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## Original Research

MnO<sub>2</sub>-Catalyzed electrocatalytic mineralization of triclosan in chlorinated wastewater

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## ABSTRACT

The rising concentrations of xenobiotic aromatic compounds in the environment pose significant risks to human and ecosystem health. Developing a universal, environmentally benign, and scalable platform for mineralizing organic pollutants before their release into the environment is therefore crucial. Electrocatalysis can be highly advantageous for wastewater treatment because it is immediately responsive upon applying potential, requires no additional chemicals, and typically uses heterogeneous catalysts. However, achieving efficient electrochemical mineralization of wastewater pollutants at parts-per-million (ppm) levels remains a challenge. Here, we report the use of manganese dioxide (MnO<sub>2</sub>), an Earth-abundant, chemically benign, and cost-effective electrocatalyst, to achieve over 99 % mineralization of triclosan (TCS) and other halogenated phenols at ppm levels. Two highly active MnO<sub>2</sub> phases— $\alpha$ -MnO<sub>2</sub>-CC and  $\delta$ -MnO<sub>2</sub>-CC—were fabricated on inexpensive carbon cloth (CC) support and evaluated for their ability to oxidatively degrade TCS in pH-neutral conditions, including simulated chlorinated wastewater, real wastewater, and both synthetic and real landfill leachates. Total organic carbon analysis confirmed the effective degradation of TCS. Electron paramagnetic resonance and ultraviolet–visible spectroscopy identified reactive oxygen species, enabling the construction of a detailed TCS degradation pathway. Upon optimization, the TCS removal rate reached 38.38 nmol min<sup>−1</sup>, surpassing previously reported rates achieved with precious and toxic metal co-catalysts. These findings highlight MnO<sub>2</sub>-CC as a promising, eco-friendly electrocatalyst with strong potential for upscaled remediation of organic pollutants in wastewater treatment.

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## 1. Introduction

Industrial and anthropogenic activities generate large amounts of harmful organic waste that seriously threatens the environment and human health [1]. In particular, halogenated organic pollutants

(HOPs) have high bioaccumulation potentials and are highly resistant to conventional biodegradation [2,3]. Thus, HOPs persist in the environment, where they can bioaccumulate and cause acute cytotoxicity in humans and other organisms [4]. For instance, triclosan (TCS) is a typical HOP widely used as an antiseptic and antimicrobial chemical in many quotidian goods, such as cosmetic, hygiene, and household cleaning products. It has been entering wastewater treatment plants (WWTPs) in a range of concentrations (0.00007–14 parts per million) for more than 30 years [5]. Over recent decades, various techniques, including physical methods

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(e.g., adsorption), oxidation processes (e.g., chlorination, photolysis/photocatalysis, ozonation, Fenton processes, and electrochemical oxidation), and biological approaches have been developed to treat triclosan-contaminated wastewater [6]. Among these, tertiary treatments, such as Fenton oxidation, electrolysis, piezo-electrocatalysis, and photocatalysis, have been extensively studied for degrading refractory contaminants in aqueous environments [7]. However, their applicability is often limited by factors such as low reaction rates, sludge stabilization and activation challenges, incomplete dehalogenation, and the high costs of stoichiometric reagents [7]. Although biodegradation and adsorption are cost-effective and environmentally friendly, they also face challenges, such as the generation of solid residues and the need for pre-treatment to convert pollutants into less toxic byproducts [8].

Primary treatment is estimated to remove approximately 28 % of TCS from wastewater, while secondary and tertiary treatments can remove over 80 % [9]. The TCS concentrations in WWTP influents and effluents worldwide are 0.0013–86.2 parts per million (ppm) and 0.0031–5.53 ppm, respectively, indicating that the complete removal of TCS from wastewater remains challenging [9].

Over the past decade, tertiary treatments, such as electrocatalytic treatments [10,11], have drawn increasing attention due to their unique ability to function immediately and robustly in ambient aqueous conditions. Moreover, several authors have reported using electrochemical or hybrid methods for TCS degradation (Table 1), and the TCS removal rates of these methods at ppm levels ranged from 5.77 to 28.8 nmol min<sup>-1</sup>. Although all of these methods operate under ambient conditions, many require elaborated electrodes, such as boron-doped diamond (BDD) [12], or even toxic elements such as lead [13], to degrade TCS. Some hybrid methods do not need elaborated or toxic materials; for example, an electrocatalytic Fenton method had a high TCS removal rate (i.e., 43.25 nmol min<sup>-1</sup>) [14]. However, it generated stoichiometric

quantities of ferrous ions to facilitate the rapid degradation of TCS with hydrogen peroxide. Other advanced oxidation processes, such as photocatalytic and bio-chemical hybrid processes, have been applied for TCS degradation and exhibited reasonable TCS removal rates (i.e., 28.8 nmol min<sup>-1</sup>) [14–16]. Nevertheless, these methods have several limitations or require specific conditions, such as costly electrodes, an organic cosolvent, or stoichiometric oxidants, thereby hindering their large-scale implementation. Therefore, there is a need to use Earth-abundant elements to fabricate active electrodes that can serve as an economical and feasible means of removing persistent TCS from wastewater.

Manganese oxide (MnO<sub>2</sub>)-based electrocatalysts are highly promising, as they are inexpensive, chemically benign, and exhibit high catalytic activities [25–27]. MnO<sub>2</sub> can exist in different structural phases, namely the  $\alpha$ -,  $\beta$ -,  $\gamma$ -,  $\delta$ -, and  $\lambda$ -phases, which consist of the same octahedral MnO<sub>6</sub> units linked in different ways. Thus far, natural birnessite MnO<sub>2</sub> has been applied in slurries for the non-electrocatalytic oxidation of organic compounds, such as phenols [28], anilines [29], fluoroquinolones [30], and antibacterial amine-oxides [31]. The high structure–activity variability in MnO<sub>2</sub> presents an excellent opportunity to develop a cost-effective electrocatalyst to remove TCS from wastewater [32].

Herein, we report a novel hydrothermal exfoliation protocol to synthesize two different nano-MnO<sub>2</sub> structures, namely  $\alpha$ -phase MnO<sub>2</sub> nanoneedles (NNs) and  $\delta$ -phase nanosheet arrays (NSAs), which were anchored on a cost-effective carbon cloth (CC) support to form MnO<sub>2</sub> electrocatalysts. Both MnO<sub>2</sub> electrocatalysts mineralized TCS at ppm levels in chlorinated wastewater simulated as WWTP effluent [33] at room temperature under open-atmosphere conditions. Reactive oxygen species (ROS) were identified using electron paramagnetic resonance (EPR) and ultraviolet–visible (UV–vis) spectroscopy, and the mechanism of TCS degradation and the intermediates involved in substrate scope studies were

**Table 1**  
Advanced oxidation methods to mineralize triclosan in aqueous environments.

| Method                  | Concentration (ppm) | Removal rate (nmol min <sup>-1</sup> )     | Degradation rate (%)                          | Conditions <sup>a</sup> | Catalyst                                                                 | Co-catalyst/Oxidant                                                                                 | Solvent                            |
|-------------------------|---------------------|--------------------------------------------|-----------------------------------------------|-------------------------|--------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------|------------------------------------|
| Electro. (this work)    | 10                  | 38.38                                      | 100                                           | Ambient                 | $\delta$ -MnO <sub>2</sub> on CC.<br>20 mA cm <sup>-2</sup>              | –                                                                                                   | pH 7 DI water                      |
| Electro [17].           | 0.8                 | 5.77                                       | N.D.                                          | Ambient (22 °C, dark)   | First: Ti/MMO<br>Second: Nb/BDD<br>2–14 mA cm <sup>-2</sup>              | –                                                                                                   | pH 8 DI water                      |
| Electro [18].           | 3                   | 830                                        | 99.9                                          | Ambient                 | Ti/SnO <sub>2</sub> –Sb/Ce–PbO <sub>2</sub> 2<br>–10 mA cm <sup>-2</sup> | 400 ppm Ce <sup>3+</sup>                                                                            | pH 3–11 DI water                   |
| Electro [19].           | 5                   | 28.8                                       | 98                                            | Ambient                 | Ti/SnO <sub>2</sub> –Sb/Gd–PbO <sub>2</sub><br>10 mA cm <sup>-2</sup>    | Gd <sup>3+</sup>                                                                                    | pH 3 urine                         |
| Photo [15,20].          | 10                  | 28.8                                       | 100                                           | Ambient                 | Anatase and rutile TiO <sub>2</sub>                                      | –                                                                                                   | pH 6.6 DI water                    |
| Fenton [16,21]          | 10                  | 28.8                                       | 100                                           | Ambient (20–50 °C)      | 1 ppm Fe <sup>3+</sup>                                                   | 25 ppm H <sub>2</sub> O <sub>2</sub>                                                                | pH 3 HNO <sub>3</sub> and DI water |
| E.-Fenton [16]          | 50                  | <sup>b</sup> k = 0.0128 min <sup>-1</sup>  | 91.39                                         | Ambient                 | Fe:Mn-SBC<br>15 mA cm <sup>-2</sup>                                      | 2600 ppm KMnO <sub>4</sub><br>748 ppm Fe <sup>2+</sup><br>5.42 ppm H <sub>2</sub> O <sub>2</sub>    | pH 3 real waste water              |
| E.-Fenton [14,22]       | 10                  | 43.25                                      | 99.9                                          | Ambient                 | 2:1 Fe:Mn-PC<br>10 mA cm <sup>-2</sup>                                   | –                                                                                                   | pH 3 DI water                      |
| Fenton [23]             | 10                  | One: 14.41 (180 min)<br>Two: 86.5 (30 min) | One: 82.7 % (180 min)<br>Two: 96.1 % (30 min) | Ambient                 | 500 ppm BiFeO <sub>3</sub> NPs                                           | One: 10 mM H <sub>2</sub> O <sub>2</sub><br>Two: 0.5 mM EDTA<br>10 mM H <sub>2</sub> O <sub>2</sub> | pH 6 DI water                      |
| Biochemical hybrid [24] | 5                   | 28.8                                       | 100                                           | Anaerobic               | 1000 p.m. Pd/Fe NPs<br>Laccase with syringaldehyde                       | –                                                                                                   | pH 5.6–7.8 DI water                |

Note: Electro., Electrocatalytic; Photo., Photocatalytic; E.-Fenton, Electrocatalytic Fenton; Conc., Triclosan concentration; N.D., Not detected; BDD, Boron-doped diamond; MMO, Mixed metal oxides; SBC, Sludge-based carbon; CB, Carbon black; PC, Porous carbon; NPs, Nanoparticles; EDTA, Ethylenediaminetetraacetic acid; DI water, Deionized water; k, rate constant.

