Advanced Treatment of Leachate Secondary Effluent by a Combined Process of MFPFS Coagulation and Sulfate Radical Oxidation

Zhanmeng Liu*, Peng Zhan, Fahui Nie
School of Civil Engineering and Architecture, East China Jiao Tong University, 808 East Shuanggang Road, Changbei District, Nanchang, 330031, China

Received: 22 March 2016
Accepted: 12 April 2016

Abstract
Leachate secondary effluent is usually unable to meet the national discharge permit in China and further treatment is a significant challenge because of the residual persistent pollutants. A novel magnetic composite coagulant of MFPFS was prepared from Fe₃O₄ nanoparticles and polymeric ferric sulfate (PFS). Landfill leachate secondary effluent was treated by MFPFS coagulation combined with sulfate radical oxidation due to Fe²⁺ activation. Higher COD and color removals were achieved using the MFPFS magnetic coagulant than by adding the coagulant PFS alone. Under the working conditions of 1:3 Fe₃O₄/PFS mass ratios, 1.2 g/L MFPFS dosage, 50 min sedimentation time, and without pH adjustment, the removals of COD and color reached up to 60% and 80%, respectively. In the oxidation process, Na₂S₂O₈ was activated by Fe²⁺ to generate sulfate radicals to oxide the residual pollutants in the effluent from MFPFS coagulation. In general, higher temperature, higher dosage, and lower pH were in favor of the oxidation efficiencies. Under the working conditions of 2.4 g/L Na₂S₂O₈ dosage, 0.3 Fe²⁺/S₂O₈²⁻ molar ratio, pH 3.0, and 60°C temperature, the COD and color removals can reach up to 75% and 95%. After being treated by the combined process, no violation of Chinese effluent standards for landfill leachate was found in respect of each individual parameter, including COD, color, and other parameters like BOD, total nitrogen, and ammonia nitrogen. The combined process was a promising alternative treatment pathway for leachate secondary effluent.

Keywords: magnetic composite coagulant, coagulation, sulfate radicals oxidation, leachate secondary effluent, activation

Introduction
The sanitary landfill has been widely accepted for municipal solid waste treatment and disposal due to its lower cost of operation and maintenance as compared to others. However, landfill leachate is considered to be a heavily polluted and a likely hazardous wastewater that is produced during the biochemical decomposition of solid waste or as a result of water infiltration through solid waste. Landfill leachate exhibits considerable spatial and temporal variations. The chemical composition of landfill leachate primarily depends on site operations, refuse characteristics, and internal processes, and the major determinants are, in general, the amount and
The experiment indicates that the addition of Fe₃O₄ can improve floc density and settling performance, thereby enhancing the removal of Microcystis aeruginosa. Fe₃O₄ nanoparticles and an organic flocculant (chitosan or polyacrylamide) are combined to prepare the inorganic organic composite flocculant that is applied to the wastewater treatment [11-12]. The combination of these materials has been found to generate a strong synergistic effect that can significantly improve coagulation performance.

Although coagulation can considerably remove COD, color, and other pollutants from wastewater, this method may not always result in satisfactory treatment efficiencies when applied alone and its effluent can often not meet the national discharge permit. Advanced oxidation processes (AOPs) have been widely applied in the treatment of wastewater containing different organic compounds that are non-biodegradable and/or toxic to microorganisms. Typical AOP systems for leachate treatment had been widely studied, including electro catalytic oxidation [13-14], wet air oxidation [15-16], ozonation [17], and Fenton [18] or Fenton-like oxidation [19] as well as their combination [20]. In recently years, Persulfate (S₂O₈²⁻) has drawn increasing attention as an alternative oxidant in oxidation of contaminants and is considered to be a promising oxidant [21]. Persulfate can be activated to form a stronger oxidant of sulfate radical by heat [22], transition metal ions [23], or UV light [24]. It has been demonstrated by related studies that a broad spectrum of organic pollutants can be rapidly degraded in surfate radical oxidation [25]. The advanced oxidation process based on sulfate radical has a strong oxidative capacity and low cost, which facilitates its potential application in wastewater treatment. However, it is rarely reported in research that a sulfate radical was applied in the treatment of leachate.

So in this work, a novel magnetic composite coagulant of MFPFS was prepared from Fe₃O₄ nanoparticles and PFS and was introduced to remove the residual pollutants from leachate secondary effluent combined with sulfate radical oxidation due to Fe²⁺. The objective of this study was to examine whether the new combined process was a feasibly integrated technique for efficiently treating leachate secondary effluent and find the best operation parameters as reference for engineering applications.

