Insights into simultaneous adsorption and oxidation of antimonite [Sb(III)] by crawfish shell-derived biochar: spectroscopic investigation and theoretical calculations

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Abstract Removal of antimonite [Sb(III)] from the aquatic environment and reducing its biotoxicity is urgently needed to safeguard environmental and human health. Herein, crawfish shell-derived biochars (CSB), pyrolyzed at 350, 500, and 650 °C, were used to remediate Sb(III) in aqueous solutions. The adsorption data best fitted to the pseudo-second-order kinetic and Langmuir isotherm models. Biochar produced at 350 °C (CSB350) showed the highest adsorption capacity (27.7 mg g⁻¹), and the maximum 78% oxidative conversion of Sb(III) to Sb(V). The adsorption results complemented with infrared (FTIR), X-ray photoelectron (XPS), and near-edge X-ray absorption fine structure (NEXAFS) spectroscopy analyses indicated that the adsorption of Sb(III) on CSB involved electrostatic interaction, surface complexation with oxygen-containing functional groups (C=O, O=C–O), π–π coordination with aromatic C=C and C–H groups, and H-bonding with –OH group. Density functional theory calculations verified that surface complexation was the most dominant adsorption mechanism, whilst π–π coordination and H-bonding played a secondary role. Furthermore, electron spin resonance (ESR) and mediated electro-chemical reduction/oxidation (MER/MEO) analyses confirmed that Sb(III) oxidation at the biochar surface was governed by persistent free radicals (PFRs) (•O₂⁻ and •OH) and the electron donating/accepting capacity (EDC/EAC) of biochar. The abundance of preferable surface functional groups, high concentration of PFRs, and high EDC conferred CSB350 the property of an optimal adsorbent/oxidant for Sb(III) removal from water. The encouraging results of this study call for future trials to apply suitable biochar for removing Sb(III) from wastewater at pilot scale and optimize the process.

Graphical abstract

Article highlights
● Crawfish-shell biochar (CSB) pyrolyzed at 350 °C showed the highest Sb(III) adsorption and oxidation.
● DFT calculations highlighted complexation, H-bonding and π–π interactions as key removal mechanisms.
● Sb(III) oxidation was mainly governed by persistent free radicals and electron transfer capacity of biochar.

Keywords Sorption · Heavy metal · Synchrotron · Density functional theory · Contaminated water

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1 Introduction

Antimony (Sb) is a metalloid that belongs to Group VA of the periodic table, and the element exhibits chemical and bio-toxicological characteristics similar to arsenic (As), where oxyanions of both the elements might transform between trivalent and pentavalent species under variable redox conditions (Nishad and Bhaskarapillai 2021; Bolan et al. 2022b). Antimony exists in four oxidation states (-III, 0, +III, and +V) of which Sb(III) and Sb(V) are the prevalent forms in the environment, with Sb(III) 10-time more toxic than Sb(V) (Xiong et al. 2020). In recent decades, natural biogeochemical release and anthropogenic activities such as mining, metallurgy, and widespread use of Sb-containing products (e.g., pigments, batteries, and flame retardants) have triggered the Sb contamination in the environment worldwide (Wei et al. 2020; Chen et al. 2022b). Considering the potential teratogenicity and carcinogenicity of Sb to human beings, the World Health Organization (WHO) has set the maximum total Sb limit in drinking water as 20 µg L\(^{-1}\) (Zhu et al. 2021). China and Japan have, however, set more stringent limits, which are 5 and 2 µg L\(^{-1}\), respectively (Xiong et al. 2020). Adsorption method using various adsorbents such as activated carbon, carbon nanotubes, and graphene has been found to be an effective and sustainable remediation strategy for the removal of metal(loid)s (Chen et al. 2022c). Nevertheless, high cost of these carbon-based materials has limited the practical potential for their large-scale applications. Therefore, it is of great importance to develop cost-efficient and eco-friendly materials for the removal of Sb anions from aquatic systems, especially for the highly toxic Sb(III) species.

Biochar is a low-cost carbonaceous material produced from the pyrolysis of biomass wastes, with high porosity and abundant surface functional groups (Altaf et al. 2021b; Wu et al. 2021). Biochar has been proven as an excellent adsorbent for the removal of toxic organic compounds (Huang et al. 2018a, b; Qin et al. 2018; Nie et al. 2021), and trace metal(loid)s such as Cd (Chen et al. 2021; Yin et al. 2021), Cr (Wei et al. 2019a; Xu et al. 2020), Pb (Wen et al. 2021; Yang et al. 2021), Hg (Altaf et al. 2021a, b; Liu et al. 2022), As (Pan et al. 2021; Yang et al. 2022), and Sb (Wan et al. 2020; Song et al. 2021; Zhu et al. 2021). Biochar can adsorb metal(loid) ions through multiple mechanisms such as pore filling, ion exchange, precipitation, and surface complexation with functional groups (Hu et al. 2020; Bolan et al. 2022a). To date, due to the limited adsorption/immobilization potential of pristine biochar for pollutants, various modification methods such as magnetization (Zhu et al. 2021), loading of zirconium (Rahman et al. 2021), and chitosan (Palansooriya et al. 2021) have been developed to strengthen biochar’s performance in pollutant removal in aquatic systems. The modification processes indeed enhance the adsorption capacity of biochar, yet the extra cost and energy consumption also inevitably rise (Rajapaksha et al. 2016). A prerequisite for the sustainable application of biochar as an effective adsorbent in wastewater treatment is its cost-effectiveness. The adsorption performance of biochar for Sb(III) removal would be closely related to its biomass feedstock source (Cui et al. 2017). Therefore, screening inexpensive and widely sourced feedstock for the preparation of biochar to effectively remove Sb(III) from water is a need of the hour.

