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## Manganese oxide-mediated halogenation of carbazole under marine-related conditions

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Polyhalogenated carbazoles (PHCZs) are emerging halogenated organic compounds detected in various environmental matrices, including marine environments. However, the origins of PHCZs are not yet fully understood. This study investigates the manganese oxide-mediated transformation of carbazole into PHCZs under marine-related conditions. Brominated and chlorinated carbazoles were produced through manganese oxide-mediated oxidation of carbazole in the presence of halide ions, with brominated carbazoles being the dominant species. Both reactive bromine species (RBS) and reactive chlorine species were detected in the reaction system. However, the detected RBS concentration was much lower than that of the observed brominated carbazoles under identical conditions. In reactions spiked with *tert*-butanol and methanol as  $\cdot\text{OH}$  scavengers, RBS production decreased from 0.56 nmol L<sup>-1</sup> to 0.12 nmol L<sup>-1</sup>, while the yield of 3-BCZ increased. These results suggest that PHCZ formation occurs through two primary pathways: (i) electrophilic substitution, where reactive halogen species generated by manganese oxide-mediated oxidation of halide ions react with carbazole and (ii) nucleophilic addition, where halide ions attack carbazole cations formed *via* manganese oxide oxidation. Our findings provide new insights into the abiotic processes contributing to the environmental load of PHCZs and suggest that manganese oxides play a critical role in the biogeochemical cycling of halogenated organic compounds in coastal and marine ecosystems.

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### Environmental significance

Polyhalogenated carbazoles (PHCZs) are emerging persistent organic pollutants frequently detected in marine and coastal environments, yet their natural formation pathways remain poorly understood. This study demonstrates that  $\delta\text{-MnO}_2$ , a ubiquitous and highly reactive manganese oxide in marine sediments, can mediate the abiotic halogenation of carbazole under environmentally relevant conditions. By revealing mineral-driven halogenation as a previously overlooked natural source of PHCZs, this work provides new insights into the geochemical processes governing the environmental occurrence of halogenated organic pollutants. These findings highlight the important role of manganese oxides in shaping the environmental fate of halogenated compounds and suggest that mineral-mediated reactions may contribute to the background levels of PHCZs in marine ecosystems.

## 1 Introduction

Polyhalogenated carbazoles (PHCZs) are a class of emerging persistent organic pollutants that have gained increasing attention due to their widespread occurrence and potential ecological risks. To date, more than 30 PHCZ congeners have been identified in diverse environmental matrices, including sediments,<sup>1–5</sup> soils,<sup>6–8</sup> air,<sup>8–10</sup> water,<sup>5,8</sup> and biological samples worldwide.<sup>8,11,12</sup> Structurally similar to polychlorinated dibenzofurans, PHCZs exhibit notable bioaccumulation potential and environmental persistence.<sup>11,13,14</sup> Moreover, toxicological studies have shown that PHCZs can induce developmental toxicity,<sup>15,16</sup> neurotoxicity,<sup>16,17</sup> and endocrine-disrupting

effects,<sup>18,19</sup> highlighting their significance in environmental and public health concerns.

Understanding the environmental sources and formation mechanisms of PHCZs is essential for assessing their environmental fate and risks. Both anthropogenic and natural processes have been proposed to explain their ubiquitous presence. Industrial activities can lead to unintended generation and release of PHCZs.<sup>20–22</sup> For example, Sun *et al.* identified and quantified PHCZ emissions from 13 major industrial sectors in China, revealing that coke production is the dominant source (9229 g per year), followed by iron ore sintering facilities (3237 g per year).<sup>23</sup> Additionally, Zhang and Lin reported that advanced oxidation processes used in coking wastewater treatment can generate PHCZs as byproducts.<sup>24</sup> However, these anthropogenic sources alone fail to explain the presence of PHCZs in sediment cores predating large-scale industrial activities,<sup>2,3,25</sup> suggesting that natural formation pathways also play a critical role. Indeed, natural halogenation

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processes have been demonstrated as a plausible source of PHCZs. Laboratory studies have shown that carbazole can undergo enzymatic halogenation catalyzed by chloroperoxidase and bromoperoxidase, resulting in the formation of a variety of chlorinated, brominated, and mixed-halogenated carbazoles.<sup>12,26</sup> These findings highlight that natural oxidative halogenation may substantially contribute to the environmental burden of PHCZs, especially in marine systems rich in halide ions.

Manganese oxides represent another important, yet less explored, natural halogenation mediator. As ubiquitous and powerful oxidants in soils and marine sediments, manganese oxides play a vital role in biogeochemical cycles of organic pollutants.<sup>27</sup>  $\text{MnO}_2$ , the most prevalent form, possesses strong redox potential ( $E_0 = 1.23$  V), enabling electron-transfer reactions with numerous organic contaminants adsorbed onto its surfaces.<sup>28–30</sup> The oxidation capability is further enhanced by reactive oxygen species (ROS) generated during  $\text{MnO}_2$  redox cycling.<sup>31–33</sup> Among manganese oxides,  $\delta\text{-MnO}_2$  is particularly reactive due to its high surface Mn(III) ( $E_0 = 1.5$  V) content.<sup>30,34,35</sup> Previous studies reported that  $\delta\text{-MnO}_2$  could also mediate the halogenation processes. For instance,  $\delta\text{-MnO}_2$  oxidizes iodide ions ( $E_0 = 0.54$  V) into reactive iodine species, subsequently leading to the formation of iodinated organic compounds.<sup>36–39</sup> Kadoya *et al.* reported that  $\delta\text{-MnO}_2$  mediates the chlorination of guaiacol (a lignin model compound) to form chloroguaiacol in the presence of chloride ions.<sup>40</sup> This process is proposed to occur by oxidizing guaiacol into cationic intermediates, which subsequently react with chloride ions to generate the chlorinated product. Such evidence suggests that  $\delta\text{-MnO}_2$ -mediated halogenation represents a potentially important abiotic pathway for the formation of PHCZs in natural environments.

We hypothesize that  $\delta\text{-MnO}_2$  can mediate the formation of PHCZs by oxidizing halide ions to reactive halogen species and by directly oxidizing carbazole to reactive intermediates that subsequently react with halide ions. To test this hypothesis, we systematically investigated  $\delta\text{-MnO}_2$ -mediated formation of PHCZs under marine-related conditions. Specifically, we evaluated the effects of  $\text{MnO}_2$  dosage and halide ion composition on PHCZ production and elucidated the underlying transformation pathways. This work advances understanding of geochemical processes contributing to natural PHCZ formation and highlights  $\delta\text{-MnO}_2$  as an abiotic driver of halogenated organic pollutant generation in marine and coastal ecosystems.