Material and Methods

Wastewater Samples

The experimental wastewater was a biological pre-treated leachate sample taken from a municipal solid waste landfill disposal plant in Jiujiang City, Jiangxi Province, PR China. The malodorous wastewater was brown and contained some suspended materials. Typical characteristics of the wastewater are presented as follows: COD 580-760 mg/L, color 280-360 times, pH 6.4-6.7, and ammonia nitrogen 62-74 mg/L.
Coagulation Experiments

With and without prior pH adjustment and for different dosages, coagulation experiments were performed by jar test. The pH values of samples were adjusted to the desired levels by the addition of appropriate amounts of H₂SO₄ or NaOH solution. The time and speed of mixing for rapid and slow mixing were set automatically by means of a controller. Briefly, 800 ml of wastewater were added with a certain amount of coagulant, after which the test was conducted by subjecting the solution to 1 min of rapid mixing followed by 10 min of slow mixing. Next, the mixed samples were allowed to settle without agitation for certain time. The supernatant was withdrawn from the beakers and used for chemical analysis and further treatment by sulfate radical oxidation.

Sulfate Radical Oxidation Experiments

With and without prior pH adjustment, sulfate radical oxidation was conducted using Na₂S₂O₈ as oxidant. 100 mL of leachate secondary effluent was added into a 250 ml Erlenmeyer flask. Certain amount of Na₂S₂O₈ and Fe²⁺ were added to the solution. Then the mixed solution was shaken at 150 rpm using a shaking water bath at a controlled temperature. Oxidation was initiated by the addition of a certain amount of Na₂S₂O₈. The oxidation time was set for 12 hours.

Analysis Methods

COD, BOD, total nitrogen, ammonia nitrogen, and color were measured according to the Manual Standard Methods for the Examination of Water and Wastewater drawn by the Chinese Environmental Protection Agency. pH was measured using a pH-Meter (model PHS-3C). The chemical reagents used here were all analytical grade purchased from Shanghai Chemical Technology Company, China.

Results and Discussion

MFPFS Coagulation Process

Effect of mass ratio of Fe₃O₄/PFS in MFPFS on coagulation performance

The MFPFS reagents with different Fe₃O₄/PFS mass ratios were prepared and coagulation experiments of the newly prepared reagents were performed under the same conditions and the results are shown in Fig. 1. As shown in Fig. 1, higher or lower mass ratios of Fe₃O₄/PFS were not facilitated regarding coagulation performance. With an increase in the mass ratio of Fe₃O₄/PFS, the COD and color removals first slowly increased and then gradually declined. The color removal rate reached its highest value of 83% when the mass ratio was 1:4. When the mass ratio was increased to 1:3, the COD removal rate reached its highest value at 62%, and the color removal rate reached 81%. As the mass ratio continued to increase, the COD and color removals became reduced to a certain degree. When the mass ratio was 1:1, the COD and color removals were only 42% and 57%, respectively. The COD removals reached up to the highest under the mass ratio of 1:3. Even if color removal was reduced slightly, the removal ratio was still high. Therefore, the optimum mass ratio of Fe₃O₄/PFS for preparing MFPFS was 1:3.

Effect of pH on Coagulation Performance

Wastewater pH is an important parameter in the coagulation process. It affects the pollutant form and the Zeta potential of colloid granule and then affects coagulation performance. In order to investigate the possible effect of initial pH and obtain the optimum pH at which the coagulation process performs most effectively, the pH of landfill leachate was adjusted to desired value. The effect of pH variation on coagulation performance was examined and the results are shown in Fig. 2. It can be clearly seen from Fig. 2 that the COD and color removals were first on the steep increase and then on the
very slow decline. When pH value was 4.0, the COD and color removal rates were the lowest – at 24% and 56%, respectively. When pH was 6.0, color removal reached its highest level at 85%. When pH increased to 6.5, COD removal reached its highest value at 60%, and color removal 80%. When pH was increased from 6.5 to 8.0, the COD and color removals slowly decreased with an increase in pH value. However, COD and color removal still reached 52% and 78%, respectively, at pH 8.0. This illustrates that the optimum wastewater pH value scope was wide. In general, the coagulation performances were good in the scope of from pH 6.0 to pH 8.0. Since the initial pH value was generally in the feasible range of coagulation and considering pH adjustment was uneconomical in practical engineering terms for an indistinct increase of COD removal, no pH adjustment was carried out during the followed trials.