<sup>a</sup> Unless specified otherwise, “ambient” refers to the conditions of 25 °C and 1 atm.

<sup>b</sup> Volume data were not available.

elucidated using gas chromatography–mass spectrometry. Total organic carbon (TOC) analyses were performed to confirm TCS mineralization.

## 2. Materials and methods

### 2.1. Materials

All chemicals were used as received. Potassium permanganate ( $\text{KMnO}_4$ ), ammonium sulfate ( $(\text{NH}_4)_2\text{SO}_4$ ), manganese sulfate monohydrate ( $\text{MnSO}_4 \cdot \text{H}_2\text{O}$ ), and 2-bromophenol (2-BrPh) were obtained from Macklin (Shanghai, China). 4-Chlorophenol (4-CP), bisphenol A (BPA), and triclosan (TCS) were obtained from the Tokyo Chemical Industry (Tokyo, Japan). 2,4,6-Trichlorophenol (2,4,6-TCP) was obtained from Thermo Scientific (Waltham, United States). Methanol (High-performance Liquid Chromatography grade), acetone, and ethanol (Analytical grade) were used. Sodium sulfate ( $\text{Na}_2\text{SO}_4$ ) and sodium chloride ( $\text{NaCl}$ ) were obtained from Fisher (Pittsburgh, United States).

### 2.2. Fabrication of $\text{MnO}_2$ nanostructures on a carbon cloth support

$\text{MnO}_2$  nanostructures were fabricated on CC via hydrothermal exfoliation (Fig. 1a). First, the CC was sonicated successively in acetone, ethanol, and deionized water to remove adsorbed impurities. For  $\alpha$ -phase  $\text{MnO}_2$  preparation, the sonicated CC (1 cm  $\times$  3 cm) was hung on the walls of a Teflon-lined autoclave while immersed in 40 mL of an aqueous solution of 220 mg  $\text{KMnO}_4$  and 60 mg  $\text{MnSO}_4 \cdot \text{H}_2\text{O}$ . Next, the autoclave was sealed, heated at 140 °C for 20 h, and cooled to room temperature overnight. As a result, a thin film of  $\alpha$ - $\text{MnO}_2$  was deposited on the CC, with some residual powder left at the autoclave bottom.  $\delta$ -phase  $\text{MnO}_2$  ( $\delta$ - $\text{MnO}_2$ ) was prepared on CC under similar hydrothermal conditions but with different precursors and heat treatment. Here, the hanging sonicated CC was immersed in 40 mL of a solution containing 126.4 mg  $\text{KMnO}_4$  and 42.8 mg  $(\text{NH}_4)_2\text{SO}_4$  and heated at 110 °C for 20 h.

All  $\text{MnO}_2$  electrodes were annealed at 350 °C for 2 h at a heating rate of 10 °C  $\text{min}^{-1}$ . The respective electrodes were denoted as  $\alpha$ - and  $\delta$ - $\text{MnO}_2$ -CC. The electrodes were stored at room temperature until use. The residual dark brown ( $\alpha$ - $\text{MnO}_2$ ) and black ( $\delta$ - $\text{MnO}_2$ ) precipitates from the autoclaves were also collected and stored under similar conditions for further characterization.

### 2.3. Electrocatalytic oxidation experiments and product analysis

All electrolysis experiments were conducted in an undivided batch reactor (100 mL) containing a  $\text{MnO}_2$  anodic working electrode (geometric area = 1  $\text{cm}^2$ ) and nickel foam as a cathodic counter electrode. All solutions contained 10 ppm of endocrine disruptors (EDCs) and were prepared in aqueous media containing 50 mM  $\text{Na}_2\text{SO}_4$  and 5 mM  $\text{NaCl}$  to provide background ionic strength and mimic the salinity of natural wastewater effluent [34]. Chronopotentiometric electrolysis was performed at 20, 40, and 80 mA  $\text{cm}^{-2}$  at room temperature ( $23 \pm 2$  °C). Aliquots of samples were collected periodically and analyzed using a high-performance liquid chromatography (HPLC, Agilent 1260 Infinity II) equipped with a photodiode array detector and a reverse-phase Zorbax XDB-C18 column (3.9  $\times$  150 mm), which was eluted isocratically with a methanol:water (70:30, v/v%) mobile phase at a flow rate of 1 mL  $\text{min}^{-1}$ . The intermediate products were detected using a gas chromatograph (GCMS, Shimadzu 2010 Ultra Gas Chromatograph–Mass Spectrometer) equipped with a capillary column (Agilent DB-5, 30 m  $\times$  0.25 mm internal diameter  $\times$  0.25  $\mu\text{m}$ ) in splitless mode at an injection temperature of 200 °C.

All samples subjected to gas chromatography analysis (100 mL each) were extracted with 4 mL of dichloromethane, followed by drying with anhydrous  $\text{Na}_2\text{SO}_4$ . Gas chromatograms were analyzed by comparison with the National Institute of Standards and Technology library and external references to identify intermediates. The extent of mineralization was determined by TOC analysis (Shimadzu, TOC-L<sub>CPH</sub>).

### 2.4. Structural phase and morphology analysis

The crystallinity and phase structure of the  $\text{MnO}_2$  electrodes and residual powders were determined by X-ray diffraction (PANalytical X'Pert X-ray diffractometer) using copper  $K\alpha$  radiation (40 kV, 30 mA). The qualitative and quantitative surface properties of the prepared electrodes were evaluated by high-resolution scanning electron microscopy (FEI–Philips XL30 Esem-FEG). Raman spectroscopy was performed using a Raman spectrometer (EnSpectr R532, EnSpectr) equipped with a 20 mW, 532 nm laser. X-ray photoelectron spectroscopy (XPS) was performed on a Thermo  $K\alpha$  photoelectron spectrometer, and monoenergetic aluminum  $K\alpha$  radiation was employed to characterize and quantify the distribution of the phases. All electrochemical experiments, i.e., linear sweep voltammetry (LSV) and cyclic voltammetry (CV), were conducted in a three-electrode setup using a CHI 660 E electrochemical station (CH Instruments, Inc., Shanghai) in 15 mL of 1 M  $\text{Na}_2\text{SO}_4$  electrolyte under ambient conditions. Unless specified otherwise, a standard bulk electrolysis reaction involved an as-prepared  $\text{MnO}_2$  electrode as the working electrode (area = 1  $\text{cm}^2$ ); a Nickel mesh and a 3.5 M silver/silver chloride ( $\text{Ag}/\text{AgCl}$ ) electrode were used as the counter and reference electrode, respectively.

#### 2.4.1. Measurement of the electrochemical active surface areas of uncoated and $\text{MnO}_2$ -Coated electrodes

The electrochemical active surface areas (ECSAs) of the electrodes were calculated from the electrochemical double-layer capacitance ( $C_{dl}$ ) of the catalytic surface obtained from double-layer charging curves, which were determined by CV at 0–0.8 V (a non-Faradaic region) at a scan rate of 20–200 mV  $\text{s}^{-1}$  with an interval of 20 mV  $\text{s}^{-1}$ . Specifically,  $C_{dl}$  was measured from the slope of the  $j$ - $\nu$  curve, where  $j$  is the non-Faradaic capacitive current obtained from a CV curve, and  $\nu$  is the scan rate. Next, the ECSA was calculated using the following equation:  $\text{ECSA} = C_{dl}/C_s$ , where  $C_s$  is the specific capacitance set to 0.02 mF  $\text{cm}^{-2}$  as previously published [35].