## 2 Materials and methods

### 2.1 Chemicals

The detailed information of carbazole and PHCZ congeners is listed in Table S1. 3,5-Dimethyl-1*H*-pyrazole (DMPZ,  $\geq 99\%$ ) and 4-bromo-3,5-dimethyl-1*H*-pyrazole (BDMPZ,  $\geq 99\%$ ) were purchased from Sigma-Aldrich (St. Louis, Missouri, USA). 4-Chloro-3,5-dimethylpyrazole (CDMPZ), 3,9'-bicarbazole (97%), 3-chloro-1,2-propanediol (3-CPD, 98%), 3-bromo-1,2-propanediol (3-BPD, 97%), and *N*-heptafluorobutyrylimidazole were purchased from Acme Biochemical (Shanghai, China). Allyl alcohol ( $10\text{ mg mL}^{-1}$ ) was purchased from BeNa Culture

Collection (Beijing, China). High-performance liquid chromatography grade acetone, *n*-hexane, acetonitrile, and methyl alcohol were purchased from Tedia (Fairfield, IA, USA), while dichloromethane was sourced from Anaqua Chemicals Supply (Houston, TX, USA). *Tert*-butanol ( $\geq 99\%$ ) was purchased from Macklin (Shanghai, China). Formic acid ( $\geq 98\%$ ) was purchased from Anpel (Shanghai, China). Analytical grade sodium bromide (NaBr), sodium nitrate ( $\text{NaNO}_3$ ), potassium permanganate ( $\text{KMnO}_4$ ), and manganese chloride tetrahydrate ( $\text{MgCl}_2 \cdot 4\text{H}_2\text{O}$ ) were purchased from Xilong Chemicals (Shantou, Guangdong, China). Guaranteed grade sodium chloride (NaCl), analytical grade sodium hydroxide (NaOH), and ascorbic acid were purchased from Sinopharm Chemical Reagent (Shanghai, China).

Salt solutions for the reactions were prepared by dissolving NaCl and NaBr solids in deionized water, and the pH was adjusted to 8.2 using  $0.01\text{ mol L}^{-1}$  NaOH. Artificial seawater was prepared following the method described by Dickson and Goyet.<sup>41</sup> Real seawater with a salinity of 35 used in the experiments was collected from the South China Sea, China.

Manganese dioxide ( $\delta\text{-MnO}_2$ ) was synthesized according to Murray's method.<sup>42</sup> Briefly, 1.64 L of deionized water was purged with nitrogen gas ( $\text{N}_2$ ) for 2 h prior to use, and 80 mL of  $0.1\text{ mol L}^{-1}$   $\text{KMnO}_4$  and 160 mL of  $0.1\text{ mol L}^{-1}$  NaOH solutions were added, followed by a dropwise addition of 120 mL of  $0.1\text{ mol L}^{-1}$   $\text{MnCl}_2$  while maintaining constant nitrogen sparging. The formed  $\delta\text{-MnO}_2$  particles were collected by centrifugation and rinsed several times with deionized water until the conductivity of the supernatant matched that of deionized water. The resulting  $\delta\text{-MnO}_2$  suspensions were stored at  $4\text{ }^\circ\text{C}$  and diluted to appropriate concentrations prior to use. To quantify the Mn content, the  $\delta\text{-MnO}_2$  particles were dissolved using ascorbic acid and subsequently analyzed on an inductively coupled plasma mass spectrometer (NexION 2000, PerkinElmer, Shelton, CT). The zeta potential of the synthesized  $\delta\text{-MnO}_2$  was measured to be  $-16.6\text{ mV}$  at pH 7.0. A portion of the synthetic  $\delta\text{-MnO}_2$  was air-dried, and its specific surface area was determined to be  $325\text{ m}^2\text{ g}^{-1}$  using the Brunauer–Emmett–Teller method based on  $\text{N}_2$  adsorption on an Autosorb-iQ-MP-C automated gas sorption analyzer (Quantachrome Instruments, Boynton Beach, FL). Scanning electron microscopy and X-ray diffraction analyses indicated that the synthetic  $\delta\text{-MnO}_2$  particles consisted of amorphous, irregular aggregates (Fig. S1).

### 2.2 $\delta\text{-MnO}_2$ oxidation experiment

The reactions were carried out in a 250 mL Erlenmeyer flask. First,  $30\text{ }\mu\text{L}$  of  $5\text{ mmol L}^{-1}$  carbazole (dissolved in methanol) was added to each flask. After the solvent evaporated, 30 mL of salt solution and an appropriate volume of  $\delta\text{-MnO}_2$  suspension were sequentially added. The mixture was homogenized to initiate the reaction. Triplicate samples were prepared for each treatment, and all samples were kept in the dark at  $25\text{ }^\circ\text{C}$  with continuous shaking at 120 rpm. At designated time intervals, triplicate samples were withdrawn, and the reaction was immediately quenched by adding an excess amount of ascorbic acid to dissolve the  $\delta\text{-MnO}_2$ . Unless otherwise noted, the



reaction was carried out at  $25 \pm 1$  °C for 24 h, with initial concentrations of  $5.0 \mu\text{mol L}^{-1}$  for carbazole,  $0.8 \text{ mmol L}^{-1}$  for  $\text{Br}^-$ , and  $100 \mu\text{mol L}^{-1}$  for  $\delta\text{-MnO}_2$ . The pH was set at 8.2 to simulate seawater conditions.

To assess the transformation of carbazole and formation of reaction products, 30 mL of the reaction solution was collected and extracted three times each with 15 mL of fresh dichloromethane. The organic phases were combined and evaporated to dryness, and the residue was redissolved in 1 mL of acetonitrile. The final samples were filtered through a PTFE filter ( $0.22 \mu\text{m}$ , Jinteng, Tianjin, China) and stored at 4 °C before instrumental analysis.

