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# Co-catalysis of trace dissolved Fe(III) with biochar in hydrogen peroxide activation for enhanced oxidation of pollutants†

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Activation of hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) with biochar is a sustainable and low-cost approach for advanced oxidation of organic pollutants, but faces the challenge of a low yield of hydroxyl radical (<sup>•</sup>OH). Herein, we hypothesize that the activation efficiency of H<sub>2</sub>O<sub>2</sub> can be enhanced through co-catalysis of trace dissolved iron (Fe) with biochar. Two biochar samples derived from different feedstock, namely LB from liquor-making residue and WB from wood sawdust, were tested in the co-catalytic systems using trace Fe(III) (0.3 mg L<sup>-1</sup>). The cumulative <sup>•</sup>OH production in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> was measured to be 3.28 times that in LB/H<sub>2</sub>O<sub>2</sub>, while the cumulative <sup>•</sup>OH production in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> was 11.9 times that in WB/H<sub>2</sub>O<sub>2</sub>. No extra consumption of H<sub>2</sub>O<sub>2</sub> was observed in LB/H<sub>2</sub>O<sub>2</sub> or WB/H<sub>2</sub>O<sub>2</sub> after addition of trace Fe(III). Consequently, the reaction rate constants (*k*<sub>obs</sub>) for oxidation of pollutants (2,4-dichlorophenoxyacetic acid and sulfamethazine) were enhanced by 3.13–9.16 times. Other iron species including dissolved Fe(II) and iron minerals showed a similar effect on catalyzing 2,4-D oxidation by biochar/H<sub>2</sub>O<sub>2</sub>. The interactions involved in adsorption and reduction of Fe(III) by biochar in which the defects acted as electron donors and oxygen-containing functional groups bridged the electron transfer. The fast regeneration of Fe(II) in the co-catalytic system resulted in the sustainable <sup>•</sup>OH production, thus the efficient oxidation of pollutants comparable to other advanced oxidation processes was achieved by using dissolved iron at a concentration as low as the concentration that can be found in natural water.

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## 1. Introduction

Biochar is a kind of carbon-rich material obtained by pyrolysis of waste biomass under oxygen-limited conditions, and has shown great potential in carbon sequestration, soil improvement, energy storage, and environmental remediation.<sup>1–3</sup> Recently, biochar has been found to be an attractive catalyst for advanced oxidation of pollutants.<sup>4–6</sup> Specifically, biochar can activate H<sub>2</sub>O<sub>2</sub>,<sup>7–10</sup> and persulfates<sup>11–13</sup> to produce reactive oxygen species (ROS) such as the hydroxyl radical (<sup>•</sup>OH), sulfate radical (SO<sub>4</sub><sup>•-</sup>) and singlet oxygen (<sup>1</sup>O<sub>2</sub>), which are powerful oxidants (e.g. *E*<sup>0</sup>(<sup>•</sup>OH/H<sub>2</sub>O) = 2.73 V) capable of mineralization of various organic pollutants.<sup>14–16</sup> As biochar is readily available from waste biomass, its application in remediation of contaminated environments is advantageous over other elaborately synthesized materials in view of cost and sustainability.<sup>17–19</sup> However, the practical application of biochar-activated systems is limited

by the low yield of ROS. For example, the cumulative yield of <sup>•</sup>OH was estimated to be only 0.09 mol per molar consumption of H<sub>2</sub>O<sub>2</sub> in a biochar/H<sub>2</sub>O<sub>2</sub> system,<sup>20</sup> and most of the H<sub>2</sub>O<sub>2</sub> was decomposed into end-products (H<sub>2</sub>O and O<sub>2</sub>), as has been observed on other carbon materials.<sup>21</sup> The low efficiency of H<sub>2</sub>O<sub>2</sub> activation by biochar leads to unsatisfactory performance on pollutant removal and excess H<sub>2</sub>O<sub>2</sub> consumption. So, biochar was more often applied as a carrier of catalysts,<sup>22–25</sup> or after special modifications<sup>26–28</sup> for use in advanced oxidation processes (AOPs).

Herein, we hypothesize that the H<sub>2</sub>O<sub>2</sub> activation by biochar can be enhanced by using trace dissolved iron (Fe) together. Biochar, being an electron-rich and conductive carbon material, can act as a reductant of Fe(III) for persulfate activation to enhance the degradation of organic pollutants,<sup>29,30</sup> and can also accelerate Fe(III)/Fe(II) cycle that is the rate-limiting step in Fenton and Fenton-like processes.<sup>31,32</sup> Thereby, the required dosage of Fe(II) in Fenton process can be diminished substantially by biochar,<sup>20</sup> which is beneficial for minimizing the production of iron sludge that is still a challenge in Fenton process.<sup>33,34</sup> In this context, it is interesting to reduce further the iron concentration to a trace level that is close to natural environment, and to explore the interactions between trace iron and biochar on co-catalyzing the oxidation of pollutant by H<sub>2</sub>O<sub>2</sub>. The

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trace dissolved iron is ubiquitous in natural water. For example, a maximum dissolved iron of  $0.163 \text{ mg L}^{-1}$  in surface water has been reported by Nagai *et al.*,<sup>35</sup> and a dissolved iron of  $0.295 \text{ mg L}^{-1}$  in groundwater was reported by Gutierrez-Zapata *et al.*<sup>36</sup> The dissolved iron may also co-exist with organic pollutants in wastewater, especially in wastewater pretreated with iron-containing flocculants.<sup>37</sup> Therefore, the in-depth investigation on the co-catalysis of trace dissolved iron with biochar in oxidation process is of significance for both remediation of contaminated environment and advanced treatment of wastewater.

In this study, trace dissolved  $\text{Fe(III)}$  was used together with biochar to activate  $\text{H}_2\text{O}_2$  for enhanced oxidation of organic pollutants. The typical  $\text{Fe(III)}$  concentration was set at  $0.30 \text{ mg L}^{-1}$ , equal to the limit of iron concentration in drinking water recommended by the World Health Organization (WHO), and close to that found in a natural water.<sup>36</sup> A common herbicide 2,4-dichlorophenoxyacetic acid (2,4-D) and a common antibiotic sulfamethazine (SMZ) were used for evaluating the performance of this novel oxidation process co-catalyzed by trace  $\text{Fe(III)}$  and biochar. Both 2,4-D and SMZ are popular model pollutants that have been frequently used in evaluating various AOPs, and their degradation pathways have been clarified in many previous studies.<sup>34,38,39</sup> Herein, we focus on the co-catalytic effect between trace dissolved  $\text{Fe(III)}$  and biochar in oxidation process, and the role of dissolved  $\text{Fe(III)}$  was highlighted by using two biochars obtained by pyrolysis of different feedstock. We will show that co-catalysis of trace  $\text{Fe(III)}$  with biochar can raise dramatically the  $\cdot\text{OH}$  yield from  $\text{H}_2\text{O}_2$  decomposition, and enhance the oxidation of pollutants consequently. The interfacial interactions between trace  $\text{Fe(III)}$  and biochar in the co-catalytic system will be investigated, so as to clarify the in-depth mechanisms. Furthermore, different iron species including dissolved  $\text{Fe(II)}$  and solid iron minerals are used instead of aqueous  $\text{Fe(III)}$ , so as to extend the knowledge about the applicability of this co-catalytic system in remediation of contaminated environment.

## 2. Experimental

### 2.1. Reagents and biochar samples

Ferric sulfate  $[\text{Fe}_2(\text{SO}_4)_3]$  and ferrous sulfate heptahydrate ( $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ ) of analytical grade used for preparing the solution of trace  $\text{Fe(III)}/\text{Fe(II)}$ , as well as aqueous solution of  $\text{H}_2\text{O}_2$  (30%, w/w) of analytical grade were purchased from the Sino-pharm Chemical Reagent Co. Ltd., China. Benzoic acid (BA) and other chemicals (*e.g.* acetonitrile, methanol, sodium hydroxide) of analytical grade, iron minerals including hematite ( $\text{Fe}_2\text{O}_3$ , 99% purity) and magnetite ( $\text{Fe}_3\text{O}_4$ , 99% purity), and the model pollutants including 2,4-D (98% purity) and SMZ (99% purity) were purchased from Aladdin Agents Co. Ltd., China. The  $18.2 \text{ M}\Omega \text{ cm}$  (Milli-Q Gradient, Millipore, USA) pure water was used for preparing the solutions.

