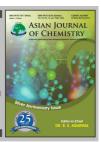
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# Removal of Chemical Contaminants of Micro-Polluted Water from Wastewater Treatment Plant Through Electrochemical Oxidation

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The electrochemical oxidation was adopted to further treat secondary effluent of wastewater treatment plants. The removal effect of major pollutants of constructed wetland effluent was mainly investigated and the main chemical reaction path of the removal of organic matter and nitrogen compounds by electrochemical method was discussed preliminarily. The results showed the best removal ratio of NH<sub>3</sub>-N, total nitrogen and total phosphorus was 94.2, 82.8 and 60.5 %, respectively after dosing NaCl on the condition of different electrolysis time and constant current 5A (namely the current density 7.4 mA/cm²). Meanwhile, the effect was better than no NaCl. If other experimental conditions remained unchanged, the major pollutants were removed faster by electrolysis. Considering power consumption, it suggested that the best operation condition was of some salt solution, the current density 7.4 mA/cm² and 0.5 h.

Key Words: Secondary effluent of wastewater treatment plants, Electrolysis time, Electrochemical removel, Constructed wetland.

#### INTRODUCTION

Electrochemical oxidation treatment of water is a relatively new type of water treatment technology, which gradually develops into a very promising technology<sup>1,2</sup>. Because of easy controlling, lesser floor areas and no secondary pollution, it has been successfully applied in the purification of wastewater from many field of water treatment<sup>3-5</sup>. In addition, certain research experience to micro-polluted water using electrochemical method has been gained in the past several years. The experience proved that the method had great removals to NH<sub>3</sub>-N and total nitrogen<sup>6,7</sup>. Secondary effluent of wastewater treatment plants is further treated exiguously. In most instances, the effluent is directly discharge to rivers, and it will bring about river pollutions marginally<sup>8</sup>. Thus, the secondary effluent of wastewater treatment plants is necessary to be treated to reduce any possible impacts on rivers<sup>9-12</sup>.

Accordingly, this study focuses on using electrochemical method for treating secondary effluent of wastewater treatment plants. The electrochemical method was applied to advanced treatment of secondary effluent, and the hybrid process which made final effluent beat the target to environmental quality standards for surface water was discussed, too.

## **EXPERIMENTAL**

**Electrolytic system:** A laboratory batch electrochemical oxidation reactor was designed and constructed to the dimensions

shown in Fig. 1. The electrolytic cell was made of synthetic glass, with a dimension of  $140 \text{ mm} \times 90 \text{ mm} \times 130 \text{ mm}$ , and the dimensions of the electrodes were  $130 \text{ mm} \times 65 \text{ mm} \times 1 \text{ mm}$ . During electrolysis, the current density was adjusted by using a DC power supply (MPS702). The reactor consisted of an undivided cell made of synthetic glass with a cathode and anode arranged in parallel. Stainless steel plates were used as cathodes while, horizontal Ti/Ru/Cr plates acted as anodes. The electrodes were arranged in the electrochemical cell of 1 L working value.

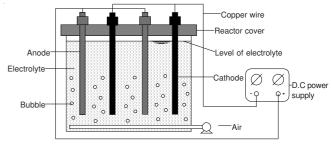


Fig. 1. Schematic diagram of the electrolytic system

**Analytical methods:** The experimental constants in electrolytic contaminants degradation were interelectrode distance, current density and initial Cl<sup>-</sup> concentration. The concentration of NaCl solution was 5.0 mol/L, the dosing ratio was 8 mL/h.

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And the removals of contaminants were changed at a time while the others were kept constant of definite salt concentration, the current density 7.4 mA/cm<sup>2</sup> and interelectrode distance 1 cm.

**Influent quality:** The raw wastewater, secondary effluent of wastewater treatment plants, was collected from wastewater treatment plant in Tangshan. The composition of the influent used in all experiments is shown in Table-1.

| TABLE-1                               |                    |                |
|---------------------------------------|--------------------|----------------|
| CHARACTERISTICS OF THE WASTEWATER     |                    |                |
| SAMPLE USED IN THE EXPERIMENTS        |                    |                |
| Parameter                             | Unit               | Concentration  |
| pH                                    | -                  | $7.2 \pm 0.2$  |
| Ammonia nitrogen (NH <sub>3</sub> -N) | mg L <sup>-1</sup> | $1.90 \pm 0.5$ |
| Total nitrogen (TN)                   | mg L <sup>-1</sup> | $3.25 \pm 0.9$ |
| Total phosphorus (TP)                 | mg L <sup>-1</sup> | $0.38 \pm 0.1$ |