To investigate the effect of  $\delta\text{-MnO}_2$  concentration,  $\delta\text{-MnO}_2$  concentrations were set at 100, 200, and  $500 \mu\text{mol L}^{-1}$ . To explore the effect of halide ion concentration,  $\text{Br}^-$  concentrations were set at 0.08, 0.2, and  $0.8 \text{ mmol L}^{-1}$  and  $\text{Cl}^-$  concentrations were set at 5.4, 54.0, and  $540.0 \text{ mmol L}^{-1}$ . To assess the effect of the molar ratio of  $\text{Cl}^-$  to  $\text{Br}^-$  ( $[\text{Cl}^-]:[\text{Br}^-]$ ), their ratios were set at 0, 10 : 1, 100 : 1, and 500 : 1, while keeping the  $\text{Br}^-$  concentration at  $0.8 \text{ mmol L}^{-1}$ . When exploring the effect of the  $[\text{Cl}^-]:[\text{Br}^-]$  ratio,  $\text{NaNO}_3$  was added to maintain a constant  $\text{Na}^+$  concentration, eliminating the potential influence of  $\text{Na}^+$  on  $\text{MnO}_2$  oxidation.

### 2.3 Identification of reactive halogen species

The generation of reactive halogen species (RHS), including reactive chlorine species (RCS) and reactive bromine species (RBS), in the  $\delta\text{-MnO}_2$  reaction system was identified using allyl alcohol and DMPZ as chemical probes. Allyl alcohol reacts with RHS to produce 3-CPD and 3-BPD,<sup>43,44</sup> while DMPZ reacts with RHS to yield 4-chloro-3,5-dimethylpyrazole (CDMPZ) and BDMPZ.<sup>45</sup>

For the allyl alcohol probe experiments, the reaction was conducted under conditions identical to those used for carbazole oxidation, except that the carbazole substrate was replaced with  $20 \mu\text{L}$  of  $10 \text{ mg mL}^{-1}$  allyl alcohol. After 24 h of reaction, 10 mL of the reaction mixture was collected, and excess  $\text{Na}_2\text{SO}_3$  and  $\text{NaHSO}_3$  were added to quench the residual RHS and dissolve the  $\delta\text{-MnO}_2$ . The sample was extracted three times with 5 mL ethyl acetate, vortexing for 60 s each time. The organic phase was collected, combined, dehydrated using anhydrous  $\text{Na}_2\text{SO}_4$ , and concentrated under nitrogen blow to approximately  $100 \mu\text{L}$ . The residue was reconstituted in 2 mL of acetonitrile, followed by the addition of  $50 \mu\text{L}$  heptafluorobutanoyl imidazole. The mixture was stirred for 60 s and heated at 70 °C in a water bath for 30 min to derivatize 3-CPD and 3-BPD into 1,2-bis(heptafluorobutyryloxy)-3-chloropropane and 1,2-bis(heptafluorobutyryloxy)-3-bromopropane, respectively. After cooling to room temperature, the sample was diluted with 5 mL of deionized water and 2 mL of saturated NaCl solution, followed by extraction with 2 mL of *n*-hexane. The extraction was repeated twice, and the combined *n*-hexane extracts were concentrated to 1 mL under a gentle nitrogen flow. The final extract was transferred to a 2 mL vial for analysis. The analysis was carried out on a gas chromatography-mass spectrometer (GCMS-QP2010 Ultra, Shimadzu, Tokyo, Japan). The

separation was achieved on a DB-5MS capillary column ( $50 \text{ m} \times 0.25 \text{ mm}$ ,  $0.25 \mu\text{m}$  film thickness). The injection ( $1.0 \mu\text{L}$ ) was in splitless mode at 250 °C, with high-purity helium (99.999%) as a carrier gas at  $1.0 \text{ mL min}^{-1}$ . The oven temperature started at 60 °C (held for 1 min), followed by heating to 120 °C at  $5 \text{ }^\circ\text{C min}^{-1}$  (held for 3 min), then increased to 250 °C at  $20 \text{ }^\circ\text{C min}^{-1}$  and held for 2 min. The MS was operated in electron impact mode (70 eV). Selected ions were  $m/z$  253, 289, and 453 for 1,2-bis(heptafluorobutyryloxy)-3-chloropropane and 197, 225, and 467 for 1,2-bis(heptafluorobutyryloxy)-3-bromopropane.

For experiments employing DMPZ as the probe, the carbazole substrate was replaced with  $30 \mu\text{L}$  of  $100 \text{ mmol L}^{-1}$  DMPZ. Triplicate samples were prepared for each treatment. At designated time intervals, triplicate samples were withdrawn and excess ascorbic acid solution was added to quench the reaction by dissolving  $\delta\text{-MnO}_2$ . The samples were then filtered through a  $0.22 \mu\text{m}$  PVDF membrane filter (Jinteng, Tianjin, China) and analyzed directly by LC-MS/MS. The derivatized BDMPZ and CDMPZ were analyzed on a liquid chromatography-triple quadrupole mass spectrometer (Agilent 1290 Infinity LC-6490 MS/MS) equipped with an electrospray ionization (ESI) source (Agilent, Palo Alto, CA, USA). The MS was operated in positive ion (ESI+) mode with multiple reaction monitoring. For the detection of BDMPZ, precursor ion  $m/z$  175 and product ions  $m/z$  95/54 were selected. The quantitative ion was  $m/z$  95. The collision energies were optimized to be 30 eV for  $m/z$  95 and 40 eV for  $m/z$  54. For the detection of CDMPZ, precursor ion  $m/z$  131 and product ions  $m/z$  54/62.8 were selected. The quantitative ion was  $m/z$  54. The collision energies were optimized to be 25 eV for  $m/z$  54 and 20 eV for  $m/z$  62.8. Other MS parameters followed those of Zhang and Lin.<sup>22</sup> The mobile phase consisted of ultrapure water (A) and acetonitrile containing 0.1% formic acid (B), with a flow rate of  $0.20 \text{ mL min}^{-1}$  and an injection volume of  $5 \mu\text{L}$ . To achieve optimal separation, the gradient elution program was 0–2 min, 5% B; 2–4 min, 5–30% B; 4–7 min, hold at 30% B; 7–9 min, 30–5% B; and 7–12 min, hold at 5% B to restore the initial conditions. To prevent salt contamination, the eluent was directed to waste for the first 5 min before switching to MS detection.