The biochar samples were derived from two different feedstock: one labeled as LB from a cereal waste after making liquor, and another labeled as WB from a pinewood sawdust, according to the method reported in our previous study.<sup>20</sup> The

temperature of  $700^\circ\text{C}$  for pyrolysis was selected for producing the biochar that has both relatively stable properties<sup>40,41</sup> and high reactivity in catalyzing Fenton process.<sup>20</sup> The carbon contents, ash contents and surface areas (Brunauer–Emmett–Teller (BET) method) were  $66.0\% \pm 0.3\%$  and  $93.9\% \pm 0.4\%$ ,  $23.4\% \pm 0.1\%$  and  $0.89\% \pm 0.03\%$ , and  $6.66$  and  $409 \text{ m}^2 \text{ g}^{-1}$ , respectively for LB and WB. More details regarding the compositions of these two biochars are provided in the ESI (Table S1 and Fig. S1†). A Fourier transform infrared spectrometer (FTIR) (Nexus, Nicolet, USA) was used to identify the functional groups of biochar with KBr as the background. Raman measurements were performed using a Renishaw inVia Raman spectrometer (UK) to evaluate the defects and graphitization of biochar. An X-ray diffractometer (XRD) (Empyrean, Panalytical, Netherlands) was used to characterize the graphite-like structure in biochar.

### 2.2. Experimental methods of oxidation

The oxidation experiments using trace  $\text{Fe(III)}$  in biochar/ $\text{H}_2\text{O}_2$  system were performed in the dark in an oscillator (180 rpm) (KS4000i, IKA, Germany). In a typical co-catalytic system containing  $100 \text{ mL}$  of water solution of trace  $\text{Fe(III)}$  ( $0.30 \text{ mg L}^{-1}$ ), the biochar ( $3.0 \text{ g L}^{-1}$ ) was added, and the initial pH ( $\text{pH}_0$ ) adjusted to  $3.0 \pm 0.1$  with dilute  $\text{H}_2\text{SO}_4$  solution. The suspension was oscillated at  $25 \pm 0.2^\circ\text{C}$  for 30 min of preliminary equilibration. After addition of pollutant (2,4-D or SMZ) with an initial concentration ( $C_0$ ) of  $20 \text{ mg L}^{-1}$ , the pH of solution was measured and re-adjusted to  $3.0 \pm 0.1$  if necessary. Then, the reaction was initiated immediately after addition of  $\text{H}_2\text{O}_2$  ( $5.0 \text{ mmol L}^{-1}$ ). After a set reaction time interval  $t$  (2–60 min),  $0.5 \text{ mL}$  of solution was sampled and immediately mixed with methanol of the same volume. The 2,4-D or SMZ concentration ( $C_t$ ,  $\text{mg L}^{-1}$ ), and total organic carbon (TOC) in the mixed solution samples was analyzed after membrane filtration ( $0.22 \mu\text{m}$ ). The mean value of triplicates was recorded for each experiment. The  $\text{Fe(III)}/\text{H}_2\text{O}_2$  and biochar/ $\text{H}_2\text{O}_2$  systems, together with those using biochar or  $\text{H}_2\text{O}_2$  alone were tested for comparison. Different  $\text{pH}_0$  (2.0–4.0), and different dosages of  $\text{Fe(III)}$  ( $0.10$ – $1.00 \text{ mg L}^{-1}$ ),  $\text{H}_2\text{O}_2$  ( $1.0$ – $9.0 \text{ mmol L}^{-1}$ ) and biochar ( $1.0$ – $5.0 \text{ g L}^{-1}$ ) were adjusted to investigate their influence on 2,4-D degradation. The different conditions used in this study are listed in Table S2.† In further comparison studies, the trace  $\text{Fe(II)}$  solution from dissolution of  $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ , and the suspensions of two iron minerals (hematite and magnetite) were used instead of  $\text{Fe(III)}$  solution to test the feasibility of various iron species. The repetitive experiments were performed to estimate the reusability of biochar by recycling it through centrifugation (8000 rpm) and freeze-drying without further purification.

### 2.3. Measurement of ROS in reaction systems

The  $\cdot\text{OH}$  production in various reaction systems was measured by using BA as a chemical probe.<sup>42,43</sup> The specific procedure is same as that in the oxidation experiment, except that a BA solution with initial concentration of  $5.0 \text{ mmol L}^{-1}$  was used instead of pollutant solution. The solution sample was analyzed after



mixing with methanol and membrane filtration. The concentration of *p*-hydroxybenzoic acid (*p*-HBA) derived from BA oxidation was referred to calculate the cumulative amount of  $\cdot\text{OH}$ . According to Tong *et al.*,<sup>42</sup> the production of 1.00 mmol of *p*-HBA is equivalent to the consumption of 5.87 mmol of  $\cdot\text{OH}$ . The production  $\cdot\text{OH}$  and superoxide anion radical ( $\text{O}_2^{\cdot-}$ ) in various reaction systems was also determined by electron paramagnetic resonance (EPR) spectrometry (FA-200, JEOL, Japan).<sup>43</sup>

#### 2.4. Instrumental analysis of solution samples

The concentration of *p*-HBA, 2,4-D or SMZ was measured using high performance liquid chromatography (LC-20A, Shimadzu, Japan). The details for analysis of *p*-HBA and 2,4-D can be found in our previous study,<sup>43</sup> and SMZ was analyzed at 275 nm using an eluent composed of acetonitrile and 0.1% acetic acid ( $v/v = 40/60$ ). The  $\text{H}_2\text{O}_2$  or  $\text{Fe(III)}$  concentration was measured using UV-vis spectrophotometry (Spectrum 1920, Shanghai, China).<sup>20</sup> The total aqueous iron was measured by inductively coupled plasma-atomic emission spectrometry (ICP-AES) (Prodigy XP, Leeman, USA), and the TOC was determined on a total carbon analyzer (Elementar vario, German).

### 3. Results and discussion

#### 3.1. Co-catalysis of trace $\text{Fe(III)}$ with biochar in oxidation of pollutants by $\text{H}_2\text{O}_2$

The co-catalysis of trace dissolved  $\text{Fe(III)}$  ( $0.30 \text{ mg L}^{-1}$ ) with biochar in oxidation of 2,4-D by  $\text{H}_2\text{O}_2$  can be found in Fig. 1. No

apparent removal of 2,4-D was observed in  $\text{Fe(III)}/\text{H}_2\text{O}_2$  (Fig. 1(a)), and only 41.8% and 35.7% of 2,4-D was removed respectively by  $\text{LB}/\text{H}_2\text{O}_2$  (Fig. 1(a)) and  $\text{WB}/\text{H}_2\text{O}_2$  (Fig. 1(b)), after 60 min of reaction. In contrast, much more 2,4-D was removed in the two systems co-catalyzed by dissolved  $\text{Fe(III)}$  and biochar, namely 89.0% in  $[\text{Fe(III)} + \text{LB}]/\text{H}_2\text{O}_2$  and 96.5% in  $[\text{Fe(III)} + \text{WB}]/\text{H}_2\text{O}_2$ , after the same reaction time. The results indicate that trace  $\text{Fe(III)}$  enhanced the pollutant removal by biochar/ $\text{H}_2\text{O}_2$ , which is further proved by kinetic analysis. The obtained *pseudo*-first-order reaction rate constants ( $k_{\text{obs}}$ ) show the much faster degradation of 2,4-D in the co-catalytic systems (Fig. 1(c)), as the  $k_{\text{obs}}$  values for 2,4-D removal in  $[\text{Fe(III)} + \text{LB}]/\text{H}_2\text{O}_2$  and  $[\text{Fe(III)} + \text{WB}]/\text{H}_2\text{O}_2$  are respectively 4.64 and 9.16 times that in  $\text{LB}/\text{H}_2\text{O}_2$  and  $\text{WB}/\text{H}_2\text{O}_2$ , and are two orders of magnitude larger than that the  $k_{\text{obs}}$  value in  $\text{Fe(III)}/\text{H}_2\text{O}_2$ . The  $k_{\text{obs}}$  values ( $0.035\text{--}0.055 \text{ min}^{-1}$ ) in these  $[\text{Fe(III)} + \text{biochar}]/\text{H}_2\text{O}_2$  systems are also larger than that observed in a Fenton process catalyzed by  $\text{FeS}^{44}$  and in a photo-catalytic process,<sup>45</sup> indicating that an efficiency comparable to other AOPs can be obtained through activation of  $\text{H}_2\text{O}_2$  using both biochar and trace  $\text{Fe(III)}$ .