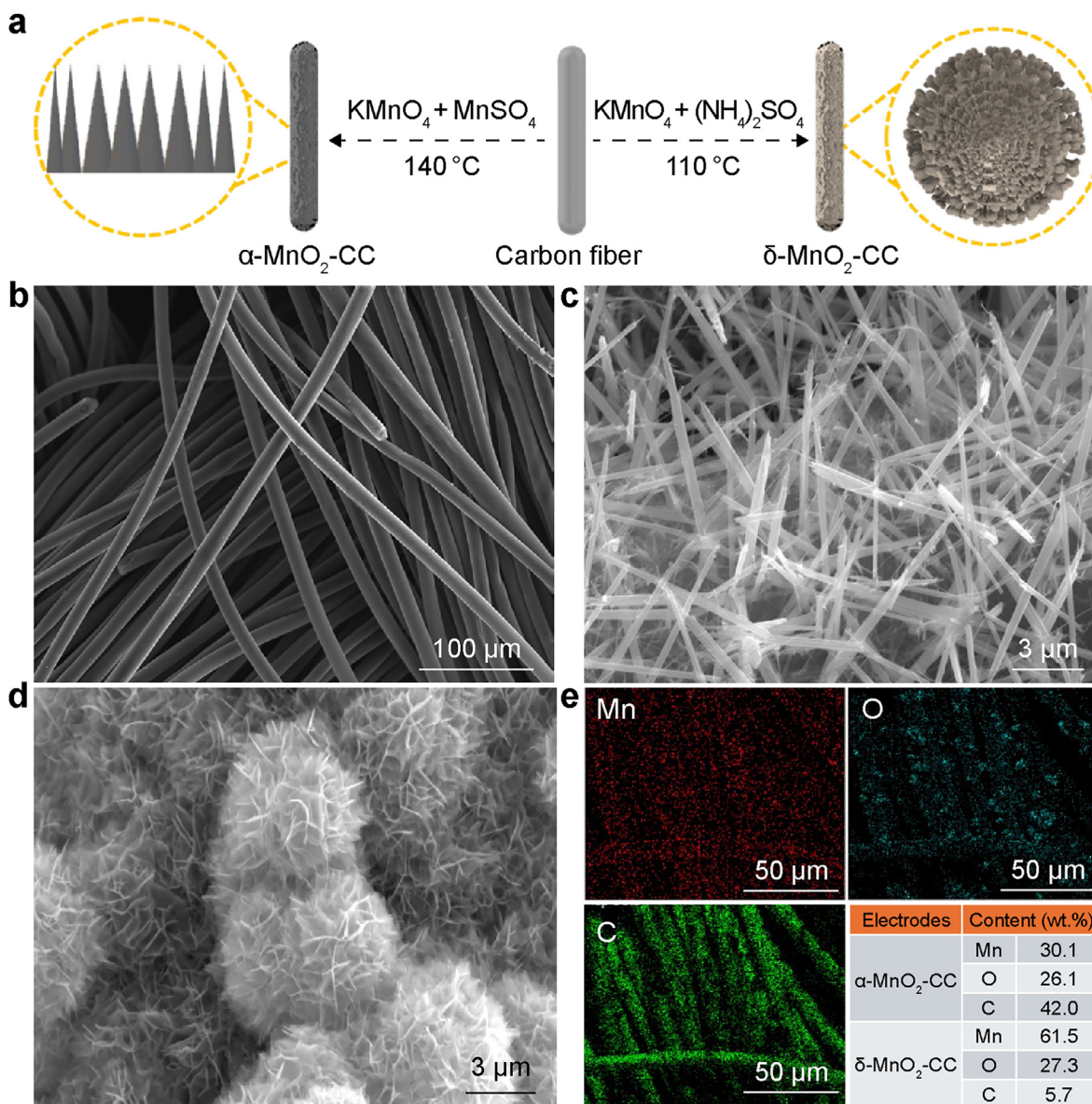
## 3. Results and discussion

### 3.1. Structural analysis of $\alpha$ - and $\delta$ - $\text{MnO}_2$ nanostructures

The nanostructures and morphologies of the  $\text{MnO}_2$  electrodes were examined using scanning electron microscopy (SEM). The bare CC had a smooth fibrous morphology (Fig. 1b). The  $\alpha$ - and  $\delta$ - $\text{MnO}_2$  electrodes uniformly covered the CC surface. The  $\alpha$ - $\text{MnO}_2$ -CC exhibited interconnected NNs on the CC surface, with an estimated thickness of 200–300 nm (Fig. 1c). The  $\delta$ - $\text{MnO}_2$ -CC exhibited interconnected NSAs that formed a micron-sized sea-urchin-like morphology (Fig. 1d). These ordered NSAs formed an open-network-like structure (Supplementary Material Fig. S1a), and the average thickness of a single nanosheet was estimated to be 22.5 nm (Supplementary Material Fig. S1b).

Energy-dispersive X-ray spectrometry (EDS) mapping analysis was performed to quantify elemental Mn, O, and C content. The Mn content (Fig. 1e) illustrates the relatively uniform Mn distribution on the surface of the  $\text{MnO}_2$  electrodes. Strong Mn and O signals were observed (Supplementary Material Fig. S2) in both phases of





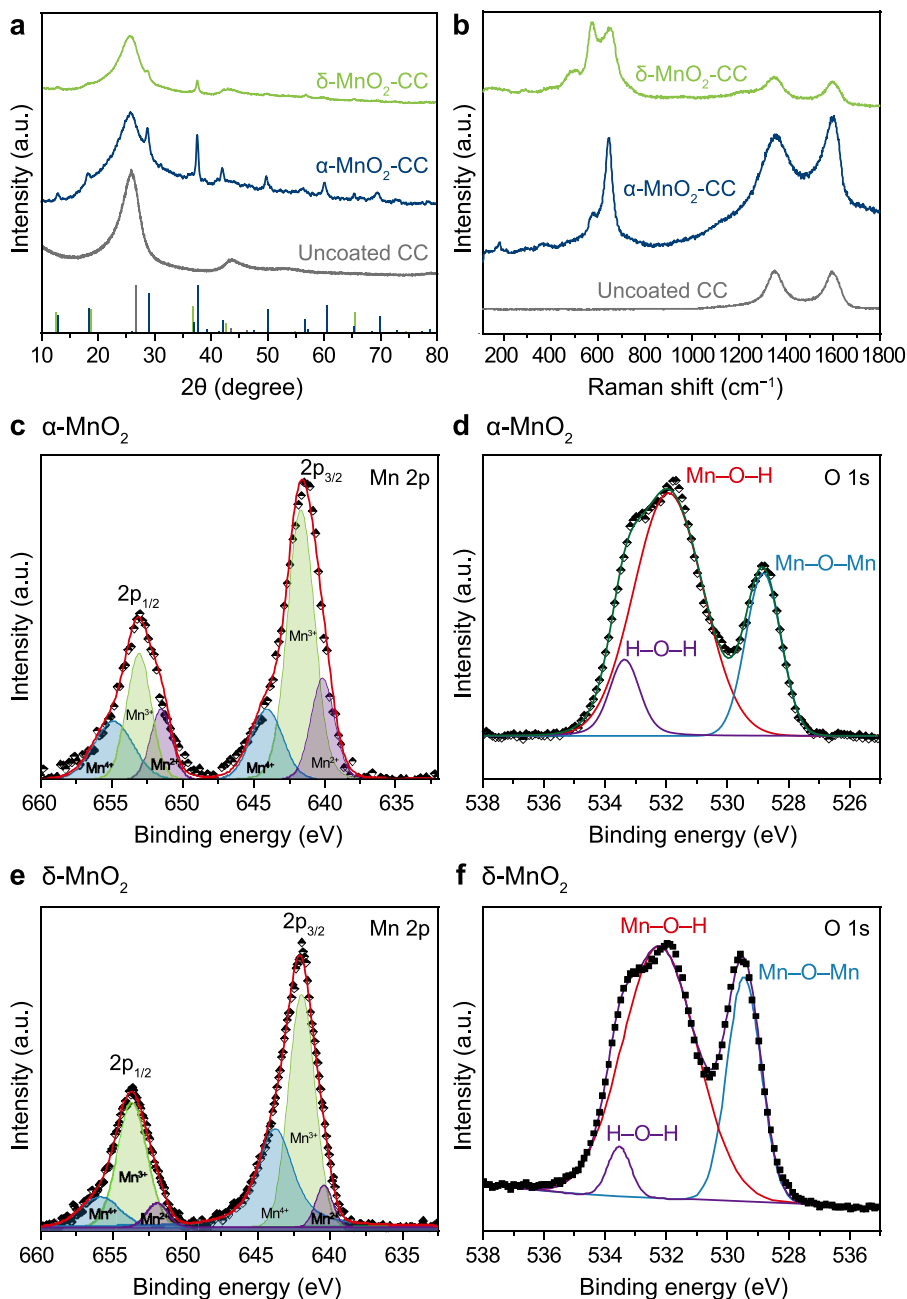
**Fig. 1.** a, Schematic illustration of the fabrication of MnO<sub>2</sub> nanostructures on a carbon cloth (CC) surface. b–d, Scanning electron microscopy images of bare CC (b), α-MnO<sub>2</sub>-CC (c), and δ-MnO<sub>2</sub>-CC (d). e, Energy-dispersive X-ray spectroscopy elemental mapping images of MnO<sub>2</sub> on the CC surface.

MnO<sub>2</sub>-CC, and the intensity was doubled on the δ-MnO<sub>2</sub>-CC (61.54 wt%, [Supplementary Material Fig. S2b](#)) compared with α-MnO<sub>2</sub>-CC (30.06 wt%, [Supplementary Material Fig. S2a](#)). In addition, the exposed carbon content of δ-MnO<sub>2</sub>-CC (5.72 wt%) was much less than that of α-MnO<sub>2</sub>-CC (42.02 wt%), indicating greater coverage of the CC surface on δ-MnO<sub>2</sub>-CC than on α-MnO<sub>2</sub>-CC ([Supplementary Material Table S1](#)).

X-ray diffractometry (XRD) was performed to examine the crystallinity of the MnO<sub>2</sub> electrodes. The XRD patterns of bare CC and of α-phase MnO<sub>2</sub> NNs and δ-phase MnO<sub>2</sub> NSAs on CC are shown in [Fig. 2a](#). Two XRD peaks at 26° and 43° were observed on the bare CC surface, consistent with the literature [36]. Neither α-MnO<sub>2</sub> nor δ-MnO<sub>2</sub> exhibited well-defined peaks, indicating that they had amorphous structures. Nevertheless, the α-MnO<sub>2</sub> peaks and the δ-MnO<sub>2</sub> peaks agree with Powder Diffraction File card #44–0141 and Joint Committee on Powder Diffraction Standards card #18–802, respectively [37]. The differences between the phase structures of the synthesized forms of MnO<sub>2</sub> were confirmed by their peak

patterns. Also, α-MnO<sub>2</sub>-CC had a predominant (211) plane of Mn, as shown by its narrow, high-intensity peaks. In contrast, δ-MnO<sub>2</sub>-CC demonstrated broader and weaker (411) and (521) peaks at 38° and 50°, respectively. The XRD patterns of residual MnO<sub>2</sub> powders collected from the autoclave following the synthesis of α- and δ-MnO<sub>2</sub> were also analyzed to determine their crystal phases ([Supplementary Material Fig. S3](#)). Their XRD patterns were consistent with those of MnO<sub>2</sub> deposited on the CC surface.

