Removal of bromate ion using powdered activated carbon

Lian Wang¹,², Jie Zhang², Jingze Liu¹,*, Hong He², Min Yang², Jianwei Yu², Zichuan Ma³, Feng Jiang⁴

¹. College of Life Science, Hebei Normal University, Shijiazhuang 050016, China. E-mail: lianwang@mail.ustc.edu.cn
². Research Center for Eco-Environmental Sciences, Chinese Academy of Sciences, Beijing 100085, China
³. College of Chemistry & Material Science, Hebei Normal University, Shijiazhuang 050016, China
⁴. Beijing YADU Indoor Environmental Protection Science and Technology Co., Ltd., Beijing 102206, China

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Abstract

Bromate ion (BrO₃⁻) removal from drinking water by powdered activated carbons (PAC₅) in bath mode was evaluated under various operational conditions. Six kinds of PACs, including wood-based carbon, fruit-based carbon, coal-based carbon, and these three carbons thermally deoxidized in a nitrogen atmosphere, were selected to investigate their capacity on BrO₃⁻ removal. With the highest zeta potential value and being richly mesoporous, coal-based carbon had a high and an excellent BrO₃⁻ adsorption efficiency. The removal content of BrO₃⁻ by per gram of coal-based carbon was 0.45 mg within 5 hr in 100 µg/L bromate solution. The surface characteristics of PACs and bromide formation revealed that both physical and chemical PACs properties simultaneously affected the adsorption-reduction process. Under acidic conditions, PACs possessed high zeta value and adequate basic groups and exhibited neutral or positive charges, promoting BrO₃⁻ adsorption-reduction on the carbon surface. Interestingly, the PACs thermally deoxidized in N₂ atmosphere optimized their properties, e.g. increasing their zeta values and decreasing the oxygen content which accelerated the BrO₃⁻ removal rate. The maximum adsorption capacity of fruit-based carbon was the highest among all tested carbons (99.6 mg/g), possibly due to its highest pore volume. Remarkably, the thermal regeneration of PACs in N₂ atmosphere could completely recover the adsorption capacity of PACs. The kinetic data obtained from carbons was analyzed using pseudo second-order and intraparticle diffusion models, with results showing that the intraparticle diffusion was the more applicable model to describe adsorption of BrO₃⁻ onto PACs.

Key words: bromate; powdered activated carbons; adsorption-reduction process; adsorption capacity

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Introduction

The presence of bromate in drinking water has attracted much attention because it is an animal carcinogen (Marhaba and Bengraïne, 2003; Wolf et al., 1998). In the process of ozonation, bromate can be formed through complex reactions on molecular ozone and hydroxyl radical (von Gunten and Hoigné, 1994). Ozonation is a disinfection method that can destroy microorganisms, reduce the color, odor, and total organic carbon, etc. In the process of ozonation, bromide is oxidized to hypobromous acid and hypobromite, which are further oxidized to bromate by ozone and/or hydroxyl radicals. In general, the concentration of bromate in drinking water ranges from 0.4 to 100 µg/L (Butler et al., 2005; Krasner et al., 1993). A maximum allowed contaminant level of bromate is 10 µg/L by the European Union (1980), while the World Health Organization set a provisional guideline value of 25 µg/L (WHO, 1996).

Up to now, three approaches have been used to reduce the concentration of bromate in water. One approach is to remove the bromate precursors, such as bromide and natural organic matter before ozonation process (Johnson and Singer, 2004; Marhaba and Bengraïne, 2003). The second one is to control the bromate formation during ozonation through pH control by adding ammonia or hydrogen peroxide, and by modifying ozonation operation (Boulard et al., 2003; Kim et al., 2007). The third approach is using physical and chemical methods to remove bromate after ozonation. Most research works have been focused on the use of activated carbons (especially granular activated carbon) to remove bromate owing to the high removal efficiency of activated carbon (Bao et al., 1999; Huang et al., 2007; Huang and Cheng, 2008; Kirisits et al., 2000; Siddiqui et al., 1996). However, the granular activated carbon capacity is carbon-specific and depends on the source water (Bao et al., 1999; Kirisits et al., 2000).