Additional experiments have been performed to prove that 2,4-D was removed through oxidation. First, the contribution of adsorption by biochar to the total removal is insignificant as few 2,4-D was removed by biochar itself within 60 min (LB in Fig. 1(a), and WB in Fig. 1(b)). Second, the TOC removal in the reaction systems using  $\text{H}_2\text{O}_2$  (Fig. 1(d)) proves that 2,4-D was removed mostly by oxidation. Much more TOC was removed in  $[\text{Fe(III)} + \text{WB}]/\text{H}_2\text{O}_2$  and  $[\text{Fe(III)} + \text{LB}]/\text{H}_2\text{O}_2$  than that in  $\text{WB}/\text{H}_2\text{O}_2$

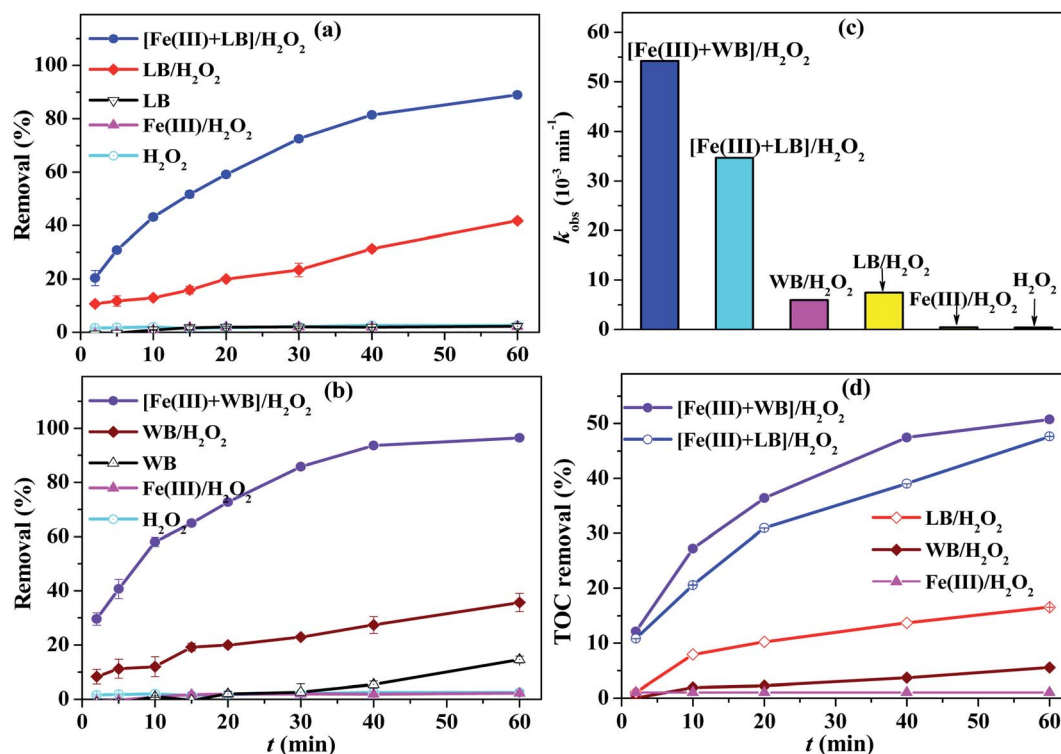


Fig. 1 (a and b) Removal of 2,4-D ( $C_0 = 20 \text{ mg L}^{-1}$ ) in the systems using different combinations of  $\text{Fe(III)}$ , biochar (LB or WB) and/or  $\text{H}_2\text{O}_2$ ; (c)  $k_{\text{obs}}$  and (d) TOC removal in different systems. Dosage:  $\text{Fe(III)} = 0.30 \text{ mg L}^{-1}$ , LB or WB =  $3.0 \text{ g L}^{-1}$ , and  $\text{H}_2\text{O}_2 = 5.0 \text{ mmol L}^{-1}$ ; and  $\text{pH}_0 = 3.0$ .

and LB/H<sub>2</sub>O<sub>2</sub>, proving the enhanced mineralization of 2,4-D in both co-catalytic systems. The release of chloride ions indicates the dechlorination of 2,4-D in the reaction systems, and the much faster dechlorination was observed in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> than in WB/H<sub>2</sub>O<sub>2</sub> (Fig. S2†). The specific degradation pathway of 2,4-D in Fenton-like oxidation by H<sub>2</sub>O<sub>2</sub> included dechlorination, hydroxylation, and opening of the aromatic ring, with the short-chain organic acids dominated the residual TOC.<sup>43</sup> Furthermore, the co-catalysis of trace Fe(III) (0.30 mg L<sup>-1</sup>) with biochar was also found in oxidation of another pollutant SMZ (Fig. S3†). The removal of SMZ in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> is much faster than in either Fe(III)/H<sub>2</sub>O<sub>2</sub> or LB/H<sub>2</sub>O<sub>2</sub>, and the *k*<sub>obs</sub> for SMZ removal in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> is calculated to be  $13.6 \times 10^{-3} \text{ min}^{-1}$ , which is 3.13 times that in LB/H<sub>2</sub>O<sub>2</sub>, and 6.07 times that in Fe(III)/H<sub>2</sub>O<sub>2</sub>. The results prove that the co-catalytic system is applicable for oxidation of different organic pollutants. It needs to be mentioned in advance here that, the difference between the two biochars (LB vs. WB) on performance for pollutant removal in the co-catalytic system will be addressed in the Section 3.5.

### 3.2. Enhanced <sup>•</sup>OH yield in the co-catalytic system

Much more <sup>•</sup>OH was produced in the two systems co-catalyzed by biochar and trace Fe(III) (0.30 mg L<sup>-1</sup>) than that in biochar/H<sub>2</sub>O<sub>2</sub>, according to results shown in Fig. 2(a). The cumulative <sup>•</sup>OH productions in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> and [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> are respectively 11.9 and 3.28 times that in WB/H<sub>2</sub>O<sub>2</sub> and LB/H<sub>2</sub>O<sub>2</sub>, and more than ten times that in Fe(III)/H<sub>2</sub>O<sub>2</sub>. The most