Raman spectroscopy of MnO<sub>2</sub>-CC electrodes was performed using a Raman spectrometer equipped with a 20 mW, 532 nm laser. Raman spectroscopy ([Fig. 2b](#)) confirmed the clear difference between two versions of MnO<sub>2</sub>. δ-MnO<sub>2</sub> had been formed, as the Raman spectrum contained representative peaks at 160.4, 504.6, 581.8, and 654.6 cm<sup>−1</sup>, which are consistent with the literature [38]. The X-ray photoelectron spectra of α-MnO<sub>2</sub>-CC and δ-MnO<sub>2</sub>-CC are shown in [Fig. 2c–f](#). Two peaks, centered on 642.4 and 654 eV representing the spin-orbit doublet states of Mn 2p<sub>3/2</sub> and Mn 2p<sub>1/2</sub>, respectively, were present in the high-resolution spectrum of Mn



**Fig. 2.** a–b, X-ray diffraction patterns (a) and Raman spectra (b) of uncoated carbon cloth (CC),  $\alpha$ -MnO<sub>2</sub>-CC, and  $\delta$ -MnO<sub>2</sub>-CC. c–f, High-resolution X-ray photoelectron spectra of Mn 2p (c, e) and O 1s (d, f), together with the corresponding spectral decomposition of  $\alpha$ -MnO<sub>2</sub>-CC (c–d) and  $\delta$ -MnO<sub>2</sub>-CC (e–f).

2p. After spectral decomposition, Mn existed in three valences: +2, +3, and +4. Both  $\alpha$ - and  $\delta$ -MnO<sub>2</sub> showed strong Mn<sup>3+</sup> peaks at 642 and 654 eV for Mn 2p<sub>3/2</sub> and Mn 2p<sub>1/2</sub>, respectively. Fig. 2d and f show the O 1s core-level spectra of  $\alpha$ -MnO<sub>2</sub>-CC and  $\delta$ -MnO<sub>2</sub>-CC, respectively. In addition to the dominant Mn–O–Mn moieties in MnO<sub>2</sub>, represented by the peak at approximately 528.8 eV, there were large proportions of Mn–O–H moieties (i.e., Mn ions bonded with hydroxyl groups) and H–O–H moieties (i.e., absorbed water), represented by peaks at approximately 531.9 and 533.4 eV, respectively. The relative surface proportions of Mn<sup>3+</sup> and Mn<sup>4+</sup> (Mn<sup>3+/4+</sup>) are often considered the main factors in the performance of Mn catalysts. For instance, a large proportion of Mn<sup>3+</sup> on the surfaces of bifunctional catalysts has been found to facilitate the oxygen evolution reaction (OER) [39–42]. Conversely, a small

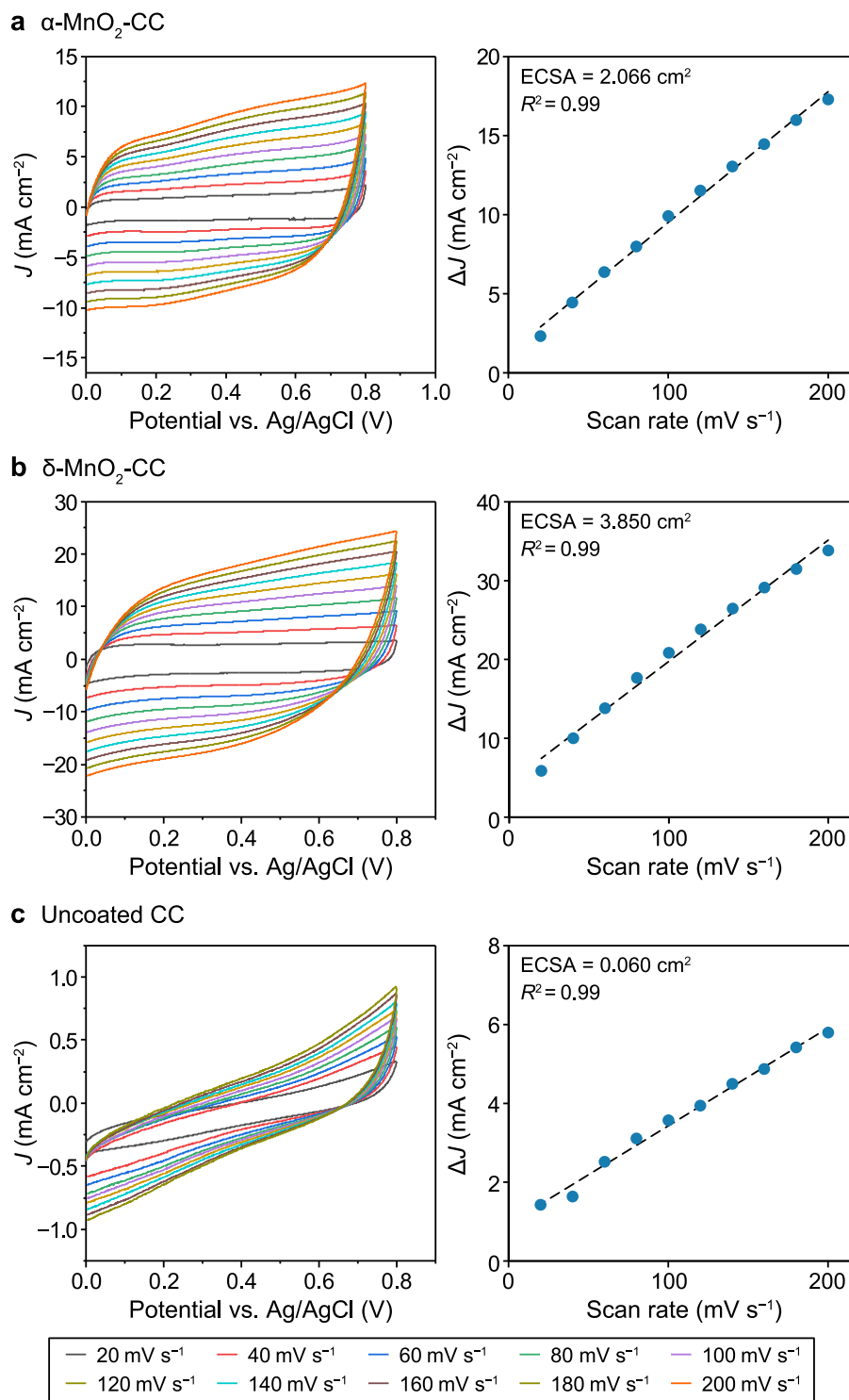
proportion of Mn<sup>4+</sup> on the surfaces of bifunctional catalysts has been suggested to enhance their adsorption of organic compounds [43]. The Mn<sup>3+/4+</sup> values for  $\alpha$ -MnO<sub>2</sub>-CC and  $\delta$ -MnO<sub>2</sub>-CC were 2.7079 and 1.6000, respectively, which matched well with the catalytic findings.

### 3.2. Determination of the ECSAs of the MnO<sub>2</sub> electrodes

The electrochemical characterization of the MnO<sub>2</sub> electrodes was conducted in a three-electrode batch system. Their ECSAs were determined via CV at 0–0.8 V in the non-Faradaic current region at a range of scan rates, i.e., 20–200 mV s<sup>-1</sup>, with data collected every 20 mV. Fig. 3 shows the cyclic voltammograms of MnO<sub>2</sub>-modified and uncoated electrodes measured in 1 M Na<sub>2</sub>SO<sub>4</sub> solution. C<sub>dl</sub> was

determined by plotting the scan rate versus the change in current density ( $\Delta J = J_{\text{anodic}} - J_{\text{cathodic}}$ ), as the resulting linear slopes equal  $2C_{\text{dl}}$ . The ECSAs of the  $\alpha$ -MnO<sub>2</sub> electrode (Fig. 3a, inset) and  $\delta$ -MnO<sub>2</sub> electrode (Fig. 3b, inset) were 33 and 62 times higher, respectively,

than that of the CC electrode (Fig. 3c, inset). Moreover, the surface area of the  $\delta$ -MnO<sub>2</sub> electrode was twice that of the  $\alpha$ -MnO<sub>2</sub> electrode, which accounts for the effectiveness of the  $\delta$ -MnO<sub>2</sub> electrode in degrading small aromatics, as described in Section 3.5.



**Fig. 3.** Cyclic voltammetry curves of  $\alpha$ -MnO<sub>2</sub>-CC (a),  $\delta$ -MnO<sub>2</sub>-CC (b), and uncoated CC electrodes (c) in the non-Faradaic region of 0–0.8 V vs. Ag/AgCl, recorded at scan rates ranging from 20 to 200 mV s<sup>-1</sup>. The subpanels on the right side of panels a–c display the corresponding electrochemical double-layer capacitance ( $C_{\text{dl}}$ ) plots derived from the cyclic voltammetry curves. CC: carbon cloth.

### 3.3. Electrocatalytic TCS-degradation performance of $\alpha$ - and $\delta$ -MnO<sub>2</sub> electrodes

The electrocatalytic performance of the as-prepared  $\alpha$ - and  $\delta$ -MnO<sub>2</sub> in the oxidative degradation of TCS was evaluated via chronopotentiometric electrolysis in an aqueous environment containing 5 mM NaCl and 50 mM Na<sub>2</sub>SO<sub>4</sub> at room temperature and atmospheric pressure. NaCl was added to provide conductivity and to mimic the typical chloride-ion concentration of wastewater [34]. The oxidative degradation of TCS by  $\alpha$ - and  $\delta$ -MnO<sub>2</sub> decreased as the current increased (Fig. 4a), indicating that the degradation efficiency was compromised by the increasing competitiveness of the OER driven by anodic water splitting. At 20 mA cm<sup>-2</sup>, the proportion of TCS degraded by  $\alpha$ - and  $\delta$ -MnO<sub>2</sub> reached 97.27 ± 2.7 % and 99.44 ± 1.4 %, respectively, but at 40 mA cm<sup>-2</sup>, this decreased to 85.18 ± 5.7 % and 88.92 ± 5.5 %, respectively. This result indicates that, compared with  $\delta$ -MnO<sub>2</sub>,  $\alpha$ -MnO<sub>2</sub> was more affected by the OER. At 80 mA cm<sup>-2</sup>, the proportion of TCS degraded by  $\alpha$ -MnO<sub>2</sub> was only 63.27 ± 7.3 %, whereas that degraded by  $\delta$ -MnO<sub>2</sub> remained reasonably high: 82.58 ± 3.1 %.

