Rejection rate and mechanisms of drugs in drinking water by nanofiltration technology

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ABSTRACT
Nanofiltration (NF) technology is a membrane-based separation process, which has been pervasively used as the high-effective technology for drinking water treatment. In this study, a kind of composite polyamide NF thin film is selected to investigate the removal efficiencies and mechanisms of 14 trace drugs, which are commonly and frequently detected in the drinking water. The results show that the removal efficiencies of most drugs are quite high, indicating the NF is an effective technology to improve the quality of drinking water. The removal efficiencies of carbamazepine, acetaminophen, estradiol, antipyrine and isopropyl-antipyrine in ultrapure water are 78.8 ± 0.8%, 16.4 ± 0.5%, 65.4 ± 1.8%, 71.1 ± 1.5% and 89.8 ± 0.38%, respectively. Their rejection rates increase with the increasing of their three-dimensional sizes, which indicates that the steric exclusion plays a significant role in removal of these five drugs. The adsorption of estradiol with the strongest hydrophobicity has been studied, which indicates that adsorption is not negligible in terms of removing this kind of hydrophobic neutral drugs by NF technology. The removal efficiencies of indomethacin, diclofenac, naproxen, ketoprofen, ibuprofen, clofibric acid, sulfamethoxazole, amoxicillin and bezafibrate in ultrapure water are 81 ± 0.3%, 86.3 ± 0.5%, 85.7 ± 0.4%, 93.3 ± 0.3%, 86.6 ± 2.5%, 90.6 ± 0.4%, 59.7 ± 1.7%, 80.3 ± 1.4% and 80 ± 0.5%, respectively. For these nine drugs, their rejection rates are better than the above five drugs because they are negatively charged in ultrapure water. Meanwhile, the membrane surface presents the negative charge. Therefore, both electrostatic repulsion and steric exclusion are indispensable in removing these negatively charged drugs. This study provides helpful and scientific support of a highly effective water treatment method for removing drugs pollutants from drinking water.

Keywords: Drinking water, drugs, Nanofiltration, Removal efficiency, Removal mechanism

1. Introduction

In recent years, it has attracted the increasing attention on the presence of drugs and their remains in the aquatic environment [1-6]. A wide range of drugs are frequently detected in the surface and ground water due to their incomplete assimilation by human and animals as well as poor removal efficiencies in wastewater treatment plants [7-8]. The long-term exposure of drugs may lead to potential threats to human health and water ecological security [9-11]. Therefore, the study on the high-effective removal techniques of drugs from water is an urgent need. In recent decades, nanofiltration (NF), a low pressure-driven membrane process, has been gaining much attention due to its high rejection rates to many low molecule weight organic pollutants [12-15]. NF was a costly technology due to the short life cycle, easily fouling and high operation cost in the past. However, with the development of research and the improved manufacturing technology, the cost of NF has decreased significantly, which makes NF become a promising application in water treatment [15-19]. For the quality and safety of supply drinking water, it is expected that NF will be widely employed into drinking water treatment in the near future.

