td>HCl</td>
<td>EDTA</td>
<td>Alkyl sulphate</td>
<td>Alpha-CT</td>
<td>NaClO</td>
<td>4Aquaclean (Aquacare)</td>
</tr>
<tr>
<td>KOH</td>
<td>HNO₃</td>
<td>SDS</td>
<td>CP-T</td>
<td>Peroxidase</td>
<td>KMnO₄</td>
<td>Divos (Diverseylever)</td>
</tr>
<tr>
<td>NH₃OH</td>
<td>H₂SO₄</td>
<td>CTAB</td>
<td>H₂O₂</td>
<td>Triclean (triton)</td>
<td></td>
<td>Triclean (triton)</td>
</tr>
<tr>
<td>Oxalic</td>
<td></td>
<td>Citric</td>
<td></td>
<td></td>
<td></td>
<td>Ultrasil/Aquaclean (Ecolab)</td>
</tr>
</tbody>
</table>

Table 3
Possible interactions between cleaning agents and foulants

<table>
<thead>
<tr>
<th>Caustic</th>
<th>Acidic</th>
<th>Sequestering</th>
<th>Oxidizing/disinfection</th>
<th>Detergent/surfactant</th>
<th>Enzymatic</th>
</tr>
</thead>
<tbody>
<tr>
<td>Organic</td>
<td>Hydrolysis</td>
<td>Hydrolysis/saponification</td>
<td>Dispersion</td>
<td>Oxidizing</td>
<td>Chelation</td>
</tr>
<tr>
<td>Inorganic</td>
<td>Solubilization/chelation</td>
<td>Solubilization/chelation</td>
<td>Dispersion</td>
<td>Oxidizing</td>
<td>Chelation</td>
</tr>
<tr>
<td>Microbial</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>Disinfecting</td>
<td>–</td>
</tr>
</tbody>
</table>

where $d_e = \left( d_0 - 2 \left( \frac{\delta}{k_1 t + \delta} \right) (k_2 t + \delta) \right)$

Particle size is decreasing which means that the effective pore diameter is increasing. Bird and Bartlett propose the following swelling–compaction relationship:

2.4.2. The Li model

Li et al. [2] modeled the chemical cleaning of dead-end ultra filtration membranes fouled by fermentation broth of glutamic acid by a simple kinetic model, assuming that the flux will increase during cleaning according to a first order equation:

$$\frac{dJ}{dt} = kJ$$

where $k$ is a rate constant related to temperature according to the Arrhenius equation:

$$k = k_0 \exp \left( -\frac{E_A}{RT} \right)$$

The advantage of the Li model is that it is simple, the protocol to determine the model parameters is straightforward and it has only two model parameters. It includes a variable to control the cleaning process (temperature) and the Arrhenius term implies cleaning is a kinetic process. However, the experimental verification of the model was not done extensively. In addition, the first-order Li model may not be able to capture cleaning behavior well, if operational setting of the cleaning process change, and dynamics become more complex.

2.4.3. Model by Zondervan

The model proposed by Zondervan et al. [34] was tested on the caustic cleaning of dead-end ultra filtration membranes fouled by surface water and is based on two component balances:

$$\frac{dx_W}{dt} = -k'J(x_{W\infty} - x_{W}) + r''_W$$

$$\frac{dx_C}{dt} = k'J(x_{C,\infty} - x_C) + nC r''_W$$

Table 4
Preferred cleaning

<table>
<thead>
<tr>
<th></th>
<th>Proteins</th>
<th>Glucanes</th>
<th>Pigments</th>
<th>Minerals</th>
<th>Hydrophobes</th>
<th>Starch</th>
<th>Tanins</th>
<th>Pectin</th>
<th>Fat</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alkaline</td>
<td>0</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Acidic</td>
<td>0</td>
<td>–</td>
<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Surfactants</td>
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<td>0</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>+</td>
<td>0</td>
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<td>0</td>
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<tr>
<td>Soil dispersers</td>
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<td>0</td>
<td>+</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Complexing agents</td>
<td>0</td>
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<td>+</td>
<td>+</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Enzymes</td>
<td>+</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>+</td>
<td>–</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Oxidizers</td>
<td>+</td>
<td>+</td>
<td>–</td>
<td>–</td>
<td>0</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>0</td>
</tr>
</tbody>
</table>