The mechanisms of BrO₃⁻ removal by activated carbon have been proposed by some researchers (Bao et al., 1999; Kirisits et al., 2000; Siddiqui et al., 1996; Studebaker, 1957). It has been widely accepted that the removal of BrO₃⁻ by activated carbon is postulated to be adsorbed first, then reduced to hypobromite (BrO⁻), and finally...
reduced to bromide (Br\textsuperscript{−}) on activated carbon surface (Siddiqui et al., 1996). Previous report found that activated carbons with less surface oxygen tend to be basic and have an anion exchange capacity because of their positive charge (Chingombe et al., 2005). It means that some reduction methods such as N\textsubscript{2} reduction treatment could decrease the surface oxygen of activated carbons, such that increase the removal rate of BrO\textsubscript{3}\textsuperscript{−} on activated carbons. Moreover, Siddiqui et al. (1996) concluded that the increase in the acidity of carbon could compromise the adsorption of BrO\textsubscript{3}\textsuperscript{−} due to unfavorable electrostatic interactions between anions and acid groups. Huang et al. (2007) reported that the removal capacity of BrO\textsubscript{3}\textsuperscript{−} increased with decreasing solution pH. Therefore, further work is needed to confirm the e

The purpose of this study was to elucidate how the physical and chemical characteristics of powdered activated carbon influence BrO\textsubscript{3}\textsuperscript{−} removal in bath mode. Six kinds of powdered activated carbons (PACs) (wood-based carbon, fruit-based carbon, coal-based carbon, and three carbons thermally deoxidized in N\textsubscript{2} atmosphere) were characterized using nitrogen adsorption, X-ray photoelectron spectroscopy (XPS), zeta potential measurement, and acid-base titrations. The adsorption capacities of the various PACs were compared and the adsorption kinetics of BrO\textsubscript{3}\textsuperscript{−} on PACs were developed.

1 Experimental

1.1 Materials

Three types of commercial PACs, wood-based carbon (PAC\textsubscript{W}) (Shanxi Xinhua Carbon Corporation, China), one fruit-based carbon (PAC\textsubscript{F}) (Tangshan Huaneng Carbon Corporation, China), and one coal-based carbon (PAC\textsubscript{C}) (Ningxia Taixi Carbon Corporation, China) were used in this study. Prior to experiments, the samples were washed using ultra-pure water and dried overnight at 110°C to remove excess water, then cooled and stored in a desiccator. In order to determine their surface characteristics, the above three carbons were thermally treated at 600°C in a vacuum condition, with a pressure lower than 3 x 10\textsuperscript{−}7 Pa. The high-resolution scans were performed over 524−544 eV (O 1s spectrums). For calibration purposes, the C 1s electron bond energy corresponding to graphitic carbon was referenced to 284.8 eV.

Acid-base titration method was applied to estimate the number of acid and base groups on the surface of activated carbon samples. Acid/base depletion from 0.1 mol/L HCl and NaOH solutions was performed on each sample to determine the acid/base characteristics of carbons (Barton, 1997).

1.3 Kinetic adsorption measurement

The experiments of BrO\textsubscript{3}\textsuperscript{−} removal by PACs were performed using a bath shaker at 120 r/min. A suitable dose of carbon was introduced into a 250-mL amber glass bottles containing 100 µg/L or 500 µg/L BrO\textsubscript{3}\textsuperscript{−} solution. The pH of BrO\textsubscript{3}\textsuperscript{−} solution was 5.6 unless otherwise stated. Then, they were placed on a shaker table for a specified time and filtered through pre-washed 0.45 µm filters, and the filtrates were analyzed by ion chromatography.

1.4 Maximum adsorption capacity measurement

The maximum adsorption capacity of PACs was obtained using Langmuir isotherm model (Eq. (1)):

\[
\frac{1}{q_e} = \frac{1}{q_m} + \frac{1}{k_q q_m} \times \frac{1}{c}
\]

where, \( q_e \) and \( c \) are the concentrations of BrO\textsubscript{3}\textsuperscript{−} in the PACs and equilibrium solution, respectively, \( q_m \) represents the maximum adsorption capacity, and \( k_q \) is the Langmuir adsorption constant that is related to the adsorption energy. To determine the isothermal (20 ± 1°C) adsorption capacity, 1000 mg/L PACs was added into BrO\textsubscript{3}\textsuperscript{−} solutions with various concentrations. After 24 hr for equilibrium adsorption, 1 mL mixture solution was extracted and filtered through pre-washed 0.45 µm filters for analyzing by ion chromatography.
1.5 Analysis