Crawfish (Procambarus clarkii), distributed worldwide, is one of the common seafood for human (Zhou et al. 2021). China’s annual production and export of crawfish in 2018 reached 1.64 million t and 10,801 t, respectively (China Crawfish Industry Development Report 2019). Large-scale consumption of crawfish in the catering industry has rendered the production of a huge volume of crawfish shell bio-waste (Chen et al. 2020b). Approximately, 580,000 t of crawfish shell waste is generated every year in China (Ma et al. 2019). Thus, converting crawfish shells to biochar could be a sustainable solution for the management of this ever-increasing waste resource (Lv et al. 2020). Previous research reported the feasibility of crawfish shell-derived biochar (CSB) for the removal of metal cations including Cd\(^{2+}\), Pb\(^{2+}\), and Cu\(^{2+}\) where cation exchange and mineral precipitation appeared as the key adsorption mechanisms (Xiao et al. 2017; Ma et al. 2021; Zhang et al. 2021). The adsorption mechanism of Sb(III) by CSB might differ as compared to metal cations, which remains least understood in the literature. A few studies also reported that biochar could oxidize Sb(III) to Sb(V), decreasing the element’s biotoxicity (Wu et al. 2019; Wei et al. 2020; Chen et al. 2022b). However, the oxidation mechanism of Sb(III) on biochar surface through electron transfer reactions, especially those involving persistent free radicals (PFRs), was not thoroughly studied before.

Since pyrolysis temperature can greatly affect the physicochemical characteristics (e.g., abundance of functional groups and electron transfer capacities) of biochar, it can influence biochar’s removal efficiency for contaminants (Xiao et al. 2017). Understanding the effect of such characteristics on Sb(III) adsorption/oxidation is crucial for assessing the applicability of CSB for Sb(III) removal from water. This study reports the synthesis of CSB pyrolyzed at 350, 500, and 650°C and its subsequent application to remove Sb(III) from aqueous solutions. The specific objectives of this study were to: (1) determine the adsorption capacity of CSB for Sb(III) under different conditions (i.e., reaction time, initial adsorbate concentration, solution pH, ionic strength, and co-existing substances); (2) unravel the Sb(III) adsorption mechanism using multiple advanced
spectroscopic techniques and theoretical calculations; and (3) elucidate the pyrolysis temperature depended electron/PFRs mediating mechanisms in the oxidative conversion of Sb(III) to Sb(V) on CSB surface.

2 Materials and methods

2.1 Chemicals and reagents

All the used chemicals in this study were of analytical grade and were purchased from Macklin Bio-Chem Technology Co., Ltd. (Shanghai, China). The chemicals were directly used without further purification. All solutions were prepared with 18.2 MΩ cm$^{-1}$ deionized water (ULPHW-I, Ulupure Co. LTD., China). In addition, the standard solution containing 1000 mg L$^{-1}$ of Sb(III) was obtained from the National Research Center for Certified Reference Materials of China.

2.2 Biochar preparation and characterization

The crawfish shells were collected from a marketplace located in Hangzhou, Zhejiang Province, China. The crawfish shells were rinsed with tap water to remove the impurities, and then oven-dried at 80°C for 24 h prior to biochar production. The CSB was produced by pyrolyzing crawfish shells in a batch pyrolysis furnace at 350, 500, and 650°C under the oxygen-limited condition, with a heating rate of 15°C min$^{-1}$, and held for 2 h once the final temperature was reached. After pyrolysis, the obtained biochar was labeled as CSB350, CSB500, and CSB650. The biochar was ground, passed through a 100-mesh sieve (0.150 mm), and stored at room temperature for characterization and further experiments. Characterization of selected biochar using multiple spectroscopic techniques including synchrotron-based micro-X-ray fluorescence (µ-XRF) and near-edge X-ray absorption fine structure spectroscopy (NEXAFS) is shown in the Supplementary Material.

2.3 Batch adsorption experiments

A stock solution containing 1000 mg L$^{-1}$ of Sb(III) was prepared by dissolving potassium antimonyl tartrate trihydrate (C$_8$H$_2$K$_2$O$_7$Sb$_2$·3H$_2$O) in deionized water. The working solutions were prepared by the dilution of the stock solution using 0.01 M NaCl solution as the background electrolyte. Batch adsorption experiments were conducted using the methods reported in our previous publication (Chen et al. 2022b). All the adsorption experiments were carried out under dark conditions and replicated three times.