### 2.4 Analysis of carbazole and reaction products

The concentrations of carbazole and PHCZs were determined using LC-MS/MS. The mobile phase comprised water (A) and acetonitrile (B) at a flow rate of  $0.25 \text{ mL min}^{-1}$ . The gradient elution program was 0–1 min, hold at 55% B; 1–8 min, a linear increase to 80% B; 8–15 min, hold at 90% B; 15–17 min, hold at 90% B; 17–17.1 min, return to 55% B; 17.01–20 min, equilibrate at 55% B. The MS was operated in negative ion (ESI-) mode with multiple reaction monitoring for the quantification of carbazole and PHCZs. The MS/MS parameters followed those of Zhou *et al.*<sup>46</sup>

For unknown reaction product analysis, LC-MS/MS was operated in full scan and product ion scan modes. In full scan mode, the  $m/z$  range was set from 60 to 800. In the product ion scans, the  $m/z$  scan range was set from 60 to  $(M + 20)$  ( $M$  denotes



the molecular weight) and collision energies were varied from 5 to 70 eV, depending on the fragment structure. Potential products were screened from full scan results, and suspected compounds were confirmed by comparing retention times and fragmentation patterns with those of authentic standards.

### 3 Results and discussion

#### 3.1 $\delta$ -MnO<sub>2</sub>-mediated transformation of carbazole into PHCZs and other products

The results showed that  $\delta$ -MnO<sub>2</sub> catalyzed the transformation of carbazole into PHCZs in solutions containing various concentrations of halide ions (Fig. 1). In the presence of Br<sup>-</sup> and Cl<sup>-</sup>, the reaction produced both mono- and di-halogenated carbazoles. Specifically, in the solution containing 0.8 mmol L<sup>-1</sup> Br<sup>-</sup> and 0.54 mol L<sup>-1</sup> Cl<sup>-</sup>, three PHCZ congeners were detected, including 3-BCZ ( $4.8 \times 10^{-1} \mu\text{g L}^{-1}$ ), 36-BCZ ( $3.0 \times 10^{-2} \mu\text{g L}^{-1}$ ), and 3-CCZ ( $1.4 \times 10^{-1} \mu\text{g L}^{-1}$ ) (Fig. 1). Notably, despite the Br<sup>-</sup> concentration being much lower than that of Cl<sup>-</sup>, brominated carbazoles dominated the product species, indicating that Br<sup>-</sup> is substantially more reactive than Cl<sup>-</sup> in the reaction system. This higher reactivity can also explain the detection of 3-BCZ in the reaction with 0.54 mol L<sup>-1</sup> Cl<sup>-</sup> alone, likely due to trace Br<sup>-</sup> impurities.

In artificial seawater, only 3-BCZ ( $1.1 \times 10^{-2} \mu\text{g L}^{-1}$ ) was detected (Fig. 1), an order of magnitude lower than that in the mixed-halide system with comparable halide concentrations. Similarly, a higher concentration of 3-BCZ ( $0.3 \times 10^{-1} \mu\text{g L}^{-1}$ ) was formed in natural seawater. This discrepancy suggests that other components in artificial and natural seawater, particularly high concentrations of metal cations, may suppress  $\delta$ -MnO<sub>2</sub>-mediated PHCZ formation. Previous studies have shown that high levels of Ca<sup>2+</sup> and Mg<sup>2+</sup> can reduce the reactivity of  $\delta$ -MnO<sub>2</sub> in two ways: (i) by adsorbing onto the negatively charged  $\delta$ -MnO<sub>2</sub> surface (the point of zero charge  $\text{pH}_{\text{pzc}} = 2.4$ ), altering surface structure, reducing colloidal stability, and inducing precipitation<sup>29,47,48</sup> and (ii) by occupying surface reactive sites, thereby blocking access for organic substrates.<sup>49,50</sup> Both effects

can inhibit the  $\delta$ -MnO<sub>2</sub>-mediated transformation of carbazole into PHCZs. The differences observed between artificial and natural seawater may also arise from variations in halide ion concentrations and the presence of natural organic matter, which influence the complex transformation process among carbazole, halide ions and MnO<sub>2</sub>.

The concentration of  $\delta$ -MnO<sub>2</sub> significantly influenced the formation of PHCZs (Fig. S2). As the  $\delta$ -MnO<sub>2</sub> concentration increased from 100  $\mu\text{mol L}^{-1}$  to 500  $\mu\text{mol L}^{-1}$ , the final production of 3-BCZ and 36-BCZ gradually decreased. This decline is likely due to the further oxidation of 3-BCZ and 36-BCZ through oxidative reactions between  $\delta$ -MnO<sub>2</sub> and these products, as supported by the results shown in Fig. S3. Like carbazole, 3-BCZ and 36-BCZ may first adsorb onto the  $\delta$ -MnO<sub>2</sub> active sites, followed by electron transfer. Higher  $\delta$ -MnO<sub>2</sub> concentrations increase the number of reactive sites, which enhances the oxidative breakdown of both carbazole and PHCZ products, ultimately reducing the final PHCZ yield.

Fig. S4A illustrates the strong oxidizing ability of  $\delta$ -MnO<sub>2</sub> toward carbazole. In the reaction system containing  $\delta$ -MnO<sub>2</sub>, carbazole removal was nearly complete, accompanied by the release of Mn<sup>2+</sup>, the reductive product of  $\delta$ -MnO<sub>2</sub>. This can be attributed to two factors: (i) carbazole's redox potential (0.93 V) is lower than that of  $\delta$ -MnO<sub>2</sub> (1.23 V),<sup>51</sup> making oxidation thermodynamically favorable and (ii) alternative transformation pathways, such as oxidative coupling, may occur.<sup>29,52</sup> Evidence for oxidative coupling was obtained *via* mass spectrometry, revealing a distinct chromatographic peak at *m/z* 330.8,

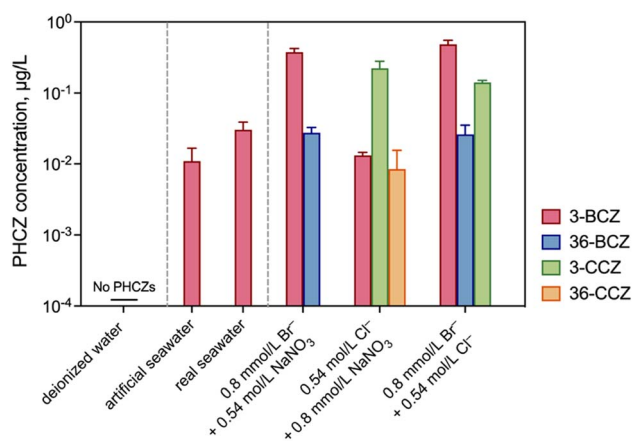


Fig. 1  $\delta$ -MnO<sub>2</sub>-mediated transformation of carbazole into PHCZs in different solutions. The initial concentrations of carbazole and  $\delta$ -MnO<sub>2</sub> were 5  $\mu\text{mol L}^{-1}$  and 100  $\mu\text{mol L}^{-1}$ , respectively. The pH of the solution was adjusted to 8.2 and the reaction was conducted at 25 °C for 24 h.