The BrO\textsuperscript{3–} measurement was conducted using an ion chromatography (ICS-1500, Dionex, Canada), which was equipped with a dual-piston pump, an UV/Vis detector (UVD-500, Dionex, Canada), an AS50 autosampler, and IonPac AS23 separator columns (UVD-500, 4 mm, Dionex, Canada). The eluent solution was 9 mmol/L Na\textsubscript{2}CO\textsubscript{3}. The chromogenic agents were 1.2 mmol/L NaNO\textsubscript{2} and the mixture solution of 1.5 mol/L KBr and 1.0 mol/L H\textsubscript{2}SO\textsubscript{4}. The minimum detection limit for BrO\textsuperscript{3–} was 1.2 µg/L. Calibrations were performed before each series of samples was analyzed. The yield of bromine was determined using an ion chromatograph consisting of a pretreatment column (IonPacAG14A-SC, 4 mm, Dionex, Canada), a separator (AS14A-SC, 4 mm, Dionex, Canada), a suppressor (ASRS-ULTRA, 4 mm, Dionex, Canada), and a pulse electronic chemical detector (Dionex, Canada). The eluent solution was 9 mmol/L Na\textsubscript{2}CO\textsubscript{3}, and IonPac AS23 separator columns (UVD-500, 4 mm, Dionex, Canada). The BrO\textsuperscript{3–} was detected at a flow rate of 1.2 mL/min and the injection volume was 25 µL.

2 Results and discussion

2.1 Physical characterization of activated carbon

Table 1 shows the porous properties of carbon samples. All the samples exhibited high surface area and possessed both mesopores and micropores. PACF carbon had the highest V\textsubscript{meso} value (0.38 cm\textsuperscript{3}/g) and pore volume (0.87 cm\textsuperscript{3}/g). For the three original carbon samples, the ranking of total pore volume was in accordance with V\textsubscript{meso} volume as PAC\textsubscript{W} < PAC\textsubscript{C} < PAC\textsubscript{F}. These differences are ascribed to the nature of the primary raw material and the manufacturing processes. After thermally reduction with N\textsubscript{2}, the pore volume and V\textsubscript{meso} volume of PAC\textsubscript{F} and PAC\textsubscript{W} carbons slightly increased, while those of the PAC\textsubscript{C} slightly decreased.

Table 1 Porous properties of the powdered activated carbons (PACs) and PACs treated with N\textsubscript{2} after saturation adsorption

<table>
<thead>
<tr>
<th>PACs</th>
<th>Surface area (m\textsuperscript{2}/g)</th>
<th>Pore diameter (nm)</th>
<th>V\textsubscript{meso} (cm\textsuperscript{3}/g)</th>
<th>V\textsubscript{micro} (cm\textsuperscript{3}/g)</th>
<th>Pore volume (cm\textsuperscript{3}/g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PAC\textsubscript{W}</td>
<td>828.0</td>
<td>976.0</td>
<td>2.46</td>
<td>2.80</td>
<td>0.24</td>
</tr>
<tr>
<td>PAC\textsubscript{F}</td>
<td>1102.0</td>
<td>940.2</td>
<td>2.55</td>
<td>2.51</td>
<td>0.38</td>
</tr>
<tr>
<td>PAC\textsubscript{C}</td>
<td>688.8</td>
<td>716.1</td>
<td>3.07</td>
<td>3.05</td>
<td>0.33</td>
</tr>
<tr>
<td>PAC\textsubscript{W} \ deoxidized with N\textsubscript{2}</td>
<td>841.1</td>
<td>880.4</td>
<td>2.81</td>
<td>2.66</td>
<td>0.31</td>
</tr>
<tr>
<td>PAC\textsubscript{C} \ deoxidized with N\textsubscript{2}</td>
<td>1270.0</td>
<td>1013.2</td>
<td>2.48</td>
<td>2.45</td>
<td>0.44</td>
</tr>
<tr>
<td>PAC\textsubscript{C} \ deoxidized with N\textsubscript{2}</td>
<td>773.1</td>
<td>711.0</td>
<td>2.95</td>
<td>3.00</td>
<td>0.22</td>
</tr>
</tbody>
</table>