Kinetic adsorption experiments were conducted by adding 0.05 g of biochar (i.e., CSB350, CSB500, and CSB650) into a 25-mL working solution containing 40 mg L$^{-1}$ of Sb(III). The mixture was shaken (180 rpm) for 24 h at 25°C. The initial Sb(III) concentration (40 mg L$^{-1}$) was based on previous studies (30–50 mg L$^{-1}$) (Jia et al. 2020; Wei et al. 2020; Chen et al. 2022b). Samples were collected at desired time intervals (5, 10, 15, 30, 60, 120, 240, 360, 480, 600, 720, 960, 1440 min) to determine the residual Sb(III) concentrations in the solution. Adsorption isotherm experiments were conducted by adding 0.05 g of biochar, weighed into a 25-mL working solution with various initial Sb(III) concentrations. The mixture was oscillated at 180 rpm for 24 h at 25°C before the quantitative measurement of residual Sb(III) in the solution. The impact of initial solution pH on adsorption was investigated with a fixed Sb(III) initial concentration (40 mg L$^{-1}$) at 25°C. The initial solution pH was adjusted in the range of 2–11 using HCl (0.1 M) and/or NaOH (0.1 M) solutions. The influence of ionic strength on Sb(III) adsorption was evaluated with NaCl concentrations varying from 0.01 to 0.25 M, with an initial Sb(III) concentration of 40 mg L$^{-1}$, at 25°C. Moreover, NO$_3^-$, Cl$^-$, SO$_4^{2-}$, and PO$_4^{3-}$ are ubiquitously found anions in the aqueous system, and thus they were chosen as the model co-existing anions to investigate their influence on Sb(III) removal. The initial concentration of these anions was selected as 40 mg L$^{-1}$, acting as a typical concentration of anions in wastewater (Wang et al. 2018).

After adsorption, all samples were filtered using a 0.45-μm polyethersulfone (PES) membrane (JINTENG Experimental Equipment Co., Ltd., China), and the concentration of Sb(III) in the supernatant was quantified using an atomic absorption spectrometer (AAS) (ZA3300, Shimadzu, Japan) equipped with an Sb hollow-cathode lamp (Shanghai Huake Experimental Equipment Co., Ltd., China) at a wavelength of 217.6 nm. The detection limit of Sb concentration (40 mg L$^{-1}$) was based on previous studies (30–50 mg L$^{-1}$) (Cui et al. 2017). The initial solution pH was recalibrated after measuring each batch of 25 samples, and the relative standard deviation of triplicate analysis was set to <5%.

The adsorption capacity of biochar and the removal efficiency of Sb(III) were calculated using Eq. [1] and Eq. [2] (Cui et al. 2017):

$$Q_e = (C_o - C_e)V/m \quad \text{(Eq. [1])}$$

$$\eta = (C_o - C_e)/C_o \times 100\% \quad \text{(Eq. [2])}$$
Table 1. Selected physicochemical characteristics of crawfish shell biochar pyrolyzed at 350°C, 500°C and 650°C.

| Sample  | C (%) | H (%) | N (%) | S (%) | pH (H₂O) | Ash content (%) | Surface alkalinity (cmol kg⁻¹) | Specific surface area (m² g⁻¹) | Pore volume (cm³ g⁻¹) |
|---------|-------|-------|-------|-------|-----------|-----------------|-------------------------------|-------------------------------|------------------------|
| CSB350  | 47.9  | 5.0   | 3.5   | 0.03  | 8.5       | 27.0            | 271                           | 3.2                           | 0.004                  |
| CSB500  | 27.4  | 1.0   | 2.3   | 0.02  | 9.2       | 38.6            | 288                           | 4.4                           | 0.031                  |
| CSB650  | 25.5  | 0.5   | 1.6   | 0.02  | 10.6      | 47.3            | 303                           | 12.1                          | 0.054                  |

Note: CSB350, CSB500, and CSB650 indicate the crawfish shell biochar pyrolyzed at 350 °C, 500 °C, and 650 °C, respectively.
Metal concentrations in the CSB showed a gradual increase with increasing pyrolysis temperature (Table S1), which was probably attributable to the volatilization losses of major elements such as C, H, and N, thereby concentrating the metallic elements (Xiao et al. 2017). The low content of heavy metals indicated that the CSB holds the potential to be applied as environmentally-friendly adsorbent.

The SEM images exhibited a denser morphology and stunted porosity of CSB350 compared to other biochars (Fig. 1). An increasing pyrolysis temperature endowed the biochar with more developed pore channels, exhibiting a fiber-like honeycomb structure (Fig. 1). These results confirmed higher surface area and pore volume in CSB650 than other two biochars (Table 1). Results of TEM analysis indicated a rough surface morphology with discernible particles on CSB surfaces (Fig. 1), suggesting the probable presence of mineral substances such as calcium carbonate. The XRD patterns confirmed that the mineral crystals in CSB were mainly CaCO₃, and a small amount of NaCl (Fig. S1) inherited from the feedstock material. The EDS spectra of CSB confirmed that higher pyrolysis temperature enhanced the contents of alkaline minerals such as Na, K, Mg and Ca in biochar (Fig. S2). As demonstrated in the TGA and DGT patterns of CSB (Fig. S3), 43%, 59%, and 60% of weight losses were noted in the case of CSB350, CSB500, and CSB650, respectively. The decomposition of CSB350 started at about 300°C with two maximum weight losses peaked at 457°C and 713°C, and the temperature of sharp mass decay was found to be at 737°C for CSB500 and 753°C for CSB650 (Fig. S3B). The first mass loss for CSB350 at 457°C represented the likely decomposition of calcium hydroxide (Habte et al. 2020). On the other hand, the weight loss of CSB above 700°C was due to the thermal destruction of the mineral contents of biochar (Park et al. 2018), i.e., calcite (CaCO₃) identified by XRD in this study (Fig. S1).

