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MOF-derived Fe-doped δ -MnO₂ nanoflowers as oxidase mimics: chromogenic sensing of Hg(II) and hydroquinone in aqueous media†

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The structure and morphology play a crucial role in enhancing the biomimetic oxidase activity of nanozymes. In this study, a facile *in situ* chemical oxidation strategy was employed to synthesize MOF-derived MnO_x, utilizing the structural features of the parent MOF to enhance oxidase-mimicking activity. We have systematically investigated the effects of phase evolution, structural modulation and morphology on the oxidase activity of MnO_x with Fe substitution. The oxidase-like activity was evaluated using the chromogenic substrate 3,3',5,5'-tetramethylbenzidine (TMB), which produced blue colored ox-TMB with an absorption peak at 652 nm upon oxidation. While all Fe-doped MnO_x nanostructures exhibited oxidase-like activity, the 10% Fe-doped sample (**10Fe-MnO_x**) demonstrated the highest performance, likely due to a synergistic effect of the structure, morphology and existing oxygen vacancies. The underlying oxidase mechanism was investigated using steady-state kinetics and electron paramagnetic resonance (EPR) analysis. In addition, a colorimetric assay was developed for detecting Hg²⁺ and hydroquinone (HQ) in real water samples collected from industrial and natural sources. The calculated detection limits of the colorimetric probe **10Fe-MnO_x** for HQ (1.74 μM) and Hg²⁺ (0.47 μM) outperformed conventional metal oxide-based nanozymes. These findings may pave the way for the development of easily synthesizable, scalable and highly sensitive MOF-derived metal oxide nanozymes with promising applications in biological and environmental settings.

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Introduction

Over the years, natural enzymes have been the subject of extensive research in various biochemical processes due to their high substrate specificity and catalytic efficacy.^{1–3} However, their catalytic activity is influenced by several factors such as pH, temperature, reaction medium, *etc.* Furthermore, their widespread application is limited by high production costs, storage challenges, instability (including denaturation and protease digestion) and purification difficulties.^{3,4} Significant efforts are thus underway to develop novel enzyme mimics exhibiting desired catalytic activity and chemical stability.

Artificial enzymes offer several advantages over their natural counterparts, including enhanced stability, greater tolerance to environmental variations, and low-cost production and storage.^{5–8} Contextually, a diverse range of synthetic materials have been documented to exhibit biomimetic activity similar to natural enzymes such as peroxidases,⁹ oxidases,¹⁰ catalases,¹¹ superoxide dismutase,¹² *etc.* To develop artificial enzymes, inorganic materials have emerged as more promising synthetic models due to their tuneable structural, electronic and redox features.¹³ Inspired by the work of Gao *et al.*, which demonstrated that Fe₃O₄ can catalyse the oxidation of the chromogenic substrate tetramethylbenzidine (TMB),¹⁴ recent studies have explored the enzymatic potential of various nanomaterials based on CuO,¹⁵ Co₃O₄,¹⁶ V₂O₅,¹⁷ MnO₂,¹⁸ Ag₂O,¹⁹ *etc.* Among them, manganese oxides (MnO_x) have gained particular attention due to their variable redox behaviour, cost-effectiveness, favourable biological safety profile and relatively higher oxidizing activity.²⁰ MnO_x can be developed in multiple phases (such as MnO, Mn₃O₄, Mn₂O₃ and MnO₂), with MnO₂ standing out due to its superior catalytic efficacy and recyclability. Moreover, MnO₂ nanostructures can be easily tuned to attain a desired morphology including nanotubes, nanowires,

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nanoplates and nanorods.²¹ For instance, Zhang *et al.* synthesized MnO₂ nanowires, MnOOH/MnO₂ nanorods, and octahedral MnO₂ nanoparticles using a hydrothermal method, finding that MnO₂ nanowires exhibited superior catalytic activity as oxidase mimics.²² Similarly, Liu *et al.* demonstrated the oxidase-mimicking activity of MnO₂ nanoparticles, showcasing their ability to catalyse the oxidation of colorless TMB to blue oxidized TMB (ox-TMB) without the need for hydrogen peroxide (H₂O₂).²³ Despite these advancements, studies on the structure–activity relationship of MnO₂ oxidase mimics remain limited. Structurally, MnO₂ exists in various crystallized forms (such as α -, β - and δ -MnO₂), each defined by its number of [MnO₆] octahedral subunit framework and tunnel size.^{24,25} Furthermore, doping of various metal ions such as V⁵⁺, Ag⁺, Co²⁺, Ni²⁺, Cu²⁺ and Fe^{+3/+4} into MnO₂ substantially improves its redox properties and the dispersion of active sites, which are crucial for catalytic applications.^{26–32} Among these, iron (Fe) is particularly beneficial as it can either substitute Mn or occupy the tunnel structures, stabilizing the material and improving catalytic efficiency.³³ Wang *et al.* reported that Fe doping influences the phase of MnO₂ nanostructures, with different Fe concentrations yielding α -MnO₂, R-MnO₂, or β -MnO₂, thereby enhancing capacitance retention and energy density.³⁴ Moreover, Fe doping introduces oxygen vacancies, which are known to significantly enhance electrocatalytic and photocatalytic activities due to the increased active sites and high surface energy.³⁵ While these reported nanozymes exhibit comparable catalytic activities to natural enzymes, their intrinsic catalytic activities are often compromised by factors such as a limited number of exposed active sites, intrinsically deficient multilevel structures and aggregation issues.^{36–38}

Among them, MnO₂ nanostructures with high porosity and a large number of exposed active sites are found to be more promising for specific catalytic applications. In this endeavour, MnO_x materials derived from metal–organic frameworks (MOFs) have emerged as a profound class of porous materials.^{39–41} In particular, MOF-derived MnO₂ nanostructures have attracted increasing research interest in catalysis due to their well-defined coordination networks and tunable porosity. The diversity of metallic nodes, linking struts, and coordination interactions in all possible directions positions MOFs as direct surrogates to develop metal oxide based artificial enzymes. The well-defined, tailorable cavities and channels of parent MOFs can be retained in the resulting metal oxides (MOs). Typically, the rationale behind the use of MOF-derived metal oxides as promising catalysts lies in their ability to retain a high surface area, enhancing catalytic activity by exposing more active sites. The MOF-to-MO conversion also enables precise control over crystal phase, particle size, and morphology. Carbon residues from the MOF structure provide stability to MOs by encapsulating the nanostructures and preventing active component loss during reactions. The porous architecture ensures efficient mass transfer and improved interaction between catalysts and reactants. Based on these advantages, several MOFs have shown potential as templates, suitable support materials, and precursors for manufacturing nanostructured materials. Compared to the existing meso-

porous materials, MOF derivatives offer a novel pathway for constructing mesoporous nanozymes. In this context, the intrinsic mesoporous properties of MOFs can be maintained, while the stability and activity of the resulting nanozymes can be significantly improved. Furthermore, the mesoporous structures and multiple channels can enable full contact with substrates and facilitate mass transport and diffusion of products in catalytic applications.^{42–45}

