

Original Research

Improved Dissolved Air Flotation Performances Using Chitosan under Different Dosing Schemes

Yu-Long Shi, Jun Ma*, Jia-Xuan Yang

State Key Laboratory of Urban Water Resource and Environment, School of Municipal and Environmental Engineering, Harbin Institute of Technology, Harbin 150090, China

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Abstract

In the present work we investigated the applications of chitosan as a bubble surface modifier in the coagulation dissolved air flotation (DAF) process for river water treatment. Zeta potentials of the residual colloidal particles in floated water and other typical water quality indexes measurements reinforced the fact that micro-bubble surface was successfully modified by dosing chitosan in a saturation system. The flotation efficiencies of both natural organic matters (NOM) and particles were obviously enhanced via chitosan-modified bubbles and the effects became more remarkable after the pH of saturation water was adjusted from 7.0 to 5.5, especially for NOM removal. Moreover, the influence of chitosan on DAF performance, as a coagulant aid, was also evaluated. The coagulation-aiding effects of chitosan were equal to acting as a bubble modifier at pH 7.0, but less efficient than the bubble modification DAF at pH 5.5. In addition, the effects of molecular weight (MW) and deacetylation degree (DD) of chitosan on its performance in both dosing strategies were discussed in detail. These findings highlight the advantages of bubble surface modification in surface water treatment and provide an alternative for DAF process optimization.

Keywords: bubble surface modification, chitosan, deacetylation degree, dissolved air flotation, molecular weight

Introduction

The application of advanced oxidation processes and biological filtration for the removal of natural organic matter is a hot research topic [1-2]. But the chemical oxidation and biodegradation of NOM will lead to the the production of carbon dioxide, which is the primary greenhouse gas and a threat to human health beyond a threshold level [3-7]. By contrast, sedimentation and flotation are the most common and economically feasible

units used to remove natural organic matters (NOM) and colloid particles in surface water treatment [8]. Flotation has been used for waste water reclamation [9], and this has the good prospects of practical applications, especially in drought areas [10]. Concerning sedimentation and flotation processes, efficient coagulation is an essential prerequisite. Over the ordinary pH range, the NOM components and particles bear a negative charge [11]. In order to accomplish a superior removal of NOM, enhanced coagulation is proposed as the best available technique by adding acid/base or increasing dosage of metal coagulants [12]. But both methods have disadvantages, such as increasing the corrosivity of treated water [13] and causing

*e-mail: majun@hit.edu.cn

human health problems related to the high aluminum residuals in tap water [14].

Recently, a new innovation has focused on coagulant optimization, which has been considered to be the third available way for enhanced coagulation [15]. Researchers prepared composited coagulants through a combination of inorganic coagulant and polymer additives with the intent of taking the advantages of both components, such as composite polyaluminum chloride (HPAC) [15], preformed polyferric chloride- polyDADMAC coagulant [16], and a novel polyaluminum-chitosan compound [17]. Although these new complex reagents exhibit superior coagulation effects over conventional metallic coagulants, most of them are still in the development stage.

Besides enhanced coagulation, there is another approach for DAF to improve its performances. As a unique participant, the enormous amounts of micro-bubbles, undertaking all tasks of floc-particles separation, recently excited the particular interest of researchers. The bubble surface zeta potential was considered to be an essential factor for its interaction with floc particles. A series of studies were conducted to optimize the bubble surface characteristics. Results proved that the negatively charged bubble surface could be positively modified using cationic chemicals [18-21]. Bubble surface positive modification may be a novel way to improve DAF performance in pollutant control. More recently, Rao et al. obtained the positively charged bubbles by dosing aromatically modified polymers in the saturator to separate algae cells [22]. Zhang and Guiraud demonstrated the utilization of positive bubble surfaces for silica nanoparticle separation, and the optimum removal rate was beyond 90% [23]. In our previous study the feasibility of bubble modification was also confirmed and the surface-modified bubbles exhibited significant potential to combine with coagulation pretreatment for river water treatment [24].