<sup>•</sup>OH production (1.11 mmol L<sup>-1</sup>) was observed in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> after 60 min of reaction, which can explain the highest *k*<sub>obs</sub> and the most TOC removal in this system (Fig. 1(c) and (d)). The ranking of various reaction systems on <sup>•</sup>OH production is generally in consistency with their order on *k*<sub>obs</sub> values (Fig. 1(c)), namely, more <sup>•</sup>OH production resulted in faster oxidation of pollutant. However, the ranking of H<sub>2</sub>O<sub>2</sub> consumption in different systems is not always consistent with <sup>•</sup>OH production. More decomposition of H<sub>2</sub>O<sub>2</sub> was observed in the two systems using LB (Fig. 2(b)), namely [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> and LB/H<sub>2</sub>O<sub>2</sub>. The H<sub>2</sub>O<sub>2</sub> consumed in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> is comparable to that in WB/H<sub>2</sub>O<sub>2</sub>, despite that the <sup>•</sup>OH production in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> was the most among all the systems (Fig. 2(a)). Therefore, the activation efficiency of H<sub>2</sub>O<sub>2</sub> should be estimated on the basis of <sup>•</sup>OH yield, namely the molar production of <sup>•</sup>OH per molar decomposition of H<sub>2</sub>O<sub>2</sub>. In fact, the <sup>•</sup>OH yield in the two biochar-activated systems is fairly low (6.19% in WB/H<sub>2</sub>O<sub>2</sub>, and 9.27% in LB/H<sub>2</sub>O<sub>2</sub>), indicating that most (>90%) of H<sub>2</sub>O<sub>2</sub> decomposed into end-products (H<sub>2</sub>O and O<sub>2</sub>).<sup>21</sup> In contrast, addition of trace Fe(III) in biochar/H<sub>2</sub>O<sub>2</sub> enhanced dramatically the <sup>•</sup>OH yield (89.8% in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub>, and 32.5% in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub>), without extra consumption of H<sub>2</sub>O<sub>2</sub> (Fig. 2(b)). Therefore, biochar facilitated the decomposition of H<sub>2</sub>O<sub>2</sub> while trace Fe(III) boosted the yield of <sup>•</sup>OH in the co-catalytic systems. The similar co-catalytic effect on enhancing the <sup>•</sup>OH yield from H<sub>2</sub>O<sub>2</sub> decomposition has been observed in the combination of biochar with iron minerals,<sup>43,46</sup> and in the Fenton-like systems with presence of

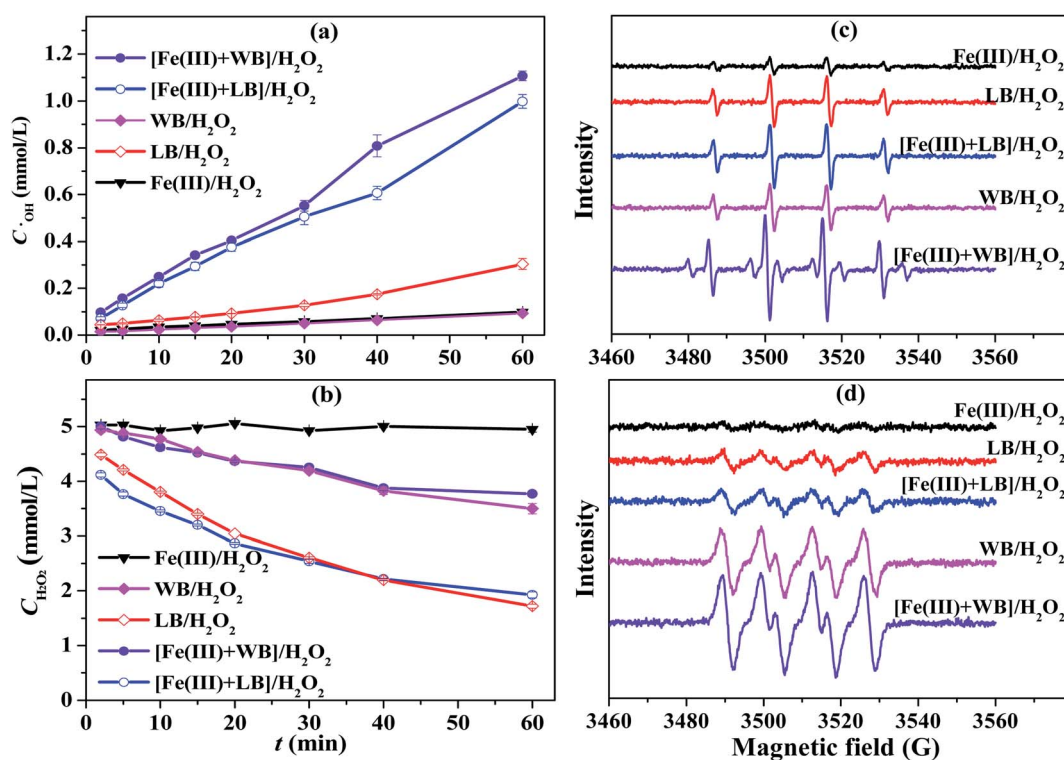


Fig. 2 (a) Cumulative <sup>•</sup>OH production, and (b) change of H<sub>2</sub>O<sub>2</sub> concentration, (c) EPR spectra of <sup>•</sup>OH trapped by DMPO, and (d) EPR spectra of O<sub>2</sub><sup>•-</sup> trapped by DMPO and methanol in different oxidation systems. Dosage: Fe(III) = 0.30 mg L<sup>-1</sup>, LB or WB = 3.0 g L<sup>-1</sup>, and H<sub>2</sub>O<sub>2</sub> = 5.0 mmol L<sup>-1</sup>; and pH<sub>0</sub> = 3.0.





activated carbon, carbon nanotubes and reduced graphene oxide.<sup>47,48</sup>

The EPR spectra confirm the production of both  $\cdot\text{OH}$  and  $\text{O}_2^{\cdot-}$  in most of the reaction systems. The  $\cdot\text{OH}$  signals with characteristic 1 : 2 : 2 : 1 pattern (Fig. 2(c)) can be ranked by relative intensity in an order similar to the  $\cdot\text{OH}$  production shown in Fig. 2(a). Namely, the stronger  $\cdot\text{OH}$  signals belonging to the two co-catalytic systems are consistent with the more production of  $\cdot\text{OH}$  measured with chemical probe. Furthermore, the degradation experiments using methanol as the  $\cdot\text{OH}$  scavenger has been performed for verifying the role of  $\cdot\text{OH}$ . The results (Fig. S4(a)†) indicated that the degradation of 2,4-D was almost ceased when methanol was added in the reaction systems, proving that  $\cdot\text{OH}$  is the dominant ROS in both  $[\text{Fe}(\text{III}) + \text{biochar}]/\text{H}_2\text{O}_2$  and  $\text{biochar}/\text{H}_2\text{O}_2$  systems. Despite this, the signals of 1 : 1 : 1 : 1 pattern shown in Fig. 2(d) indicates that there was  $\text{O}_2^{\cdot-}$  in the reaction systems except  $\text{Fe}(\text{III})/\text{H}_2\text{O}_2$ . Chloroform ( $\text{CHCl}_3$ ) was used as the scavenger of  $\text{O}_2^{\cdot-}$  according to a previous study by Diao *et al.*<sup>49</sup> The experimental results indicate that  $\text{O}_2^{\cdot-}$  had contributed to 2,4-D degradation as well, because addition of chloroform in  $[\text{Fe}(\text{III}) + \text{LB}]/\text{H}_2\text{O}_2$  slowed down the 2,4-D degradation (Fig. S4(b)†). As  $\text{O}_2^{\cdot-}$  is a reductant ( $E^0(\text{O}_2/\text{O}_2^{\cdot-}) = -0.16 \text{ V}$ ), it contributed to 2,4-D degradation most likely by serving as the intermediate for production of  $\cdot\text{OH}$ .<sup>43,50</sup>

### 3.3. Effect of $\text{Fe}(\text{III})$ concentration and iron species

The influence of  $\text{Fe}(\text{III})$  concentration on production of  $\cdot\text{OH}$  in  $[\text{Fe}(\text{III}) + \text{LB}]/\text{H}_2\text{O}_2$  is shown in Fig. 3(a). As can be seen, more  $\text{Fe}(\text{III})$  resulted in production of more  $\cdot\text{OH}$  and subsequently faster oxidation of 2,4-D (Fig. 3(b)). Specifically, the 2,4-D

degradation rate was enhanced sharply when the  $\text{Fe}(\text{III})$  concentration increased from 0 to  $0.30 \text{ mg L}^{-1}$ . But after then, the enhancement of degradation rate was no longer so distinct. The dissolved iron may also exist in water in ferrous species ( $\text{Fe}(\text{II})$ ), especially in anaerobic environment. For this case, a comparison study using  $\text{Fe}(\text{II})$  instead of  $\text{Fe}(\text{III})$  was conducted to degrade 2,4-D under the same reaction conditions. The results (Fig. 3(c)) indicate that 2,4-D was degraded in  $[\text{Fe}(\text{II}) + \text{LB}]/\text{H}_2\text{O}_2$  with a reaction rate ( $k_{\text{obs}}$ ) a little higher than that in  $[\text{Fe}(\text{III}) + \text{LB}]/\text{H}_2\text{O}_2$ , but much higher than that in the Fenton process without biochar ( $\text{Fe}(\text{II})/\text{H}_2\text{O}_2$ ). Therefore, the trace iron of both valence ( $\text{Fe}(\text{III})$  and  $\text{Fe}(\text{II})$ ) is applicable in the co-catalyzed oxidation of pollutant by  $\text{H}_2\text{O}_2$ .