Recently, several studies have demonstrated that MOF-derived manganese oxides offer precise structural control and significantly improve their catalytic properties. For example, Lu *et al.* synthesized a hollow MnFeO oxide from a PBA@PBA core–shell structure *via* thermal treatment, achieving both oxidase- and peroxidase-like activities.⁴⁶ Similarly, Fe₃O₄@C@MnO₂ composites, formed *via* an interfacial reaction between pyrolyzed Fe-MIL-88A and KMnO₄, exhibited appreciable biomimetic activities.³⁸ Gai *et al.* developed MnO_x@NC with dual-enzyme properties by calcining a Mn-TTPCA MOF.⁴² While these studies highlight the potential of MOF-derived MnO₂ nanozymes, their thermal synthesis often results in aggregation or structural collapse, limiting their catalytic efficiency. A more controlled *in situ* chemical oxidation strategy is thus essential to optimize the structure and morphology of such materials. To the best of our knowledge, the impact of Fe doping on the phase, structure, and morphology of MnO_x nanostructures and its influence on oxidase activity remains largely unexplored.

Herein, we introduce a new cost-effective strategy to prepare MOF-derived Fe-doped δ -MnO₂ nanoflowers *via* a straightforward solvent treatment of the MOF template. The as-prepared MnO_x successfully retained the morphology of the parent MOF while exhibiting phase transitions influenced by varying levels of Fe doping. Notably, at 10% Fe doping, a complete conversion into a pure δ -MnO₂ phase was observed. The oxidase activities of MOF-derived manganese oxides were assessed by monitoring the oxidation of TMB in the absence of H₂O₂. All the synthesized materials demonstrated oxidase activity; however, 10% Fe-doped MnO_x (referred to as **10Fe-MnO_x**) exhibited the highest catalytic activity, attributed to its phase structure, nanoflower-like morphology and oxygen vacancies. The catalytic mechanism was investigated using steady-state kinetics and an EPR study. Furthermore, under optimized conditions, we demonstrated a straightforward colorimetric method for sensitive detection of hydroquinone and Hg²⁺ ions.

Experimental section

Preparation of the MnBTC MOF and Fe-doped MnBTC MOFs

The MnBTC MOF was prepared using a procedure reported elsewhere.⁴⁷ Briefly, 2.0 g of PVP ($M_w = 30\,000$) was dissolved in a 100 mL solvent mixture of ethanol and water (2 : 1, v/v) and stirred for 10 min at 400 rpm. Thereafter, 2.0 mmol of Mn(CH₃COO)₂ was added to the above solution to form a colourless transparent solution A. In a separate beaker, 4.73 mmol

(0.9 g) of 1,3,5-benzenetricarboxylic acid (BTC) was dissolved in a 100 mL ethanol–water (2 : 1, v/v) mixture to obtain colorless solution B. The two solutions A and B were then mixed and stirred for 1 hour at 400 rpm, and the resulting mixture was left to rest overnight at room temperature. A white-colored powder was collected by centrifugation, washed thrice with ethanol/water and finally dried at 60 °C for 4 hours.

A series of Fe-doped MnBTC MOFs were synthesized following a procedure similar to that described above, with varying Fe molar percentages (1%, 5% and 10%) relative to Mn, while keeping the total metal precursor amount ($\text{Mn}(\text{CH}_3\text{COO})_2$ and FeCl_3) at 2.0 mmol. The resulting products were labelled as 1Fe-MnBTC, 5Fe-MnBTC and 10Fe-MnBTC, respectively.

Preparation of MOF derived MnO_x and Fe-doped MnO_x

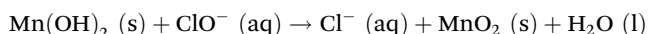
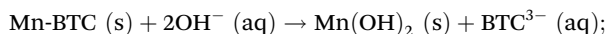
MOF-derived manganese oxides were synthesized using a facile chemical oxidation method. An alkaline NaOCl solution was prepared by mixing 100 mL of a 0.5 M NaOH aqueous solution with 100 mL of a 0.3 mmol NaOCl aqueous solution at room temperature. Subsequently, 0.05 g of the synthesized MnBTC MOF was added to the alkaline solution of NaOCl and stirred overnight to obtain a black suspension of MnO_x . The resulting suspension was centrifuged, washed three times with ethanol/water, and dried under vacuum to yield a black-colored powder of MnO_x .

Fe-doped MnO_x samples were prepared using the same chemical oxidation method applied to 1Fe-MnBTC, 5Fe-MnBTC and 10Fe-MnBTC, resulting in **1Fe-MnO_x**, **5Fe-MnO_x**, and **10Fe-MnO_x**, respectively.

Results and discussion

Structural description

Fe-doped MnO_x samples were synthesized through a simple chemical oxidation of Fe-MnBTC MOFs under alkaline conditions using NaOCl. Weak Mn(II)–O bonds in Mn-BTC MOFs facilitate their easy transformation into metal oxides *via* solvent treatment. Upon exposure to NaOH, OH^- ions diffuse into the porous MOF structure, displacing BTC ligands through a thermodynamically favorable process and forming an unstable $\text{Mn}(\text{OH})_2$ intermediate. This intermediate subsequently reacts with ClO^- ions in solution, resulting in the formation of mixed-phase MnO_x . The reaction process can be represented as follows:



The phase structure of the parent Fe-MnBTC MOFs was first examined using PXRD analysis, as shown in Fig. S1.† The PXRD patterns closely matched the reported and simulated patterns, confirming the successful synthesis of the MOFs. With increasing Fe doping, a slight peak shift toward a higher angle was observed, suggesting minimal structural alterations. This shift suggests a gradual substitution of Mn by Fe while

preserving the original crystal structure. To investigate the structural and phase evolution with varying Fe-doping levels, the PXRD patterns of MOF-derived manganese oxides were analyzed (Fig. 1). The undoped MnBTC MOF-derived manganese oxide exhibited a mixed-phase composition, primarily consisting of the tetragonal Mn_3O_4 phase (JCPDS: 24-0734) and the monoclinic $\delta\text{-MnO}_2$ phase (JCPDS: 80-1098). The weak diffraction peaks indicated a predominantly amorphous nature, while the broad peaks suggested a nanoscale morphological structure. Although the phase composition of MnO_x remained largely unchanged upon 1% and 5% Fe doping, a significant improvement could be realized in the crystallinity as evidenced by enhanced peak intensities. The resulting higher crystallinity indicates that the incorporation of small amounts of Fe may promote better ordering within the crystal structure without altering the crystal phases.