The greater tendency of  $\alpha$ -MnO<sub>2</sub> than  $\delta$ -MnO<sub>2</sub> to exhibit the OER at high currents is consistent with a previous electrochemical observation that the  $\alpha$  phase has a lower overpotential for OER than the  $\delta$  phase; this implies that  $\alpha$ -MnO<sub>2</sub> will trigger OER before  $\delta$ -MnO<sub>2</sub> [44]. The OER activity of  $\alpha$ -MnO<sub>2</sub> is superior to that of  $\delta$ -MnO<sub>2</sub> because the former has a greater Mn<sup>3+/4+</sup> ratio on its surface, as demonstrated by XPS (Fig. 2c). Mn<sup>3+</sup> favors the occurrence of the OER because the single electron occupying its  $\sigma^*$ -orbital (eg) is transferred to its O–O  $\pi^*$ -orbital when Mn<sup>3+</sup> is oxidized to Mn<sup>4+</sup> [45]. The model supports our XPS-based analysis of the surface, which revealed that Mn<sup>3+/4+</sup> ratios for the  $\alpha$ - and  $\delta$ -phases were 2.7079 and 1.6000, respectively (Fig. 2c–e). Thus, compared with the  $\delta$ -MnO<sub>2</sub> catalyst, the  $\alpha$ -MnO<sub>2</sub> catalyst had a larger proportion of Mn<sup>3+</sup>, which shifted its activity from TCS degradation toward the OER.

In addition to  $\delta$ -MnO<sub>2</sub> having a wider electrochemical window for the OER than  $\alpha$ -MnO<sub>2</sub>,  $\delta$ -MnO<sub>2</sub> adsorbed more TCS (Supplementary Material Fig. S4). Specifically, in an open-circuit TCS adsorption control experiment,  $\delta$ -MnO<sub>2</sub> adsorbed 13.7 % of the TCS to which it was exposed, whereas  $\alpha$ -MnO<sub>2</sub> adsorbed only 8.3 %. This is attributable to the ECSA of  $\delta$ -MnO<sub>2</sub> being greater than that of  $\alpha$ -MnO<sub>2</sub> (3.847 cm<sup>2</sup> versus 2.066 cm<sup>2</sup>; Fig. 3a and b, panels on the right). Moreover, the electrocatalytic performance of as-synthesized MnO<sub>2</sub> electrodes for some mono- and di-aromatic structures were tested under the same conditions (Fig. 4b).

During the bulk electrolysis of TCS, trace amounts of chlorinated mono-aromatic intermediates were detected. Therefore, to elucidate the degradation pathways and obtain comprehensive mechanistic insights, TCS and the chlorinated mono-aromatics 4-CP and 2,4,6-TCP were subjected to time-resolved electrolysis on  $\alpha$ -MnO<sub>2</sub> and  $\delta$ -MnO<sub>2</sub> to investigate the relative rates of degradation by each catalyst. Two additional EDCs structurally similar to TCS were also examined under these conditions, namely BPA, due to its di-aromatic structure, and 2-BrPh, due to its halogenated structure (Fig. 4c and d). All electrolysis was performed at 20 mA cm<sup>-2</sup> to minimize the possibility of competition with the OER. A pseudo-first-order kinetic analysis was conducted to calculate the degradation rate constants ( $k$ , min<sup>-1</sup>; Fig. 4c and d, insets).

Under the initial conditions ( $j = 20$  mA cm<sup>-2</sup>),  $\alpha$ -MnO<sub>2</sub> performed slightly better than  $\delta$ -MnO<sub>2</sub>, as all of the  $k$  values of the former were greater than those of the latter. All degradation patterns obeyed the pseudo-first-order kinetic model, as the data exhibited a good fit, i.e., the average coefficients of determination were 0.88 and 0.93 for the  $\alpha$ -MnO<sub>2</sub> and  $\delta$ -MnO<sub>2</sub> datasets, respectively, in linear regression using  $\ln(C/C_0) = -kt$ . The  $k$  of TCS

degradation was 22 min<sup>-1</sup> on  $\alpha$ -MnO<sub>2</sub> and 9.6 min<sup>-1</sup> on  $\delta$ -MnO<sub>2</sub>.  $\alpha$ -MnO<sub>2</sub> also degraded BPA at a greater rate than did  $\delta$ -MnO<sub>2</sub>, indicating the universal applicability of  $\alpha$ -MnO<sub>2</sub>. Halogenated aromatic species that appeared during TCS degradation were also examined. 2,4,6-TCP was degraded by both  $\alpha$ -MnO<sub>2</sub> ( $k = 16.4$  min<sup>-1</sup>) and  $\delta$ -MnO<sub>2</sub> ( $k = 6.6$  min<sup>-1</sup>), with its degradation by the former being slightly more efficient than its degradation by the latter. However, the rate constant for 4-CP degradation by  $\alpha$ -MnO<sub>2</sub> ( $k = 40.5$  min<sup>-1</sup>) was almost thrice that for 4-CP degradation by  $\delta$ -MnO<sub>2</sub> ( $k = 14.9$  min<sup>-1</sup>). We performed a control comparison using 2-BrPh and observed a similar rate enhancement for  $\alpha$ -MnO<sub>2</sub> over  $\delta$ -MnO<sub>2</sub> ( $k = 32.5$  min<sup>-1</sup> versus 14.9 min<sup>-1</sup>), which indicated that the rapid degradation observed on  $\alpha$ -MnO<sub>2</sub> was not due to the chlorine substituent of 4-CP but rather its monomeric ring structure and mono-halogenation. Control experiments (Fig. 4c and d) on open-circuit adsorption showed that  $\alpha$ -MnO<sub>2</sub> and  $\delta$ -MnO<sub>2</sub> had similar adsorption capacities (Supplementary Material Fig. S4) for 4-CP and 2,4,6-TCP, which indicated that the superior catalytic activity of  $\alpha$ -MnO<sub>2</sub> was attributable to its intrinsic properties, i.e., the fact that it had a greater Mn<sup>3+/4+</sup> ratio (2.7079) than  $\delta$ -MnO<sub>2</sub> (1.6000).

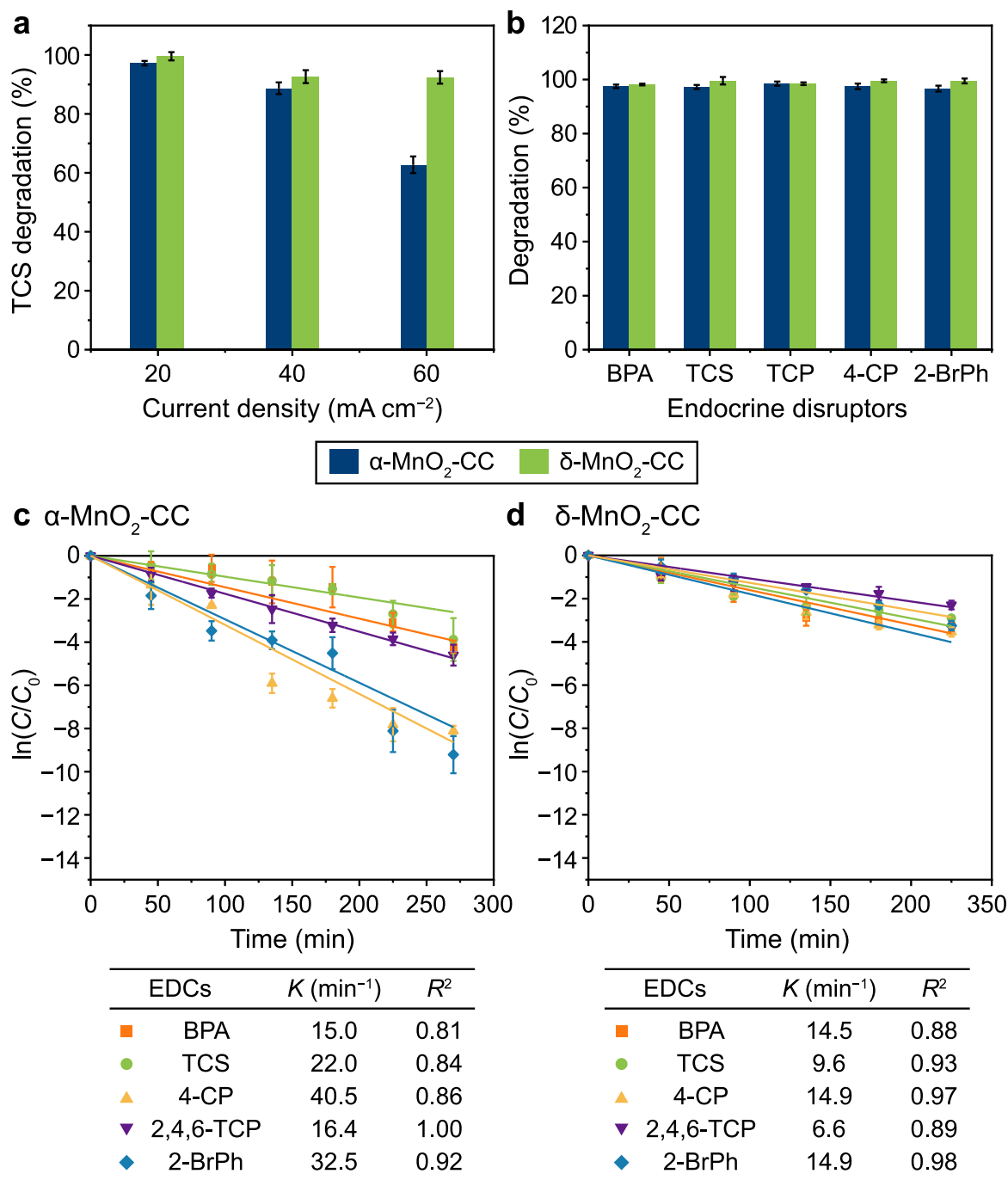
Overall, the degradation trend for  $\alpha$ -MnO<sub>2</sub> was approximately 4-CP, 2-BrPh > 2,4,6-TCP, TCS > BPA. The exceedingly high degradation of 2-BrPh and 4-CP by  $\alpha$ -MnO<sub>2</sub> indicates that it excels at degrading halogenated mono-aromatics and does not appear to be restricted by the location of the halogen. An open-circuit adsorption control experiment was also performed using 4-CP and 2-BrPh and both  $\alpha$ -MnO<sub>2</sub> and  $\delta$ -MnO<sub>2</sub> catalysts. The results showed that  $\delta$ -MnO<sub>2</sub> was slightly more effective than  $\alpha$ -MnO<sub>2</sub> at adsorbing organic compounds, which excluded the possibility that favorable adsorption accounted for the high reactivity of  $\alpha$ -MnO<sub>2</sub>. Thus, we attribute the better performance of  $\alpha$ -MnO<sub>2</sub> to the fact that it has a greater Mn<sup>3+/4+</sup> ratio than  $\delta$ -MnO<sub>2</sub>, which enhances the oxidative catalytic performance of  $\alpha$ -MnO<sub>2</sub>. The degradation trend for  $\delta$ -MnO<sub>2</sub> was approximately 4-CP, 2-BrPh > BPA, TCS > 2,4,6-TCP. This suggests that the accessibility of C–H sites on the aromatic ring of an EDC plays a key role in its oxidative degradation by  $\delta$ -MnO<sub>2</sub>. Specifically, 4-CP and 2BrPh have four accessible aryl C–H sites at which oxidation can occur; BPA and TCS also have four such sites per ring, but these sites may be sterically hindered by the molecules' bulky biphenyl structure; and 2,4,6-TCP only has two accessible aryl C–H sites.