Iron minerals have been used as the iron source for catalyzing the Fenton-like processes,<sup>51,52</sup> so we hypothesize that the solid iron species can also be used together with biochar for co-catalyzing the oxidation of pollutants by  $\text{H}_2\text{O}_2$ . In this study, two common iron minerals, namely hematite ( $\text{Fe}_2\text{O}_3$ ) and magnetite ( $\text{Fe}_3\text{O}_4$ ), were used together with WB for activation of  $\text{H}_2\text{O}_2$  at  $\text{pH}_0$  of 3.0. The results in Fig. 3(d) show the enhanced 2,4-D removal by both  $[\text{Fe}_2\text{O}_3 + \text{WB}]/\text{H}_2\text{O}_2$  and  $[\text{Fe}_3\text{O}_4 + \text{WB}]/\text{H}_2\text{O}_2$ , with the removal efficiencies apparently higher than the simple sum of  $\text{WB}/\text{H}_2\text{O}_2$  and corresponding heterogeneous Fenton system ( $\text{Fe}_2\text{O}_3/\text{H}_2\text{O}_2$  or  $\text{Fe}_3\text{O}_4/\text{H}_2\text{O}_2$ ). In addition, the dissolved iron was measured to be  $0.073$  and  $0.26 \text{ mg L}^{-1}$  after reaction in  $[\text{Fe}_2\text{O}_3 + \text{WB}]/\text{H}_2\text{O}_2$  and  $[\text{Fe}_3\text{O}_4 + \text{WB}]/\text{H}_2\text{O}_2$ , respectively. The 2,4-D degradation rate by  $[\text{Fe}_3\text{O}_4 + \text{WB}]/\text{H}_2\text{O}_2$  is comparable to that obtained in  $[\text{Fe}(\text{III}) + \text{WB}]/\text{H}_2\text{O}_2$  with a dissolved  $\text{Fe}(\text{III})$  of  $0.30 \text{ mg L}^{-1}$ , while the degradation rate by  $[\text{Fe}_2\text{O}_3 + \text{WB}]/\text{H}_2\text{O}_2$  is close to that obtained with a dissolved  $\text{Fe}(\text{III})$  of  $0.10 \text{ mg L}^{-1}$  (Fig. S5†). Therefore, it is likely that the iron minerals catalyzed

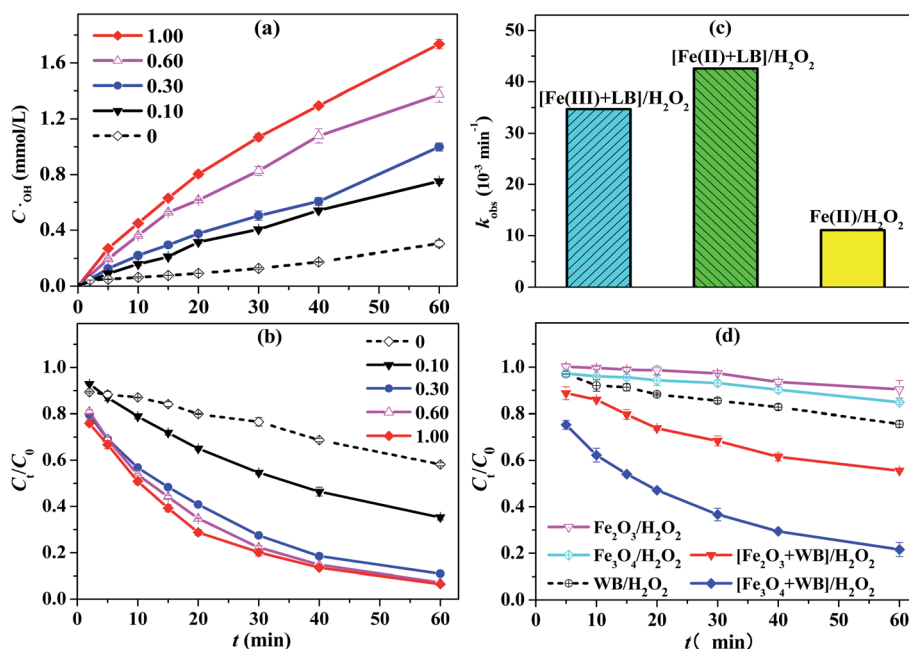


Fig. 3 (a) Cumulative  $\cdot\text{OH}$  production, and (b) removal of 2,4-D ( $C_0 = 20 \text{ mg L}^{-1}$ ) by  $[\text{Fe}(\text{III}) + \text{LB}]/\text{H}_2\text{O}_2$  with different  $\text{Fe}(\text{III})$  concentrations; (c)  $k_{\text{obs}}$  of 2,4-D removal in  $[\text{Fe} + \text{LB}]/\text{H}_2\text{O}_2$  using  $\text{Fe}(\text{II})$  instead of  $\text{Fe}(\text{III})$ , and (d) removal of 2,4-D ( $C_0 = 20 \text{ mg L}^{-1}$ ) in the oxidation systems using WB ( $1.0 \text{ g L}^{-1}$ ) and iron minerals ( $1.0 \text{ g L}^{-1}$ ). Dosage: LB =  $3.0 \text{ g L}^{-1}$  (if present), and  $\text{H}_2\text{O}_2 = 5.0 \text{ mmol L}^{-1}$ ; and  $\text{pH}_0 = 3.0$ .

the degradation of pollutants in biochar/ $\text{H}_2\text{O}_2$  through releasing dissolved iron at acidic pH.<sup>43,53,54</sup> The more and in-depth investigations are underway to clarify the mechanisms involved in the activation of  $\text{H}_2\text{O}_2$  by both biochar and iron minerals. Herein, the preliminary investigation validated the same role of dissolved iron from solid minerals on catalyzing the oxidation of pollutant in biochar/ $\text{H}_2\text{O}_2$ . These results, together with that obtained by using both  $\text{Fe(III)}$  and  $\text{Fe(II)}$  species, are significant for remediation of contaminated environment by utilizing the ubiquitous iron in water and soil. Furthermore, the co-catalysis of trace dissolved iron with biochar for activation of  $\text{H}_2\text{O}_2$  is also advantageous for treatment of wastewater, because the production of iron sludge can be minimized at an iron concentration as low as  $0.3 \text{ mg L}^{-1}$ . For validating this, the solution after reaction in  $[\text{Fe(III)} + \text{LB}]/\text{H}_2\text{O}_2$  was separated out and its pH was adjusted from acidic to neutral (6.5). The solution maintained to be clear after the pH adjustment (Fig. S6<sup>†</sup>), proving that the iron sludge was hardly produced.