Recently, some researchers investigated the removal performance of drugs by NF technology [20-25]. Three major drug removal mechanisms are proposed, including steric exclusion, electrostatic repulsion and adsorption. Steric exclusion is considered as one of the removal mechanism in removing all kinds of drugs, especially for drugs having no charge, which are called as neutral...
drugs. The molecular weight cut-off (MWCO) of NF and the molecular weight of drug are employed to evaluate the removal efficiency. However, it is inadequate and inaccurate to adopt the MWCO to evaluate the removal performance of NF as well as use the molar weight to weigh the size of drug molecule [20]. In addition, increasing environmental awareness coupled with more stringent regulation standards has triggered various industries to challenge themselves in seeking appropriate wastewater treatment technologies [21]. Note that the removal efficiency is not consistent with the MWCO and drug molecular weight in NF. Regarding this, this paper is focused on the influence of the molecular dimensions of three-dimensional structure on the rejection of non-absorption neutral drugs by NF [22-23]. The removal mechanism can be further elaborated instead of only focus on the NF MWCO and drug molecular weight. Besides, hydrophobic interaction is also noticeable between some neutral drugs and NF in the experiments. However, hydrophobic interaction may cause the different consequences, which need more researchers’ efforts to investigate [24-26]. On the one hand, hydrophobic interaction causes the membrane fouling and makes the membrane pore size become smaller, which can increase the removal efficiency. On the other hand, the hydrophobicity makes the drug absorbed onto and partition into membrane surface, which will facilitate the drug diffusion to go through the NF membrane. The previous research showed that most of NF membranes were negatively charged at the condition of neutral or alkaline solution due to the dissociation of functional groups (e.g., carboxylic acid groups) on the membrane surface [27]. The amount of negative charges on membrane surface increases with the increase of the pH value of the solution due to the acid group on membrane surface is gradually dissociated. Thus, the amount of negative charges on membrane surface would be changed with the pH value of the solution [20-23, 28-29]. Note that drugs containing acid functional groups (e.g., the carboxylic group, phenolic hydroxyl group and et al.) have several different forms based on the pH value of feed solution. These drugs are regarded as the negatively charged drugs. Usually, negatively charged drugs exist in two kinds of specifications (negative ionic and neutral forms) when the dissociation constant of the drug (pKa) is smaller than the pH value of solution. Therefore, the electrostatic repulsion can occur between negatively charged drugs and negative NF membrane in certain solution conditions. As known, the removal mechanism for negatively charged drugs is completely different from that of the neutral drugs. Thereby, the pH value of the solution plays a significant role in removing charged drugs by NF. Some studies have reported that the role of electrostatic repulsion is very crucial in removing the charged organic solutes [20-22, 27-29]. Based on our literature survey, the role of steric exclusion for removal the charged organic solutes is still not clear. So, it is interesting to know the removal mechanisms for the charged drugs. The cost of NF membrane water treatment plant varies and is dependent on the production capacity, type of treatment involved, design criteria, characteristics of land and building, etc. [30-31]. NF technology will have the tremendous market requirement in the near future with the increase of water quality standard and the decrease of NF cost. The rejection rate for NF membrane are the most important aspect for the design of membrane filtration plant as it is a direct measure of productivity, operating pressure (energy requirements) and amount of membrane required (membrane area) [32]. Hence, studying the rejection rate for drugs in the drinking water is capable to provide acceptable economic estimation for the water treatment by NF membrane.

Fourteen drugs (i.e. carbamazepine, indomethacin, diclofenac, anti-pyrene, naproxen, ketoprofen, ibuprofen, clofibric acid, acetaminophen, sulfamethoxazole, amoxicillin, estradiol, bezafibrate and isopropyl-antipyrine) were selected as the objective water pollutants due to their high exposure levels in water environment. These drugs also stand for a broad range of physical and chemical properties (i.e., size, pKa, hydrophillicity, and water solubility) and their different rejection rates are expected to achieve by the NF. In summary, this paper will employ a popular commercial NF membrane to remove those selected fourteen drugs and the removal efficiency and mechanism will be systematically investigated. This study will provide the helpful and scientific supports for the development of high-effective water treatment technology (NF) in the application of removing drug pollutants from water.

2. Materials and Methods

2.1. Membrane and Chemicals

A composite NF membrane thin film named NF-270 was selected in this study and the properties of NF-270 are presented in Table 1 [29]. The physic-chemical properties of the selected 14 drugs are summarized in Table 2. All of the drugs were purchased from Sigma-Aldrich. The purities of all drugs were more than 98%. The stock solution was prepared by dissolving each drug 20 mg/L in methanol solvent and stored at -4°C [24, 29]. All chemicals were at least reagent grade. It is worth pointing out that based on the drugs distribution curve, five drugs (acetaminophen, carbamazepine, anti-pyrene, isopropyl-antipyrine and estradiol) are neutral non-charged solutes and the other nine drugs are negatively charged solutes while the solution is ultra pure water (i.e. the pH value is about 7.01).