In this work, based on the conventional coagulation-DAF process (*conv*.DAF) coagulation with polyaluminum chloride (PACl), the performance of bubble positive modification DAF (*posi*.DAF) was investigated during river water treatment. Chitosan, a cationic natural biopolymer, was selected as the bubble surface modifier due to its non-toxicity, high biodegradability, and ecological friendliness [25]. The primary purpose of this study was to determine the optimum dosing strategy for this *posi*.DAF system. For comparison, the flotation efficiency using chitosan as flocculation aids was also examined. In addition, the influences of the DD and MW on chitosan flotation aiding effects were evaluated in these two different dosing schemes.

Materials and Methods

Materials

All chemicals used were of analytical grade or higher, except the PACl (Al₂O₃ content 30.5%), which was produced by Gongyi Fuyuan Water Purification Material Co., Ltd., China. Three native chitosans with medium or high molecular weight (MMW or HMW) and medium or high deacetylation degree (MDD or HDD) were provided by Jinan Haidebei Marine Bioengineering Co., Ltd., China. The molecular weight of chitosan is determined in terms of the Mark-Houwink equation, and the intrinsic viscosity was measured by a Ubbelohde viscometer. The degree of deacetylation was determined by acid-base titration. The charge density of cationic polymers was quantified by the colloid titration method based on the procedure proposed by Kam and Gregory using potassium polyvinyl sulphate (PPVS) calibrated by standard reagent CTAB [26]. The properties of chitosans are summarized in Table 1. Stocking solution of Chitosan 1 g/L was prepared with 0.025 M HCl and overnight stirring at room temperature (25°C) before dosing. Other reagents were purchased from Sinopharm Chemical Reagent Co., China.

Experimental Set-up

A column DAF was employed in this work, located at a waterworks along the Huai River (HR). The unit was mainly composed of a multiphase pump (20FPD05 Nikuni, Japan), a stainless steel air-liquid separation vessel, a recycled water tank, and four transparent acrylic columns (Fig. 1). Each column installed a blade stirrer with automatic timer and mixing speed control.

The experiments of *posi*.DAF via chitosan were carried out as follows:

1. Coagulation/Flocculation – Tests were performed with 4 L raw water transferred into each column. A given volume of PACl working solutions were added in columns using a pipette, and immediately the stirring apparatus were operated as the following procedures: rapid mixing at 500 rpm for 1 min, then gentle stirring at 100 rpm for 5 min.
2. Bubble generation – Filtered water of the waterworks was transferred into the DAF recycle water tank and the pH value (e.g., 7.0±0.1, 5.5±0.1) was adjusted using 0.2 M HCl solutions. The chitosan working solution was added to the recycle tank and mixed evenly. Then the DAF pump started up. The recycled water was saturated with air at pressure of 500 kPa for

Table 1. Characteristics of chitosans used in the tests.

Polymer	Molecular weight (g/mol)	Deacetylation degree (%)	Charge density (meq/g)
Ch.HMW HDD	510,900	87.8	4.5 (pH 5.5) 2.3 (pH 7)
Ch.HMW MDD	507,200	76.5	3.6 (pH 5.5) 1.8 (pH 7)
Ch.MMW MDD	218,000	75.9	3.2 (pH 5.5) 1.5 (pH 7)

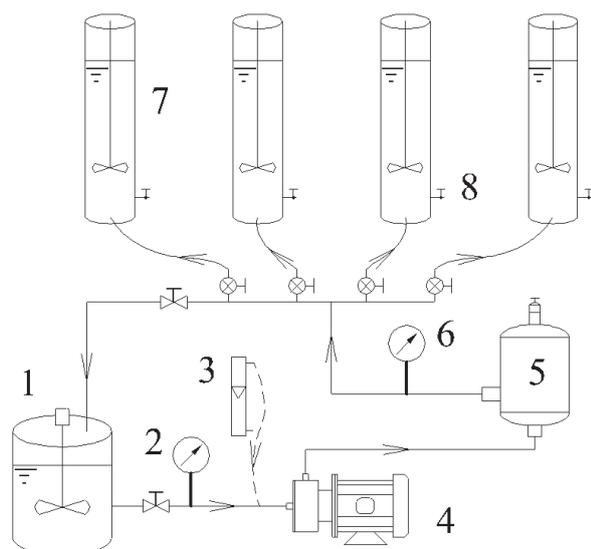


Fig. 1. Schematic diagram of DAF pump system: 1) recycled water tank, 2) vacuum pump, 3) air rotameter, 4) multiphase pump, 5) air-liquid separation vessel, 6) manometer, 7) flotation column, 8) sampling point.