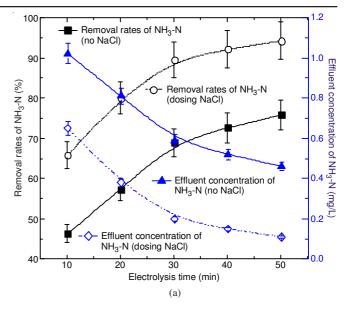
# RESULTS AND DISCUSSION

Removal of nitrogenous compounds: As shown in Fig. 2, when current density was 7.4 mA/cm<sup>2</sup> and the interelectrode distance was 1 cm, the NH<sub>3</sub>-N and total nitrogen in constructed wetland were removed remarkably, too. Because of the lowconcentration NH<sub>3</sub>-N, the removals of NH<sub>3</sub>-N with direct electrolysis were lower than the dosing NaCl solution. The maximum rates were 75.8 and 94.2 %, respectively. On the contrast, the maximum removal rates of total nitrogen were 71.4 and 82.8 %, the concentration of total nitrogen could attain 0.56 mg/L after dosing NaCl solution, which could satisfy the environmental quality standards for surface water to full advantage. In addition, the removals of the electrolytic oxidation of NH<sub>3</sub>-N and total nitrogen increase generally with the retention time, too. Meanwhile, the removal principles of NH<sub>3</sub>-N and total nitrogen is that dosing NaCl solution could accelerate the rates of generating oxidizability substances, so it had higher removals to certain contaminants. The oxidizing ability of electrodes attained the relative limit when the electrolysis time was 0.5 h. At that time, the reduced oxidizability substances made the oxidation rate slacken remarkably, so the removals of NH<sub>3</sub>-N and total nitrogen increased tardily. But the effluent concentration had been satisfied our demand. The processes not only ensured the removal effects, but also saved the energy consumption, too.

Removal of total phosphorus: As is shown in Fig. 3, when current density was 7.4 mA/cm<sup>2</sup> and the interelectrode distance was 1 cm, the total phosphorus was also removed remarkably. The removals of NH<sub>3</sub>-N with direct electrolysis were lower than the dosing NaCl solution. The maximum rates were 60.5 and 42.1 %, respectively. The concentration of total phosphorus could attain 0.15 mg/L after dosing NaCl solution. According above stated, when the electrolysis time exceeded 0.5 h, the removals of total phosphorus increased tardily, too. In addition, the removals of phosphorus may be related to the electro-flocculation.

Course of chemical reaction: During electrolytic reaction,  $O_2$  was generated by the anode and dissolved oxygen in wastewater generated  $H_2O_2$  at the cathode.

$$O_2 + e^- \longrightarrow O_2^- \tag{1}$$



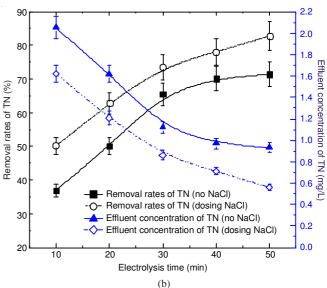


Fig. 2. Removal effect of nitrogenous compounds was showed when the electric current was constant with the electrolysis time prolonging

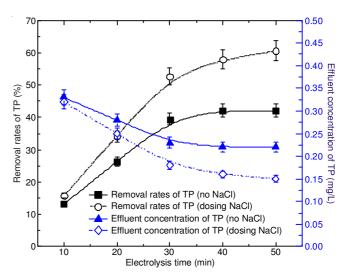


Fig. 3. Removal effect of total phosphorus was showed when the current density was constant with the electrolysis time prolonging

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$$O_2^- + H^+ \longrightarrow H_2O^{\bullet}$$
 (2)

$$2H_2O^{\bullet} \longrightarrow H_2O_2 + O_2 \tag{3}$$

Another course of chemical reaction may possibly happen,

$$O_2^- + H_2O^{\bullet} \longrightarrow O_2 + H_2O^-$$
 (4)

$$H_2O^- + H^+ \longrightarrow H_2O_2$$
 (5)

and HO<sup>•</sup>, hydroxyl free radical was generated further by H<sub>2</sub>O<sub>2</sub>.

$$H_2O_2 + e^- \longrightarrow HO^- + HO^{\bullet}$$
 (6)

HO\* could oxidize organic and nitrogenous compounds and the overall reaction occurring in the anodic solution between hypochlorite and ammonia can be expressed as follows:

$$2NH_4^+ + 3ClO^- \longrightarrow N_2 + 3H_2O + 2H^+ + 3Cl^-$$
 (7)

## Conclusion

The experimental results indicated that the best operating condition was of some salt solution, the current density 7.4 mA/cm² and electrolysis time 0.5 h. In addition, the removals of contaminants with direct electrolysis were lower than the dosing NaCl solution. Moreover, the results showed the best removal ratio of NH<sub>3</sub>-N, total nitrogen and total phosphorus were 94.2, 82.8 and 60.5 %, respectively after dosing NaCl on the condition of different electrolysis time and constant current 5A (namely the current density 7.4 mA/cm²). The main indexes

of electrolysis effluent could satisfy the environmental quality standards for surface water to a great extent.

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