Effect of Sedimentation Time on Coagulation Performance

The sedimentation process aims to separate the solid (mainly flocs) from the wastewater and produce the treated supernatant. The effect of sedimentation time on the coagulation performance is shown in Fig. 3. Both COD and color removals were on a sharp upward trend with the sedimentation time increasing from 10 to 50 min, while keeping nearly constant after that. Sedimentation time is an important factor affecting the quality of treated supernatant. The fine solid particle left in the supernatant caused by scant sedimentation time is a contribution not only to the pollutants in supernatant but also to the burden of the latter treatments. Excessive sedimentation time may benefit better coagulation performance. However, this needed a larger footprint and then results in increasing much more capital construction cost investment. Considering the COD and color removal variations in general, it was suggested that the optimum sedimentation time was about 40 to 50 minutes under the given conditions.

Coagulation Performance of PFS and MFPFS at Different Dosages

The coagulation performance of PFS and MFPFS at different dosages is shown in Fig. 4, where coagulation performance of MFPFS was obviously better than that of PFS under the same conditions. Coagulation performance had been greatly improved by the introduction of Fe3O4 nanoparticles. As the dosages of PFS and MFPFS increase, COD and color removals tended to initially increase quickly followed by a slow decline. When the dosage is 0.9 g/L, COD removal with the MFPFS coagulant reached its maximum at 56%. By contrast, COD removal with PFS alone was only 45%. COD removal when MFPFS was added was 11% higher than when only PFS was added. At this dosage, color removal of MFPFS and PFS were 82% and 69% respectively. Color removal when MFPFS was added was 13% higher than when only PFS was added. The reason is that, firstly, Fe3O4 nanoparticles with larger specific surface area and adsorption capacity can become the cores of floc in the coagulation process, playing the role of heterogeneous nucleation [10] and, consequently, the density and settling property of the coagulation body were improved, and the floc formation, aggregation, and settling time were shortened, thereby enhancing the coagulation of PFS [9]. The second (with increased dosage) Fe3O4 became easier to combine with the coagulation body of the coagulant to form a more compact magnetic flocculation due to its magnetic property. Moreover, mutual attraction between magnetic particles was enhanced, thus increasing the number of particles that aggregate into large aggregates and ultimately leading to a good coagulation effect [26].

As shown in Fig. 4, when the dosage of MFPFS increased from 2.0 to 2.5 g/L, the COD and color removals decrease from 62% to 46%, and from 78% to 71%, respectively. This indicated that overdosing of coagulant would deteriorate the coagulation performance. The reason is that an increased dosage of MFPFS causes a greater dosage of Fe3O4. Thus, self coagulation would
occur between Fe$_3$O$_4$ nanoparticles, which would no longer combine with the coagulation body of the coagulant to form a more compact magnetic coagulation. As such, these nanoparticles can no longer facilitate heterogeneous nucleation, resulting in poor coagulation performance. In addition, according to the charge neutralization theory, hydrolyzed products of the added coagulant can interact with negative colloids and neutralize their charges, which cause colloid destabilization and removal. With more than the appropriate dosage than the amount required for coagulation, the colloids can absorb the cations then become positively charged, resulting in renewed stability because of electrical repulsion, thus leading to poor coagulation performance [27]. Overdosing the coagulant not only weakened coagulation efficiency but also increased the cost. With MFPFS dosage in the scope from 0.9 g/L to 2.0 g/L, the COD and color removals still reached up to 56% and 80%. These illustrate that the optimum MFPFS dosage scope was wide at the experimental conditions, which makes dosage control and coagulant popularization easier.

On the basis of the above analysis, the optimum conditions for the treatment of landfill leachate by MFPFS coagulation are at 1:3 mass ratio of Fe$_3$O$_4$ /PFS in MFPFS, approximately 1.2 g/L MFPFS dosage, and approximately pH 6.5 (without adjusting pH). Under these conditions, the removal rates of COD and color reached up to 62% and 85%, respectively.

### Sulfate Radical Oxidation Process

After treatment by MFPFS coagulation, the typical parameters of the effluent were: COD 235-282 mg/L, color 70-100 times, and pH 6.1-6.5. The effluent of MFPFS coagulation was further subject to treatment by sulfate radical oxidation due to Fe$^{2+}$ activation.