2.2 Chemical characterization of activated carbons

The analysis of the contents of oxygen and carbon composition as well as deconvolution results of O 1s core-level XPS spectra of PACs samples are presented in Fig. 1 and Table 3. The O1s core-level spectra indicated that the C=O (about 531.5 eV) and C–O (about 533 eV) groups existed in the samples, whose appearance might be due to the existence of carboxylic acids, lactones, or phenolic species (Pigamo et al., 2002). Clearly, the amounts of oxygen in C=O and total oxygen in PAC\textsubscript{F} were the lowest among all original carbons. Additionally, total oxygen...
Fig. 1  XPS survey spectra of the PACs samples (a), O1s core-level XPS spectra of the PACs samples (b). The dots represent the actual experimental data; the black lines are the fitting curves and baselines fitted by the Shirley function. The dash lines represent the fitting peaks of C=O and C–O.
Table 3  Distribution of oxygen-bearing structures (at.%) from O 1s core-level XPS spectra (Fig. 1) and the atomic ratios of O and C on PACs surfaces

<table>
<thead>
<tr>
<th>PACs</th>
<th>C=O</th>
<th>C–O</th>
<th>H2O adsorbed</th>
<th>Total O (at.%)</th>
<th>C (at.%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PACW</td>
<td>4.03</td>
<td>2.84</td>
<td>2.07</td>
<td>8.94</td>
<td>91.06</td>
</tr>
<tr>
<td>PACF</td>
<td>2.86</td>
<td>2.23</td>
<td>1.40</td>
<td>6.49</td>
<td>93.51</td>
</tr>
<tr>
<td>PACC</td>
<td>3.38</td>
<td>3.04</td>
<td>1.77</td>
<td>8.19</td>
<td>91.81</td>
</tr>
<tr>
<td>PACW deoxidized with N2</td>
<td>2.28</td>
<td>1.70</td>
<td>0.60</td>
<td>4.58</td>
<td>95.42</td>
</tr>
<tr>
<td>PACF deoxidized with N2</td>
<td>1.88</td>
<td>1.22</td>
<td>0.19</td>
<td>3.29</td>
<td>96.71</td>
</tr>
<tr>
<td>PACC deoxidized with N2</td>
<td>3.36</td>
<td>1.81</td>
<td>0.01</td>
<td>5.18</td>
<td>94.59</td>
</tr>
</tbody>
</table>

Contents, as well as the contents of C=O and C–O groups of PACs deoxidized carbons with N2 at 600°C declined largely, which means PACs could be deoxidized efficiently through N2 treatment.

2.3 Adsorption kinetic and adsorption volume studies

Figure 2 presents batch kinetic removal curves of adsorption capacity versus contact time for the six carbons. The adsorption capacity of BrO₃⁻ by activated carbons varied significantly with the type of carbons, ranging from 0.01 mg/g for PACW to almost complete adsorption (0.43 mg/g) for PACC within 5 hr, and PACF showed a moderate removal rate. As compared with the published results (Siddiqui et al., 1996), PACC used in this study showed an excellent efficiency for removal of BrO₃⁻. The quickest adsorption of PACC might be due to the highest zeta potential, high Vmeso, and less C=O groups, since surface properties and Vmeso volume are reported to be important to the adsorption of carbon (Huang et al., 2007; Huang and Cheng, 2008; Siddiqui et al., 1996). The adsorption rate of PACW in this study was the lowest, which was contrary with the previous reported results (Huang and Cheng, 2008), this might be owing to the different volume of mesopore, zeta potential, and oxygen content on account of different manufacturing processes. Surely, our reduction method of optimization on the nature of carbons was significant for the study on the adsorption capacity of BrO₃⁻. The thermally deoxidized carbons greatly promoted the adsorption rate of carbons through the increases of zeta potential and the obvious decrease of oxygen content.

Figure 3 shows the maximum adsorption capacity of PACs calculated with Langmuir isotherm model. The maximum adsorption capacity of PACF was about 99.6 mg/g, which was the highest saturation adsorption capacity because of its highest pore volume among all tested carbons. It is surprising to note the maximum adsorption capacity of the thermally deoxidized carbons largely decreased. Considering pore volume values of activated carbons, it could be found that the pore volume and Vmeso volume were somewhat related to their maximum adsorption capacity, However, the reason need to be clarified by further work. Regeneration experiments of carbon samples were also performed. The maximum adsorption capacity of recovered PACs was almost the same to their original state, indicating that the adsorption capacity of PACs could be recovered by thermal regeneration and the recovered carbons could be recycled.