### 3.4. Interactions between dissolved $\text{Fe(III)}$ and biochar

The enhanced activation of  $\text{H}_2\text{O}_2$  in the co-catalytic system is related to the interfacial interactions between dissolved iron and biochar, which was investigated using a  $\text{Fe(III)}$  concentration of  $1.00 \text{ mg L}^{-1}$ , because the lower concentration (*e.g.*  $0.30 \text{ mg L}^{-1}$ ) makes it difficult to measure accurately the concentration of regenerated  $\text{Fe(II)}$ . First, the results shown in Fig. 4 indicate that a significant fraction of aqueous  $\text{Fe(III)}$  was transformed into  $\text{Fe(II)}$  during the oxidation of 2,4-D in  $[\text{Fe(III)} + \text{biochar}]/\text{H}_2\text{O}_2$ . There was hardly any  $\text{Fe(II)}$  detected in  $\text{Fe(III)}/\text{H}_2\text{O}_2$ , while the  $\text{Fe(II)}$  concentrations in  $[\text{Fe(III)} + \text{LB}]/\text{H}_2\text{O}_2$  and  $[\text{Fe(III)} + \text{WB}]/\text{H}_2\text{O}_2$  reached to  $0.451$  and  $0.608 \text{ mg L}^{-1}$  after 60 min, respectively. Therefore, nearly 45% and 80% of aqueous iron were present in  $\text{Fe(II)}$  species in  $[\text{Fe(III)} + \text{LB}]/\text{H}_2\text{O}_2$  and  $[\text{Fe(III)} + \text{WB}]/\text{H}_2\text{O}_2$ , respectively, if calculated on the basis of total aqueous iron measured at 60 min. Second, the total aqueous iron in  $[\text{Fe(III)} + \text{WB}]/\text{H}_2\text{O}_2$  ( $\sim 0.75 \text{ mg L}^{-1}$ ) is apparently fewer than the added  $\text{Fe(III)}$  ( $1.00 \text{ mg L}^{-1}$ ), and the total aqueous iron in  $[\text{Fe(III)} + \text{LB}]/\text{H}_2\text{O}_2$  is also fewer than the sum of added

$\text{Fe(III)}$  ( $1.00 \text{ mg L}^{-1}$ ) and the released  $\text{Fe}$  ( $\sim 0.3 \text{ mg L}^{-1}$ ) from LB. The results indicate that nearly 25% of aqueous iron was adsorbed by biochar.

The reduction and adsorption of aqueous  $\text{Fe(III)}$  by other carbon materials in the Fenton-like process have been reported previously. Qin *et al.*<sup>55</sup> found that  $\text{Fe(III)}$  ions were adsorbed onto the surface of hydrothermal carbon (HTC) to form complexes with hydroxyl groups. Then  $\text{Fe(III)}$  was reduced into  $\text{Fe(II)}$  through electron transfer from persistent free radicals (PFRs) in HTC with the mediation of  $\text{Fe(III)}$  complexes. Peng *et al.*<sup>56</sup> suggested that the reduction of  $\text{Fe(III)}$  by carbon nanotubes (CNTs) occurred through direct electron transfer or mediated by adsorption of both  $\text{Fe(III)}$  and  $\text{H}_2\text{O}_2$  onto CNTs surface. Yang *et al.*<sup>57</sup> proposed a new pathway for the fast reduction of  $\text{Fe(III)}$  in a Fenton-like process with presence of functionalized multi-walled carbon nanotubes (FCNT-H), namely  $\text{Fe(III)}$  was reduced by  $\text{H}_2\text{O}_2$  through a mediation of  $\text{Fe(III)}$  complexes with carboxyl groups on the FCNT-H surface. Like these carbon materials, biochar has shown its potential in electron donation and transfer in AOPs, but the specific routes are different for the biochars prepared at different conditions. For example, Wang *et al.*<sup>29</sup> and Hong *et al.*<sup>34</sup> have proposed that PFRs in the biochar prepared at  $500^\circ\text{C}$  were the dominant electron donors to reduce  $\text{Fe(III)}$  for activation of persulfates. However, our previous study has shown that the biochar prepared at  $700^\circ\text{C}$  was more reactive on facilitating the  $\text{Fe(III)}/\text{Fe(II)}$  cycle in Fenton process,<sup>20</sup> although Yang *et al.*<sup>58</sup> and Zeng *et al.*<sup>30</sup> have reported that there are few PFRs in the biochar prepared at  $700^\circ\text{C}$ . Therefore, PFRs are not likely to be the major electron donors for reduction of  $\text{Fe(III)}$  in this study. Recently, Ouyang *et al.*,<sup>59</sup> and Wang and Chen<sup>60</sup> proposed that the defects in the biochar prepared at high temperature ( $700\text{--}900^\circ\text{C}$ ) acted as electron donors for activation of persulfates. In addition, the graphite-like structure in biochar facilitated the electron transfer.<sup>61–63</sup> In the two biochars (LB and WB) used in this study, the defects was proved by the relatively high intensity ratios of D-band ( $1350 \text{ cm}^{-1}$ ) to G-band ( $1570 \text{ cm}^{-1}$ ) ( $I_D/I_G$ ) in Raman spectra (Fig. 5(a)), and the graphite-like structure was confirmed by the two bands at  $2\theta = 23.5$  and  $43.5^\circ$  in XRD patterns (Fig. 5(b)). So, it can be supposed

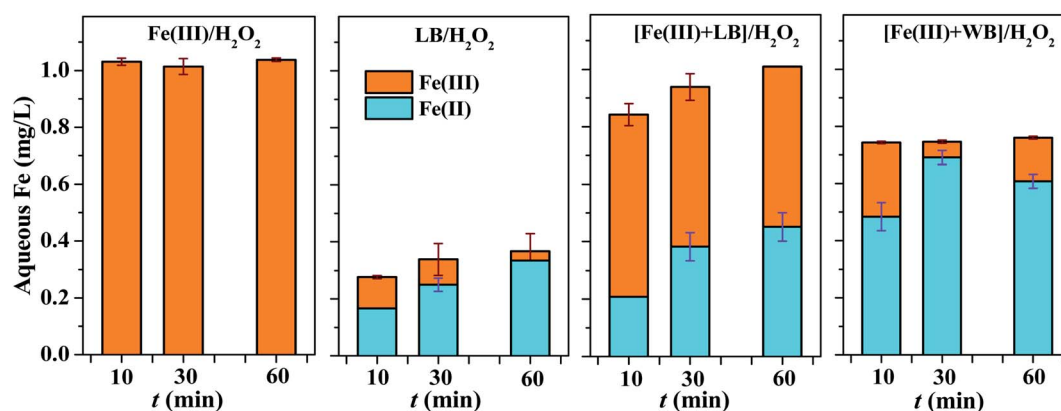
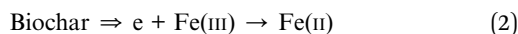
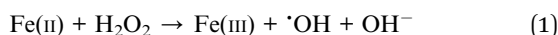


Fig. 4 Change of total Fe and  $\text{Fe(II)}$  concentrations in the solutions of different oxidation systems. Dosage:  $\text{Fe(III)} = 1.00 \text{ mg L}^{-1}$ , LB or WB =  $3.0 \text{ g L}^{-1}$ , and  $\text{H}_2\text{O}_2 = 5.0 \text{ mmol L}^{-1}$ ; and  $\text{pH}_0 = 3.0$ .



that the defects and graphite-like structure in LB and WB contributed to the electron donation/transfer for direct reduction of Fe(III) by biochar. Such direct reduction is further proved by mixing biochar in aqueous Fe(III) solution (Fig. 5(d)). As expected, Fe(II) accounted for 73% and 84% (at 60 min) of aqueous iron in the suspension of Fe(III) + LB and Fe(III) + WB, respectively. Furthermore, the surface hydroxyl groups were observed by FTIR in both biochars (Fig. 5(c)), and carboxyl groups were found in the biochars recycled after reaction (Fig. S7†). These oxygen-containing functional groups can form complexes with Fe(III),<sup>55,57</sup> which bridged the electron transfer from biochar to reduce Fe(III). Thus, the reduction of Fe(III) by biochar makes the regeneration of Fe(II) fast enough for catalyzing the sustainable  $\cdot\text{OH}$  production (eqn (1) and (2)), so that the efficient oxidation of pollutant was achieved even at a trace level of iron supply in the co-catalytic system. However, the trace iron used in this study makes it difficult to obtain more specific information about the Fe(III) complexation on biochar.