XRD analysis was conducted on all the MnO<sub>2</sub> catalysts after electrolysis, and the results showed that they exhibited good retention of their respective phases (Supplementary Material Fig. S5). Thus, the  $\alpha$ -MnO<sub>2</sub> peaks at 28.74° and 37.58° and the key  $\delta$ -MnO<sub>2</sub> peak at 38.3° retained suitable shapes after exposure to all the EDC substrates.

### 3.4. Generation mechanism and role of ROS

EPR and UV–vis spectroscopy were used to identify the ROS. The following text describes a possible mechanism based on the spectroscopic results. First, the anodic oxidation of two chloride ions (Cl<sup>-</sup>) affords molecular chlorine (equation (1)) [46], which then combines with H<sub>2</sub>O to yield hypochlorous acid (HClO; equation (2)) [47]. The presence of HClO was confirmed by UV–vis spectroscopy, as it showed an increase in the size of the peak at ~290 nm, which matches the reported wavelength of HClO in UV–vis spectra (Fig. 5g and h). Next, HClO reacts with either a chlorine radical (Cl<sup>•</sup>; equation (3)) [48] or a hydroxyl radical (HO<sup>•</sup>; equation (4)) [49] to yield chlorine monoxide (ClO<sup>•</sup>), which subsequently reacts with a hydroxide ion to afford a superoxide anion (O<sub>2</sub><sup>•-</sup>; equation (5)) [50]. Studies have suggested that ClO<sup>•</sup> can react with HO<sup>•</sup> to yield a chlorite ion [51], which is commonly used as an oxidizing agent. It can accept electrons from other substances, leading to the





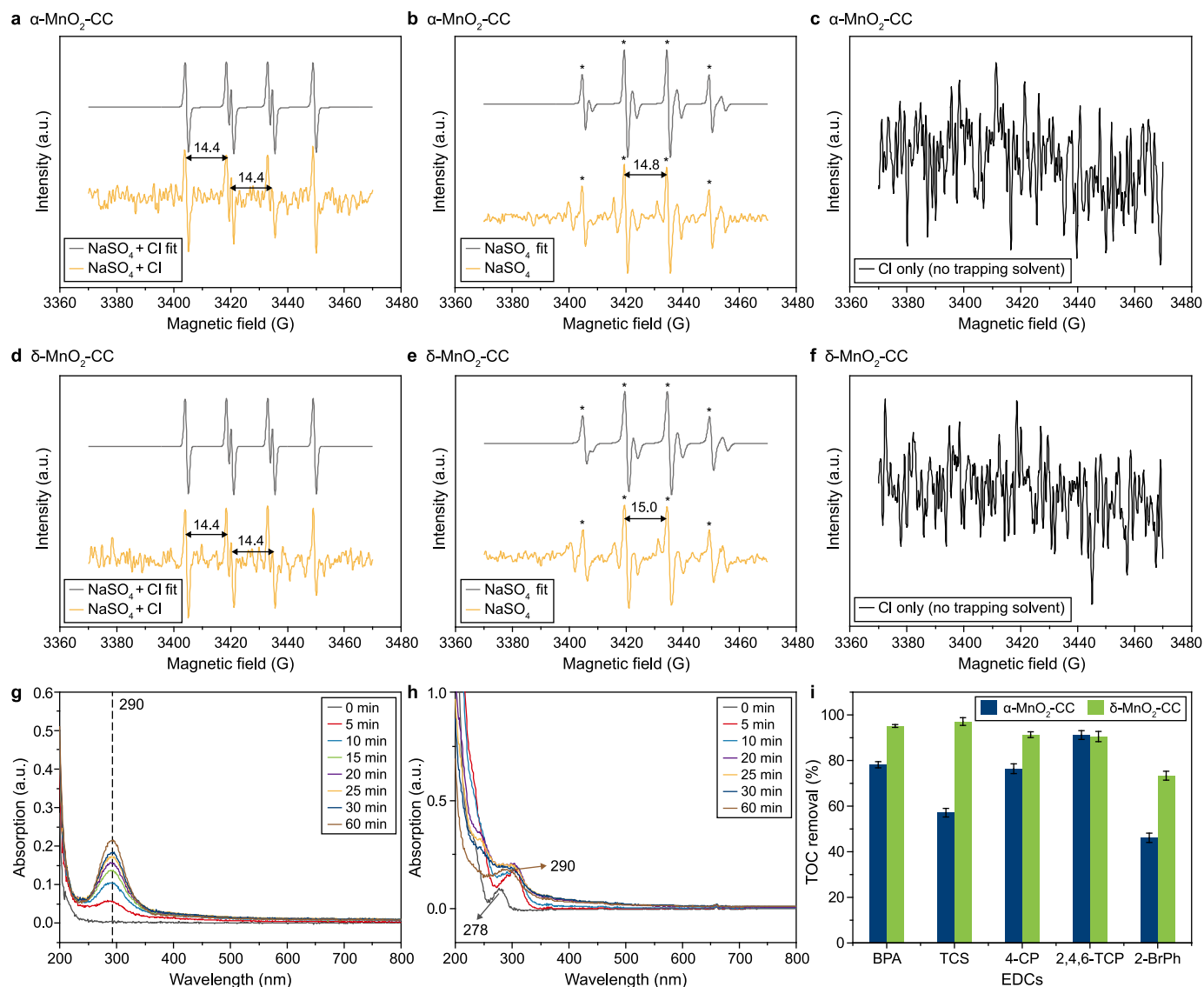
**Fig. 4.** **a**, Triclosan (TCS) degradation efficiencies of  $\alpha\text{-MnO}_2\text{-CC}$  and  $\delta\text{-MnO}_2\text{-CC}$  at different current densities. **b**, Electrocatalytic oxidative activity of  $\alpha\text{-}$  and  $\delta\text{-MnO}_2\text{-CC}$  for endocrine disruptors (EDCs, 10 ppm) at room temperature ( $25^\circ\text{C}$ ) at 20 mA (4.5 h). **c–d**, Plots of  $\ln(C/C_0)$  versus time indicating the first-order behavior of  $\alpha\text{-MnO}_2\text{-CC}$  (**c**) and  $\delta\text{-MnO}_2\text{-CC}$  (**d**) at a current density of 20 mA (where  $C = C_t$  and  $C_0 = C_i$ ). The tables below show the corresponding first-order parameters for TCS and related EDCs. The average of triplicate data and standard deviation is given in Supplementary Materials [Tables S3–S6](#). CC: carbon cloth; EDC: endocrine disruptors; BPA: bisphenol A; TCP: trichlorophenol; 4-CP: 4-chlorophenol; 2-BrPh: 2-bromophenol.

formation of chloride ions and chlorine dioxide.