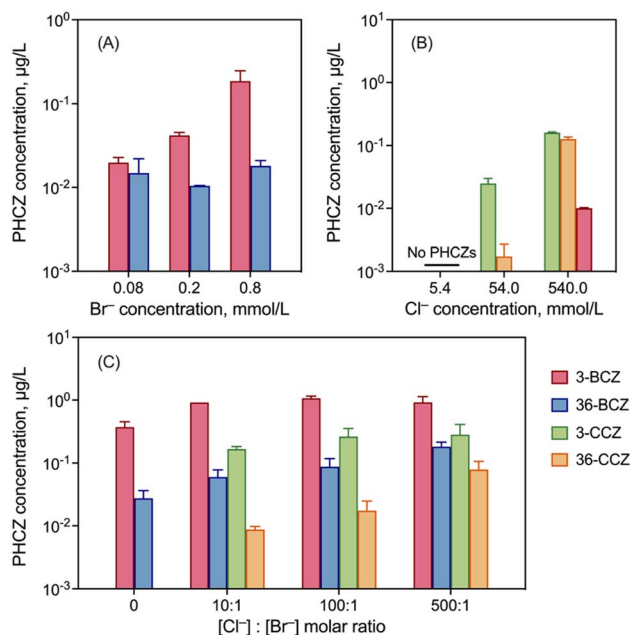


Fig. 2 Effect of Br<sup>-</sup> concentration (A), Cl<sup>-</sup> concentration (B), and Cl<sup>-</sup>/Br<sup>-</sup> ratio (C) on PHCZ production. The initial concentrations of carbazole and  $\delta$ -MnO<sub>2</sub> were 5  $\mu\text{mol L}^{-1}$  and 100  $\mu\text{mol L}^{-1}$ , respectively. The initial concentration of Br<sup>-</sup> was 0.8 mmol L<sup>-1</sup> in panel (C), and NaNO<sub>3</sub> was added to maintain a constant Na<sup>+</sup> concentration to eliminate the potential influence of Na<sup>+</sup> on MnO<sub>2</sub> oxidation. The pH of the solution was adjusted to 8.2 and the reaction was conducted at 25 °C for 24 h.



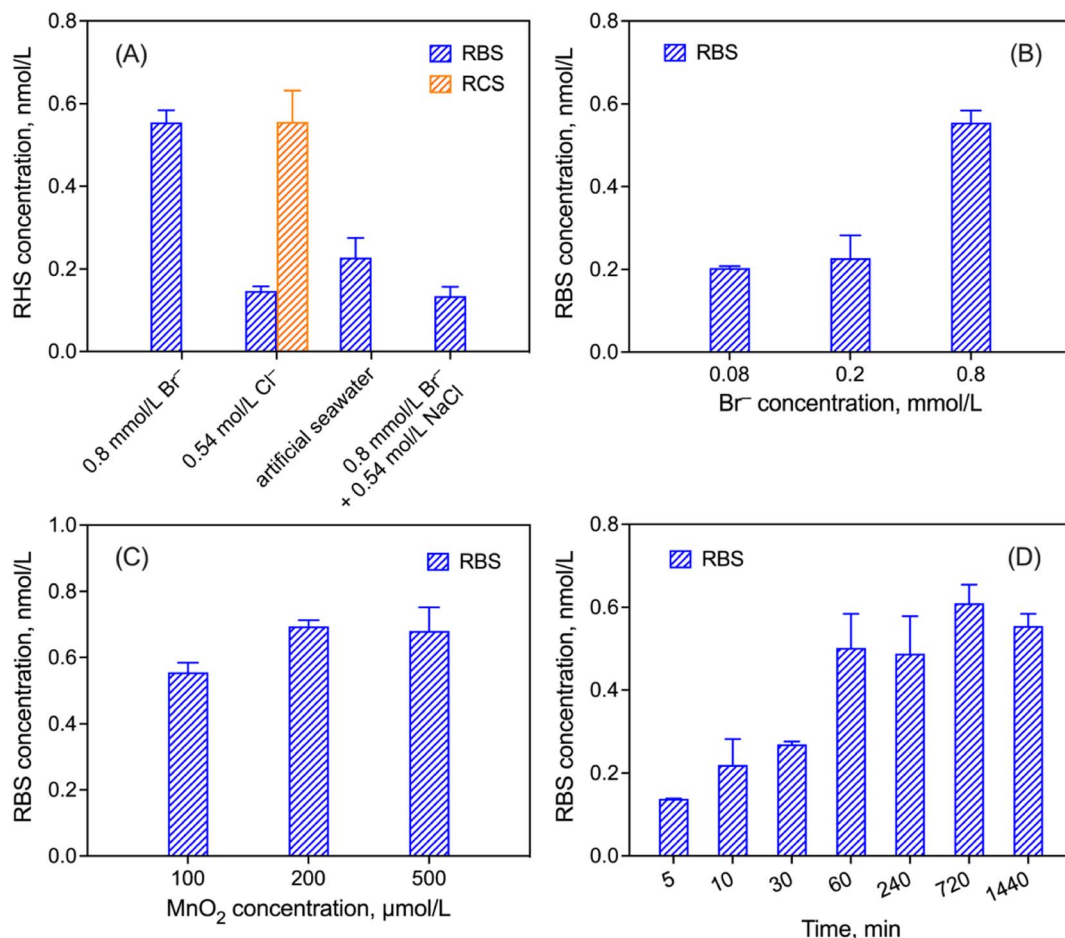


Fig. 3 Effect of reaction solutions (A), Br<sup>-</sup> concentration (B), MnO<sub>2</sub> concentration (C), and reaction time (D) on the generation of reactive halogen species (RHS). Unless specified, the reaction solution contained 0.8 mmol L<sup>-1</sup> NaBr and 100 μmol L<sup>-1</sup> δ-MnO<sub>2</sub> and the reaction time was 24 h. The solution pH was adjusted to 8.2.

corresponding to a carbazole dimer (Fig. S4B). The major fragment ion at *m/z* 165.9 corresponded to the carbazole monomer. Based on retention time and mass spectral comparison with those of an authentic standard, the unknown compound was identified as 3,9'-bicarbazole (Fig. S4C).