2 min. Finally, microbubbles were introduced into the columns via a releasing device under an equivalent recycling ratio of 10%. After 15 min flotation, samples were collected from the bottom of the column for further analysis. If chitosan was used as a coagulant aid, its dosing point changed to the flotation columns under different sequences with PACl. Other procedures remained unchanged.

Raw Water

Source waters for DAF tests were collected from the HR, located in central China between the Yangtze and Yellow rivers. The raw water dissolved organic carbon (DOC) level was generally around 5.4 mg/L with UV_{254} absorbance of 0.12 cm^{-1} , both of which determined a relatively lower value of the specific ultraviolet absorbance (SUVA) of about 2.2. This indicated that the organic matters in HR were a mixture of aromatic humic substances and aliphatic non-humic matters [11]. The raw water showed a moderate to high alkalinity level, maintaining above $105 \text{ mg/L as CaCO}_3$. The pH values varied between 7.3 and 7.5. In addition, during this work the raw water turbidity remained under 70 NTU at 22.8°C on average.

Analytical Methods

Total organic carbon (TOC) was analyzed using prepared reagents based on the persulfate oxidation photometric principle (HACH Standard Method 10129), with the assistance of a HACH DR5000 UV-Vis spectrophotometer and Digital Reactor Block 200. DOC was analyzed according to the same method after samples were filtrated through a polycarbonate membrane of

$0.45 \mu\text{m}$ pore size. UV_{254} was determined by DR5000 after samples were filtrated through a $0.45 \mu\text{m}$ syringe filter. Turbidity was measured by a HACH 2100N turbidimeter. pH was measured using a PHS-3C pH meter (Leici Shanghai, China). Zeta potential of suspended particles in floated water was measured using a JS94H micro-electrophoresis instrument (POWEREACH, China). All tests were performed in duplicate and the average error between replicates was always under 5%.

Results and Discussion

Chitosan Modification DAF with Coagulation Pretreatment

Effect of [CH]:[Al] Ratios

The flotation efficiencies for HR water treatment at different chitosan-to-aluminum concentration ratios ([CH]:[Al]) are shown in Fig. 2. The pH of recycled water was kept at 7 ± 0.1 during the experiments. As can be seen compared with the *conv.*DAF, the introduction of chitosan-modified bubbles in *posi.*DAF obviously enhanced DOC and turbidity removal. The promotion effects decreased with the increase of PACl dosage (in term of Al concentration). For example, at [CH]:[Al] ratio of 0.4 and with a low Al concentration (1 mg/L), the removal rates of turbidity and DOC were improved by about 20.7% and 8%, respectively. While with moderate Al concentration (4 mg/L) their increase dropped to 7.3% and 4.3%. Under the same Al concentration, with the [CH]:[Al] ratio rising from 0.2 to 0.4, there was a positive correlation between the DOC removal rate and chitosan dosage. But for turbidity control there was no remarkable improvement.

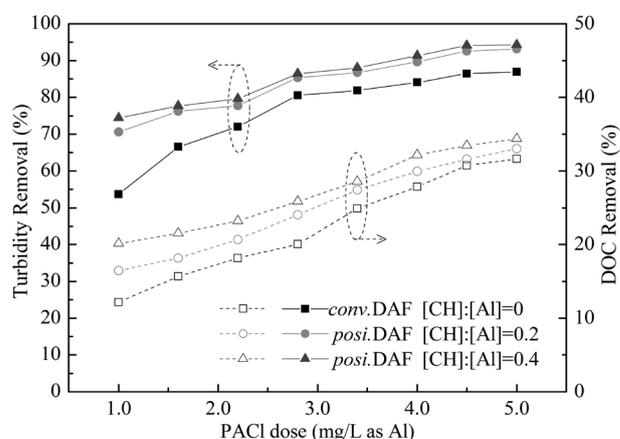


Fig. 2. Turbidity and DOC removal by *conv.*DAF and *posi.*DAF at different [CH]:[Al] ratios (raw water quality: DOC 5.32 mg/L, UV_{254} 0.131 cm^{-1} , turbidity 54.8 NTU, 23.2°C ; recycled water pH: 7 ± 0.1).