(+): positive cleaning effect; (−): negative cleaning effect; (0): positive, neither negative effect. Reference: Ecolab Membrane Cleaning Congress 2005, Dusseldorf.
constant, $x_{W,\infty}$ is the fouling state at infinite cleaning time and $n_C$ is a pseudo-stoichiometric constant, $r_W$ is the cleaning rate, defined as:

$$r_W'' = -k'' n_C (x_W - x_{W,\infty})$$  \hspace{1cm} (10)

where $k''$ is a cleaning rate parameter which may be dependent on the cleaning temperature.

The fouling state is defined in terms of trans-membrane pressure:

$$x_W(t) = \frac{\Delta P(t)}{\Delta P_0}$$  \hspace{1cm} (11)

where $\Delta P_0$ is the trans-membrane pressure at the beginning of the cleaning and $\Delta P(t)$ is the trans-membrane pressure during cleaning. As trans-membrane pressure during cleaning cannot directly be measured, cleaning dynamics are captured by turbidity measurements. The cleaning concentrate is collected at the membrane outlet, and turbidity is subsequently measured. The following relationship between the trans-membrane pressure during cleaning and the measured turbidity is suggested:

$$\Delta P(t) = 1 - \epsilon_{\infty} \int_0^t \frac{E}{\int_0^t E \, dt}$$  \hspace{1cm} (12)

$E$ is the turbidity measured at the outlet of the membrane unit and $t_c$ is the duration of the chemical cleaning procedure. The integral of the turbidity reflects the amount of irreversible fouling that has been removed. The integral is normalized to one and scaled by means of the trans-membrane pressure-based cleaning effectiveness:

$$\epsilon_{\infty} = \frac{\Delta P_0 - \Delta P_{\infty}}{\Delta P_0 - \Delta P_m}$$  \hspace{1cm} (13)

where $\Delta P_{\infty} \approx \Delta P(t_c)$ and $\Delta P_m$ is the trans-membrane pressure of a clean membrane determined at a specified clean water flux. The cleaning effectiveness is the ratio of the decrease in trans-membrane pressure during a chemical cleaning procedure and the increase in trans-membrane pressure during a production cycle.

The cleaning state can be determined by means of pH measurements and is defined according to:

$$x_C(t) = \frac{10^{-\text{pH}(t)}}{10^{-\text{pH}^*}}$$  \hspace{1cm} (14)

where pH(t) is the pH measured at the outlet of the membrane unit and pH* is a reference pH.

To characterize the effect of a cleaning agent we introduce two important evaluation criteria, the instant cleaning rate, defined as:

$$r_i = \frac{dx_W}{dt} \bigg|_{t=0}$$  \hspace{1cm} (15)

And the overall cleaning effectiveness, $\eta$:

$$\eta = \frac{x_W(0) - x_W(t_c)}{x_W(0) - x_{W,\infty}}$$  \hspace{1cm} (16)

where $x_W(0)$ is the fouling state at the beginning of the cleaning and $x_W(t_c)$ is the fouling state at the end of the cleaning process.

The model by Zondervan is simple, has three model parameters, namely $k'$, $k''$ and $n_C$, that can be determined by a straightforward experimental protocol (based on pH and turbidity measurements). The model also includes control variables as cleaning flow and cleaning agent concentration. Even temperature may be incorporated, by determining the relationship between temperature and the cleaning rate constant. The model structure is based on component balances and is suitable for process optimization. From the model two intuitive criteria can be derived easily for quantification of the overall cleaning effect.