**Effect of Na$_2$S$_2$O$_8$ Dosage on Oxidation Performance**

Oxidant dosage is an important parameter in the sulfate radical oxidation process. The effect of Na$_2$S$_2$O$_8$ dosage on oxidation performance is shown in Fig. 5, which shows that a high Na$_2$S$_2$O$_8$ dosage generally favored COD and color removals. Generally, the trends of the COD removal and color removal were similar to Na$_2$S$_2$O$_8$ dosage, and increasing Na$_2$S$_2$O$_8$ dosage would improve oxidation performance. Observing the COD removal curve, it can be seen that COD removal significantly increased with Na$_2$S$_2$O$_8$ dosage from 0.4 to 2.4 g/L. When the oxidant dosage was 2.4 g/L, the COD removal rate reached up to 72%. When the Na$_2$S$_2$O$_8$ dosage was continually increased from 2.4 to 3.2 g/L, the COD removal percentage did not significantly increase and remained relatively steady. The sulfate radical oxidation had a strong capacity of decoloration from leachate secondary effluent. When the oxidant dosage was 1.6 g/L, the color removal rate reached up to 90%, and in the range of Na$_2$S$_2$O$_8$ dosage from 2.0 to 3.2 g/L, the effluent color nearly completely disappeared and the color removals were all more than 95%.

**Effect of Fe$^{2+}$/S$_2$O$_8^{2-}$ Molar on Oxidation Performance**

A suitable Fe$^{2+}$/S$_2$O$_8^{2-}$ molar ratio is necessary in the sulfate radical oxidation due to Fe$^{2+}$ activation. Too high and too low Fe$^{2+}$/S$_2$O$_8^{2-}$ molar ratios are unfavorable to oxidation performance. As shown in Fig.6, a suitable Fe$^{2+}$/S$_2$O$_8^{2-}$ molar ratio favored oxidation performance. COD removal first remarkably increased with Fe$^{2+}$/S$_2$O$_8^{2-}$ molar ratio from 0.1 to 0.3 and then declined with Fe$^{2+}$/S$_2$O$_8^{2-}$ molar ratio from 0.3 to 0.8. The peak removal rate of COD appeared at Fe$^{2+}$/S$_2$O$_8^{2-}$ molar ratio 0.3, with 72% COD removal. When the Fe$^{2+}$/S$_2$O$_8^{2-}$ molar ratio was increased to 0.6, the COD removal rate declined to 56%. This likely occurred because the Fe$^{2+}$ dosage added was greater than the amount required for activation. Observing the removal curve, it can be seen that the color removal rate first
increased and then slightly declined with the $\text{Fe}^{2+}/\text{S}_2\text{O}_8^{2-}$ molar ratio. Furthermore, it should be mentioned that the effluent color was always lower than 40 times in the whole $\text{Fe}^{2+}/\text{S}_2\text{O}_8^{2-}$ molar ratio and met the discharge permission (lower than 40 times) established by the Chinese Environmental Protection Agency. In consideration of COD removal and color removal, it was suggested that the application of $\text{Fe}^{2+}/\text{S}_2\text{O}_8^{2-} = 0.3$ would be appropriate for wastewater treatment.

**Effect of pH on the Oxidation Performance**

pH is also an important parameter in treating wastewater by sulfate radical oxidation. As shown in Fig. 7, in general the trends of COD removal and color removal were similar to pH, and heightening pH would weaken oxidation performance. Both the COD removal and color removal were the highest at pH 3, and the removal rates reached 75% and 98%, respectively. Observing the COD removal curve, it can be seen that the COD removal rate first slowly declined with pH from 3 to 4 and then rapidly declined with pH from 4 to 7. When the pH value increased to 7.0, the COD removal rate declined to 39%. As for color removal, the removal rate slightly declined with pH from 3 to 5 and then remarkably declined with pH from 5 to 7. It had been reported that heightening pH has a dual effect (positive and restrictive) on pollutant removal in sulfate radicals oxidation based on thermal activation. On the one hand, heightening pH is in favor of Na$_2$S$_2$O$_8$ activation generating sulfate radicals and then enhancing oxidation capacity. On the other hand, heightening pH can elevate the alkalinity level, which results in enlarging the side reactions of sulfate radicals with CO$_3^{2-}$ and HCO$_3^{-}$, thus weakening the oxidation efficiencies. As a whole, it is reflected (as shown in Fig. 5) that the second parameter played a key role and heightening pH would deteriorate the oxidation efficiencies in the treatment of the wastewater. The result was similar to that provided in a study conducted by Shen to evaluate the oxidation efficiency of diphenylamine by persulfate [28]. Considering COD removal and color removal, it was suggested that applying pH 3-4 would be appropriate for wastewater treatment.

**Effect of Temperature on Oxidation Performance**

It can be seen from Fig. 8 that high temperatures generally favor oxidation performance. The oxidation efficiencies were better at higher temperatures than those at low temperatures under the same conditions. When the temperature was 25°C, COD removal and color removal were only 43% and 80%, respectively. When the temperature was increased to 45°C, COD and color removals reached up to 65% and 95%, respectively. However, it could be found that although the higher temperature resulted in higher COD removal and higher color removal, oxidation efficiency improvement due to temperature elevation was gradually weakened. Improvement due to the elevation activation temperature became poor when continuously increasing the activation temperature from 45°C to 75°C. As a consideration of COD removal and color removal, it was suggested that the application of 45-60°C would be appropriate for wastewater treatment.