As shown in Fig. S4, CSB350 exhibited an advantage in the diversity of surface functional groups over the other two biochars, including O–H stretching vibrations of alcoholic hydroxyl group (3410 cm⁻¹) (Wei et al. 2020), stretching vibrations of C–H bonds (–CH2– at 2922 cm⁻¹ and –CH3– at 2850 cm⁻¹), carbonyl C=O (1580 cm⁻¹) (Chen et al. 2022a), O–C=O of the carbonates (1410 cm⁻¹) (Ma et al. 2021), aromatic ether C–O–C bonds (1040 cm⁻¹), and vibrations of aromatic C–H out-of-plane groups (875 and 713 cm⁻¹) (Xu et al. 2020). This phenomenon suggested that CSB350 might hold the potential to adsorb Sb(III) via chemisorption process. As for the CSB500 and CSB650, the stretching vibrations of O–H bond, C–H bond, and C=O disappeared, whereas enhanced bands for O–C=O, aromatic ether C–O–C, and aromatic C–H bonds were observed (Fig. S4). The high temperature-induced degeneration of O–H vibration peak was attributable to the decomposition of the hydroxyl groups bonded by hydrogen and oxygen bonds (Park et al. 2018). The disappearance of C–H and C=O peaks was caused by the transformation of O-alkylate groups to volatile or fixed carbon (e.g., C–O, C–O–C) with the increase in pyrolysis temperature (Xu et al. 2020; Ma et al. 2021). Enhanced peaks centered at 875 and 731 cm⁻¹ indicated the promotion of “biochar’s” aromaticity under high-temperature pyrolysis (Cui et al. 2017). In addition, the peaks observed around 605 and 567 cm⁻¹ in CSB were associated with O–P–O bonds (Park et al. 2018).

The XPS spectra verified that the increasing pyrolysis temperature led to higher proportions of O 1s, Ca 2p, N 1s, and Na 1s in biochar, and a lower C 1s peak proportion (Fig. S5). The C 1s spectra (Fig. 2 A) of CSB350 could be deconvoluted into three peaks, i.e., 284.8, 285.2, and 288.9 eV, which were assigned to the C–C (50.0%), C=C (42.6%), and HO–C=O groups (7.4%), respectively (Zeng et al. 2019; Palansooriya et al. 2021). For O 1s, CSB350 displayed two peaks at 531.6 and 532.4 eV (Fig. 2D), which represented C–O and C=O, respectively (Liu et al. 2019). As the pyrolysis temperature increased, C=C peak in CSB350 disappeared, whereas HO–C=O peak proportion increased from 7.4% in CSB350 to 12.8% in CSB500, and 18.7% in CSB650 (Fig. 2 A-C). A new C–O–C peak in CSB500/650 was observed, and its peak intensity positively responded to the increasing pyrolysis temperature (Fig. 2B, C), which was consistent with the results in FTIR spectra (Fig. S4). The peak intensity and location shift in C 1s and O 1s of CSB indicated that the pyrolysis temperature was
Biochar adsorption process could be divided into two linear stages. At the first fast-stage (0–60 min), the higher $K_1$ value represented the high adsorption rate, implying the rapid occupancy of Sb(III) on the available active sites, which was mainly due to the film diffusion where Sb(III) was diffused towards the surface of biochar (Rusmin et al. 2015; Xiong et al. 2020). As the majority of adsorption sites were occupied by Sb(III), the adsorption process entered the second slow stage (60-1440 min), i.e., an intra-particle diffusion governed Sb(III) adsorption into the internal pores of biochar (Liu et al. 2019).

### 3.2 Sb(III) adsorption and desorption

#### 3.2.1 Adsorption kinetics

Kinetic data revealed that the adsorption of Sb(III) by CSB increased rapidly in the first 30 min, accounting for 92–96% of the total adsorption; then the adsorption rate gradually slowed down until equilibrium reached (Fig. 3 A). The kinetic adsorption data for Sb(III) were fitted to the pseudo-first-order, pseudo-second-order (Fig. 3 A), and intra-particle diffusion models (Fig. 3B), and the associated fitting parameters are presented in Table S2. The pseudo-second-order model best described the kinetic data with the highest $R^2$ values (0.84–0.97) (Table S2). The theoretically calculated $Q_e$ value also agreed closely with the obtained experimental data, indicating that chemical interactions occurred between functional groups on the biochar surface and Sb(III) anions (Xiong et al. 2020).

The intra-particle diffusion model fitting results suggested that more than one rate-limiting steps were involved during the Sb(III) adsorption process, as the curve of $Q_t$ versus $t^{0.5}$ was multi-linear with an intercept of $C \neq 0$ (Fig. 3B; Table S2), as indicated by Rusmin et al. (2015). The Sb(III) adsorption process could be divided into two linear stages. At the first fast-stage (0–60 min), the higher $K_1$ value represented the high adsorption rate, implying the rapid occupancy of Sb(III) on the available active sites, which was mainly due to the film diffusion where Sb(III) was diffused towards the surface of biochar (Rusmin et al. 2015; Xiong et al. 2020). As the majority of adsorption sites were occupied by Sb(III), the adsorption process entered the second slow stage (60-1440 min), i.e., an intra-particle diffusion governed Sb(III) adsorption into the internal pores of biochar (Liu et al. 2019).

