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Influence of Different Factors on the Removal of Ammonia by Electrolysis with Three-Dimensional Electrode

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Electrolytic removal of ammonia was investigated by using zeolite packed three dimensional electrode reactor under batch mode. The results showed that 6-100 mg N/L ammonia could be successfully removed within 1-7 h under various conditions of 30-300 mg/L chloride ion concentration, 0.5-2 A current and pH 3-9. Current density and initial ammonia concentration had significant effect on the ammonia removal rate, while the influence of chloride concentration and pH was weak under batch mode in this research. For the treatment of municipal sewage, the ammonia concentration was reduced from 27.8 mg N/L to 0.5 mg N/L after 1.5 h electrolysis. Moreover, 72 and 61 % removal of the total nitrogen and COD could be achieved after continue electrolysis for 2.25 h.

Keywords: Three-dimensional electrode, Electrochemical oxidation, Ammonia, Zeolite.

INTRODUCTION

Nowadays eutrophication is one of most serious environmental problems in China. In general, excess discharge of nutritious elements such as nitrogen and phosphorus are recognized as the main reason for algae blooming. In wastewater, ammonia is one of the most important forms of nitrogen contamination due to its characteristics of oxygen consumption, promotion to algae growth, toxicity to fish *etc.* Low limits of 0.02 and 0.1 mg/L of unionized ammonia (NH_3) are suggested for chronic and acute toxicity, respectively¹. Deep removal of ammonia from the secondary effluent of sewage discharge facility has become a challenge issue.

As one of the advanced oxidation methods, electrochemical oxidation has attracted more and more attention due to its effectiveness, environmental friendly products, easy operation and maintenance. Many researchers have reported that electrochemical process can be used for removal of refractory organics, nitrate, ammonia, heavy metals, *etc.*²⁻⁴. For example, significant chemical oxygen demand (COD) and ammonia removal as well as decolorization have been observed by Rao *et al.*⁵ when treating tannery wastewater with Ti/Pt, Ti/PbO₂, and Ti/MnO₂ as anodes and Ti plate as cathode.

The electrolytic removal of contaminants from wastewater was found to be related to various factors such as electrode materials, electrolyte, current, pH, type of reactor, *etc.*⁶. Previous studies have summarized that higher current density and electrolyte concentration, neutral pH condition could promote

the reaction rate of ammonia. In addition, the electrode materials were the most important factor for the ammonia removal rate. As the electrode materials changed, the ammonia removal rate altered. Moreover, ammonia was proved to have obvious effects on the phenol removal efficiency in chloride mediated system during electrochemical oxidation with RuO₂-Pt/Ti as anode and Ti plate as cathode⁷. In recent years, 3D electrode reactors had been successfully applied to the treatment of landfill leachate, tannery wastewater, textile wastewater *etc.*⁸⁻¹⁰, which was proved to significantly enhance the mass transfers as well as contaminants removal.

In this research, 3D electrode reactor was designed for deep and fast removal of ammonia by electrolysis. Different influencing parameters including current, chloride ion concentration, pH and initial ammonia concentration were investigated for electrocatalytic ammonia removal by using bipolar 3D electrode reactor. Typical secondary effluent from sewage discharge facility was treated to evaluate the application of the system in real situation.

EXPERIMENTAL

(NH₄)₂SO₄, NaCl, Na₂SO₄, H₃BO₃, H₂SO₄ and NaOH were all of analytical grade purchased from Sinopharm Chemical Reagent Co., Ltd (Shanghai, China). Distilled water was used as the solvent to prepare synthetic wastewater with ammonia ($\text{NH}_3 + \text{NH}_4^+$), chloride ion (Cl^-) and suitable electro conductivity. The final pH of the synthetic solution was adjusted by using H₂SO₄ and NaOH with the help of a pH meter. 30 %

solution of boric acid was prepared as adsorption solution for ammonia eluting.

Set-up and procedure: Batch electrolysis system was employed to investigate the influence of different factors on electrolytic removal of ammonia by three dimensional electrode. The experimental set-up was composed of DC power supply, circulating pump, gas washer and collection bag and 3D electrode reactor with RuO_2/Ti anode, stainless steel cathode and 2-3 mm natural zeolite particle electrode. The wastewater containing ammonia was passed through the 3D electrode reactor with the force of the pump at a constant flow rate of 150 mL/min. During electrolysis, samples were taken and analyzed immediately, and the eluting ammonia in the gases was adsorbed in the gas washer and analyzed after the experiment.