### 3.2 Effect of halide ions

Halide ions act as halogen donors for the halogenation process, directly influencing PHCZ yield. As shown in Fig. 2A, increasing Br<sup>-</sup> concentration from 0.08 mmol L<sup>-1</sup> to 0.8 mmol L<sup>-1</sup> led to nearly a tenfold increase in 3-BCZ. Under conditions containing 5.4 mmol L<sup>-1</sup> of Cl<sup>-</sup>, no chlorinated carbazoles were detected. However, when the Cl<sup>-</sup> concentration was elevated from 54.0 mmol L<sup>-1</sup> to 540 mmol L<sup>-1</sup>, the concentrations of 3-CCZ and 36-CCZ rose from  $2.5 \times 10^{-2}$  μg L<sup>-1</sup> and  $1.7 \times 10^{-3}$  μg L<sup>-1</sup> to  $1.6 \times 10^{-1}$  μg L<sup>-1</sup> and  $1.3 \times 10^{-1}$  μg L<sup>-1</sup>, respectively (Fig. 2B). These results clearly show that δ-MnO<sub>2</sub>-mediated PHCZ formation is halide concentration-dependent, with higher Br<sup>-</sup> and Cl<sup>-</sup> levels promoting increased yields of their respective halogenated carbazoles.

Given that Br<sup>-</sup> and Cl<sup>-</sup> often coexist in marine environments, we further examined the effect of their molar ratio on

PHCZ formation. The molar ratios of Cl<sup>-</sup> to Br<sup>-</sup> ([Cl<sup>-</sup>]:[Br<sup>-</sup>]) were varied at 0, 10 : 1, 100 : 1, and 500 : 1, while keeping the Br<sup>-</sup> concentration at 0.8 mmol L<sup>-1</sup>. The results demonstrated that the increasing proportion of Cl<sup>-</sup> enhanced the formation of both chlorinated and brominated carbazoles (Fig. 2C). This effect may be explained by two factors: (i) higher Cl<sup>-</sup> availability facilitates its adsorption to reactive δ-MnO<sub>2</sub> sites, directly promoting 3-CCZ and 36-CCZ formation and (ii) greater Cl<sup>-</sup> surface occupation may partially suppress further oxidative degradation of halogenated carbazoles by δ-MnO<sub>2</sub>, thereby improving their final yield.

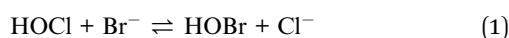
### 3.3 Production of RHS

To further probe how halide ions influence PHCZ formation, the generation of RHS in the reaction system was investigated. RHS are key intermediates in the formation of halogenated organic compounds; therefore, their detection was essential for understanding the reaction mechanism. Phenolic chemical probes commonly used for RHS detection (*e.g.*, 2,6-dichlorophenol) can be readily oxidized by δ-MnO<sub>2</sub>, leading to potential interference. In contrast, allyl alcohol and DMPZ are inert to δ-MnO<sub>2</sub>, making them suitable probes in this study.



Since the allyl alcohol derivatives 3-CPD and 3-BPD can undergo further reactions with allyl alcohol,<sup>43,44</sup> allyl alcohol was used only for qualitative identification of RHS. DMPZ, however, reacts stoichiometrically (1 : 1 molar ratio) with RCS and RBS to form CDMPZ and BDMPZ, respectively,<sup>45</sup> and was therefore used for quantitative analysis.

The results consistently confirmed RHS generation in solutions containing either Cl<sup>-</sup> or Br<sup>-</sup> (Fig. 3 and S5). In the reaction system with 0.54 mol L<sup>-1</sup> Cl<sup>-</sup>, 0.55 nmol L<sup>-1</sup> RCS and 0.15 nmol L<sup>-1</sup> RBS were detected, with the formation of RBS primarily attributed to trace bromide Br<sup>-</sup> impurities in the NaCl reagent. In both the mixed-halide system and the artificial seawater system, only RBS was detected while no RCS was observed. This result can be explained by the fact that both high Br<sup>-</sup> concentrations and the generated RBS can convert RCS into other RBS, such as HOBr and BrOCl, as shown in eqn (1) and (2).<sup>53,54</sup>



Both the Br<sup>-</sup> concentration and  $\delta$ -MnO<sub>2</sub> concentration significantly influenced RBS generation (Fig. 3B and C). Increasing the Br<sup>-</sup> concentration from 0.08 mmol L<sup>-1</sup> to 0.8 mmol L<sup>-1</sup> increased RBS yield from 0.2 nmol L<sup>-1</sup> to 0.56 nmol L<sup>-1</sup>, likely due to more Br<sup>-</sup> binding to reactive  $\delta$ -MnO<sub>2</sub> sites and promoting MnO<sub>2</sub>-mediated RBS formation. Increasing the  $\delta$ -MnO<sub>2</sub> concentration from 100  $\mu$ mol L<sup>-1</sup> to 200  $\mu$ mol L<sup>-1</sup> raised RBS production from 0.56 nmol L<sup>-1</sup> to 0.69 nmol L<sup>-1</sup>; however, further increasing  $\delta$ -MnO<sub>2</sub> to 500  $\mu$ mol L<sup>-1</sup> did not significantly enhance RBS generation. This result suggests that once  $\delta$ -MnO<sub>2</sub> reaches an excess level, Br<sup>-</sup> availability becomes the limiting factor for further RBS production.

The production of RBS gradually increased with reaction time before eventually reaching a plateau (Fig. 3D). This trend might be attributed to the continuous release of Mn<sup>2+</sup>, the reduction product of  $\delta$ -MnO<sub>2</sub>, into the reaction system as the reaction progresses.<sup>29,55,56</sup> On the one hand, Mn<sup>2+</sup> readily adsorbs on the  $\delta$ -MnO<sub>2</sub> surface, occupying reactive sites that

would otherwise be available for Br<sup>-</sup> binding. The adsorption of Mn<sup>2+</sup> and its inhibitory effect on the oxidative capacity of  $\delta$ -MnO<sub>2</sub> have been extensively documented in studies focusing on  $\delta$ -MnO<sub>2</sub>-catalyzed oxidation of hydroquinone, phenols, and anilines.<sup>30,55,57</sup>



The Nernst equation

$$E_{\text{h}} = E_0 + 0.0296 \lg \frac{[\text{MnO}_2][\text{H}^+]^4}{[\text{Mn}^{2+}]} \quad (4)$$

predicts that the accumulation of Mn<sup>2+</sup> would lower the redox potential of  $\delta$ -MnO<sub>2</sub>. Together, Mn<sup>2+</sup> surface adsorption and the decrease in  $\delta$ -MnO<sub>2</sub> redox potential reduce its ability to generate RHS, ultimately leading to the stabilization of RBS production over time.