#### 3.2.2 Adsorption isotherm

The Langmuir, Freundlich, and Temkin models were fitted to the adsorption isotherm data of Sb(III) (Fig. 3 C), and their corresponding parameters are listed in Table S3. As compared to the Freundlich fitting results, the Langmuir model provided higher $R^2$ values ranging from 0.97 to 0.99, indicating that Sb(III) adsorption was much closer to a monolayer adsorption, rather than a multilayer adsorption (Chen et al. 2022b). As predicted from the Langmuir model, the maximum adsorption capacity ($Q_m$) was found to be 27.7, 18.2, and 16.1 mg g$^{-1}$ for CSB350, CSB500, and CSB650, respectively (Table S3), indicating that the increase of pyrolysis temperature negatively affected the removal of...
we select biochar for modification to obtain greater removal of Sb(III).

3.2.3 Influence of ionic strength and co-existing substances on Sb(III) adsorption

The adsorption of Sb(III) onto CSB was insignificantly \((P > 0.05)\) affected by the solution ionic strength ranging from 0.01 to 0.2 M (Fig. 4 A). This phenomenon indicated inner-sphere complex formation between Sb(III) and functional groups on biochar surface (Rahman et al. 2021). The high ionic strength (0.25 M) suppressed the adsorption capacity of CSB350, CSB500, and CSB650, and adsorption was found to decrease by 15%, 14%, and 11%, respectively, as compared to the 0.01 M treatment (Fig. 4 A). The elevated degree of charge screening caused by increased Na\(^+\)
Biochar (Xi et al. 2013). Moreover, the introduced humic acid (HA) significantly ($P < 0.05$) inhibited the adsorption of Sb(III) on CSB by 46.4–52.3%, as compared to the control (Fig. 4B). This could be attributable to the blockage of surface reactive sites on biochar by the large-molecular HA (Wei et al. 2020).

3.2.4 Influence of initial solution pH on Sb(III) adsorption

The highest adsorption of Sb(III) on CSB350 was noted at pH 2, up to 3.2 mg g$^{-1}$ (Fig. 4C). The response of Sb(III) adsorption to the initial solution pH might be due to changes in speciation of Sb(III) at different pH conditions (Xiong et al. 2020). Under strongly acidic conditions at pH of 2,
the dominating species of Sb(III) would be electropositive Sb(OH)$_2^{+}$, and electrostatic interaction might occur between the electronegative surface of CSB and electropositive Sb(OH)$_2^{+}$, thus promoting Sb(III) adsorption (Wan et al. 2020). When the initial solution pH ranged from 3 to 6, the amount of Sb(OH)$_2^{+}$ decreased, which suppressed the electrostatic interaction between biochar and Sb(OH)$_2^{+}$, thus decreasing Sb(III) adsorption (Cui et al. 2017). To be specific, at initial solution pH of 6, the adsorption of Sb(III) on CSB350, CSB500 and CSB650 decreased by 30.5%, 21.7%, and 21.8%, respectively, as compared to the adsorption at pH 2 (Fig. 4 C). At a pH range of 6 to 11, Sb(III) adsorption on CSB appeared not to be affected significantly by the solution pH, as the adsorption amount of Sb(III) only slightly varied. These results could be explained by the fact that the predominant form of Sb(III) was Sb(OH)$_3^{0}$ at this specific pH range, which was not readily adsorbed by biochar due to its electroneutrality (Xiong et al. 2020).

### 3.2.5 Desorption and regeneration of biochar

As expected, desorption of Sb(III) from CSB350, CSB500, and CSB650 increased with the increase of desorption time, and the desorption process tended to be stable after 4 h (Fig. S6A). After 24 h of desorption, CSB350 retained 88% of the maximum adsorption capacity of Sb(III), slightly lower than that of CSB500 (90%) and CSB650 (90%). The desorption results indicated a robust resistance of CSB to desorb Sb(III) anions. Moreover, the Sb-adsorbed CSB350 was desorbed using NaOH solution (0.5 M) to test its reusability. The regeneration results demonstrated that Sb(III) adsorption efficiency of CSB350 was maintained 83.1% of the maximum adsorption capacity after 3 adsorption-desorption cycles (Fig. S6B), suggesting that CSB held potential reusability in practical application.
3.3 Qualitative analysis on biochar surface after Sb(III) adsorption

The μ-XRF-based elemental mapping of CSB after Sb(III) adsorption confirmed the presence of Sb and mineral elements (i.e., K and Fe) on the biochar surface, with a heterogeneous distribution (Fig. 5). The greater density and brightness of Sb dots were observed in the Sb-laden CSB350 (Fig. 5 A), indicating its higher adsorption capacity than CSB500 and CSB650. Additionally, the Sb distribution area was not consistent with that of Fe distribution area, which might suggest that the Fe-Sb complex formation was not a key Sb(III) adsorption mechanism in this study. The HR-TEM based elemental mapping revealed the distribution change of non-metal elements (i.e., C, O, N, P) in biochar (Fig. S7). The weakened intensities for C and O might indicate the contribution of C/O-containing groups in the Sb(III) adsorption process. The CSB350 containing more C and O groups showed higher Sb(III) adsorption than CSB650, confirming the μ-XRF and HR-TEM results.