### 3.5. Comparison between two biochars

The different performance on pollutant removal between the two biochars, LB and WB, was observed in different oxidation

systems. First, the faster degradation of 2,4-D was observed in LB/H<sub>2</sub>O<sub>2</sub> than that in WB/H<sub>2</sub>O<sub>2</sub> (both without added Fe(III)) (Fig. 1(c)), despite the higher adsorption of 2,4-D by WB than that by LB, according to the adsorption isotherms obtained after 24 h of equilibration (Fig. S8†). The faster 2,4-D degradation in LB/H<sub>2</sub>O<sub>2</sub> corresponds to the relatively more  $\cdot\text{OH}$  production in LB/H<sub>2</sub>O<sub>2</sub> than that in WB/H<sub>2</sub>O<sub>2</sub> (Fig. 2(a)). The results indicate the relatively higher reactivity of LB for activation of H<sub>2</sub>O<sub>2</sub>, which should be ascribed to the different compositions of biochar. Huang *et al.*<sup>64</sup> and Tao *et al.*<sup>65</sup> have reported that the inorganic components such as metals in biochar contributed significantly to activation of H<sub>2</sub>O<sub>2</sub>. In this study, there are more inorganic components in LB than in WB, as indicated by the higher ash content (23.4% vs. 0.89%) and lower carbon content (66.0% vs. 93.9%) in LB (Table S1†). The iron leaching was observed in LB/H<sub>2</sub>O<sub>2</sub> (Fig. 4), indicating there are trace protogenetic iron species in LB. Besides, nitrogen element (3.36% by weight) derived from the biomass feedstock was detected in LB, indicating some pyrogenic N species in LB. Both Fe and N species inherent in biochar have been reported to be favorable for enhancing the activation of peroxides.<sup>26,64</sup> Despite this, the yield of  $\cdot\text{OH}$  in LB/H<sub>2</sub>O<sub>2</sub> is still very low (9.27%) when calculated on the basis of much consumption of H<sub>2</sub>O<sub>2</sub> (Fig. 2(b)). So, there should be some components in LB that impeded the transformation of H<sub>2</sub>O<sub>2</sub> into  $\cdot\text{OH}$ , except those favored the production of  $\cdot\text{OH}$ .

Different from that observed in biochar/H<sub>2</sub>O<sub>2</sub>, the higher  $k_{\text{obs}}$  for oxidation of 2,4-D and more TOC removal were obtained in

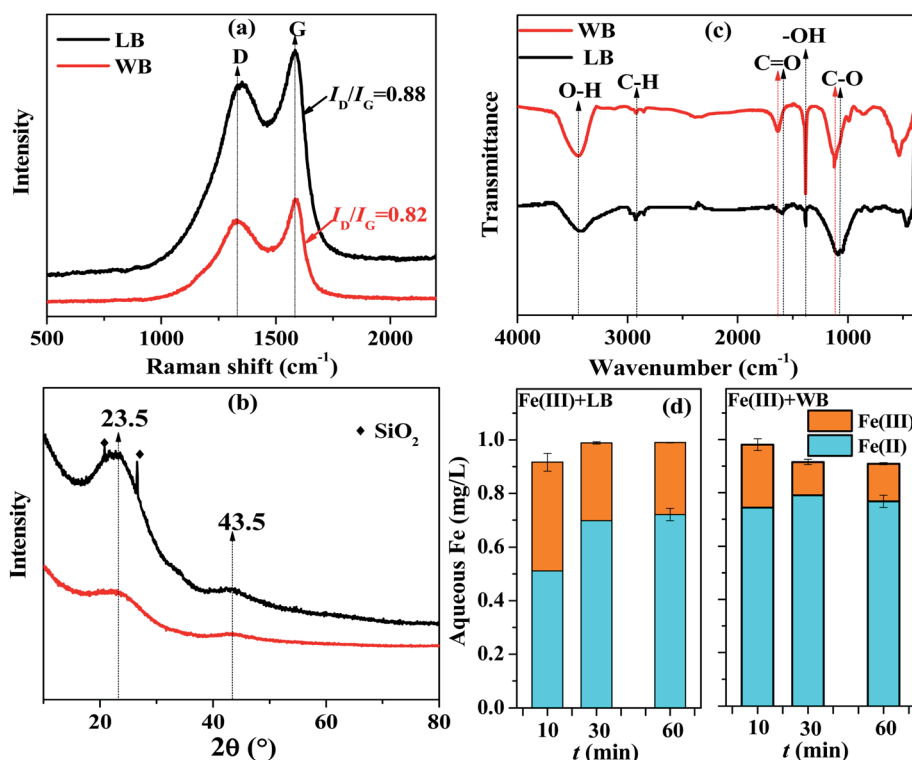
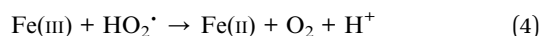
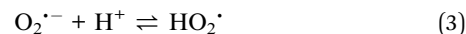


Fig. 5 (a) Raman spectra, (b) XRD patterns, and (c) FTIR of LB and WB samples; and (d) change of total Fe and Fe(II) concentrations in the mixtures of biochar (3.0 g L<sup>-1</sup>) and Fe(III) solution (1.00 mg L<sup>-1</sup>) at pH<sub>0</sub> = 3.0.



[Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> than those in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> (Fig. 1(c) and (d)), which is in consistency with the more  $\cdot$ OH production in the former system (Fig. 2(a)). Furthermore, the apparently higher  $\cdot$ OH yield in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> (89.8%) was obtained, accompanying with apparently less consumption of H<sub>2</sub>O<sub>2</sub> than in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> (Fig. 2(b)). As there is hardly any inorganic component in WB, the enhanced  $\cdot$ OH yield should be derived from the interfacial interactions between biochar and the added trace Fe(III). Fig. 4 shows that the aqueous Fe(II) concentration in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> is apparently higher than that in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub>, namely more regenerated Fe(II) in the former system. So, the performance of the co-catalytic system on pollutant removal relies heavily on regeneration of Fe(II), as Fe(II) is much more efficient on catalyzing the production of  $\cdot$ OH than biochar. The relatively purer carbon matrix of WB, as indicated by its higher carbon content (93.9%), makes it a better conductor for electron transfer to reduce Fe(III). The high surface area of WB also favors electron transfer to Fe(III) by providing more reactive sites. Besides the direct reduction of Fe(III) by biochar, another possible route (eqn (3) and (4)) involving O<sub>2</sub><sup>•−</sup> should also contribute to the regeneration of Fe(II) in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub>, because more O<sub>2</sub><sup>•−</sup> was produced in [Fe(III) + WB]/H<sub>2</sub>O<sub>2</sub> than that in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> according to the EPR analysis (Fig. 2(d)). O<sub>2</sub><sup>•−</sup> was produced from electron transfer from biochar to O<sub>2</sub>, the byproduct of H<sub>2</sub>O<sub>2</sub> decomposition, according to that reported by Fang *et al.*<sup>66</sup> Therefore, the direct and indirect reduction of Fe(III) by biochar dominated the performance of the co-catalytic system on pollutant removal, and the specific routes of electron transfer from biochar (the electron donor) involved in adsorption of Fe(III) by biochar and mediation of H<sub>2</sub>O<sub>2</sub>/O<sub>2</sub>.