A drastic transformation occurred at 10% Fe doping, where the phase composition shifted to pure $\delta\text{-MnO}_2$. The PXRD pattern of **10Fe-MnO_x** exhibited distinct peaks dedicated solely to $\delta\text{-MnO}_2$, confirming a complete transition from mixed-phase MnO_x to $\delta\text{-MnO}_2$ at this doping concentration. This phase change suggests that 10% Fe doping disrupts the stability of Mn_3O_4 under oxidizing conditions, favouring the formation of pure $\delta\text{-MnO}_2$. In this process, Fe acted as both a structure-directing agent and a catalytic promoter, enabling the oxidation of Mn_3O_4 to MnO_2 .

Though several research groups have attributed sharp peaks at similar 2θ values to the $\alpha\text{-MnO}_2$ phase with an orthorhombic crystal structure, a few reports suggest the Mn_3O_4 phase with a tetragonal structure. We resolved this ambiguity using Rietveld refinement analysis, confirming the peaks correspond to Mn_3O_4 . Rietveld refinement analysis was performed on PXRD data of **5Fe-MnO_x** and **10Fe-MnO_x** nanostructures using GSAS-II software⁴⁸ to confirm the initially identified phases and quantify their percentages (Fig. 2). Table S1† presents the lattice parameters and crystallographic

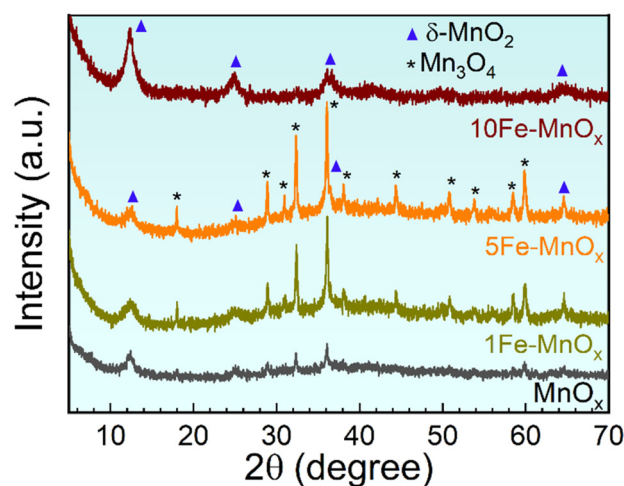


Fig. 1 PXRD spectra of MOF-derived MnO_x , **1Fe-MnO_x**, **5Fe-MnO_x** and **10Fe-MnO_x**.

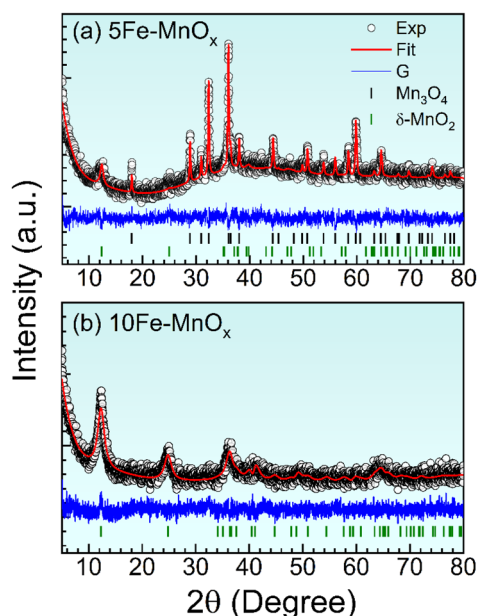


Fig. 2 Rietveld refinement plots of XRD patterns for (a) **5Fe-MnO_x** and (b) **10Fe-MnO_x** nanostructures. The experimental data are shown as open black circles, the theoretical fit as solid red lines, and the Bragg positions for **Mn₃O₄** and **δ-MnO₂** crystal structures are indicated by black and olive bars, respectively. Blue solid lines at the bottom reflect the difference between experimental and theoretical data.

angles derived from the best-fit data, along with relevant fitting parameters. The XRD pattern for the **5Fe-MnO_x** nanostructures was best modelled as a mixture of tetragonal **Mn₃O₄** (78%, space group *I4₁/amd*) and monoclinic **δ-MnO₂** (22%, space group *C2/m*). In contrast, the **10Fe-MnO_x** nanostructures were found to be purely **δ-MnO₂** with a monoclinic crystal structure (space group *C2/m*). The Rietveld analysis also confirmed that in both samples, Fe occupies some of the Mn sites in the manganese oxide crystal lattice. These analyses provide insights into the influence of iron doping concentration on phase formation, crystal structures, and microstructures, which will be discussed in subsequent sections.

FE-SEM and TEM analyses were conducted to investigate the morphology and microstructures of the as-prepared Fe-MnBTC MOFs and **Fe-MnO_x**. The SEM images of Mn-MOF revealed the presence of urchin/flower-like architectures composed of thin nanosheets (Fig. 3a). FE-SEM micrographs of **MnO_x** (Fig. 3b) showed that the flower-like morphology of the parent MOF is well preserved in the respective **MnO_x**.

Further comparison of SEM images of the **MnO_x** samples revealed the impact of Fe doping on morphology. The undoped **MnO_x** exhibited a mixed phase of **Mn₃O₄** and **δ-MnO₂**, characterized by an amorphous morphology resembling flower-like structures. In contrast, a notable transformation in morphology was observed at 5% Fe doping. The sample exhibited a mixed morphology comprising both rod-like and flower-like structures (Fig. 3c). These results corroborate the PXRD analysis exhibiting improved crystallinity in **5Fe-MnO_x**. At 10% Fe-doping, the morphology largely transformed

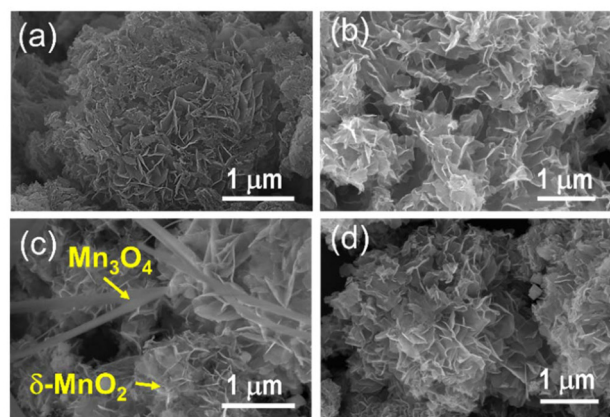


Fig. 3 FE-SEM micrographs of (a) Mn-MOF; (b) **MnO_x**; (c) **5Fe-MnO_x**; and (d) **10Fe-MnO_x**.

into flower-like structures (Fig. 3d). This transition indicates that the higher levels of Fe-doping favour the establishment of a more uniform flower-like morphology. It is worth mentioning that the incorporation of Fe into the framework likely influences the nucleation and growth trend during the synthesis of MOF derived metal oxides. 10% Fe dopants can significantly alter the coordination environment and surface energy of growing crystallites, thus promoting a more controlled and homogeneous self-assembly process. The results from the FE-SEM analysis aligned well with those obtained from PXRD analysis, highlighting that variation in Fe levels not only influences the phase composition and crystallinity but also significantly affects the morphological characteristics of **MnO_x**.