Electrolytic experiments: Batch mode was employed to evaluate the electrolytic removal efficiency of ammonia by bipolar 3D electrode. The possible influencing factors were investigated by using simulated domestic sewage. Single factor approach was used to test the effect of initial ammonia concentration, current density, chloride concentration and pH. Each time, only one experiment factor was changed while other factors were kept same as shown in Table-1. Actual wastewater from Dongqu wastewater treatment plant was taken to evaluate the efficiency of electrolytic ammonia removal in real situation.

TABLE-1 SINGLE FACTOR EXPERIMENTAL CONDITION			
$\text{C}_{\text{NH}_3\text{-N}}$ (mg/L)	Current density (mA/cm ²)	C_{Cl^-} (mg/L)	pH
10, 30, 50, 100	13.7	300	7
30	3.8, 7.6, 11.4, 13.7	300	7
30	13.7	30, 120, 225, 300	7
30	13.7	300	3, 7, 9

Detection method: Ammonia was measured according to standard methods¹¹. COD was determined by low range COD reagent (HACH, Loveland, Colo.). pH was measured by a pH meter (PHS-2C, Leici Company, Shanghai, China). Total nitrogen was measured by a TOC/TN analyzer (Multi N/C 3100, Analytikjena Company, Germany).

RESULTS AND DISCUSSION

After each experiment, the concentration of ammonia in gas washer was measured and the result showed that ammonia volatilization was less than 1 %. So the amount of ammonia removed through volatilization can be ignored.

Current density: In order to investigate the effects of ammonia removal by zeolite through ion-exchange and adsorption, synthetic wastewater containing 29.5 mg N/L ammonia and 300 mg/L chloride was circulated inside the experimental setup. The change of ammonia was shown in Fig. 1, which indicated that in the beginning 15 min rapid adsorption resulted in 50 % removal of ammonia. Slow adsorption of ammonia was observed in the following 3 h and the final ammonia concentration was measured to be 6.8 mg N/L. To evaluate the effect of current densities, the same synthetic wastewater were electrolyzed under the currents of 0.5 A, 1.0 A, 1.5 A

and 1.8 A, respectively. As shown in Fig. 1, more than 99 % of ammonia could be removed in 3, 2, 1.5 and 1.5 h, respectively, while 12, 7, 4, and 2.5 h of electrolytic time were needed to achieve similar results under the same conditions when using traditional 2D electrode in previous research¹². The efficiency of ammonia removal by 3D electrode reactor increased significantly as compared with 2D electrode. Moreover, it was obviously that higher contaminant removal rate was obtained when higher current density was applied, which was similar with the results obtained by Rosilda and his co-workers¹³ studied the ammonium ion removal from produced water by a batch single-stage electrolytic process using $\text{RuO}_2 + \text{TiO}_2/\text{Ti}$ as anode and pure titanium plate as cathode. From previous research with RuO_2/Ti ¹² and IrO_2/Ti ¹⁴ anode, electrolytic ammonia oxidation followed pseudo-zero order kinetics at different currents. This difference could be attributed to the combination reaction of ion-exchange and electrochemical oxidation in 3D electrode reactor. Although high ammonia removal rate was observed under high current, energy cost would be significantly increased.

Fig. 1. Effect of current density on the electrolytic removal of ammonia

Chloride ion concentration: Chloride ion concentration in the system was very important for ammonia oxidation because of the formation of HClO or ClO^- . As shown in Fig. 2, the electrolytic removal of ammonia was investigated under different chloride ion concentrations between 30 and 300 mg/L. After 1.5 h electrolysis, the ammonia concentration decreased from 30 mg N/L to less than 1 mg N/L under all chloride concentrations, which was different from the result obtained by 2D electrode. Vanlangendonck *et al.*¹⁵ and Li & Liu¹² concluded that high chloride concentration had positive influence on ammonia oxidation rate because higher concentration was beneficial for hypochlorite production. The improved removal in this experiment might due to ammonia adsorption on zeolite and simultaneously electrocatalytic regeneration of zeolite during electrolysis with 3D electrode.