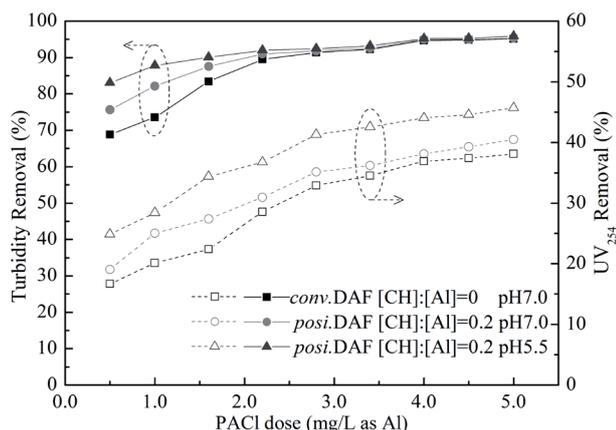


Fig. 3. Turbidity and UV_{254} removal by *conv.DAF* and *posi.DAF* with recycled water at pH 7.0 and 5.5 (raw water quality: DOC 5.20 mg/L, UV_{254} 0.119 cm^{-1} , turbidity 29.3 NTU, 23.5°C).

Effects of the Recycled Water pH Values

As illustrated in Fig. 3, *posi.DAF* performed more effectively at pH 5.5 than at pH 7.0, especially for the organic matters separation. The change tendency accorded well with the results of the previous modifiers selection tests [24]. Compared with UV_{254} , the optimal removal of turbidity was first obtained. At the Al concentration of 4mg/L, the maximum turbidity removal of $95\pm 0.3\%$ was successively reached for all dosing schemes, beyond which the residual turbidity of the floated water remained unchanged. By contrast, there was an obvious improvement of UV_{254} reduction with its average removal percentage increasing from 32.4% at pH 7.0 to 38.1% at pH 5.5.

The pH dependence of *posi.DAF* on NOM removal was closely related to the pH sensitive charge density of chitosan. The protonation degree of $-NH_2$ groups along the backbone of chitosan determined its positive charge

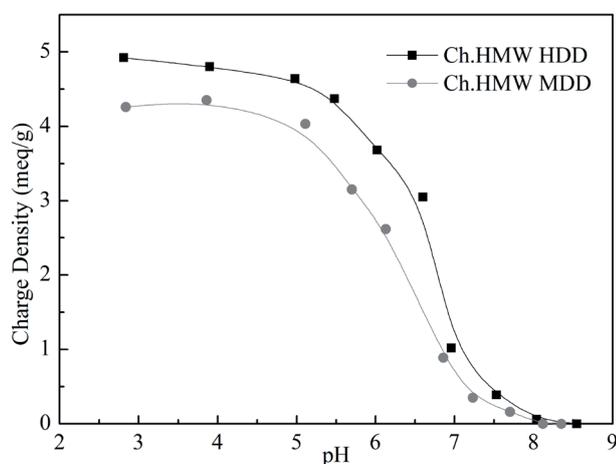


Fig. 4. Effect of pH on charge density of high molecular weight chitosan (HMW) with medium/high deacetylation degree (MDD or HDD).

density and also the bubble interfacial properties covered with it. The pKa value of the amino group was about 6.3 [25]. It can be considered that a small proportion or most of the amino groups were protonated at pH 7.0 and 5.5, respectively. As seen in Fig. 4, when pH is below 5.0 the cationic charge of chitosans sustained high levels, then sharply decreased with pH increasing from 5.5 to 7.0 and became almost zero at pH around 8.0. Thus the adjustment of pH from 7.0 to 5.5 facilitated the electrostatic interactions between the oppositely charged modification bubbles and organic matter. In addition, the different removal characteristic between UV_{254} and turbidity also revealed that charge neutralization played an important role in organic matter separation.