Furthermore, the morphology, microstructure and composition of the as-prepared **5Fe-MnO_x** and **10Fe-MnO_x** were evaluated using TEM analysis (Fig. 4). The TEM images of **5Fe-MnO_x** (Fig. 4a) revealed the presence of nanorod-like morphology. This observation was consistent with the SEM analysis. High-resolution TEM (HR-TEM) micrographs showed a lattice fringe spacing (*d*) of ~0.49 nm, corresponding to the (101) planes of the tetragonal **Mn₃O₄** structure (Fig. 4b).

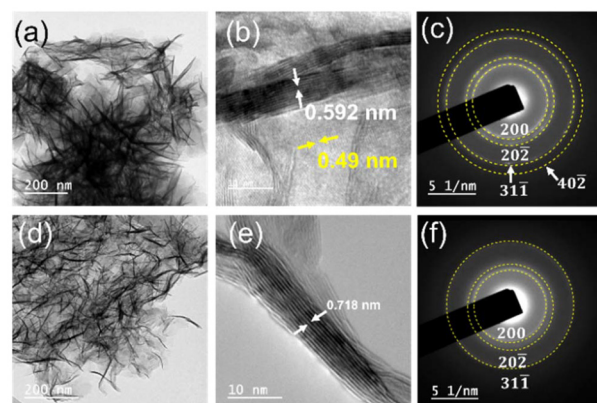


Fig. 4 High resolution TEM images and SAED patterns of **5Fe-MnO_x** (a–c) and **10Fe-MnO_x** (d–f).

Additionally, a flower-like morphology was observed, featuring a stripe spacing of ~ 0.592 nm, a characteristic fingerprint of birnessite-type δ - MnO_2 . The SAED image in Fig. 4c indicated polycrystallinity and the rings are in good agreement with the mixed phase pattern of Mn_3O_4 and δ - MnO_2 . Similar to the SEM analysis, a complete transfer of morphology to nanoflowers could be seen when 10% Fe was doped (Fig. 4d). These nanoflowers exhibited stripe spacings of ~ 0.72 nm, attributed to the presence of interlayer water molecules and cations that stabilize the layered sheets. This d -spacing corresponds to the (001) reflection planes of δ - MnO_2 (Fig. 4e), which could also be observed in the PXRD pattern at $2\theta = 12.6^\circ$. The layer thickness of 10Fe-MnO_x was determined to be ~ 3.9 nm, which is consistent with the crystallite size measured from the PXRD pattern (Fig. S2[†]). This atomic-level thickness and layered morphology likely enhance the exposure of active sites, facilitating electron transfer and thereby contributing to the superior catalytic activity. The SAED pattern of 10Fe-MnO_2 exhibited small spots along the rings, indicating its polycrystalline nature (Fig. 4f). All the rings closely matched the d -values associated with δ - MnO_2 . This structural evolution highlights the significant impact of Fe doping on the morphology and crystal structure of MnO_x , promoting the transition from a mixed morphology to a nanoflower-like structure with enhanced layer stability. Furthermore, the EDS elemental mapping of both 5Fe-MnO_x and 10Fe-MnO_x demonstrated a homogeneous distribution of elements throughout the nanostructure, with C, O, Mn and Fe as the main elements (Fig. S3 and S4[†]). The EDX spectra of 5Fe-MnO_x and 10Fe-MnO_x confirmed %Fe doping levels of 6 and 9.4, as shown in the inset table of Fig. S5[†].

To analyze the chemical state and composition of the MOFs and their derived manganese oxides, X-ray photoelectron spectroscopy (XPS) was conducted. The XPS survey spectra (Fig. S6[†]) confirmed the presence of Mn, O and Fe, with no detectable impurity peaks. The Mn 2p spectra exhibited two main peaks at ~ 642.6 eV and ~ 654.0 eV, corresponding to Mn 2p_{3/2} and Mn 2p_{1/2}, respectively. High-resolution spectra were deconvoluted using XPSPEAK41 software. For all manganese oxide samples, deconvolution of the Mn 2p_{3/2} peak revealed three distinct oxidation states: Mn^{4+} (~ 644.1 eV), Mn^{3+} (~ 642.7 eV) and Mn^{2+} (~ 641.1 eV) (Fig. 5). However, the relative contributions of Mn^{2+} , Mn^{3+} and Mn^{4+} varied significantly among samples. Notably, the 10Fe-MnO_x sample exhibited a higher surface concentration of Mn^{3+} and Mn^{4+} compared to the 0% and 5% Fe-doped samples. Additionally, shifts in the 2p_{3/2} and 2p_{1/2} peak positions of Mn could be noticed at higher binding energies with increasing Fe-doping, along with changes in the $\text{Mn}^{2+}/\text{Mn}^{3+}/\text{Mn}^{4+}$ ratios, indicating the modulation of oxidation states along with related spin-orbit coupling upon Fe incorporation. The deconvolution of the asymmetric O 1s spectra for all MOF-derived manganese oxides revealed three distinct components with binding energies approximately at 529.6 eV, 530.7 eV and 532.8 eV (Fig. 5). Based on the existing literature, these peaks correspond to lattice oxygen (O_{latt} , Mn–O–Mn), oxygen vacancies (O_V), and surface-adsorbed oxygen (O_{ads} , Mn–O–H), respectively. The oxygen vacancy concen-