Furthermore, the presence of crystallographic phase of Sb on the biochar surface was investigated using XRD analysis (Fig. 6 A). In addition to the diffraction reflection of calcium carbonate, several new reflections occurred in the Sb-laden CSB. For instance, based on the Powder Diffraction File (PDF) database, antimony sulfate $\text{Sb}_2\left(\text{SO}_4\right)_3$ (Card #34-1097) was detected in Sb-laden CSB650, while potassium antimony oxide ($\text{KSB}_2\text{O}_3$) (Card #31–0973) and $\text{Sb}_2\left(\text{SO}_4\right)_3$ were simultaneously observed in Sb-laden CSB500 (Fig. 6 A). Although CSB350 exhibited the highest Sb(III) adsorption, the Sb-laden CSB350 showed no Sb-related crystal phases, indicating the presence of Sb on CSB350 surface as an amorphous phase. This could be ascribed to the low mineral content in CSB350, as revealed by the EDS spectra (Fig. S2), which confined the formation of antimony crystal phases on this adsorbent.

3.4 Potential adsorption mechanisms

3.4.1 FTIR, XPS and NEXAFS analyses

Noticeable changes of FTIR band features occurred in CSB after interaction with Sb(III) (Fig. 6B). The CSB350 sample was used as a reference to study the role of surface functional groups of biochar in Sb(III) adsorption. Firstly, the band for O–H groups disappeared (Fig. 6B), highlighting the role of hydrogen bonding in Sb(III) adsorption. The hydroxyl groups could serve as hydrogen donors to bond with oxygen atoms from Sb(III), as suggested by Xiong et al. (2020). Secondly, a weakened intensity of aromatic C–H out-of-plane vibrations was noted (Fig. 6B), indicating that the Sb(III)-π interaction occurred during Sb(III) adsorption (Cui et al. 2017). A stable combination of a π from aromatic C–H with another π from Sb(III) anions could enhance the adsorption onto biochar (Xiong et al. 2020). Thirdly, the stretching vibrations of oxygen-containing groups such as C=O and C–O were also weakened with slight shift in peak position (Fig. 6B). These results suggested that the oxygen-containing functional groups on biochar also participated in Sb(III) adsorption via surface complexation reaction (Jia et al. 2020). Instead, a new band centered at 743 cm$^{-1}$ was noted, which could be assigned to the stretching vibrations of Sb–O–Sb (Wei et al. 2020), as also demonstrated by XPS data, confirming the presence of Sb(III) on the biochar surface.

High-resolution XPS spectra of C 1s and O 1s for Sb-laden CSB are shown in Fig. 7 A-F. The C 1s spectra of CSB350 illustrated that C=C peak proportion decreased from 42.6 to 21.7% after Sb(III) adsorption (Fig. 7 A), implying a π–π coordination between aromatic C=C bonds and Sb(III) anion (Chen et al. 2022b). As for the C 1s spectra of CSB500, the ratio of C=C peak decreased from 71.7 to 60.8% after Sb(III) adsorption (Fig. 7B). In addition, in the Sb-laden CSB650, the carboxyl peak in C 1s spectra shifted from 290.0 to 289.5 eV (Fig. 7 C), with a proportional decrease of 4%. The proportion change and peak shift of carboxyl stressed the key factor of surface complexation during CSB650-Sb(III) interaction. In the O 1s spectra of the three CSB, Sb adsorption had a noticeable impact on the peak shift and proportion change of C–O and C=O groups, to positions with higher binding energy (Fig. 7D-F). This indicates the contribution of oxygen-containing...
groups during the adsorption process, such as the formation of C–O–Sb bonds (Chen et al. 2022b). For instance, in the O 1s spectra of CSB350, a decrease of peak proportion for C–O from 74.6 to 59.2% occurred, which highlighted its crucial role in Sb(III) adsorption on the biochar (Fig. 7 F).

The NEXAFS-based C 1s spectroscopic analysis was employed to further identify the C species of CSB350 before and after Sb adsorption (Fig. 7 G, H). The assignment of C 1s deconvolved peaks was obtained from previous publications (Wei et al. 2019a; Li et al. 2020). The C 1s NEXAFS spectra illustrated that alkyl C (G1, 32.6%), O-alkyl C (G4, 30.8%), carbonyl-related C (G3 + G5, 18.7%), and aromatic C (G1 + G6, 17.1%) were the dominant C species on CSB350 (Table S5). After interaction with Sb, the proportion of aromatic C = C decreased from 15.3 to 8.3%, and the peak for aromatic C–C disappeared (Fig. 7 H; Table S5). Concomitantly, Sb association resulted in a pronounced decrease in the O-alkyl C peak, from 30.8 to 20.5% (Table S5). These feature changes strongly suggested the interaction of Sb ions with aromatic C = C/C–C and C–OH groups on the CSB350 surface, which coincided with the results of FTIR and XPS analyses.