pH: In aqueous solution, the form of ammonia is significantly affected by pH¹⁶. Under pH 9.25, NH_4^+ accounts for 50 % and NH_3 takes the other half. Under pH 7, NH_4^+ rises from 50 % to 99.4 percent, which was possible to affect the electrolytic removal of ammonia by 3D electrode. As shown in Fig. 3, electrolytic removal of ammonia were studied under pH 3, 7 and 9, the results showed that the ammonia concentrations

Fig. 2. Effect of chloride ion concentration on the electrolytic removal of ammonia

were 1.5, 0.3 and 0.6 mg N/L, respectively, after 1.5 h electrolysis. When pH was 3, the removal rate of ammonia was slightly lower than results obtained under pH 7 and 9. This suggested that pH had little influence on ammonia oxidation so the removal of ammonia by electrochemical oxidation can adapt to relatively wide pH range. It was consistent with the results obtained by Mahvi *et al.*¹⁷ who observed that the contaminant removal efficiency had no obvious difference over the entire range of pH values from 5 to 10. Moreover, Vanlangendonck *et al.*¹⁵ demonstrated that pH values between 5.5 and 10 had no effect on ammonia oxidation rate when using stainless steel as cathode and $\text{TiO}_2 + \text{RuO}_2/\text{Ti}$ as anode to treat power plants wastewater.

Fig. 3. Effect of pH on the electrolytic removal of ammonia

Initial ammonia concentration: In actual municipal sewage, the ammonia concentration usually fluctuated around 30 mg N/L, and possibly increase if it was mixed with agriculture runoff or industrial wastewater. In this research, the performance of 3D electrode was evaluated using synthetic ammonia wastewater with concentrations ranging from 6 to 99 mg N/L. The results showed that either under low or high ammonia concentrations, ammonia could be completely removed given sufficient electrolysis time. As shown in Fig. 4, 0.5, 1.5, 2.5 and 6.5 h of electrolysis time were needed to

reach an effluent ammonia concentration lower than 1 mg N/L for initial ammonia concentrations of 6, 30, 50, 99 mg N/L, respectively. Under the same condition, 1, 2.5, 4.5 and 8.5 h of electrolysis time were needed to achieve the same effect when using traditional 2D electrode reactor¹². So the ammonia removal rate by 3D electrode was significantly improved compared with 2D electrode.

Fig. 4. Effect of initial ammonia concentration on the electrolytic removal of ammonia

Performance of 3D electrode in treating actual municipal sewage: The actual wastewater was taken from Dongqu wastewater treatment plant in Shanghai, which was used to evaluate the efficiency of electrolytic for ammonia removal under the condition of 1.8 A current, pH 7 and 300 mg/L chloride ion concentration. As shown in Fig. 5, the changes of COD, total nitrogen and ammonia were determined. After electrolysis for 1.5 h, the ammonia concentration decreased from 27.8 mg N/L to 0.5 mg N/L, which met the demand of The First Grade of Environment Quantity Standards for surface water in China. In addition, the total nitrogen concentration decreased from 30.6 to 8.6 mg N/L within 2.25 h, which accounted for 72 % ammonia removal from aqueous phase. As for organic compounds, the COD concentration decreased from 38 mg/L to 15 mg/L after 2.25 h electrolysis, which accounted for 61 % removal. These results indicated that bipolar 3D electrode was a promising method for deep and rapid removal of ammonia in domestic sewage.

Fig. 5. Ammonia removal in municipal wastewater treated by 3D electrode

Conclusion

Enhanced ammonia removal was observed by electrolysis with bipolar 3D electrode reactor with RuO₂/Ti as anode, stainless steel as cathode and natural zeolite as particle electrode. Under various conditions, faster ammonia removal rates were achieved when compared with traditional 2D electrode reactor. Current density and initial ammonia concentration was found to have significant effect on the ammonia removal rate while the effect of chloride ion concentration and pH was weak. Moreover, 3D electrode reactor showed good performance for ammonia removal in treating actual municipal sewage. 27.8 mg N/L ammonia could be reduced to 0.5 mg N/L within 1.5 h, which met the demand of The First Grade of Environment Quantity Standards for surface water in China.

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