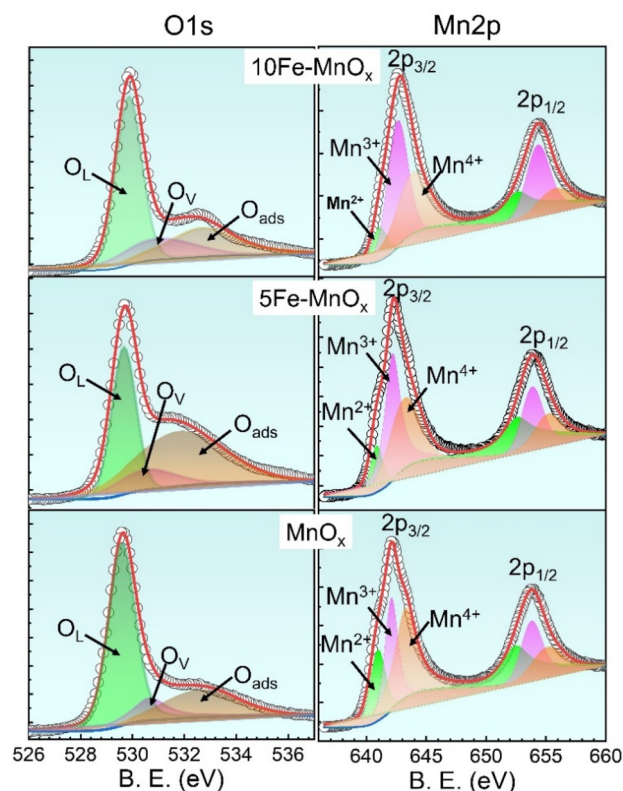


Fig. 5 Deconvoluted XPS spectra of O 1s and Mn 2p for MnO_x , 5Fe-MnO_x and 10Fe-MnO_x .

tration was quantified using the ratio $[\text{O}_V/(\text{O}_L + \text{O}_V + \text{O}_{\text{ads}})]$. The Fe doping level significantly influenced oxygen vacancy concentrations, with the 10Fe-MnO_x sample exhibiting the highest value (19%), followed by 5Fe-MnO_x (15%) and 0Fe-MnO_x (12%). An increased oxygen vacancy concentration correlates with a greater number of active oxygen sites, which can eventually enhance catalytic oxidation performance.

High-resolution Fe 2p spectral analysis of the 5Fe-MnO_x and 10Fe-MnO_x samples is presented in Fig. S7[†]. In both samples, the Fe spin-orbit peaks 2p_{1/2} and 2p_{3/2} were observed at binding energies of 724.7 eV and 710.6 eV, respectively. The deconvolution of these spin-orbit peaks revealed three distinct peaks: two at approximately 710.2 eV and 711.5 eV, corresponding to Fe^{3+} , and a smaller peak at 713.5 eV, attributed to Fe^{2+} . This indicates the presence of both Fe^{3+} and Fe^{2+} in the MnO_x lattice, with varying Fe doping levels. Notably, the intensity of the Fe 2p peaks increases with higher Fe doping. Additionally, satellite peaks have been observed at higher binding energies, specifically at 719.3 eV and 732.6 eV. The synthesized 10Fe-MnO_x shows promising potential for oxidase-mimic catalysis, attributed to the solid-state redox couples of $\text{Mn}^{3+}/\text{Mn}^{4+}$, $\text{Fe}^{3+}/\text{Fe}^{2+}$ and the presence of oxygen vacancies.

Oxidase-like activity of manganese oxides

The synthesized manganese oxides have been employed as catalysts to explore their potential in biomimetic oxidase-like

activity. It is well known that the reaction conditions and environmental factors significantly influence the oxidase-like activities of both natural and artificial enzymes. In order to evaluate the impact of Fe doping in MnO_x , key catalytic parameters including incubation time, pH, temperature, and catalytic dosage of 10Fe-MnO_x were optimized for the oxidation of a chromogenic substrate, TMB. A UV-Vis study was conducted to monitor the activity of 10Fe-MnO_x for a period of 20 min, where the absorbance at 652 nm rapidly increased and reached saturation within 15 min (Fig. S8a[†]). Although 10Fe-MnO_x exhibited the highest activity at 15 °C, most likely due to the increased solubility of dissolved oxygen at lower temperatures, all subsequent experiments were conducted at room temperature for practical relevance (Fig. S8b[†]). pH optimization is also crucial in enzyme-based catalysis, as biological enzymes often exhibit varying activity across different pH levels. The catalytic activity of 10Fe-MnO_x was thus tested in different buffer solutions with pH ranging from 2.0 to 10.0. As illustrated in Fig. S8c[†], the oxidase activity of 10Fe-MnO_x increased between pH 2.0 and 4.0, reaching the highest response at pH 4.0. A further increase in pH led to a decline in catalytic efficiency, and therefore, pH 4.0 was selected as the optimal value. Additionally, to determine the ideal catalyst loading, various concentrations of 10Fe-MnO_x were evaluated for TMB oxidation. As shown in Fig. S8d[†], the highest catalytic efficiency was achieved at an optimal dosage of $6 \mu\text{g mL}^{-1}$.

Under optimized conditions, the oxidase-like activity of all MOF-derived MnO_x samples was systematically compared. A notable trend emerged, wherein progressive Fe-doping led to a steady enhancement in catalytic performance, with 10Fe-MnO_x exhibiting the highest activity. This was further validated through UV-Vis absorption studies. To quantitatively assess the oxidase-like efficiency of both undoped and Fe-doped MnO_x , the absorbance change of TMB at 652 nm was monitored. Among all samples, 10Fe-MnO_x demonstrated the most pronounced absorption peak at 652 nm, confirming its superior catalytic activity (Fig. 6a).

The relative activity of different catalysts was calculated using the formula as described in the Experimental section. The catalytic efficiency of the MOF-derived MnO_x samples followed the order $\text{MnO}_x < 1\text{Fe-MnO}_x < 5\text{Fe-MnO}_x < 10\text{Fe-MnO}_x$ (Fig. 6b). The exceptional performance of 10Fe-MnO_x can be attributed to its unique phase structure and nanoflower-like morphology, which features interlayer channels and cavities facilitating superior substrate diffusion, thereby enhancing catalytic activity. Fe^{3+} doping into the MnO_x lattice, which primarily contains Mn^{4+} , induced the formation of oxygen vacancies and promoted the presence of redox-active $\text{Mn}^{3+}/\text{Mn}^{4+}$ and $\text{Fe}^{3+}/\text{Fe}^{2+}$ couples, both of which are critical for oxidation reactions. XPS analysis further confirmed that 10Fe-MnO_x exhibited the highest concentration of oxygen vacancies among all tested catalysts, reinforcing its superior catalytic efficiency.