The PACC, PACF, PACC deoxidized with N2, and PACF deoxidized with N2 were optimally selected in the following experiments considering their high adsorption rates and short-time removal of BrO₃⁻. Figure 4 shows pH effects on the removal of BrO₃⁻. The removal rate of BrO₃⁻ increased with the decrease of pH value of BrO₃⁻ solutions. The negative zeta potential of PACs (Table 2) in high pH solutions was corresponding to the presence of negatively charged carboxylate anionic surface functional groups on the PACs (Siddiqui et al., 1996). In solution with lower pH values, the weakly acidic and basic functional groups were protonated, accompanying with the zeta potential of carbons turning into positive value. As a result, the electrostatic interaction between the carbon and bromate was promoted. The point of zero charge varies among carbons, and the literature describes carbons with points of

Figure 2  Effect of the PACs type on BrO₃⁻ removal. Initial concentration of BrO₃⁻ and PACs are 100 µg/L and 200 mg/L, respectively.

Figure 3  Maximum adsorption capacity of the PACs and PACs after thermal regeneration for BrO₃⁻ adsorption.
Fig. 4 Effect of pH on BrO$_3^-$ removal by PAC$_F$ (a) and PAC$_C$ (b). Initial concentration of BrO$_3^-$ and PACs are 100 µg/L and 200 mg/L, respectively.

Fig. 5 BrO$_3^-$ adsorption/reduction by the selected PACs and the formation volume of Br$^-$. Initial concentration of BrO$_3^-$ and PACs are 500 µg/L and 1000 mg/L, respectively.

The reduction contribution on the BrO$_3^-$ removal by PACs was evaluated. In the batch experiment with PACs, BrO$_3^-$ reduction accompanying with bromide formation were observed (Fig. 5). After 96 hr contact with PACs, 3.9 µmol/L BrO$_3^-$ was almost completely removed and 0.6 µmol/L bromide formed, indicating that approximately 20% of bromate was reduced to bromide and 80% of bromate adsorbed by PACs in the process of BrO$_3^-$ removal. Furthermore, the bromide concentration did not linearly increase as a function of time, which might be due to the adsorption process of bromide onto PACs (Bao et al., 1999). The release and diffusion of bromide in the bromide adsorption process were interactional and competitive, leading that bromide could not immediately release into the solution once formed.

In the process of BrO$_3^-$ removal, the adsorption rate was high at the initial stage and then quickly decreased to a pseudo platform stage (5–96 hr), which was similar to the process of bromide release. On the basis of this, it could be supposed that the adsorption and reaction processes reached pseudo homeostasis in the platform stage. The process of BrO$_3^-$ removal was complex, including the surface and inner diffusion of bromate, adsorption of bromate, formation of bromide, and the release and diffusion of bromide, and so on. The kinetics of adsorption of BrO$_3^-$ on the selected carbons was described using intraparticle diffusion models and pseudo second-order in this work. The validity of the two models could be determined using the linear plots of $q_t$ versus $t^{1/2}$ (Eq. (2)) and $t/q_t$ versus $t$ (Eq. (3)).

$q_t = k_q t^{1/2}$

where, $k_q$ is the intraparticle diffusion rate constant, $q_t$ is the amounts of BrO$_3^-$ adsorbed at time $t$ (Wu et al., 2001).

\[
\frac{t}{q_t} = \frac{1}{k_2 q_e^2} + \frac{1}{q_e} t
\]

where, $k_2$ is the equilibrium rate constant of pseudo second-order adsorption, $q_e$ is the amounts of BrO$_3^-$ adsorbed at equilibrium.

In order to more accurately describe the kinetics, the quick adsorption stage (before 5 hr) and the pseudo platform stage (5–96 hr) of the adsorption curves were simulated separately. Tables 4 and 5 present the correlation coefficient, standard deviation (SD), and standard error (SE) to compare quantitatively the fitness of the models. The values of SD and SE for pseudo second-order kinetics evidently exceed those in the intraparticle diffusion model. Furthermore, the values of $q_e$ (before 5 hr) and $q_e'$ (5–96 hr) calculated with pseudo second-order kinetics were not consistent with the experimental values (Fig. 5). In addition, the intraparticle diffusion model was strongly consistent with the experimental data, suggesting that the adsorption of BrO$_3^-$ on powdered activated carbon was diffusion-controlled.

zero charge ranging from approximately 3 to 9 (Karanfil et al., 1998; Siddiqui et al., 1994). Therefore, the electrostatic attraction between bromate and the carbon depend not only on the pH of solution but also on the type of carbon.
Table 4 Kinetic parameters of intraparticle diffusion model for the adsorption of BrO$_3^-$ on the selected PACs