To confirm the formation of chlorite ions in our reaction medium, we examined the reaction using spin-trap-free EPR spectroscopy, and the results showed that the reaction did not follow such a pathway (Fig. 5c–f). Specifically, the EPR spectrum of the reaction electrolyte (which contained both  $\text{Na}_2\text{SO}_4$  and  $\text{NaCl}$ ) contained no signals for  $\text{HO}^\bullet$  because this species had been scavenged by  $\text{Cl}^-$  <sup>46</sup> and the increasing amount of  $\text{HClO}$  (equation (4)). We

separately examined the reaction using  $\text{Cl}^-$ -free EPR spectroscopy, which confirmed that  $\text{HO}^\bullet$  could be formed by both  $\text{MnO}_2$  electrodes (Fig. 5b–e). The presence of  $\text{O}_2^{\bullet-}$  was also confirmed by EPR spectroscopy, which showed that the splitting pattern corresponded to 5,5-dimethyl-1-pyrroline *N*-oxide–superoxide with a hyperfine coupling constant of 14.4, which is consistent with the published value of 14.53 [52,53] (Fig. 5a). Finally, the reaction of  $\text{HClO}$  with  $\text{O}_2^{\bullet-}$  forms additional  $\text{HO}^\bullet$  and  $\text{Cl}^-$ , which are





**Fig. 5.** a–f, Electron paramagnetic resonance analysis of electrochemical oxidative mineralization of triclosan (TCS) by MnO<sub>2</sub> electrocatalysts [ $\alpha$ -MnO<sub>2</sub>-CC (a–c) and  $\delta$ -MnO<sub>2</sub>-CC (d–f)] in different electrolytes: Na<sub>2</sub>SO<sub>4</sub> + Cl (a, d); Na<sub>2</sub>SO<sub>4</sub> (b, e); and Cl environment only (c, f). g–h, Formation of hypochlorous acid in the absence (g) and presence (h) of TCS. i, Total organic carbon removal by the electrochemical oxidative mineralization of TCS and structurally similar EDCs by the MnO<sub>2</sub> catalysts. The average triplicate data and the standard deviation are given in [Supplementary Material Table S7](#). CC: carbon cloth; EDCs: endocrine disruptors; BPA: bisphenol A; TCP: trichlorophenol; 4-CP: 4-chlorophenol; 2-BrPh: 2-bromophenol; \*: corresponds to hydroxy radical signal.

subsequently converted to various reactive chlorine species (RCS), such as ClO<sup>•</sup> and Cl<sup>•</sup> (equation (6)) [46–49,51,54,54].



Based on the spectroscopic analysis, HClO appears to be the ROS

responsible for the degradation of TCS. In the presence of TCS, a slight redshift of the  $\lambda_{\text{max}}$  from ~290 to 304 nm indicated the consumption of HClO by TCS. After the complete mineralization/disappearance of TCS, the  $\lambda_{\text{max}}$  returned to ~290 nm, indicating the reformation of HClO. In the absence of Cl<sup>−</sup>, however, HO<sup>•</sup> was the dominant ROS (Fig. 5b–e), as HClO could not form, and the degradation of TCS was significantly slower ([Supplementary Material Fig. S6](#)). This demonstrated that HO<sup>•</sup> could not degrade TCS efficiently.

TOC was measured to quantify the extent of organic content mineralization resulting from oxidative degradation (Fig. 5i). Before measuring TOC content after the degradation of EDCs, TOC content after electrolysis of both MnO<sub>2</sub> electrodes without compounds were measured as a control experiment ([Supplementary Material Table S2](#)). The  $\delta$ -MnO<sub>2</sub> degraded TCS and its structurally similar EDCs very efficiently. Over 90 % of TOC was removed after electrochemical treatment of TCS, BPA, and halogenated phenols. In

comparison,  $\alpha$ -MnO<sub>2</sub> was less efficient at complete mineralization because as the TOC decreased, the electrochemical oxidative degradation of organics occurred via the OER rather than by mineralization. Meanwhile, although the catalytic performance of  $\delta$ -MnO<sub>2</sub> was less active, it was less prone to exhibit the OER and thus achieved better mineralization. Only 2,4,6-TCP underwent more than 90 % degradation on both  $\alpha$ - and  $\delta$ -MnO<sub>2</sub> catalysts, which indicates that such highly chlorinated species can be degraded effectively by these catalysts, regardless of the nature of their surfaces.

### 3.5. TCS mineralization pathway and toxicity evaluation

Fig. 6 and S8 (Supplementary Material) depict a possible TCS-degradation pathway based on the literature findings and products detectable by mass spectrometry [55,56]. The observed fragments were categorized based on their structural complexity and appearance during electrolysis.

Based on the detectable fragments, we propose that TCS initially undergoes chlorination and dichlorination to form T1 (*m/z* 253.99), T2 (*m/z* 324), and T3 (*m/z* 358). Subsequently, cleavage of the aryl ether C–O bond produces various mono-aromatic products: T4 (2,4,6-TCP, *m/z* 195), T5 (*m/z* 161.95), T6 (*m/z* 143.95), T7 (*m/z* 161.95), T8 (*m/z* 149.95), T9 (*m/z* 94), T10 (*m/z* 108), and T11 (4-CP, *m/z* 128).

Further, we conducted two control experiments to confirm the ring-opening pathway of transformation products. First, a control experiment was conducted to electrolyze T4 (2,4,6-TCP) separately to determine whether multi-chlorinated aromatic monomers are degradable under our conditions. The reaction produced T4a (*m/z* 179), T6, and T7. As T6 and T7 were also formed during TCS electrolysis, this confirmed that T4 underwent degradation during the TCS degradation trials. We propose that the electrode oxidizes T4–T11 to yield T11a (*m/z* 108), T11b (*m/z* 158), and T11c (*m/z* 191). Although T11a–c were not observed directly during TCS electrolysis, we posit that they are formed during TCS degradation because the oxidative treatment of mono-aromatics is known to yield quinones and ring-opened carboxylic acids [33]. We thus conducted a second control experiment in which we electrolyzed T11 (4-CP) under the same conditions and obtained detectable amounts of T11a–c. The absence of T11a–c during TCS degradation is attributable to the fact that only low concentrations of T11 were formed.

We hypothesize that T11a–c is oxidized and mineralized to carbon dioxide and H<sub>2</sub>O. The cleavage of dimers occurred at the ether C–O bonds, but this cleavage occurred non-selectively on either side. Chlorination also occurred, confirming that RCS were formed during electrolysis (equation (6)). Nevertheless, the mineralization of aromatic compounds continued, as confirmed by the TOC analysis. After electrochemical treatment, the amount of organic carbon substantially decreased. Small amounts of a hydroxylated product (i.e., T6) were observed during the TCS degradation and the control T4 trials, which confirmed that HO• had been formed (equation (6)).

The acute and chronic toxicity of TCS and its individual transformation product on fish and daphnids is predicted by the Ecological Structure Activity Relationships (ECOSAR) software (Supplementary Material Table S10). The tabulated dataset provides a comprehensive overview of TCS and its transformation products, detailing their chemical structures through simplified molecular input line entry system (SMILES), molecular weights (*m/z*), aquatic toxicity parameters (LC50 and chronic values, ChV), test organisms (fish or daphnids), exposure durations (96 h for fish acute studies, 48 h for daphnid acute studies, and preferable durations for chronic endpoints), and flags indicating solubility- or log *K*<sub>ow</sub>-based considerations. Triclosan displays relatively high toxicity to both fish and daphnids, evidenced by acute LC50 values of 0.965 and 0.701 mg L<sup>−1</sup>, respectively, and corresponding low chronic thresholds, indicating a heightened hazard potential. By contrast, most identified intermediates (T4a, T4b, T4c, T4d, T4e, T11a, T11b, and T11c) exhibit substantially lower toxicity, with the notable exceptions of T2 and T3. Both T2 and T3 show a “no effects at saturation” flag due to their high log *K*<sub>ow</sub>, which indicates that under standard aquatic conditions, their solubility may be too low to reach the concentrations required for acute or chronic effects in water. Consequently, these findings suggest that electrochemical oxidation can mitigate TCS-related ecological risks, as most resultant byproducts pose less harm than the parent compound.

### 3.6. Assessing the performance and scalability of MnO<sub>2</sub> electrodes in complex environments

The performances of the  $\alpha$ - and  $\delta$ -MnO<sub>2</sub> electrodes were examined in a scaled-up (300 mL) electrolysis of TCS in different environments. The performances of the prepared MnO<sub>2</sub> electrodes

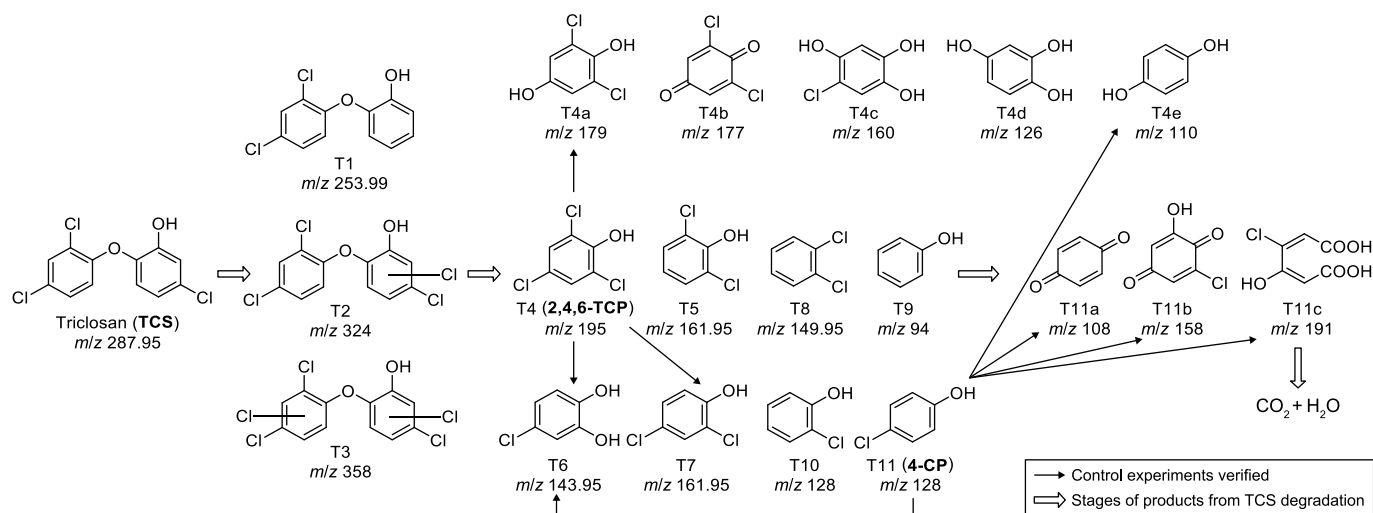


Fig. 6. Possible pathway of electrochemical oxidative mineralization of triclosan in an aqueous environment based on products that can be observed by mass spectrometry.