For comparison, the oxidase-like activity of the parent MOF and 10Fe-MnO_x synthesized *via* a co-precipitation method

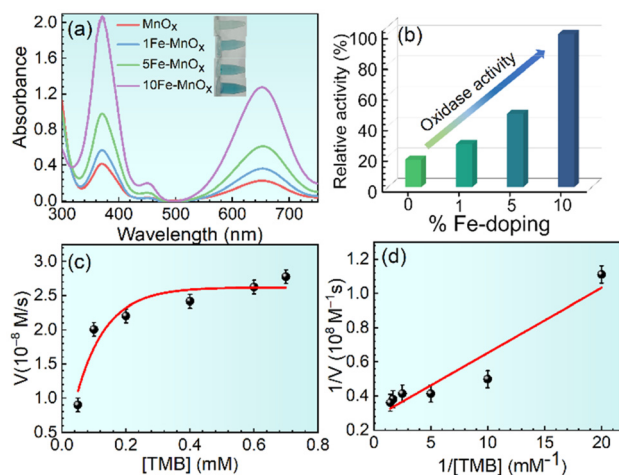


Fig. 6 (a and b) Comparative oxidase-like activity of different MOF-derived MnO_x catalysts (assay conditions: catalyst loading $6.6 \mu\text{g mL}^{-1}$; pH = 4.0; r. t.; TMB = 0.1 mM). Inset: photograph of the TMB-buffer system with different catalysts. Steady-state kinetic analysis using (c) Michaelis–Menten curves and (d) Lineweaver–Burk plots for catalytic oxidation of TMB using 10Fe-MnO_x .

was evaluated. The detailed synthesis procedure is provided in the ESI,[†] and the PXRD pattern (Fig. S9[†]) of 10% Fe-doped MnO_x confirms the formation of $\delta\text{-MnO}_2$. As shown in Fig. S10,[†] MOF-derived 10Fe-MnO_x exhibits significantly enhanced activity, highlighting the advantage of the MOF-templated approach in generating catalytically active Fe-doped $\delta\text{-MnO}_2$.

Kinetic studies of oxidase mimicking activity

A steady-state kinetic analysis was conducted under optimized conditions by varying TMB concentrations (0–100 μM), while maintaining all other parameters constant. The formation rate of the oxidized product was monitored by tracking the absorbance increase at 652 nm. Using the molar extinction coefficient ($\epsilon = 3900 \text{ M}^{-1} \text{ cm}^{-1}$), the corresponding concentrations of ox-TMB were calculated. Michaelis–Menten curves presented in Fig. 6c show the relationship between V_0 and TMB concentration. The Lineweaver–Burk plots (Fig. 6d) were obtained by plotting the double reciprocal $1/V$ vs. $1/[\text{TMB}]$, in which the slope of the double reciprocal curve corresponds to the K_m/V_{max} ratio, and the intercept of the double reciprocal curve corresponds to $1/V_{\text{max}}$. The K_m value represents the affinity between the substrate TMB and the catalyst, with a lower K_m value indicating a strong affinity of the substrate for the catalyst.⁴⁹ As summarised in Table S2,[†] compared to HRP and other reported bioinspired enzymes, 10Fe-MnO_x exhibited a relatively lower K_m value (0.15 mM), indicating its higher catalytic activity. The superior activity of 10Fe-MnO_x was mainly due to the large number of catalytically active sites generated *via* oxygen vacancies. Meanwhile, the good dispersive effect of MOF-derived metal oxides avoids the shortcoming of easy agglomeration, thus increasing the effective substrate–catalyst interactions.

Oxidase mechanism

Synthetic enzyme mimics typically generate reactive oxygen species (ROS), which play a critical role in the oxidation of chromogenic substrates such as TMB. The oxidase-like activity of these enzymes relies on dissolved oxygen, which acts as an electron acceptor to generate superoxide radical anions ($O_2^{\cdot-}$). These active species can spontaneously convert into H_2O_2 , which further decomposes to produce hydroxyl radicals ($\cdot OH$) or singlet oxygen (1O_2). As previously discussed, $10Fe-MnO_x$ contains a high density of oxygen vacancies and redox-active metal couples (Fe^{3+}/Fe^{2+} and Mn^{3+}/Mn^{4+}), both of which facilitate efficient ROS generation. These ROS oxidize TMB to its blue-colored oxidized form, confirming the oxidase-mimicking activity of $10Fe-MnO_x$. To verify the *in situ* generation of ROS, we employed DPPH (2,2-diphenyl-1-picrylhydrazyl), a stable free radical, as a detection probe. The presence of ROS was confirmed by a decrease in the absorbance of DPPH near 520 nm, indicating ROS-mediated radical scavenging (Fig. S11†). To specifically confirm the role of $O_2^{\cdot-}$ in TMB oxidation, we performed the reaction in the presence of *p*-benzoquinone, a known radical scavenger for $O_2^{\cdot-}$ radical anions. As shown in Fig. 7a, the relative oxidase-like activity of $10Fe-MnO_x$ gradually decreased with increasing concentrations of *p*-benzoquinone, confirming the *in situ* generation of $O_2^{\cdot-}$ to catalyze TMB oxidation.

To further confirm the ROS formation during oxidase-like activity, EPR studies were conducted with and without $10Fe-MnO_x$ in ethanol (Fig. 7b). A strong hyperfine signal was observed in the EPR spectra within the 280–360 gauss range when 5.0 mg of $10Fe-MnO_x$ was dispersed in ethanol. In contrast, no signal was detected in the control experiment

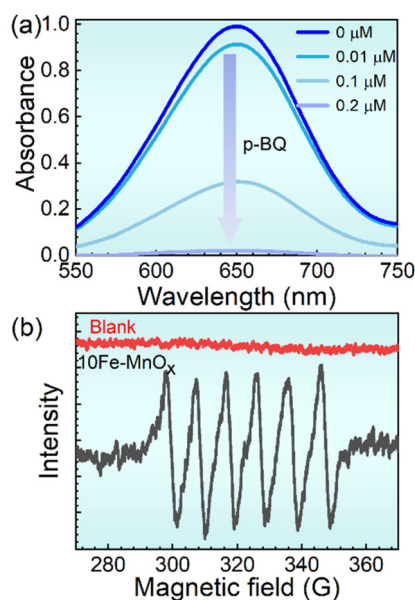
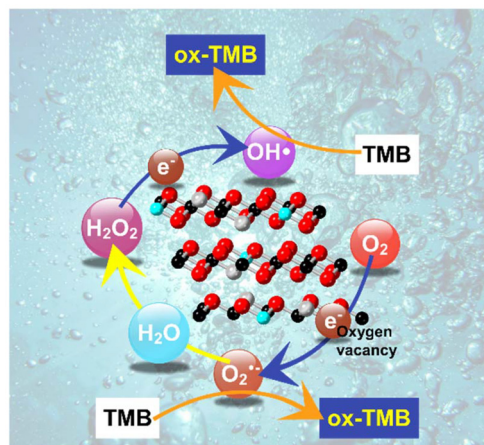


Fig. 7 (a) Variation in the absorbance of $10Fe-MnO_x$ /TMB system upon adding *para*-benzoquinone (*p*-BQ) and (b) EPR spectra of ethanolic solution with and without $10Fe-MnO_x$.