Zeta Potentials of Floated Samples

The zeta potentials of the residual particles after flotation are shown in Fig. 5. For *conv.DAF*, the zeta potential increased steadily following the charge neutralization between hydrolyzed Al species and pollutants. The extra addition of chitosan as bubble surface modifiers in *posi.DAF* at pH 7.0 did not obviously elevate the zeta potential. When dosing chitosan in a recycled water tank at pH 5.5, the main difference was that the introduction of acidic white water would lower the pH of finished water. Although this decrease was not distinct (0.3-0.6 pH units), it might theoretically lead to an increase of the cationic charge of chitosan and amplify the effect of residual chitosan on the zeta potentials. This supposition, however, wasn't proved by test results. We can see that the zeta potential after the acidic recycle water leading-in had no obvious promotions and followed nearly the same profile as the other two dosing plans. This indicated that the majority of chitosan can be tightly bonded with bubbles and only a little remained in the bulk water, which agreed well with the previous results that chitosan strongly bound onto the bubble surfaces under various pH conditions through electrostatic

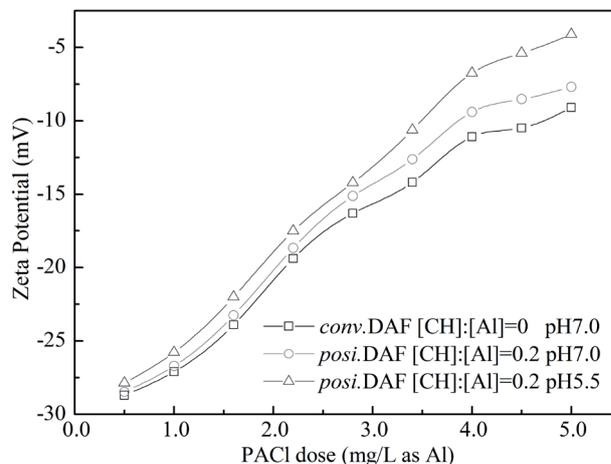


Fig. 5. Zeta potentials of floated water for *conv.DAF* and *posi.DAF* with recycled water at pH 7.0 and 5.5 (raw water quality: DOC 5.20 mg/L, UV_{254} 0.119 cm^{-1} , turbidity 29.3 NTU, 23.5°C).

interaction, hydrophobic attraction, and hydrogen bonding [24].

Influence of MW and DD on Chitosan Performance

Molecular weight (Ch.MMW MDD versus Ch.HMW MDD)

Molecular weight and deacetylation degree are two important characteristics of chitosan playing a subtle role in its interaction with dissolved and colloidal pollutants. In order to study the effects of MW and DD of chitosan on *posi*.DAF performances, a complementary experiment was conducted. In all trials, the pH of recycled water and [CH]:[Al] ratio were kept at 5.5 ± 0.1 and 0.2, respectively. Results showed that a much higher MW of chitosan was favorable for bubble flotation. For example, the removal rate of UV_{254} increased from 26.6% to 29.7% with the MW increase from 2.18×10^5 g/mol to 5.11×10^5 g/mol at PACl dosage of 2 mg/L. As is well known, increasing polymer MW can strengthen its bridging adsorption ability. But when chitosan was used as a bubble surface modifier, there was another important derivative effect that the higher the chitosan MW, the more readily it adsorbed onto the bubble surface.