Table 1. The Properties of NF Membrane Used in the Experiments

| Material | Pure water permeability (L/m²/h) | MgSO₄ rejection(%) | NaCl rejection(%) | Contact angle(°) | Applied pH range |
|----------|----------------------------------|------------------|------------------|-----------------|-----------------|
| Polyamide| 77 ± 3                            | 99.38            | 51.71            | 20.6 ± 2        | 3.0-11.0        |

a) Experimental conditions: 5 mmol/L MgSO₄ in ultra pure water, 5 bar, pH = 7.00; MgSO₄ rejection based on Flame Atomic Absorption Spectrometry measurement (FAAS).
b) Experimental conditions: 10 mmol/L NaCl in ultra pure water, 5 bar, pH = 7.00; NaCl rejection based on FAAS measurement.
c) The contact angle was measured by the static captive bubble method.

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Table 2. Physic-chemical Properties of Selected Fourteen Drugs (n.d.: not determined)

| Drug                     | MW (g/mol) | log$\text{k}_{\text{ow}}$ | pKa  | chemical formula | structure | Water solubility (mg/L) | Charge at pH 7.00 |
|--------------------------|------------|--------------------------|------|------------------|-----------|------------------------|------------------|
| Acetaminophen            | 151.16     | 9.5                      | 0.46 | C$_8$H$_9$NO$_2$  |           | $1.4 \times 10^4$       | neutral          |
| Antipyrine               | 188.22     | 1.5                      | 0.38 | C$_{11}$H$_{12}$N$_2$O |           | $1.0 \times 10^6$       | neutral          |
| Carbamazepine            | 236.27     | 13.9                     | 2.45 | C$_{13}$H$_{15}$N$_2$O |           | 17.66                  | neutral          |
| Estradiol                | 272.39     | 10.7                     | 4.01 | C$_{18}$H$_{24}$O$_2$ |           | 81.97                  | neutral          |
| Isopropyl-antipyrine     | 230.31     | n.d.                     | 1.94 | C$_{14}$H$_{18}$N$_2$O |           | n.d.                   | neutral          |
| Indomethacin             | 357.79     | 4.5                      | 4.27 | C$_{19}$H$_{16}$ClNO$_4$ |           | 0.94                   | negatively charged |
| Diclofenac               | 318.13     | 4.15                     | 4.51 | C$_{14}$H$_{18}$ClNaO$_2$ |           | 2.37                   | negatively charged |
| Naproxen                 | 230.26     | 4.15                     | 3.18 | C$_{13}$H$_{18}$O$_3$ |           | 15.9                   | negatively charged |
| Ketoprofen               | 254.28     | 4.45                     | 3.12 | C$_{16}$H$_{14}$O$_3$ |           | 51                     | negatively charged |
| Ibuprofen                | 206.28     | 4.91                     | 3.97 | C$_{13}$H$_{18}$O$_2$ |           | 21                     | negatively charged |
| Clofibric acid           | 214.65     | 3.0                      | 2.57 | C$_{10}$H$_{11}$ClO$_3$ |           | 582.5                  | negatively charged |
| Sulfamethoxazole         | 253.27     | 5.7, 1.8                 | 0.89 | C$_{10}$H$_{11}$N$_2$O$_4$S |           | 3,942                  | negatively charged |
| Amoxicillin              | 365.41     | 2.7                      | 0.87 | C$_{18}$H$_{13}$N$_2$O$_5$S |           | n.d.                   | negatively charged |
| Bezafibrate              | 361.82     | 3.44                     | 4.25 | C$_{19}$H$_{20}$ClNO$_4$ |           | 15                     | negatively charged |
In this study, we employed the parameter of three-dimensional structure to describe the molecular dimensions of non-absorption neutral drugs, so that steric exclusion effect on the removal efficiency of this kind of drug can be validated in more details. The molecular length was defined as the distance between the two most distant atoms of a particular molecule [26]. The average dimension was employed and defined as the geometric mean of length, width and depth, so that the molecular size can be fully evaluated from the three different dimensions. This mathematical equation about the relation of the average dimension, length, width and depth is shown as follows.