(absence of the catalyst), confirming the formation of ROS in the colloidal suspension of $10Fe-MnO_x$. Additionally, the oxidation of TMB under aerobic and anaerobic conditions was examined to assess the role of dissolved oxygen or surface hydroxyls in the oxidase-like activity of $10Fe-MnO_x$. A significant reduction in absorbance at 652 nm was observed when N_2 gas was bubbled into the $10Fe-MnO_x$ /TMB system, indicating the crucial effect of dissolved oxygen on oxidase activity (Fig. S12†). The possible mechanism for oxidase-like activity shown by $10Fe-MnO_x$ is illustrated in Scheme 1.

Evaluation of antioxidant properties

Given the strong oxidase-like activity of $10Fe-MnO_x$, it can be effectively employed to assess the oxidation resistance and, in particular, the ROS scavenging capabilities of various antioxidants. The antioxidant properties of $10Fe-MnO_x$ have been evaluated against various antioxidants including cysteine (Cys), guanine (Gua), dopamine (Dop), arginine (Arg), and glutathione (GSH) (Fig. S13a†). The addition of these antioxidants to ox-TMB resulted in varying degrees of color reaction suppression, confirming their antioxidant functionality. Cys and GSH exhibited the most significant color and spectral changes, likely due to the presence of their sulfhydryl ($-SH$) groups.^{50,51} The color fading trend followed the order $GSH > Cys > Dop > Gua > Arg$ (Fig. S13b†), indicating that Cys and GSH possess superior ROS scavenging abilities. Furthermore, increasing concentrations of GSH led to a more pronounced decrease in absorbance at 652 nm (Fig. 8a), further validating its antioxidant efficiency. These results demonstrate that the $10Fe-MnO_x$ -TMB colorimetric system provides a reliable method for antioxidant evaluation (Fig. 8b). Recyclability and reusability are key parameters to determine the reliability and practical utility of a catalytic system. In this effort, the recyclability test has been performed by recovering the catalyst *via* centrifugation after each sensing cycle. Notably, $10Fe-MnO_x$



Scheme 1 Possible mechanism for oxidase-like activity shown by $10Fe-MnO_x$. Schematic representation for the crystal structure of $10Fe-MnO_x$ is shown in the middle. The balls with black, cyan, red and grey colours represent Mn, Fe, O and oxygen vacancy (Ov), respectively.

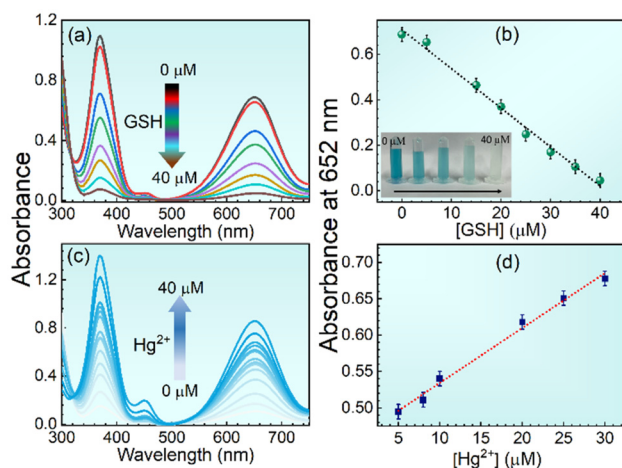


Fig. 8 (a) The absorption spectra of $10\text{Fe-MnO}_x/\text{TMB}$ system in the presence of GSH with different concentrations (0–40 μM); (b) calibration plot for the detection of GSH. Inset: photograph of $10\text{Fe-MnO}_x/\text{TMB}$ system with different concentrations of GSH; (c) the absorption spectra of $10\text{Fe-MnO}_x/\text{TMB}$ system in the presence of different concentrations of Hg^{2+} (0–40 μM) and (d) calibration plot for the detection of Hg^{2+} .

retained its catalytic activity over three cycles, making it a promising and reliable catalytic/sensing platform for further applications (Fig. S14[†]).

Colorimetric detection of Hg^{2+}

Mercury ions are recognized as highly toxic heavy metal pollutants, prevalent in the natural environment, and a primary contributor to water contamination.⁵² Hg^{2+} can be readily transformed into more hazardous organic mercury compounds through microbial activity, leading to bioaccumulation and biomagnification. The Hg^{2+} accumulation in the human body can trigger a range of chronic ailments, including Minamata disease, acrodynia, kidney dysfunction, cognitive impairment, and even damage to the central nervous system.^{53–55} Given these health risks, regulatory bodies have established stringent limits for Hg^{2+} in drinking water: the United States Environmental Protection Agency (EPA) sets a maximum contaminant level of $2.0 \mu\text{g mL}^{-1}$, while the World Health Organization (WHO) defines a limit of $6.0 \mu\text{g mL}^{-1}$. Consequently, the development of sensitive and selective analytical methods for monitoring trace levels of Hg^{2+} in aqueous environments and biological samples is of paramount importance.

As previously discussed, 10Fe-MnO_x demonstrates antioxidant properties in the presence of GSH, and the colorimetric signal diminishes significantly due to the ROS scavenging ability of the $-\text{SH}$ group. Under optimized conditions, increasing the concentration of GSH leads to a decrease in absorption intensity at 652 nm (Fig. 8a). Conversely, the addition of varying concentrations of Hg^{2+} to the $10\text{Fe-MnO}_x/\text{TMB-GSH}$ system inhibits this decrease in absorbance (Fig. 8c). Therefore, we have employed this system for colorimetric detection of Hg^{2+} (0–40 μM). Notably, $10\text{Fe-MnO}_x/$

TMB-GSH was found to be highly selective for Hg^{2+} , with a limit of detection (LOD) determined to be $0.47 \mu\text{M}$ (Fig. 8d). The colorimetric Hg^{2+} sensing can be attributed to its strong binding capability with GSH, ultimately resulting in a concomitant deactivation of antioxidant function. As a result, ROS accumulation restores the colorimetric signal of ox-TMB at 652 nm. The interference study was also conducted in the presence of various competing metal ions such as Ag^+ , Cd^{2+} , K^+ , Na^+ , Mg^{2+} , Ca^{2+} , Zn^{2+} , Cu^{2+} , Co^{2+} , Fe^{2+} , and Pb^{2+} (Fig. S15[†]), which clearly showed the exceptional selectivity of $10\text{Fe-MnO}_x/\text{TMB-GSH}$ towards Hg^{2+} .