The generation of RBS may occur *via* two potential pathways. First, the redox potentials of  $\delta$ -MnO<sub>2</sub> ( $E_0 = 1.23$  V) and the Mn(III) species ( $E_0 = 1.50$  V) within  $\delta$ -MnO<sub>2</sub> are higher than that of Br<sup>-</sup> ( $E_0 = 1.09$  V), enabling direct oxidation of Br<sup>-</sup> to Br<sub>2</sub>. Second, previous studies have demonstrated that when molecular oxygen (O<sub>2</sub>) adsorbs vertically onto the surface oxygen vacancies in MnO<sub>2</sub> and is bound to adjacent Mn(III) atoms, single-electron transfer from Mn(III) to O<sub>2</sub> occurs, leading to sustained production of O<sub>2</sub><sup>•-</sup>.<sup>31,32,58</sup> This superoxide can be converted to other reactive oxygen species (ROS), such as <sup>•</sup>OH, through a series of radical reactions (eqn (5)–(7)). Wang *et al.* detected <sup>•</sup>OH generation in MnO<sub>2</sub> with different crystallinities and confirmed significant contribution of <sup>•</sup>OH to the mineralization of dissolved organic matter.<sup>33</sup> Given that <sup>•</sup>OH is the most powerful oxidative ROS ( $E_0 = 2.8$  V), it can further oxidize Br<sup>-</sup>, thereby facilitating the production of RBS.

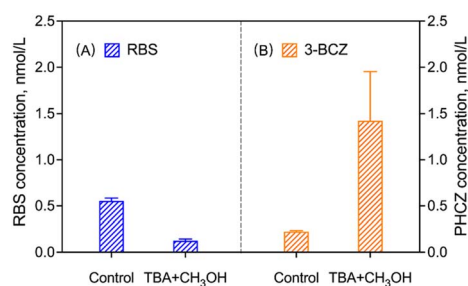
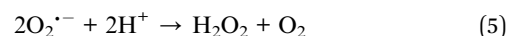


Fig. 4 Effect of *tert*-butanol (TBA) and methanol on the generation of RBS (A) and PHCZs (B). Conditions in panel (A): the reaction was carried out in 0.8 mmol L<sup>-1</sup> NaBr solution with pH adjusted to 8.2. The initial concentrations of DMPZ and  $\delta$ -MnO<sub>2</sub> were both 100  $\mu$ mol L<sup>-1</sup>. The initial concentrations of TBA and CH<sub>3</sub>OH were both 200 mmol L<sup>-1</sup> in experimental groups. No TBA or CH<sub>3</sub>OH was added in control groups. The reaction time was 24 h. Conditions in panel (B): after 24 h of reaction in the system shown in panel (A), 5  $\mu$ mol L<sup>-1</sup> of carbazole was added, and the mixture was allowed to react for an additional 24 h.

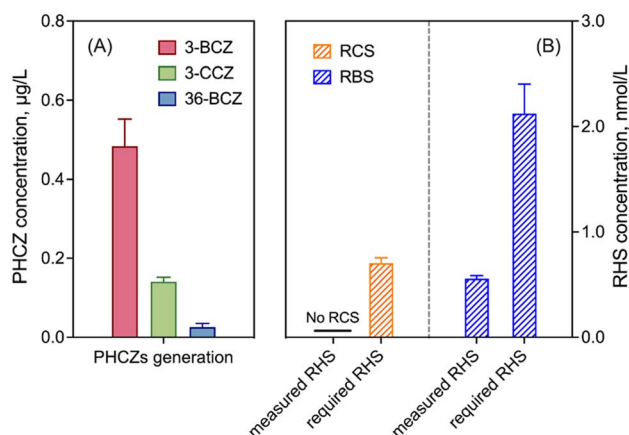
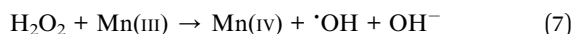
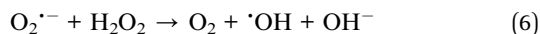


Fig. 5 The generation of PHCZs (A), RHS, and the required amount of RHS to generate PHCZs (B) in the mixed halide solution. The initial concentration of carbazole and  $\delta$ -MnO<sub>2</sub> was 5  $\mu$ mol L<sup>-1</sup> and 100  $\mu$ mol L<sup>-1</sup>, respectively with a reaction time of 24 h. For the investigation of RHS production, the substrate carbazole was replaced with 100  $\mu$ mol L<sup>-1</sup> of DMPZ.





To investigate whether  $\cdot\text{OH}$  participates in RBS generation, 200 mmol  $\text{L}^{-1}$  *tert*-butanol and 200 mmol  $\text{L}^{-1}$  methanol were added into the reaction systems as  $\cdot\text{OH}$  scavengers. The addition of these scavengers significantly reduced RBS production from 0.56 nmol  $\text{L}^{-1}$  to 0.12 nmol  $\text{L}^{-1}$ , while simultaneously increasing the yield of 3-BCZ (Fig. 4). This pronounced reduction of RBS confirms that RBS formation predominantly depended on ROS, particularly  $\cdot\text{OH}$ . The residual RBS detected in the system likely originated from the direct oxidation of  $\text{Br}^-$  by  $\text{MnO}_2$  and  $\text{Mn(III)}$ . The higher 3-BCZ yields likely resulted from the inhibition of its further oxidative degradation due to  $\cdot\text{OH}$  scavenging.

Since RCS was not detected in real seawater, it is unlikely that RCS form under typical environmental conditions. Thus,

the formation mechanism of RCS is not discussed in detail. Nonetheless, trace amounts of RCS could potentially be generated through pathways analogous to those of RBS formation, likely mediated by  $\text{Mn(III)}$  and ROS-driven oxidation, considering the relatively high redox potential of  $\text{Cl}_2/\text{Cl}^-$  (1.36 V), which makes its direct oxidation by  $\delta\text{-MnO}_2$  less favorable than that of  $\text{Br}^-$ .

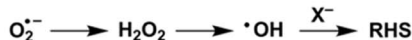
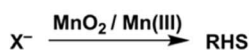
### 3.4 Potential PHCZ formation pathways

The potential pathways for PHCZ formation are outlined in Fig. 6. In brief,  $\delta\text{-MnO}_2$ -mediated transformation of carbazole into PHCZs likely occurs *via* two distinct pathways: (i) electrophilic substitution involving  $\delta\text{-MnO}_2$ -generated RHS (pathway I) and (ii) nucleophilic addition of halide ions to the carbazole cation formed through  $\delta\text{-MnO}_2$  oxidation (pathway II).