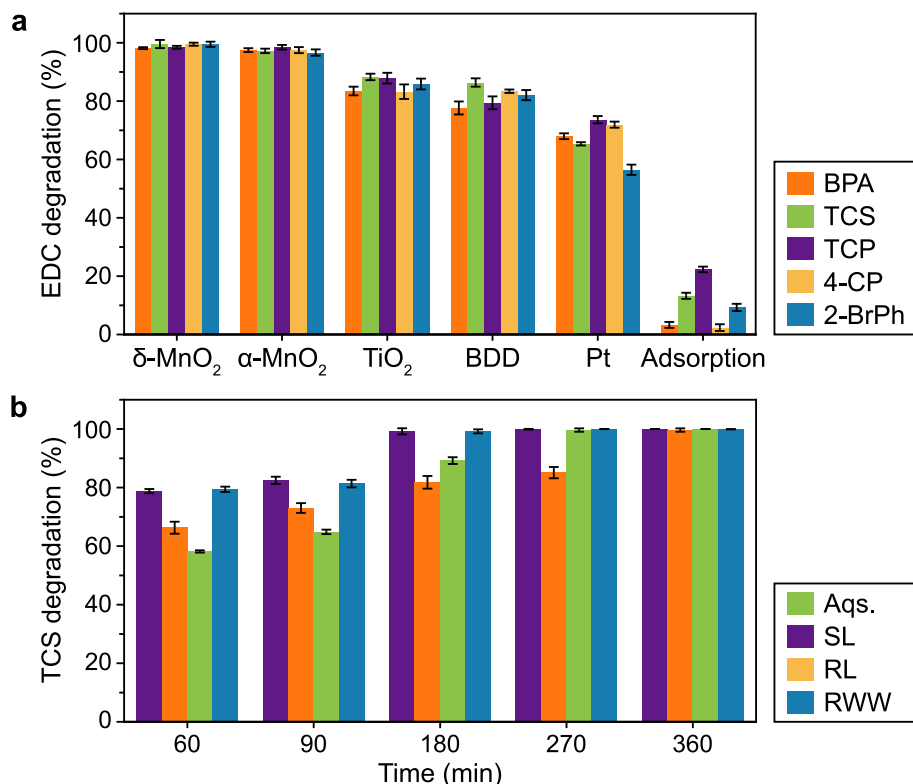
were compared with those of other common anodes, such as platinum mesh and BDD electrodes. Titania-CC was also prepared via a hydrothermal method, and its catalytic efficiency was compared with the other electrodes.  $\alpha$ - and  $\delta$ -phase  $\text{MnO}_2$  exhibited the greatest catalytic degradation of the five EDCs (Fig. 7a). An adsorption study in an open-circuit setting was also conducted using the  $\text{MnO}_2$  electrodes, and the results confirmed that the removal of EDCs was achieved electrochemically (Supplementary Material Fig. S4). Thus, coupled with the results of the TOC analysis, we concluded that EDC degradation was achieved electrocatalytically rather than by surface physisorption. In this study, we also tested the performance of  $\delta$ - $\text{MnO}_2$  electrodes for degradation of TCS in four different environments (aqueous [Aqs], real wastewater [RWW], synthetic [SL], and real landfill leachates [RL]). The performance of the  $\delta$ - $\text{MnO}_2$  electrodes was examined in a scaled-up (300 mL) electrolysis of TCS in all of these environments. The degradation of organic pollutants present in landfill leachates and industrial wastewater is one of the critical issues in environmental pollution control, as they include many micropollutants and high concentrations of inorganic compounds, thereby hindering the overall degradation process. In this study, we successfully degraded TCS in all possible complex environments (RWW, SL, and RL) and maintained greater than 75 % degradation in just 3 h (Fig. 7b). After the degradation reaction was completed in a complex environment (RL and RWW), samples were collected and extracted with DCM and then subjected to GCMS analysis to quantify the leftover TCS in the solution (Supplementary Material Fig. S8).

Fig. 7b shows that TCS was degraded significantly in both simple and complex environments. Compared with the standard aqueous environment (simple environment), the degradation efficiency was

slightly lower in the SL and RL environments, but 98 % TCS degradation was nevertheless achieved in 6 h, which indicated that the presence of many wastewater species did not influence TCS degradation (Fig. 7b). Moreover, TCS degradation was achieved in real wastewater that was comparable to the TCS degradation in the standard aqueous environment. These results confirm that noble metal catalysts, such as gold or Ag, can be replaced with earth-abundant material-based catalysts (Mn), as the latter exhibit remarkable catalytic performances that approach practical requirements. This opens an avenue for future wastewater remediation and represents significant progress toward closing the anthropogenic carbon cycle to realize global sustainability.

### 3.7. Environmental implications

Removal of EDCs from wastewater through advanced oxidation processes is currently challenged by the use of toxic oxidants and chemicals to initiate the oxidation reaction, the large amounts of noxious residuals that result from incomplete mineralization, and excessive energy input. Current endeavors to achieve the electrocatalytic mineralization of various EDCs via two highly active  $\text{MnO}_2$  phases,  $\alpha$ - $\text{MnO}_2$  and  $\delta$ - $\text{MnO}_2$ , were fabricated on a cost-effective CC surface. The resulting  $\text{MnO}_2$  catalysts were mostly utilized to accelerate mineralization in real wastewater streams (chlorinated wastewater, real wastewater, and synthetic and real landfill leachates) without the addition of oxidants for ROS generation. Moreover, the performance of the  $\alpha$ - and  $\delta$ - $\text{MnO}_2$  electrodes was also examined in a scaled-up (300 mL) electrolysis of TCS in a synthetic leachate (SL) environment; they maintained 98 % TCS degradation, which indicated that the presence of over 10 other common wastewater species did not influence TCS mineralization.



**Fig. 7.** Comparison of electrocatalytic performances of common electrode and  $\text{MnO}_2$ -based electrodes for degradation of several endocrine disruptors (EDCs, **a**) and degradation of triclosan (TCS) by  $\delta$ - $\text{MnO}_2$  electrodes in various environments on a large scale (300 mL) at 80 mA (**b**). BDD: boron-doped diamond; BPA: bisphenol A; TCP: trichlorophenol; 4-CP: 4-chlorophenol; 2-BrPh: 2-bromophenol. Aqs.: aqueous; SL: synthetic leachates; RL: real landfill leachates; RWW: real wastewater on a large scale (300 mL). The average of triplicate data and standard deviation is given in Supplementary Materials Tables S8–S9.

Significantly, the  $\text{MnO}_2$ -CC electrocatalyst performs robustly in real environmental matrices without additional oxidants or chemicals, produces no residual byproducts, and exhibits oxidative performance comparable to that of precious metal catalysts. These favorable features enabled  $\text{MnO}_2$ -electrocatalyst (Earth-abundant) to be a promising, sustainable, efficient, and scalable solution toward developing catalyst-driven technologies for completely mineralizing multiple organic pollutants.

#### 4. Conclusion

In summary, two highly active  $\text{MnO}_2$  phases,  $\alpha$ - $\text{MnO}_2$  and  $\delta$ - $\text{MnO}_2$ , were fabricated on a cost-effective CC surface, and the resulting  $\text{MnO}_2$  catalysts were comprehensively characterized. The  $\text{MnO}_2$  catalysts achieved nearly complete EDC degradation in a pH-neutral chlorinated aqueous environment similar to typical wastewater.  $\alpha$ - $\text{MnO}_2$ -CC was found to possess unique nanostructures that facilitate the degradation of small aromatic compounds and to contain more  $\text{Mn}^{4+}$  than  $\text{Mn}^{3+}$  on its surface, endowing it with enhanced catalytic performance before it reaches the onset potential of the OER. Compared with  $\alpha$ - $\text{MnO}_2$ ,  $\delta$ - $\text{MnO}_2$  delivered a more stable electrocatalytic performance overall and performed better at high-current flows. These catalytic differences allow the treatment of targeted pollutants, such as mono-aromatic or poly-aromatic organic pollutants. ROS identification by EPR and UV-vis spectroscopy confirmed the presence of various highly reactive intermediates, such as  $\text{HClO}$  and  $\text{O}_2^{\cdot-}$ . TOC analysis confirmed that aromatic pollutants, such as TCS, were effectively mineralized by the catalysts. Finally,  $\alpha$ - and  $\delta$ - $\text{MnO}_2$  both exhibited good performance in a scaled-up setting and were able to degrade TCS efficiently in an SL environment. By fabricating highly active  $\alpha$ - $\text{MnO}_2$  and  $\delta$ - $\text{MnO}_2$  phases on cost-effective CC surfaces, we developed robust electrocatalysts that efficiently achieve near-complete mineralization of endocrine disruptors in real wastewater streams without additional oxidants, thereby overcoming the challenges of toxic initiators, incomplete mineralization, and high energy inputs associated with traditional advanced oxidation processes. Thus, we have developed an efficient Earth-abundant metal catalyst that exhibits oxidative performance comparable to that of precious metal catalysts.

#### CRediT authorship contribution statement

**Asma Batool:** Writing – review & editing, Writing – original draft, Visualization, Validation, Methodology, Investigation, Formal analysis, Data curation, Conceptualization. **Shan Shao:** Visualization. **Kartick Chandra Majhi:** Visualization. **Azeem Mushtaq:** Visualization. **Yi Jiang:** Writing – review & editing, Visualization. **Wingkei Ho:** Writing – review & editing, Visualization. **Yiu Fai Tsang:** Writing – review & editing, Visualization. **Yuhe He:** Writing – review & editing, Visualization. **Kenneth Mei Yee Leung:** Writing – review & editing, Visualization. **Jason Chun-Ho Lam:** Writing – review & editing, Visualization, Supervision, Investigation, Conceptualization.

#### Data availability

No data was used for the research described in the article.

#### Declaration of competing 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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#### Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.es.2025.100559>.

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