To further validate the practical applicability of the developed 10Fe-MnO_x system, the colorimetric detection of Hg^{2+} was performed in real water samples spiked with varying concentrations of Hg^{2+} . This approach results in excellent recovery rates in the range of 97–105% (Table S3[†]). These findings confirm the reliability of the developed colorimetric assay for Hg^{2+} detection in real water samples.

Colorimetric detection of hydroquinone

Hydroquinone, also known as 1,4-dihydroxybenzene, is widely used in various industrial sectors, including cosmetics, dyes, and pesticides. However, its inherent toxicity and limited biodegradability pose significant environmental concerns.^{56–58} Excessive exposure to HQ can lead to a range of adverse health effects, such as fatigue, tachycardia, liver dysfunction, and even kidney damage.⁵⁹ Therefore, developing accurate, sensitive, rapid, and cost-effective methods for HQ detection is crucial for environmental monitoring and public health protection. Various analytical techniques, including high-performance liquid chromatography (HPLC),⁶⁰ capillary electrophoresis,⁶¹ fluorescence spectroscopy,⁶² and electrochemical methods,⁶³ have been well employed for detecting and quantifying HQ. While these techniques offer good sensitivity and specificity, they often suffer from limitations such as time-consuming procedures, reliance on expensive and complex instrumentation, or the need for electrode modification. In contrast, colorimetric methods offer a promising alternative for on-site detection due to their simplicity, speed, and low cost, making them particularly suitable for naked-eye analysis.⁶⁴ The selective detection of HQ using a simple colorimetric approach is a meaningful but challenging task in environmental monitoring, with the key requirement being the identification of a suitable chromogenic probe capable of inducing a distinct color change in the presence of HQ.

Here, we employed the $10\text{Fe-MnO}_x/\text{TMB}$ colorimetric sensing platform for HQ detection. Under optimized conditions, varying concentrations of HQ (0–50 μM) were introduced into the ox-TMB system. As shown in Fig. 9b, the absorbance peak corresponding to ox-TMB at 652 nm decreased with increasing HQ concentration. This enabled the detection of HQ at concentrations as low as $1.74 \mu\text{M}$, with a correlation coefficient (R^2) of 0.99 (Fig. 9c). The underlying mechanism involves the transfer of two electrons from HQ to ox-TMB due to the strong reducing power of HQ. This process leads to the conversion of HQ into its *p*-benzoquinone tautomer, while the

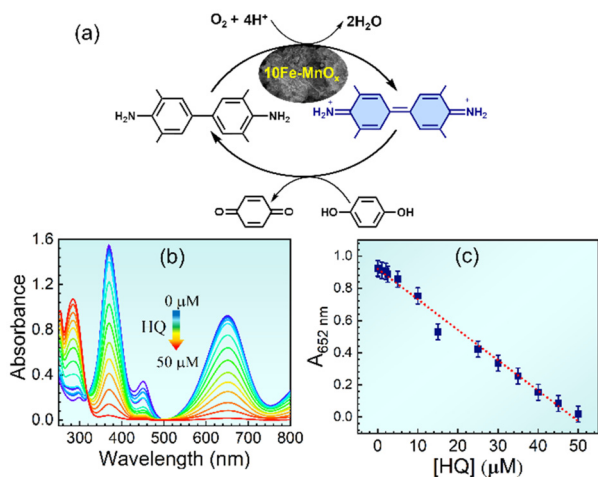


Fig. 9 (a) Schematic for TMB oxidation and subsequent detection of HQ; (b) UV-Vis spectra of the **10Fe-MnO_x**/TMB system in the presence of various concentrations of HQ (0–50 μM) and (c) calibration curve for the detection of HQ using **10Fe-MnO_x**/TMB.

blue ox-TMB is reduced to colorless TMB, as illustrated in Fig. 9a.

To assess the selectivity of the **10Fe-MnO_x**/TMB colorimetric system for HQ detection, we tested its response to a range of common interfering species, including Na⁺, K⁺, Ca²⁺, Mg²⁺, Cu²⁺, Co²⁺, Zn²⁺, Fe³⁺, Pb²⁺, Pd²⁺, Ni²⁺, Al³⁺, phenol, oxalic acid and ethanol. As shown in Fig. S16,[†] these interferents induced only weak absorbance changes at 652 nm compared to the significant decrease observed in the presence of HQ. These results clearly indicate that the **10Fe-MnO_x**/TMB colorimetric system exhibits high selectivity for HQ detection.

To further demonstrate the practical utility of the probe in real samples, water samples from different sources were collected and pre-treated. The resulting samples were spiked with different concentrations of HQ. The resultant data are summarized in Table S4.[†] The **10Fe-MnO_x** based colorimetric method demonstrated accuracy for determining HQ in real samples, with recoveries between 96% and 106%.

Stability and scalability tests of **10Fe-MnO_x**

The commercial viability of the **10Fe-MnO_x** heterogeneous catalyst is significantly dependent on its stability and scalability. After being stored for one month, the catalyst retained its efficiency, demonstrating good environmental tolerance and surpassing the performance of natural enzymes (Fig. S17a[†]). Furthermore, bulk preparation of the catalyst did not diminish its high efficiency in the oxidation of TMB, highlighting its strong potential for commercial applications (Fig. S17b[†]).

Conclusions

In summary, we successfully synthesized a pure phase of Fe-doped δ-MnO₂ nanoflowers *via* straightforward chemical oxi-

dation of Fe-doped MnBTC MOF. Systematic Fe-doping into the MOF resulted in phase evolution and structural as well as morphological modulation, which significantly influenced oxidase-like activity. Among the synthesized materials, 10% Fe-doped MnO_x exhibited the highest oxidase-like activity, which could be attributed to the synergistic effect of phase structure, oxygen vacancies, and nanoflower morphology. Under the optimized conditions, **10Fe-MnO_x** has been effectively utilized for the colorimetric detection of Hg²⁺ and hydroquinone in aqueous media. The as-developed assay demonstrated remarkable sensitivity, with detection limits of 0.47 μM for Hg²⁺ and 1.74 μM for HQ, outperforming conventional nanozymes. Furthermore, mechanistic insights into the catalytic pathways responsible for oxidation reactions were investigated using steady-state kinetics, scavenging study and EPR spectroscopy.

Data availability

The data will be made available on request.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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