Deacetylation Degree (Ch.HMW MDD versus Ch.HMW HDD)

The deacetylation degree of chitosan was intimately associated with its rheological behavior and also determined the distribution of cationic sites along its backbone. The higher DD of chitosan meant that more amino groups could be activated. When the pH of the recycled water was decreased to 5.5, the removal efficiency of UV_{254} was obviously improved by the increase of DD. For Ch.HMW, with its DD rising from 76.5% to 87.8%, the UV_{254} removal rates increased by 4.7% at Al concentration

of 5 mg/L. The very reason for the more efficient performance of Ch.HDD than Ch.MDD can be found in Fig. 4. The Ch.HDD exhibited a higher positive charge density than Ch.MDD, especially for the pH values less than the pKa of chitosan.

DAF with Dual Coagulants Strategy

Dosing Sequence of PACl and Chitosan

The primary concern of the dual-coagulant system was the dosing sequence of PACl and chitosan. There were also different opinions about this issue [27-28]. Focusing on this, a series of experiments were carried out under various raw water quality conditions at [CH]:[Al] ratio of 0.2 following three dosing orders, such as adding chitosan or PACl first and adding them at the same time. For organic matter removal, an accordant conclusion could be drawn from the test results that dosing chitosan prior to Al salts was the most efficient. The DOC flotation rates according to other two dosing sequences were approximate to each other, and adding them at the same time was a little better than adding Al first. However, the situations became complicated for turbidity control. Many other factors – including suspended solid load of the source water, mixing intensity and its duration, as well as flocs breakage and re-growth during this process, etc. – must also be taken into consideration. But one thing was certain: dosing chitosan after PACl was conducive to the aggregation of small size floc particles.

This research particularly emphasized organic matter removal by flotation. Thus, the subsequent dual-coagulant tests were performed by adding chitosan first, and then mixed intensively for 1 min prior to PACl dosing.

Effect of [CH]:[Al] Ratios

Fig. 6 shows the removal percentage of DOC and turbidity for HR water after flotation in a dual-coagulant strategy at different chitosan-to-aluminium concentration ratios. Using PACl alone, the removal of 84.5% turbidity was obtained at Al concentration of 4.5 mg/L; beyond that it only slightly changed with further increase of Al dosage. There was a significant improvement in turbidity removal with the addition of chitosan, especially at low Al concentration. The coagulation aiding effects of chitosan on turbidity control were close to each other as the [CH]:[Al] ratios varied from 0.2, 0.4, to 0.6. At the [CH]:[Al] ratio of 0.6, when the Al concentration exceeded 2.8 mg/L the residual turbidity of floated water picked up slightly. Thus the lowest [CH]:[Al] ratio of 0.2 was enough for a desirable particle separation.

With or without chitosan addition, the same change tendency was observed for DOC removal. At any specific Al concentration, the flotation efficiency increased gradually with the [CH]:[Al] ratio growing from 0.2 to 0.6. For example, at [CH]:[Al] ratio of 0.4, the removal rate of DOC increased by 5.2% and 7.7% with a low Al

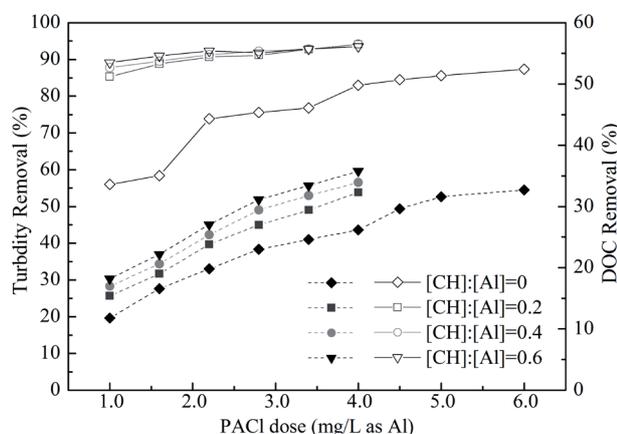


Fig. 6. Flotation efficiency of turbidity (solid line) and DOC (dashed line) in dual-coagulant strategies at different [CH]:[Al] ratios (raw water quality: DOC 5.16 mg/L, UV_{254} 0.110 cm^{-1} , turbidity 42.5 NTU, 24.1°C; recycled water pH: 7 ± 0.1).

concentration (1 mg/L) and a moderate dose (4 mg/L), respectively.