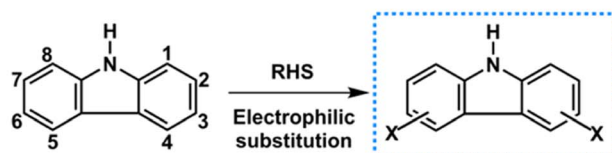
In the electrophilic substitution pathway,  $\delta\text{-MnO}_2$  generates RHS, such as  $\text{Br}_2$ ,  $\text{HOBr}$ , and their chlorinated analogs, which

#### (I) RHS-mediated the formation of PHCZs

##### (I-i) RHS formation:

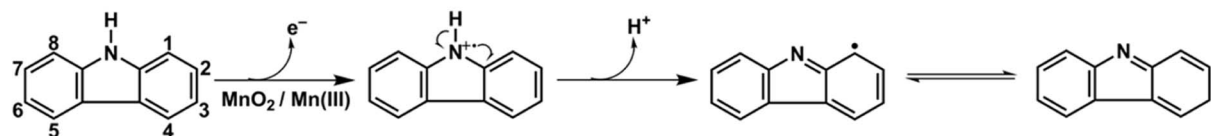


##### (I-ii) PHCZs formation:

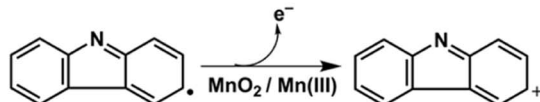


#### (II) Carbazole cation-mediated the formation of PHCZs

##### (II-i) Carbazole radical formation:



##### (II-ii) Carbazole cation formation:



##### (II-iii) PHCZs formation:

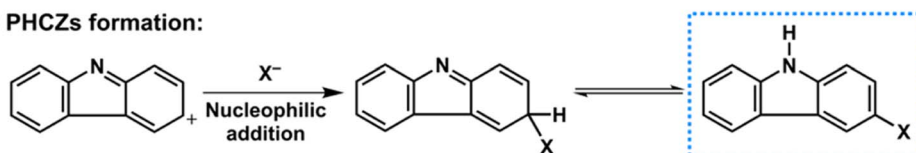


Fig. 6 Proposed reaction pathways for  $\text{MnO}_2$ -mediated transformation of carbazole into PHCZs.



subsequently attack the electron-rich carbazole ring. In aqueous systems,  $\delta$ -MnO<sub>2</sub> oxidizes halide ions (e.g., Br<sup>-</sup> and Cl<sup>-</sup>) either directly or *via* intermediate ROS, producing RHS with strong electrophilic reactivity. Carbazole, with its high electron density at the C-3 position due to resonance effects from the nitrogen atom, is particularly susceptible to electrophilic attack.<sup>59</sup> The RHS species preferentially substitute at this C-3 position, forming halogenated carbazole derivatives such as 3-BCZ and 3-CCZ. This pathway is supported by the detection of PHCZs in  $\delta$ -MnO<sub>2</sub>/halide systems, even in the absence of other known halogenating agents, highlighting the important role of  $\delta$ -MnO<sub>2</sub>-generated RHS in initiating halogenation.

Although electrophilic substitution by  $\delta$ -MnO<sub>2</sub>-generated RHS represents a key pathway for PHCZ formation, additional mechanisms are likely involved. Two observations support this: (i) in the presence of <sup>•</sup>OH scavengers, the yield of 3-BCZ exceeded the measured RBS concentration (Fig. 4) and (ii) 3-CCZ was detected despite the absence of measurable RCS in a mixed-halide solution containing 0.8 mmol L<sup>-1</sup> Br<sup>-</sup> and 0.54 mol L<sup>-1</sup> Cl<sup>-</sup>, with the RBS concentration theoretically required for the detected brominated carbazoles being much higher than the measured value (Fig. 5). Similarly, a previous study demonstrated that  $\delta$ -MnO<sub>2</sub> can oxidize guaiacol to a phenoxyl radical, which undergoes further one-electron oxidation to form a phenoxenium cation. This cation reacts with chloride or bromide to form chloroguaiacol and bromoguaiacol.<sup>40</sup> Given carbazole's electron-rich aromatic structure and relatively low redox potential, it is prone to electron transfer with  $\delta$ -MnO<sub>2</sub>, making a similar pathway plausible. We propose that direct  $\delta$ -MnO<sub>2</sub>-mediated transformation of carbazole into PHCZs involves the following steps: (i) radical formation: the nitrogen atom's lone electron pair facilitates electron transfer to  $\delta$ -MnO<sub>2</sub>, generating a carbazole radical. The unpaired electron delocalizes within the conjugated system, localizing at the C-3 position; (ii) cation formation: the carbazole radical undergoes further oxidation by  $\delta$ -MnO<sub>2</sub> or Mn(III) to yield a carbazole cation; (iii) nucleophilic addition: halide ions attack the carbazole cation at C-3, followed by isomerization to form 3-BCZ or 3-CCZ.

## 4 Conclusions

This study demonstrates that  $\delta$ -MnO<sub>2</sub> can mediate the transformation of carbazole into PHCZs under marine-related conditions. The formation of PHCZs likely proceeds through two concurrent pathways: (i) electrophilic substitution, where carbazole reacts with reactive halogen species generated by the oxidation of halide ions by  $\delta$ -MnO<sub>2</sub>, and (ii) nucleophilic addition, where halide ions attack carbazole cations produced *via*  $\delta$ -MnO<sub>2</sub>-induced oxidation. Elevated halide ion concentrations, particularly Br<sup>-</sup>, significantly enhance PHCZ formation. Given the widespread presence of halide ions, manganese oxides, and carbazole in marine environments, these findings suggest that  $\delta$ -MnO<sub>2</sub>-mediated halogenation could be an important natural pathway for PHCZ formation. This study provides new insights into the abiotic processes contributing to the environmental load of PHCZs and highlights  $\delta$ -MnO<sub>2</sub> as a key geochemical

driver of halogenated organic compound formation in coastal and marine ecosystems.

## 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.

## Data availability

The data supporting this article have been included as part of the supplementary information (SI). Supplementary information: Table S1 and Fig. S1–S5. See DOI: <https://doi.org/10.1039/d6em00064a>.

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