\[ D = (A \times B \times C)^{1/3} \]  

where \( D, A, B \) and \( C \) are the average dimension, length, width and depth, nm, respectively.

2.2. Filtration Protocol and Equipment

In all filtration experiments, the basic solution contained NaCl (10 mmol/L) and phosphate buffer (5 mmol/L). The concentration of drugs was 200 ug/L. The pH of the solution was achieved by adjusting with addition of either sodium hydroxide (0.1 mol/L) or hydrochloric acid (0.1 mol/L) solution. Ultra pure water was used for the preparation of all stock solutions and filtration experiments.

A laboratory-scale, cross-flow membrane filtration test unit is consisted of feed tank (1 L), diaphragm pump, flow meter and membrane cell (shown in Fig. 1). Temperature of the feed tank was maintained at 25 ± 0.5°C by a thermostat (DC-0515, made in China). All experiments were carried out in a circulation mode, with both permeate and concentrate recycling back to the feed tank so as to hold the constant feed concentration. To estimate the removal efficiencies of NF on 14 drugs, the removal experiments were firstly performed with ultra pure water as a reference. For each experiment, the membrane sheet was soaked in ultra pure water for at least 48 h in order to remove impurities from the manufacturing process. Approximately, 0.5 mL of feed and permeate samples were analyzed three times in parallel. The drug concentrations of the influent and effluent were basically steady corresponding to a constant rejection over time. Rejection of drugs was calculated according to Eq. (2)

\[ R(\%) = (1 - C_p / C_f) \times 100\% \]  

where \( R \) is the rejection rate (%), \( C_p \) and \( C_f \) are the drug concentrations in the permeate and feed tank, ug/L, respectively.

2.3. Analytical Methods

The concentrations of drugs were analyzed by high performance liquid chromatography (HPLC, Agilent 1200, USA) equipped with a UV detector and a Poroshell 120 EC-C18 column (4.6 × 50 mm, 2.7 um). The detection wavelengths of carbamazepine, indomethacin, diclofenac, antipyrine, naproxen, ketoprofen, ibuprofen, clofibric acid, acetaminophen, sulfamethoxazole, amoxicillin, estradiol, bezafibrate and isopropyl-antipyrine were 219, 228, 246, 268, 230, 279, 228 and 266 nm, respectively. Acetonitrile, methanol and ultra pure water (60:30:10, v:v) were used as the mobile phase, delivered at a constant flow rate of 0.5 mL/min. The injection volume was 10 uL. The value of pH was obtained by using a Sartorius Professional pH meter (PP-50). The lower limit of detecting ability for all drugs was approximately 5 ug/L. The relative coefficients (R²) of standard curves reached 0.999.