Influence of DD and MW on Chitosan Performance

Molecular weight (Ch.MMW MDD versus Ch.HMW MDD)

As illustrated in Fig. 7, MW and DD of chitosan played different roles in the dual-coagulant system. It is generally believed that polymers with a higher MW had the more remarkable bridging adsorption ability [29]. On the contrary, the present results showed that the increasing MW caused decreased flotation efficiency. This may be ascribed to the stronger affinity and more rapid interaction between Ch.MMW molecules and colloidal particles compared with the Ch.HMW ones. Similar results in previous research have also verified that the number of particles destabilized by low MW chitosan was higher than the high MW chitosan in the early stage after their addition [30].

Deacetylation Degree (Ch.HMW MDD versus Ch.HMW HDD)

Likewise, the stimulating effects of chitosan DD increasing from 76.5% to 87.8% on flotation efficiency was not as good as expected. Only marginal elevation of turbidity and UV_{254} removal was obtained.

By comparison, this indicated that when chitosan was used as a coagulant aid, the MW rather than the DD had a more pronounced effect on its flocculation performances. This finding agreed well with the results obtained by Chen et al. [31]. Notably, this didn't mean the insignificance of DD for chitosan, but instead indicated the complex raw water chemistry – especially the large quantities of blend ions that interfered with the effectiveness of increasing

DD [32]. Besides, in the neural or weakly basic raw water, the protonating of amino groups was restricted by the shortage of H^+ . Without the addition of available H^+ , merely increasing $-NH_2$ would only intensify the competition among plenty of amino groups and enhance the formation of intramolecular hydrogen bonds.

Conclusions

For conventional coagulation-DAF process, the introduction of chitosan, either as a coagulant aid or bubble surface modifier greatly improved the flotation efficiency of DOC, UV_{254} and turbidity. There was marginal difference of the DAF performances with chitosan used as coagulant aid and bubble modifier with recycled water pH at 7.0. The increase of DOC removal percentage varied between 4.3% and 8.0%. The [CH]:[Al] ratio of 0.2 was enough for desirable turbidity control. After the pH of the recycled water was adjusted to 5.5, the *posi*.DAF performed most efficiently. An obvious increase of UV_{254} removal was obtained from 32.4% at pH 7.0 to 38.1% at pH 5.5.

Recycled water pH value was a key external parameter for chitosan. The acidic recycled water with pH lower than the pKa of chitosan was favorable for *posi*.DAF. On the other hand, both the MW and DD were two important internal parameters for chitosan. Chitosan with high MW could be strongly bound onto bubble surfaces and led to high bubble modification efficiency. However, when chitosan was used as coagulant aids, medium MW chitosan performed more efficiently. By contrast, the influence of DD on chitosan flotation-aiding effects was complex and vulnerable to blended ions in raw water.

Acknowledgements

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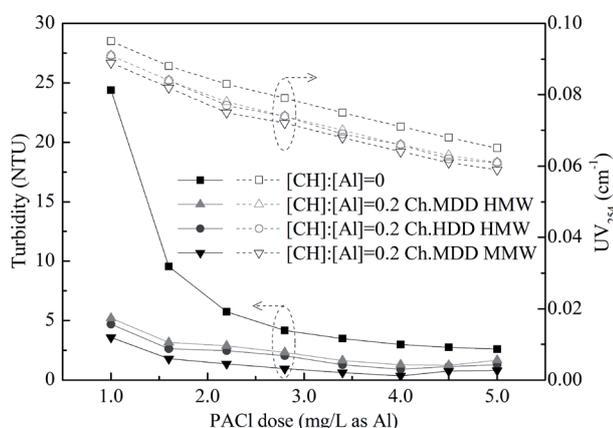


Fig. 7. Residual turbidity and UV_{254} after flotation in dual-coagulant strategies using chitosan with medium/high molecular weight (MMW or HMW) and deacetylation degree (MDD or HDD) (raw water quality: DOC 5.58 mg/L, UV_{254} 0.125 cm^{-1} , turbidity 62.3 NTU, 24.6°C; recycled water pH: 7 ± 0.1).

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