The above results demonstrated that sulfate radical oxidation based on Fe$^{2+}$ activation could adequately remove refractory organics from leachate secondary effluent. This indicated that sulfate radical oxidation can readily degrade organic matter in wastewater.

**Removal Efficiencies of Pollutants by the Combined Process of MFPFS Coagulation and Sulfate Radical Oxidation**

The pollutants in the leachate secondary effluent are not only indicated as COD and color, but also BOD$_5$, ammonia nitrogen, and total nitrogen. The removal efficiencies of these parameters were also investigated under the optimized conditions. In this experiment, the coagulant dosage of MFPFS was fixed at 1.2 g/L in the
coagulation process; the conditions in the oxidation process were: \( \text{Na}_2\text{S}_2\text{O}_8 \) dosage = 2.0 g/L, \( \text{Fe}^{2+}/\text{S}_2\text{O}_8^{2-} = 0.3 \), \( \text{pH} = 3.0 \), and \( T = 60^\circ\text{C} \). After treatment by the combined process, the typical parameters of the wastewater are listed in Table 1.

As shown in Table 1, the combined processes can result in satisfactory treatment efficiencies. The concentration of COD decreased from 624 mg/L to 261 mg/L by MFPFS coagulation and further decreased from 261 mg/L to 74 mg/L. It can be seen that the reduction of color was significant and the color completely disappeared (after oxidation), which indicated that the combined processes have a notable capacity for removing color. Concentration changes of ammonia nitrogen and total nitrogen before and after treatment were investigated. The coagulation process hardly removed ammonia nitrogen and total nitrogen while sulfate radical oxidation exhibited good capacity for ammonia nitrogen and total nitrogen removals. The concentration of ammonia nitrogen and total nitrogen decreased to 18 mg/L and 35 mg/L, respectively. Certainly, it should be mentioned that after being treated by the combined process no violations of Chinese effluent standards for landfill leachate were found in respect to each individual parameter. The encouraging results demonstrated that the combined process could remove most target compounds and provide a viable alternative to the treatment of leachate secondary effluent.

### Conclusion

In this work, the application of the combined process of MFPFS coagulation and sulfate radical oxidation based on \( \text{Fe}^{2+} \) activation for leachate secondary effluent was studied, and it proved to be an efficient and promising method.

Coagulation performance is significantly better when the novel magnetic coagulant MFPFS was prepared by introducing \( \text{Fe}_3\text{O}_4 \) nanoparticles into PFS as applied in treating leachate secondary effluent when PFS is used alone. \( \text{Fe}_3\text{O}_4 \) nanoparticles can improve floc density and settling performance, thereby enhancing pollutant removal from the wastewater. The optimum mass ratio of \( \text{Fe}_3\text{O}_4/\text{PFS} \) for preparing MFPFS is 1:3. Without adjusting \( \text{pH} \), at a dosage of 1.2 g/L the removal rates of COD and color reach up to 60% and 80%, respectively.

The oxidation efficiencies were affected by the factors of \( \text{Na}_2\text{S}_2\text{O}_8 \) dosage, \( \text{Fe}^{2+}/\text{S}_2\text{O}_8^{2-} \) molar ratio, wastewater \( \text{pH} \), and temperature. With a 2.0 g/L \( \text{Na}_2\text{S}_2\text{O}_8 \) dosage, 0.3 \( \text{Fe}^{2+}/\text{S}_2\text{O}_8^{2-} \) molar ratio, \( \text{pH} = 3.0 \), and \( T = 60^\circ\text{C} \) temperature as the optimum working conditions for the treatment of effluent from MFPFS coagulation, sulfate radical oxidation due to \( \text{Fe}^{2+} \) activation could efficiently remove the residual contaminants, and the COD and color removals can reach up to 73% and 95%, respectively.

Apart from good efficiencies of COD and color obtained, the combined process can also efficiently remove other parameters like BOD, total nitrogen, and ammonia nitrogen. After being treated by the combined process, no violations of Chinese effluent standards for landfill leachate were found in respect to each individual parameter. The encouraging results demonstrated that the combined process could remove most target compounds and provide a viable alternative to the treatment of leachate secondary effluent.

### Acknowledgements

This work was supported by the National Natural Sciences Foundation of China (No. 51468016), the Natural Sciences Foundation of Jiangxi (No. 20142BAB203027), and the Technology Development Research Project of Jiangxi (No. 20151BBG70020).

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