3. Results and Discussion

3.1. Rejection Fourteen Different Types of Drugs by NF Technology

3.1.1. The removal efficiencies of fourteen drugs by NF technology

Fig. 2 shows that NF achieves very high rejection for the most of 14 drugs and the rejection rates of nine negatively charged drugs are higher than those of the neutral drugs. The highest rejection rate by NF is found for ketoprofen (93.3 ± 0.3%). However, acetaminophen is removed with relatively poor efficiency (16.4 ± 0.5%). It is obvious that the rejection rates of neutral drugs are lower in comparison to the negatively charged drugs based on the Fig. 2. As we know, the rejection rates of NF are quite high for most of pollutants even for the small molecules. However, the rejection rates of fourteen drugs are definitely different ranging from the lowest (16.4 ± 0.5%) to the highest (93.3 ± 0.3%). Sieving exclusion is an important removal mechanism in the process of membrane separation. Compared to neutral drugs, the relatively higher degree of rejection of the negatively charged drugs may due to the electrostatic repulsion aside from the steric exclusion mechanism. Thus, it is worth further investigating the reasons of considerable difference in these drugs. Therefore, the selected 14 drugs are divided into two categories (i.e., neutral drugs and negatively charged drugs) based on the drug speciation distribution curve. It is also noted that the material balance is performed to investigate the amount of drugs adsorbed on the membrane shell after the end of separation experiment. 60.16 ug of estradiol is absorbed on the membrane shell, whereas the absorption amount of other 13 drugs is less than 10 ug. The absorption effect of the estradiol will be specially studied in details in section 3.4.
3.1.2. Molecule weight effect on the drug rejection

It is obvious that the relationship between the rejection and the molecule weight of 14 drugs is not linear based on the Fig. 3. The linear correlation coefficient ($R^2$) is only 0.1499. It is prominent that although neutral carbamazepine has higher molecular weight than that of the neutral isopropyl-antipyrine (shown in Table 2), the rejection rate of carbamazepine is obviously lower than that of isopropyl-antipyrine. Thus, it seems that the rejection rate of drug is irrelevant with their molecule weights. The molecular weight cut-off of NF membrane is about 300 g/mol. However, the rejection rates of some drugs whose molecule weight are larger than 300 g/mol are still below 90%. Thus, MWCO cannot accurately predict the removal efficiencies of drugs by NF membrane [23-24, 33-34]. The lowest rejection of acetaminophen among these drugs might be explained by two causes as follows. On the one hand, there is no electrostatic interaction between the neutral non-charged acetaminophen molecule and the membrane surface. It seems that only the steric exclusion can play a part (have effect or play an important role) in this kind of drugs rejection. On the other hand, it has the greater affinity towards water due to the higher water solubility (shown in Table 2), so that it can easily across the membrane structure into the permeate.

3.2. The Removal Mechanisms of the Neutral Drugs

3.2.1. Influence of pH on neutral drugs rejection

The rejection rates of four neutral drugs basically keep constant while pH value ranges from acidic condition (3.48) to alkaline condition (10.01) unlike the trend of acetaminophen shown in Fig. 4. As the pH value below 7.50, acetaminophen, carbamazepine, antipyrine and isopropyl-antipyrine mainly exist as neutral species. However, acetaminophen is negatively charged when the pH value above 7.50, while the other three drugs are still neutral substance. The amount of charges on the membrane surface is strengthened as the increase of the pH value. However, the rejection of carbamazepine, antipyrine and isopropyl-antipyrine still keep constant. Thus, the removal mechanism of neutral drugs may only attribute to the sieving exclusion other than the electrostatic interaction. It is noted that at pH = 10.01, the rejection rate of acetaminophen significantly increases to 73.7%. The reason is that phenolic group of acetaminophen can be deprotonated resulting in a more negatively charged, and then the electrostatic repulsion dominates to remove acetaminophen. The steric exclusion and electrostatic repulsion simultaneously exist in the alkaline solution, so that the rejection rate of acetaminophen cannot be remarkably increased by changing the solution's pH into alkaline condition.