### 3.6. Effect of other reaction conditions and reusability of biochar

The [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> system was chosen for further investigations on the influence of other reaction conditions including pH<sub>0</sub>, biochar and H<sub>2</sub>O<sub>2</sub> dosages on the degradation rate of 2,4-D. One reason is that LB derives from the liquor waste that is seldom utilized, and another reason is related to its poor adsorption to 2,4-D, which is favorable for highlighting the contribution of oxidation to 2,4-D removal. First, pH is the most influential factor determining the oxidation efficiency of organic pollutants in AOPs. In this study, the faster degradation of 2,4-D was recorded at pH<sub>0</sub> = 2.5–3.0 in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> (Fig. 6(a)), which is same as that observed in ordinary Fenton process. pH monitoring during the reaction indicates that there was not apparent fluctuation from the initial pH<sub>0</sub>, except at pH<sub>0</sub> = 4.0 when the degradation was very slow (Fig. S9†). The stable pH during reaction implies that the release of OH<sup>−</sup> along with  $\cdot$ OH production (eqn (1)) was neutralized by release of H<sup>+</sup> from interactions between Fe(III) and biochar, such as complexation of Fe(III) with hydroxyl and carboxyl groups in biochar. The pH higher than 4.0 leads to the precipitation of Fe(III) hydroxides, so that the regeneration of Fe(II) will be suppressed. In contrast, the excess acidity may suppress the formation of Fe(III)

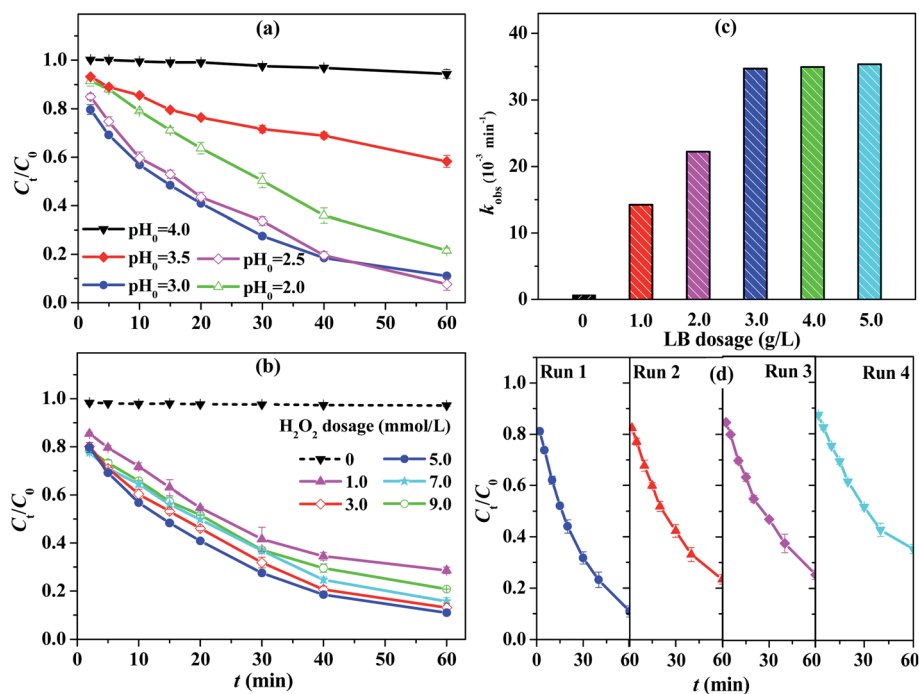


Fig. 6 Removal of 2,4-D ( $C_0 = 20 \text{ mg L}^{-1}$ ) at (a) different pH<sub>0</sub>, and (b) different H<sub>2</sub>O<sub>2</sub> dosage; (c) Change of  $k_{\text{obs}}$  with LB dosage; and (d) 2,4-D removal in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> with repetitive use of LB. Typical conditions (except the variable): Fe(III) = 0.30 mg L<sup>−1</sup>, LB = 3.0 g L<sup>−1</sup>, and H<sub>2</sub>O<sub>2</sub> = 5.0 mmol L<sup>−1</sup>, and pH<sub>0</sub> = 3.0.



complexes, which is not favorable for regeneration of Fe(II) either. Overall, the performance of the co-catalytic system on oxidation of pollutant still relies heavily on acidic pH, which needs improvement with additional measures in future work. For example, using natural chelating reagents may enhance the efficiency of oxidative removal of organic pollutants at neutral pH.<sup>67</sup> Second, the degradation reaction in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> is not as sensitive to H<sub>2</sub>O<sub>2</sub> dosage as to pH (Fig. 6(b)). The 2,4-D degradation remained at a relatively fast rate when the H<sub>2</sub>O<sub>2</sub> dosage was as low as 1.0 mmol L<sup>-1</sup>, although the fastest one was observed at 5.0 mmol L<sup>-1</sup>. Less H<sub>2</sub>O<sub>2</sub> dosage led to insufficient supply of H<sub>2</sub>O<sub>2</sub> and slowed down the degradation at latter reaction stage, while oversupply of H<sub>2</sub>O<sub>2</sub> also resulted in slower pollutant degradation due to its reaction with <sup>•</sup>OH.<sup>68</sup> In this study, the high <sup>•</sup>OH yield from H<sub>2</sub>O<sub>2</sub> decomposition reduced the H<sub>2</sub>O<sub>2</sub> dosage required in the co-catalytic system, which is beneficial for reducing the cost in water treatment. Third, the degradation rate of 2,4-D in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> became faster with the increase of LB dosage from 1.0 to 3.0 g L<sup>-1</sup> (Fig. 6(c) and Fig. S10†). The *k*<sub>obs</sub> was enhanced from 14.2 × 10<sup>-3</sup> to 34.7 × 10<sup>-3</sup> min<sup>-1</sup>, and the removal efficiency was lifted from 63.3% to 89.0% after 60 min of reaction. The biochar of higher dosage can provide more reactive sites for regeneration of Fe(II), which leads to the faster oxidation of pollutants. However, further increase of biochar dosage above 3.0 g L<sup>-1</sup> did not lead to significant change on the degradation rate of 2,4-D. The reason should be due to the trace amount of Fe(III) used in this study, which limited the total amount of regenerated Fe(II).

Finally, the reusability of biochar in the co-catalytic system for removal of 2,4-D was evaluated. As can be seen in Fig. 6(d), the oxidation of 2,4-D in [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> gradually slowed down after each sequential run. The declining reactivity of biochar in repetitive experiments is supposed to be resulted from surface contamination by the intermediates (such as short-chain organic acids) derived from incomplete 2,4-D degradation, because the biochar was reused directly without any purification. Another reason may be related to the surface oxidation of biochar.<sup>69</sup> The more carboxyl groups were observed by FTIR (Fig. S7†) on the recycled biochar, confirming the change of surface properties of biochar after reaction. The decrease of biochar's surface area after reaction (e.g. from 409 m<sup>2</sup> g<sup>-1</sup> to 360 m<sup>2</sup> g<sup>-1</sup> for WB) implies fewer reactive sites available for repetitive use. However, no other apparent change can be found by XRD analysis (Fig. S11†) and SEM observation (Fig. S12†) of the recycled biochar, indicating there is not structural change occurred on biochar during the reaction. Further extension of reaction time may help transform the degradation intermediates more completely and enhance the reusability of biochar. For example, 2,4-D was removed completely by [Fe(III) + LB]/H<sub>2</sub>O<sub>2</sub> after 180 min of reaction, accompanied with an enhanced TOC removal to 71.2%.

## 4. Conclusions

The co-catalysis of trace dissolved Fe(III) with biochar significantly enhanced the activation of H<sub>2</sub>O<sub>2</sub>, and consequently accelerated the oxidation of organic pollutants with a reaction

rate comparable to other AOPs. Replacement of dissolved Fe(III) with aqueous Fe(II) and iron minerals (hematite and magnetite) also showed the similar catalytic effect on pollutant degradation by biochar/H<sub>2</sub>O<sub>2</sub>. The interfacial interactions between dissolved Fe(III) and biochar resulted in fast regeneration of Fe(II) in the co-catalytic system. The defects in biochar acted as the major electron donors, and the graphite-like structure and oxygen-containing functional groups in biochar facilitated the electron transfer. The aqueous iron at trace level (0.3 mg L<sup>-1</sup>) prevented the precipitation of iron sludge, and biochar is advantageous over other elaborately synthesized materials in view of cost and sustainability, so their cooperation provides an efficient way to enhance the performance of AOPs on wastewater treatment. Furthermore, as dissolved iron can be found in natural water, its potential on enhancing the activation of H<sub>2</sub>O<sub>2</sub> by biochar may be utilized for remediation of contaminated environment. The future work should address the influence of co-solutes in water on performance of this novel co-catalytic oxidation system, and its applicability at neutral pH should be improved with additional measures.

## Author contributions

Dongqing Feng: methodology, investigation, writing – original draft. Jianxin Shou: conceptualization, investigation, writing – review & editing, funding acquisition. Sen Guo: investigation. Mengna Ya: validation. Jianfa Li: methodology, investigation, writing – review & editing, funding acquisition. Huaping Dong: investigation, funding acquisition. Yimin Li: supervision.

## Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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