3.4.2 DFT calculations

The Sb(III) adsorption capacities of CSB at different reaction sites were verified using DFT calculations. The optimized structure segment of CSB with 5 functional groups (–OH, C–O–C, –CH₃, C = O, and O = C–O) is presented in Fig. 8 A. As demonstrated in Fig. 8 B-F, the calculated adsorption energy (AE) values of O = C–O (-2.94 eV) and C = O (-1.81 eV) were remarkably higher than those of –OH (-0.49 eV), C–O–C (-0.10 eV) and –CH₃ (-0.04 eV). Additionally, O–Sb bond distances in various reaction sites were obtained, showing the order C = O (2.00 Å) < O = C–O (2.16 Å) < –OH (2.35 Å) < –CH₃ (2.78 Å) < C–O–C (3.64 Å). Active sites with large negative AE and short bond length were identified as promising features for Sb(III) adsorption on biochar (Zhang et al. 2019a; Chen et al. 2022b). These results further confirmed the strong Sb(III) adsorption capacity of O = C–O and C = O groups, highlighting that surface complexation was the key adsorption mechanism in the removal of Sb(III) by CSB. The DFT studies also suggested that hydrogen bonding and π–π coordination, respectively, triggered by hydroxyl and aromatic methyl might play a secondary role in the Sb(III) adsorption due to the low calculated AE (Fig. 8 B, D).

Overall, FTIR, XPS, and NEXAFS analyses and theoretical calculations confirmed the key role of functional groups in the Sb(III) adsorption process. CSB350 with abundant surface functional groups such as C = O, O = C–O, hydroxyl, and aromatic C–H has shown the greatest adsorption capacity for Sb(III). Therefore, it is recommended as the most suitable sorbent in the removal of aqueous Sb(III).
3.5 Oxidation of Sb(III) on biochar surface

The oxidation state of Sb was examined following Sb(III) adsorption on biochar by the deconvolution of Sb 3d photoelectron spectra. Given the narrow scan of Sb 3d_{3/2} overlapping with the O 1s region, the Sb 3d_{3/2} spectra were applied to determine the oxidation state of Sb on CSB, and the deconvolution results are shown in Fig. 9 A. As demonstrated, Sb(V) proportion in the CSB decreased in the order of CSB350 (78%) > CSB650 (61%) > CSB500 (45%). It was reported that Sb(III) oxidation by O$_2$ in the nature is a slow process (Cui et al. 2017; Wei et al. 2020). Hence, the oxidation phenomenon was most likely influenced by the biochar addition.

Biochar-derived persistent free radicals (PFRs) may favor the oxidation of Sb(III) (Wei et al. 2020). Electron spin resonance (ESR) analysis was employed to identify the presence of PFRs on selected biochar samples. All three CSBs generated pronounced ESR signals, and the concentration of PFRs decreased in the following order: CSB500 > CSB350 > CSB650 (Fig. 9B-1). The intensity of ESR signal increased from CSB350 to CSB500, suggesting that the increasing pyrolysis temperature favored the formation of PFRs in biochar (Cui et al. 2017). The decrease of ESR signals in CSB650 could be ascribed to the destruction and reorganization of some organic structures in the biochar (Yang et al. 2016). Moreover, the ESR signal decreased after Sb(III) adsorption by CSB350 (Fig. 9B-2), suggesting the involvement of PFRs in biochar during the biochar-Sb interaction process. Therefore, an ESR analysis using DMPO (a typical spin trapping agent) was employed to identify the specific free radical species. Results showed that DMPO-•OH and DMPO-•O$_2^-$ were detected as the major active free radical species, and the intensity of •OH signal was higher than that of •O$_2^-$ (Fig. S8). Therefore, the oxidation of Sb(III) might be induced by CSB-derived PFRs, i.e., •OH was the major oxidative radical species whilst •O$_2^-$ also contributed to the oxidation process (Huang et al. 2018a, b; Li et al. 2019). However, the oxidation capacities of CSBs (CSB350 > CSB650 > CSB500) were not in accordance with their corresponding concentration of PFRs (CSB500 > CSB350 > CSB650). This phenomenon indicated that Sb(III) oxidation by biochar was not solely governed by PFRs.

It was thus hypothesized that Sb(III) oxidation could also be affected by the electron transfer property of biochar, i.e., the electron-donating capacity (EDC) and electron-accepting capacity (EAC). The values of EDC and EAC were calculated (Fig. S9) and are presented in Fig. 10 A. As the pyrolysis temperature increased, CSB displayed decreasing EDC from 0.128 to 0.085 mmol e$^-$ (g biochar)$^{-1}$, while the EAC increased from 0.055 to 0.130 e$^-$ (g biochar)$^{-1}$. The electron exchange capacity (EEC) was obtained by summing up EAC and EDC. An increasing pyrolysis temperature caused a decrease of EDC/EEC and an increase of EAC/EEC (Fig. 10B), indicating that CSB350 possessed the strongest reducing capacity and CSB650 had the highest oxidizing capacity (Klüpfel et al. 2014; Zhang et al. 2019b;