3.2.2. Effect of molecular dimensions on the rejections of drugs

Previous studies have shown that the NF membrane has an average pore radius of 0.42 nm [29]. As discussed in Section 3.2.1, molecular weight cannot be used to stand for the precise molecular dimensions and predict the drug rejection rate. The average dimension, molecular length, width and depth are employed to describe the molecular dimensions of drugs. The molecular dimensions of the three neutral
drugs are summarized in Table 3. It can be seen that the rejection rates of neutral drugs by NF membrane vary from 16.4% to 89.8%. The rejection rates of neutral drugs appeared dramatically in accordance with the molecular length, width, depth and average dimension. This is the reason that the removal efficiencies of neutral drugs increase with the molecular dimensions. This phenomenon might be attributed to their removal mechanism mainly determined by steric exclusion. It seems that the rejection rates of neutral non-adsorptive drugs increase with the average dimension. In addition, the three-dimensional models of antipyrine and isopropyl-antipyrine are presented in Fig. 5. It is apparent to see that the additional functional group of an isopropyl exists in isopropyl-antipyrine molecule structure other than the structure of antipyrine molecule, which indicates that the molecular dimensions of isopropyl-antipyrine are larger than antipyrine. Therefore, the rejection of isopropyl-antipyrine is higher than that of antipyrine attributed to the steric exclusion.

### 3.3. Removal Mechanisms of Negatively Charged Drugs by NF Membrane

Naproxen, bezafibrate and ibuprofen are employed to study the removal mechanisms of negatively charged drugs in this section due to the apparent different physico-chemical properties with the same special functional group of carboxyl acid group. The properties of these three drugs are expected to be sensitive to the pH value due the carboxyl acid group. The speciation distribution curves of three drugs are given in the Fig. 6.

The surface chemical characteristics of NF membrane are determined by the polyamide layer. This polyamide layer contains both carboxylic and amine functional groups that can ionize in the aqueous solution [27]. The degree that NF membrane surface has been negatively charged can be clearly reflected and characterized by using the rejection rates of sodium chloride at the different pH values shown in Fig. 7. As seen in Fig. 7, the rejection rate of sodium chloride increases from 51.71% to 99.38%, which means the surface of NF membrane has been negatively charged as a function of pH, calculated based on their pKa values.

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**Table 3. The Relationship between Molecular Dimensions and the Removal Efficiencies of Acetaminophen, Antipyrine and Carbamazepine**

| Drug            | R (%) | Molecule weight (g/mol) | AFM (nm) | molecular depth (nm) | molecular width (nm) | molecular length (nm) | average dimension (nm) |
|-----------------|-------|-------------------------|----------|----------------------|----------------------|-----------------------|------------------------|
| acetaminophen   | 16.4  | 151.2                   | 105.7    | 0.42                 | 0.68                 | 1.14                  | 0.53                   |
| antipyrine      | 71.1  | 188.2                   | 66.6     | 0.56                 | 0.78                 | 1.17                  | 0.66                   |
| carbamazepine   | 78.8  | 236.3                   | 84.5     | 0.58                 | 0.92                 | 1.00                  | 0.73                   |

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**Fig. 5.** Three-dimensional models of antipyrine and isopropyl-antipyrine.
and the degree that the NF membrane surface has been negatively charged increases along with pH value.

Fig. 8 shows the influence of solution’s pH on the rejection of three negatively charged drugs. It is observed that for NF membrane, as the pH of the solution increased, the rejection of drugs is also increased. This is because the pH value determines the species distribution of drugs in the solution and changes the degree that NF membrane surface has been negatively charged. The percentage of negatively charged drugs increases as the pH value of the solution. Thus, the electrostatic repulsion force between drugs and membrane has been gradually increased, resulting in the increase of the rejection rates obviously. It is noted that the rejection rates of naproxen and ibuprofen, in contrast to that of bezafibrate, are significantly impacted by pH value of the solution. At pH = 3.48, the rejection rates of naproxen and ibuprofen are 25.7% and 11.7%, respectively. Based on the speciation distribution curves, it can be seen that at pH above 5.53, the naproxen and ibuprofen are almost completely dissociated, indicating both of them exist in the form of the negatively charged substance. Thus, the rejection rates of naproxen and ibuprofen are sharply increased to 94% as the pH of the solution is higher than 5.53, due to the increase of electrostatic repulsion force between the drugs and the negatively charged membrane. When the pH is lower than 3.48, the amount of negatively charged of membrane surface is decreased, and the dissociated and negatively charged drugs regain the proton, and then is changed to the neutral substance. Therefore, the elimination of charge repulsion between drugs and membrane is not beneficial to the removal efficiencies of drugs. In this situation, the rejection rates of the drugs are only determined by steric exclusion and adsorption between the drugs and membrane. Besides, the rejection of bezafibrate is also influenced by electrostatic repulsion between the membrane and drugs. However, the rejection rate of bezafibrate increases from 73% to 90% as the pH value ranges from 3.48 to 9.97. Although the degree that membrane surface has been negatively charged sharply increases when the pH value changes from acidic into alkaline condition, the increment of rejection rate of bezafibrate is smaller than those of naproxen and ibuprofen. It seems that the electrostatic repulsion mechanism is more dominated in removal of naproxen and ibuprofen than the removal of bezafibrate.

