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## ORIGINAL ARTICLE

# Adsorption characteristics of sulfur solution by acticarbon against drinking-water toxicosis

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## KEYWORDS

Acticarbon;  
Drinking-water toxicosis;  
Fe<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>;  
Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub>;  
Na<sub>2</sub>SO<sub>3</sub>

**Abstract** Sulfur and ammonia nitrogen are rich nutrient pollutants, after entering water can cause algal blooms, cause eutrophication of water body, the spread of them will not only pollute the environment, destroy the ecological balance, but also harm human health through food chain channels, especially drinking-water toxicosis. Acticarbon can adsorb harmful substances, it was beneficial for people's health. In order to figure out the optimal adsorption condition and the intrinsic change of acticarbon, five chemicals were adsorbed by acticarbon and analyzed by FT-IR. The optimal adsorption condition of Fe<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>, Na<sub>2</sub>SO<sub>4</sub>, Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub>, S and Na<sub>2</sub>SO<sub>3</sub> was 9 g/1000 g at 80 min, 21 g/1000 g at 20 min, 15g/1000 g at 20 min, 21 g/1000 g at 60 min and 21 g/1000 g at 100 min, respectively. FT-IR spectra showed that acticarbon had eight characteristic peaks, such as S-S stretch, H<sub>2</sub>O stretch, O—H stretch, —C—H stretch, C=O or C=C stretch, CH<sub>2</sub> bend, C—H were at 3850 cm<sup>-1</sup>, 3740 cm<sup>-1</sup>, 3435 cm<sup>-1</sup>, 2925 cm<sup>-1</sup>, 1630 cm<sup>-1</sup>, 1390 cm<sup>-1</sup>, 1115 cm<sup>-1</sup>, 600 cm<sup>-1</sup>, respectively. For FT-IR spectra of Fe<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>, the peaks at 3850 cm<sup>-1</sup>, 3740 cm<sup>-1</sup>, 2925 cm<sup>-1</sup> achieved the maximum with 9 g/1000 g at 20 min. For Na<sub>2</sub>SO<sub>4</sub>, the peaks at 2925 cm<sup>-1</sup>, 1630 cm<sup>-1</sup>, 1390 cm<sup>-1</sup>, 1115 cm<sup>-1</sup>, 600 cm<sup>-1</sup> achieved the maximum with 21 g/1000 g at 120 min. For ones of Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub>, the peaks at 3850 cm<sup>-1</sup>, 3740 cm<sup>-1</sup>, 1390 cm<sup>-1</sup>, 1115 cm<sup>-1</sup>, 600 cm<sup>-1</sup>, achieved the maximum with 2 g/1000 g at 80 min. For ones of S, the peaks at 3850 cm<sup>-1</sup>, 3740 cm<sup>-1</sup>, 2925 cm<sup>-1</sup> achieved the maximum with 19 g/1000 g at 100 min, the peaks at 1390 cm<sup>-1</sup>, 1115 cm<sup>-1</sup>, 600 cm<sup>-1</sup> achieved the maximum with 19 g/1000 g at 20 min. For FT-IR spectra of Na<sub>2</sub>SO<sub>3</sub>, the peaks at 1630 cm<sup>-1</sup>, 1390 cm<sup>-1</sup>, 1115 cm<sup>-1</sup>, 600 cm<sup>-1</sup> achieved the maximum with 2 g/1000 g at 100 min. It provided that acticarbon could adsorb and desulphurize from sulfur solution against drinking-water toxicosis.

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## 1. Introduction

Sulfur and ammonia nitrogen mainly come from chemical fertilizers, processed meat, leather and other industry emissions of industrial waste water, and city life sewage and farmland irrigation and drainage (He et al., 2008; Mao et al., 2000; Chai et al., 2010). Sulfur and ammonia nitrogen are rich nutrient pollutants, after entering water can cause algal blooms, cause eutrophication of water body, the spread of them will not only pollute the environment, destroy the ecological balance, but also harm human health through food chain channels, such as drinking-water toxicosis. Ecological effects of acid precipitation can be determined from the timing of changes in lake chemistry or acid-sensitive micro fossils and metallic pollutants in sediments. This is thought to be a result of a few factors: increased construction of large power plants and smelters with tall smokestacks coupled with a decrease in use of coal for home heating, converting the local air pollution problem into along-range, transboundary one; emissions of  $\text{NO}_x$  and other pollutants that aid in the oxidation of sulfur and nitrogen oxide have increased; and it took years for lakes, streams and their catchments to lose their buffering capabilities, so that lower pH levels were not recognized until sometime after the precipitation became acidic. Anthropogenic emissions are comparable to natural emissions on the global level, but regionally over 90% of sulfur deposited from the atmosphere is anthropogenic (Schindler, 1988).