3. Materials and methods

3.1. Experimental setup

Fouling and subsequent cleaning experiments were performed with a laboratory scale dead-end ultra filtration unit. The setup consisted of a 1 in. ultra filtration membrane module (100 fibers of 30 cm in a module housing with diameter = 1 in.), a filtration and a backwash pump, flow meters to monitor fluxes, thermometers to record temperature and correct trans-membrane pressure. Pressure meters to determine trans-membrane pressure. In this setup, trans-membrane pressure can only be measured during filtration. During a cleaning experiment, concentrate is collected in fractions, at the permeate outlet, and from these fractions pH and turbidity ($E$) are determined. A simplified diagram of the setup is shown in Fig. 1.

Fig. 1. Simplified scheme of the experimental setup. During a fouling experiment, surface water is pumped from the feed tank through the membrane and is collected in the permeate tank. During a backwash, permeate water is flushed in opposite direction through the membrane and is drained from the system at the concentrate outlet. During chemical cleaning, chemicals are flushed through the membrane and drained at the concentrate outlet.
Twente canal surface water is used to foul the UF membranes. In Table 5 water quality parameters are summarized.

No flocculant is added to the surface water. The water has an average pH of 7.5, an average conductivity of 580 μS/cm, an average O₂ concentration of 8–10 mg/l and an average turbidity of 2–20 FNU. The average water temperature was approximately 15°C. Twente canal water is known for its high organic content (DOC/TOC values around 10 mg/l). Surface water is stored in a 20 l feed tank, which is refreshed every 5 h. Ten different cleaning agents were used to clean the fouled membrane. The membrane used in the experiments is a Norit-Xiga FSU hollow fiber ultra filtration module with a membrane surface of $A_m = 0.07 \text{ m}^2$.

### 3.2. Experimental procedure

The experimental protocol followed for fouling and cleaning of the membrane is shown in Fig. 2. First a filtration procedure with a flux of 100 l/(h m²) is performed until the temperature corrected trans-membrane pressure of the membrane has increased from $P_m = 350 \text{ mbar}$ to approximately 500 mbar. Subsequently a backwash procedure is executed, in which a volume of 500 ml permeate is flushed back with a backwash flux of 150 l/(h m²). Then a volume 1000 ml of the selected cleaning agent is flushed back through the membrane with a flux of 150 l/(h m²) at a concentration of 0.05 mol/l. (For the commercial blends, the instructions for cleaning solution preparation were applied.) At the permeate outlet, the concentrate is collected and pH and turbidity are measured. After the chemical cleaning, remaining chemicals are flushed out of the system with an additional backwash of 500 ml performed at a flux of 150 l/(h m²). Trans-membrane pressure is measured before and after the chemical cleaning procedure by a short filtration at a flux of 100 l/(h m²) with clean water. The processed cleaning agent volume can be converted to time $t$ (min) by $t = V/(A_m J)$, where $A_m = 0.07 \text{ m}^2$ and $J = 150 \text{ l/(h m}^2)$ = 2500 ml/(min m²). The system has a dead time of $\tau_d = 1.14 \text{ min}$ which corresponds to a dead volume of approximately $V_d = 200 \text{ ml}$.

Trans-membrane pressure and turbidity measurements are used to monitor the fouling state, while the pH is used as an indicator for the cleaning agent state. In Zondervan et al. [34] the proposed relationship between the model variables ($x_W$ and $x_C$) and the measurable variables (trans-membrane pressure, pH and turbidity) is explained in more detail.
Fig. 3. Modeling and experimental results for different cleaning agents; the upper figures are the fouling states and the lower figures are the cleaning agent states. From left to right: hydrochloric acid (a), sulphuric acid (b), citric acid (c), 4Aquaclean Fer 12 (d) (imminodisuccinic salt) and P3 Aquaclean Sal (e).
Fig. 4. Modeling and experimental results for different cleaning agents; the upper figures are the fouling states and the lower figures are the cleaning agent states. From left to right: P3 Ultrasil 70 (phosphoric acid) (a), Kleen MTC 411 (sodium borate) (b), P3 Ultrasil 115 (potassium hydroxide) (c), sodium hypochlorite (d) and hydrogen peroxide (e).
4. Results and discussion

4.1. Modeling results for different cleaning agents

Ten different cleaning agents were evaluated, covering the overall range of available cleaning agents. In Table 6 an overview of the tested cleaning agent is presented. Also the estimated model parameters and the fits are listed in Table 6. It is noted that $n_C$ is evaluated as a fit parameter and does not have physical relevance.