![Fig. 9 The Sb 3d_{3/2} XPS spectra of Sb-adsorbed crawfish shell biochar (CSB) pyrolyzed at 350°C (CSB350), 500°C (CSB500), and 650°C (CSB650); ESR spectra of CSB before adsorption (B-1); CSB350 in the blank solution and after adsorption of Sb(III) (B-2)](image-url)
Our results demonstrated that crawfish shell-derived biochar (CSB) can effectively remove Sb(III) from aqueous solution, and simultaneously reduce the toxicity of Sb by oxidizing Sb(III) to Sb(V). Therein, CSB pyrolyzed at 350°C (CSB350) possessed the highest adsorption capacity for Sb(III); and the maximum adsorption was found to be 27.7 mg g⁻¹. Batch adsorption and characterization results revealed the mechanisms for Sb(III) removal by CSB350 involved electrostatic interaction between Sb(OH)₂⁺ and negative biochar surface under strongly acid condition, surface complexation with oxygen-containing functional groups, π–π coordination with aromatic groups, and hydrogen bonding with hydroxyl groups. Density functional theory calculations showed that O=\text{C}–\text{O} and C=\text{O} groups had the highest adsorption energy (AE = -2.94 eV and AE = -1.81 eV), highlighting the key role of surface complexation in Sb(III) adsorption. Moreover, CSB350 exhibited the strongest oxidation for Sb(III), and 78% of Sb(III) was oxidized to Sb(V), which might be due to its higher concentration of persistent free radicals and electron-donating capacity (0.128 mmol e⁻ (g biochar)⁻¹). Desorption and regeneration experiments indicated that CSB350 was a feasible adsorbent with robust stability for Sb(III) and great reusability performance. Overall, crawfish shell-derived biochar can be an environmentally-friendly and efficient adsorbent for the remediation of Sb(III) contaminated water. Future studies are needed to investigate the potential influence of crawfish shell biochar on the bioavailability and phytoavailability of Sb in the more complex soil-plant systems.

Cui et al. (2017) reported that the electron-donating phenolic moieties in biochar firstly might have reduced solution-derived molecular oxygen to reactive oxygen species (ROS, •O₂⁻), then the generated ROS facilitated the Sb(III) oxidation in aqueous solution, and finally Sb(V) was adsorbed on the biochar surface. We infer that the high EDC of CSB350 was the most plausible reason for its high Sb(III) oxidation capacity through an indirect process. The CSB650 showed a stronger Sb(III)oxidation capacity as compared to CSB500 (Fig. 9 A), which might be related to its high EAC and oxidative metallic elements. First, CSB650 possessed more electron-accepting moieties (mainly carbonyl and quinone), which could capture the electrons in Sb(III) during the interaction between biochar and Sb, thus directly promoting the oxidation. Second, the higher contents of Fe and Mn were noted in CSB650 (Table S1), which might be another reason for its stronger oxidation capacity for Sb(III), as compared to CSB500. In a previous study, the presence of Fe and Mn in biochar was found to be responsible for As(III) oxidation, as reported by Dong et al. (2014), which might have a similar contribution to the Sb(III) oxidation in this study.

As discussed above, Sb(III) oxidation was mainly influenced by the PFRs and EEC of biochar. The reason for the greatest oxidation capacity of low-temperature derived CSB350 could be its highest EDC through indirect oxidation. Another possible pathway of direct oxidation to Sb(V) by •O₂⁻ and •OH cannot be excluded. As for the high-temperature pyrolyzed biochar, CSB650 had the highest OEC and concentrations of Fe and Mn, and the Sb(III) oxidation mechanism was majorly governed by the direct oxidation process.

4 Conclusions

Our results demonstrated that crawfish shell-derived biochar (CSB) can effectively remove Sb(III) from aqueous solution, and simultaneously reduce the toxicity of Sb by oxidizing Sb(III) to Sb(V). Therein, CSB pyrolyzed at 350°C (CSB350) possessed the highest adsorption capacity for Sb(III); and the maximum adsorption was found to be 27.7 mg g⁻¹. Batch adsorption and characterization results revealed the mechanisms for Sb(III) removal by CSB350 involved electrostatic interaction between Sb(OH)₂⁺ and negative biochar surface under strongly acid condition, surface complexation with oxygen-containing functional groups, π–π coordination with aromatic groups, and hydrogen bonding with hydroxyl groups. Density functional theory calculations showed that O=\text{C}–\text{O} and C=\text{O} groups had the highest adsorption energy (AE = -2.94 eV and AE = -1.81 eV), highlighting the key role of surface complexation in Sb(III) adsorption. Moreover, CSB350 exhibited the strongest oxidation for Sb(III), and 78% of Sb(III) was oxidized to Sb(V), which might be due to its higher concentration of persistent free radicals and electron-donating capacity (0.128 mmol e⁻ (g biochar)⁻¹). Desorption and regeneration experiments indicated that CSB350 was a feasible adsorbent with robust stability for Sb(III) and great reusability performance. Overall, crawfish shell-derived biochar can be an environmentally-friendly and efficient adsorbent for the remediation of Sb(III) contaminated water. Future studies are needed to investigate the potential influence of crawfish shell biochar on the bioavailability and phytoavailability of Sb in the more complex soil-plant systems.

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Availability of data and materials The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

Declarations

Conflict of interest The authors have no conflicts of interest to disclose, financial or otherwise.

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