Acticarbon can use almost any type of carbon materials, such as wood (Wang et al., 2009a), sawdust (Zhang et al., 2010), coal (Ahmadpour and Do, 1996), shells (Chandra et al., 2009), the stone of the fruit (Jumasiah et al., 2005), bagasse, oil waste, waste plastics (Zhou et al., 2007), paper and leather scrap (Yuan et al., 2004), waste tires and urban waste (Wang, 2004). Acticarbon with highly developed porous structure and huge specific surface area (Ding et al., 2002), good chemical stability and thermal stability, high mechanical strength and surface contains a variety of oxygen containing functional groups (Yu et al., 2005 and Pu and Jiang, 2005). What's more, acticarbon, which contained potassium, calcium and other minerals, could have adsorption and filtration of extractives, oil, other matters (Peng et al., 2013a,b,c, 2014a, 2012, 2011; Xiao et al., 2013; Wang et al., 2013; Peng and Le, 2012; Liu et al., 2008; Zhang et al., 2008; Qi et al., 2012). The fabric inhibited bacterial metabolism causing fewer allergic skin reactions than other fibers sterilized with antimicrobial agents. Because the trait was due to the highly porous structure of the bamboo fabric, it could absorb sulfur-based compounds, nitrogen-based compounds and so on (Milena

et al., 2003; Ikuo et al., 2001; Masakazu et al., 2003; Kei et al., 1994; Wang et al., 2006; Xue et al., 2014; Cui et al., 2014). In order to figure out the optimal adsorption condition and the intrinsic change of the acticarbon, five chemicals were adsorbed by acticarbon and were analyzed by FT-IR.

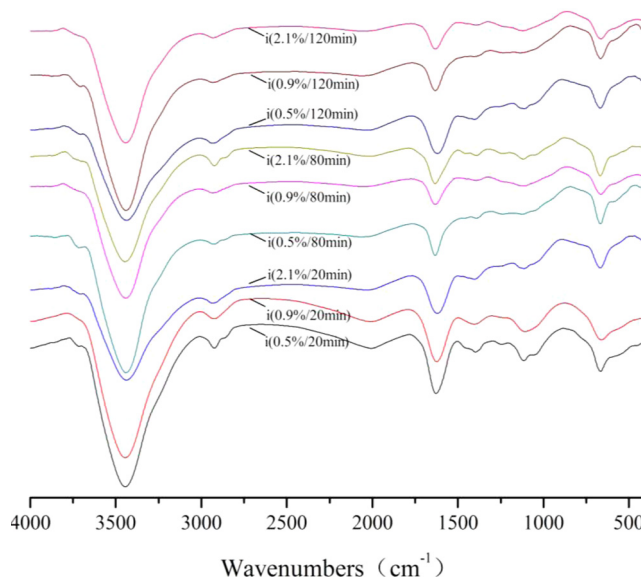
## 2. Materials and methods

### 2.1. Materials

Acticarbon,  $\text{Fe}_2(\text{SO}_4)_3$ ,  $\text{Na}_2\text{SO}_4$ ,  $\text{Na}_2\text{S}_2\text{O}_8$ , S, and  $\text{Na}_2\text{SO}_3$  were purchased from the market.

### 2.2. Methods

The  $\text{Fe}_2(\text{SO}_4)_3$  powder was weighed in quantities of 5 g, 9 g, and 21 g. These powder and 4 g over dry acticarbon were put into beaker which equipped with 1000 ml water, respectively. It was stirred in a beaker for 20 min, 80 min and 120 min. The  $\text{Na}_2\text{SO}_4$  powder was weighed in quantities of 11 g, 19 g, and 21 g. These powder and 4 g over dry acticarbon were put into beaker which equipped with 1000 ml water, respectively. It was stirred in a beaker for 20 min, 60 min and 120 min, respectively. The  $\text{Na}_2\text{S}_2\text{O}_8$  powder was weighed



**Figure 1** FT-IR spectra of acticarbon during adsorption of  $\text{Fe}_2(\text{SO}_4)_3$  solution.

**Table 1** Adsorption results.

SC (%)	Fe <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub>			SC (%)	Na <sub>2</sub> SO <sub>4</sub>			SC (%)	Na <sub>2</sub> S <sub>2</sub> O <sub>8</sub>			SC (%)	S			SC (%)	Na <sub>2</sub> SO <sub>3</sub>		
	Stir time (min)				Stir time (min)				Stir time (min)				Stir time (min)				Stir time (min)		
	20	80	120		20	60	120		20	80	100		20	60	100		20	40	100
0.5	3.76	2.49	1	1.1	1.25	0.75	2.51	0.2	0.5	1.79	1.03	0.2	7.75	4.52	4.73	0.2	0	2.74	1.75
0.9	1.75	23.7	0.5	1.9	3.52	1.99	2.26	1.1	8.25	3.23	2.23	1.9	37.3	58.6	11.5	1.1	2.74	2.74	2.26
2.1	17.5	18.5	2.75	2.1	8.21	1.75	2.99	1.5	20.1	12.7	2.19	2.1	37.7	80.3	32.8	2.1	1.25	2.26	4.26

Note: SC – Concentration of sulfur solution.

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