The experimental data and estimated models are graphically represented in Figs. 3 and 4, showing the fouling state $x_W$ (upper row) and cleaning agent $x_C$ (lower row) state as a function of time. In general the cleaning model fit is 70% or more. In some cases, however, the model fit is not adequate, especially for citric acid (64%), hydrogen peroxide (60.3%) and Kleen MTC (51%). In these cases, the measurement signal (turbidity) was small compared to the measurement noise, leading to increased error. It is also noted that concentrations of cleaning agents containing detergents (such as the Ultrasil products) should be kept minimal. Overdosing with detergents leads to an increase in membrane resistance instead of a decrease. The pores may be covered with a film of detergent, resulting in an increased resistance. However, the loss of performance is not permanent, and the membrane resistance may be restored by extensive rinsing with clean water. Lower dosings (in the range of 0.05–0.1%) were tried until the desired cleaning effects took place.

In Figs. 5 and 6 the instant cleaning rate and overall cleaning effectiveness are calculated for the different cleaning agents, according to Eqs. (15) and (16). A reference cleaning with sodium hydroxide was performed to correct for changes in water quality. In general, caustic and oxidizing agents were found to have higher instant cleaning rates and overall cleaning effectiveness. This is in good agreement with the fact that the surface water is characterized as containing higher concentrations of organic content.

Fig. 5. Instant overall cleaning rate and overall cleaning effectiveness for different cleaning agents.

In Fig. 6 overall cleaning effectiveness is plotted versus the instant cleaning rate for the different cleaning agents. The figure shows that faster cleaning may yield higher cleaning effectiveness.

5. Conclusions

The cleaning model and experimental protocol proposed by Zondervan et al. can be effectively used to capture cleaning dynamics for a considerable number of cleaning agents. For two cleaning agents the model fit was less adequate due to sensitivity to measurement error. Data was collected from turbidity and pH measurements. The response for the turbidity measurements was in some cases small, compared to the measurement noise, leading to this error. By introduction of the instant cleaning rate $r_i$ and the overall cleaning effectiveness $\eta$, the overall effect of a cleaning agent can be effectively quantified. For membranes fouled by Twente canal surface water (characterized as water with high organic content) it can be concluded that caustic and oxidizing cleaning agents are the most effective.

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Nomenclature

List of symbols

- $A_m$: membrane surface ($m^2$)
- $d_0$: initial pore diameter ($mm$)
- $d_e$: effective pore diameter ($mm$)
- $E_A$: activation energy (J/mol)
$J$ cleaning flux

$k$ rate constant (1/min)

$k_0$ Arrhenius constant (1/min)

$k_{1,2}$ swelling/compaction rate constant (1/min)

$k'$ flushing rate constant (1/min)

$k''$ cleaning rate constant (1/min)

$l$ pore length (mm)

$n_C$ pseudo stoichiometric constant

$\Delta P$ trans-membrane pressure (mbar)

$r_i$ instant cleaning rate (1/min)

$r_W$ cleaning rate (1/min)

$R$ gas constant (J/(mol K))

$R_c$ resistance as result of surface fouling (1/m)

$R_{cp}$ resistance as result of concentration fouling (1/m)

$R_m$ membrane resistance (1/m)

$R_p$ resistance as result of in-pore fouling (1/m)

$R_t$ total resistance (1/m)

$t$ time (min)

$T$ temperature (K)

$V$ volume (m$^3$)

$\lambda_C$ cleaning agent state

$\lambda_{C,\text{in}}$ cleaning agent state at the inlet

$\lambda_W$ irreversible fouling state

$\lambda_{W,\infty}$ irreversible fouling state at infinite cleaning time

Greek symbols

$\delta$ swelling/compaction parameter (mm)

$\varepsilon$ porosity

$\eta$ cleaning effectiveness

$\mu$ viscosity (Pa s)

$\tau$ tortuosity

$\tau_d$ dead time (min)

References