<table>
<thead>
<tr>
<th>PACs</th>
<th>$k_p$ (mg/(g·hr$^{1/2}$))</th>
<th>SE</th>
<th>SD</th>
<th>$k'_p$ (mg/(g·hr$^{1/2}$))</th>
<th>SE</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>PAC_F</td>
<td>0.049</td>
<td>0.012</td>
<td>0.013</td>
<td>0.013</td>
<td>0.014</td>
<td>0.004</td>
</tr>
<tr>
<td>PAC_C</td>
<td>0.19</td>
<td>0.027</td>
<td>0.029</td>
<td>0.013</td>
<td>0.013</td>
<td>0.004</td>
</tr>
<tr>
<td>PAC_F deoxidized with N$_2$</td>
<td>0.32</td>
<td>0.079</td>
<td>0.084</td>
<td>0.0037</td>
<td>0.0017</td>
<td>0.010</td>
</tr>
<tr>
<td>PAC_C deoxidized with N$_2$</td>
<td>0.51</td>
<td>0.11</td>
<td>0.011</td>
<td>0.0036</td>
<td>0.0010</td>
<td>0.0063</td>
</tr>
</tbody>
</table>

Table 5 Kinetic parameters of second-order kinetic model for the adsorption of BrO$_3^-$ on the selected PACs

<table>
<thead>
<tr>
<th>PACs</th>
<th>$q_e$ (mg/g)</th>
<th>$k_2$ (g/(mg·hr))</th>
<th>SE</th>
<th>SD</th>
<th>$q_e'$ (mg/g)</th>
<th>$k_2'$ (g/(mg·hr))</th>
<th>SE</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>PAC_F</td>
<td>2.33</td>
<td>0.39</td>
<td>0.73</td>
<td>0.97</td>
<td>0.73</td>
<td>0.40</td>
<td>0.058</td>
<td>4.27</td>
</tr>
<tr>
<td>PAC_C</td>
<td>3.93</td>
<td>0.46</td>
<td>0.20</td>
<td>0.27</td>
<td>0.71</td>
<td>0.45</td>
<td>0.038</td>
<td>2.82</td>
</tr>
<tr>
<td>PAC_F deoxidized with N$_2$</td>
<td>10.09</td>
<td>0.49</td>
<td>0.14</td>
<td>0.19</td>
<td>3.88</td>
<td>0.46</td>
<td>0.022</td>
<td>1.63</td>
</tr>
<tr>
<td>PAC_C deoxidized with N$_2$</td>
<td>32.67</td>
<td>0.47</td>
<td>0.021</td>
<td>0.029</td>
<td>1.92</td>
<td>0.49</td>
<td>0.0031</td>
<td>2.29</td>
</tr>
</tbody>
</table>

3 Conclusions

The adsorption capacity and kinetics of BrO$_3^-$ on powdered activated carbon revealed that the characteristics of activated carbon influenced their capacity and rate for the adsorption-reduction process. The efficiency of PACs for BrO$_3^-$ removal depended on the type of PACs, contact time, and pH of BrO$_3^-$ solution. Analysis of various properties of PACs indicated that the carbons with rich mesopores, higher zeta potential value, and lower oxygen content could rapidly and effectively remove the more BrO$_3^-$.

Among the selected carbons, coal-based carbon had a high and an excellent efficiency on adsorption of BrO$_3^-$.

The PACs thermally deoxidized in N$_2$ atmosphere possessed higher zeta values and contained lower oxygen content than original carbons, leading to the rapid adsorption of BrO$_3^-$. The reproduction adsorption capacity of PACs could be completely recovered by thermal treatment in N$_2$ atmosphere and the recovered PACs could be recycled. Under acidic conditions, the PACs possessed high zeta values and adequate basic groups exhibited neutral or positive charges, which had a higher BrO$_3^-$ adsorption capacity.

The kinetic data obtained in this study using pseudo second-order and intraparticle diffusion models indicated that the intraparticle diffusion was more applicable model to describe the removal process and the rate determining step was apparently the rate of diffusion of BrO$_3^-$ through the laminar layer surrounding the surface of activated carbon. The combination of zeta potential value, mesopore volume, and oxygen content could be used as the effective indicators for the selection of activated carbon for the removal of BrO$_3^-$.  

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