In contrast to the solution with the lower pH value (< 5.53), the higher rejection rates of naproxen, bezafibrate and ibuprofen by the NF membrane are observed in the solution with the pH value ranging from 5.53 to 9.97. It is noted that steric exclusion may also contribute to the removal mechanism of these three negatively charged drugs. Generally speaking, electrostatic repulsion and steric exclusion are simultaneously responsible for the removal of negatively charged drugs by NF membrane. Therefore, both electrostatic repulsion and steric exclusion are dominated the rejection of negatively charged drugs by NF membrane.

3.4. Adsorption of NF Membrane for Estradiol (E2)

The contact angle of clean NF was measured as 20.6°, which shows that the clean NF had contact angle within the range of the hydrophilic material (Vogler, 1999). Adsorption has also an important influence on the rejection of hydrophobic drugs by using NF treatment. The hydrophobic drug can be easily adsorbed onto the NF membrane surface and then dissolved and permeated into the membrane. Therefore, hydrophobic interaction plays a key role between the drug solute and the membrane phase, and often results in a lower rejection rate [8]. Estradiol, which is a hydrophobic neutral solute with the highest logKow among the selected fourteen drugs, is employed to study the absorption effect during the NF membrane separation. The result from our previous experimental data in ultra pure water showed that adsorption accounted for 30% in the rejection of the estradiol based on the material balance. There is no electrostatic interaction between the estradiol and NF membrane at the condition of pH = 7.01. The rejection of the estradiol only can be determined by steric exclusion and adsorption between the estradiol and the NF membrane.

During the separation process of removing estradiol, the two eminent phenomena are observed shown in the Fig. 9. One is that the concentration of estradiol in the influent decreases due
Fig. 9. Variation of estradiol's concentration in both the effluent and influent via time.

Fig. 10. Freundlich adsorption isotherms of the amount of estradiol in the unit of membrane area with the concentration of estradiol.

NF membrane has very high rejections for the most of fourteen drugs due to the different size, pKa and charge. Thus, the NF can be a high-effective technology to improve the quality and safety of drinking water. The NF membrane in removal of different types of drugs exhibits the different removal mechanisms. And the experiment results show that the rejection rates of nine negatively charged drugs are commonly higher than those of the neutral drugs. For neutral drugs, the steric exclusion is a major removal mechanism and the rejection rate remarkably increases with the increase of 3-D structure size. For negatively charged drugs, both electrostatic repulsion and the steric exclusion are the major removal mechanisms. And electrostatic repulsion has stronger influence on the rejection of negatively charged drugs in contrast to the steric exclusion. The rejection rate of acetaminophen, which is one of the neutral drugs, can be improved by adjusting the solution into the alkalinic condition because electrostatic repulsion also works on removing acetaminophen other than steric exclusion. It is worth to point out that the adsorption effect cannot be neglected in removing the E2. The rejection rate of E2 can be strengthened by the adsorption effect. In summary, the steric exclusion, electrostatic repulsion and adsorption effect are simultaneously existed in the removing drugs by using NF membrane separation.

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