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## Holistic functional biomimetics: a key to make an efficient electrocatalyst for water oxidation†

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Water oxidation is the holy grail reaction of natural and artificial photosynthesis. How to design an efficient water-oxidation catalyst remains a long-term challenge for solar fuel production. The rate of water oxidation in photosystem II by the oxygen-evolving complex (OEC)  $\text{Mn}_4\text{CaO}_5$  cluster is as high as  $100\text{--}400\text{ s}^{-1}$ . Mimicking the structures of the OEC is a straightforward strategy to design water-oxidation catalysts. However, the high efficiency of the OEC relies on not only its highly active site but also its holistic system for well-organized electron transfer and proton transport. Lacking such a holistic functional system makes  $\delta\text{-MnO}_2$  a poor water-oxidation catalyst, although the local structure of  $\delta\text{-MnO}_2$  is similar to that of the  $\text{Mn}_4\text{CaO}_5$  cluster. Electrocatalysts simultaneously imitating the catalytically active sites, fast electron transfer, and promoted proton transport in a natural OEC have been rarely reported. The significance of the synergy of a holistic system is underrated in the design of water-oxidation catalysts. In this work, we fabricated holistic functional biomimetic composites of two-dimensional manganese oxide nanosheets and pyridyl-modified graphene ( $\text{MnO}_x\text{-NS/py-G}$ ) for electrocatalytic water oxidation.  $\text{MnO}_x\text{-NS/py-G}$  simultaneously imitates the synergy of catalytically active sites, fast electron transfer, and promoted proton transport in a natural OEC, resulting in overall 600 times higher activity than that of typical  $\delta\text{-MnO}_2$ . This work demonstrates the significance of holistic functional biomimetic design and guides the development of highly active electrocatalysts for small molecule activation related to solar energy storage.

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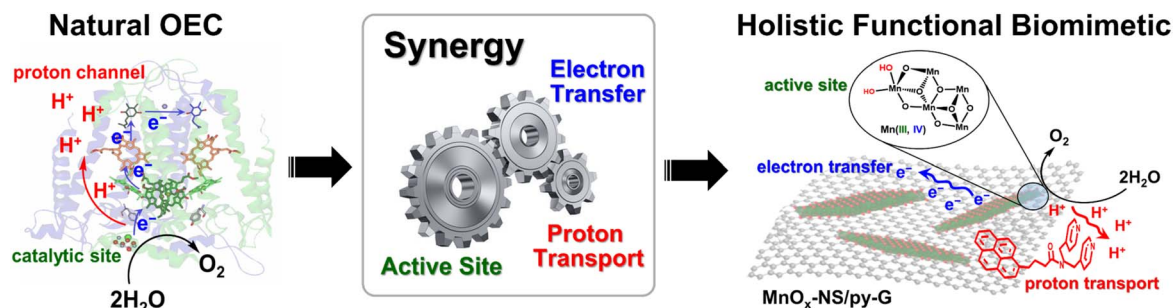
## Introduction

The development of efficient water-oxidation catalysts is of significance to realize high-performance artificial photosynthesis for solar energy conversion and fuel production.<sup>1,2</sup> In photosystem II (PSII), water is oxidized by the oxygen-evolving complex (OEC), a  $\text{Mn}_4\text{CaO}_5$  cluster surrounded by amino acid residues, two  $\text{Cl}^-$  ions, and many  $\text{H}_2\text{O}$  molecules.<sup>3–5</sup> The  $\text{Mn}_4\text{CaO}_5$  cluster, which can be regarded as a nanosized Mn oxide in a protein environment, catalyzes water oxidation with a low overpotential of around 160 mV and a high rate of  $100\text{--}400\text{ s}^{-1}$ .<sup>6</sup> Mimicking the structure of the OEC in PSII is

a straightforward strategy to design water-oxidation catalysts.<sup>2,7</sup> A successful example is the introduction of carboxylic groups into molecular catalysts, leading to the breakthrough of water oxidation performance.<sup>8</sup> However, the structural similarity to the natural OEC structure is not always related to the catalytic performance.  $\delta\text{-MnO}_2$  possess similar structural properties to the  $\text{Mn}_4\text{CaO}_5$  cluster in PSII, *e.g.*, the cabane-like local structure, and the mixing of  $\text{Mn}^{3+}$  and  $\text{Mn}^{4+}$  oxidation states, therefore, can be considered structural models for the OEC.<sup>9–11</sup> But the performance of typical  $\delta\text{-MnO}_2$  for water oxidation is poor, far less than that of natural enzymes.

The high efficiency of the OEC relies on its well-organized holistic system, which includes not only highly active sites for the formation of reactive high-valent metal-oxo and O–O bonds but also effective electron transfer pathways *via* nearby tyrosine residues<sup>12</sup> and proton transport channels *via* the  $\text{Ca}^{2+}$  ion,<sup>13</sup> namely the synergy of the catalytically active site, electron transfer, and proton transport (Scheme 1). However,  $\delta\text{-MnO}_2$  only imitates the core structure of the  $\text{Mn}_4\text{CaO}_5$  cluster but cannot imitate the OEC's secondary environment.<sup>9</sup> Although its local structural unit is close to that of the  $\text{Mn}_4\text{CaO}_5$  cluster, the water-oxidation performance of  $\delta\text{-MnO}_2$  is heavily limited due to the lacking exposure of catalytic sites and the components that facilitate electron transfer and proton transport.<sup>10</sup> In

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**Scheme 1** Concept of holistic functional biomimetics inspired by the natural OEC's synergy of the catalytically active site, electron transfer, and proton transport.

addition to the structure of the active site, the inspiration that the design of water-oxidation catalysts can take from a natural OEC is the synergy of its holistic system.<sup>11</sup> To date, the significance of the synergy of a holistic system is underrated in the design of water-oxidation catalysts. Electrocatalysts simultaneously imitating the catalytically active sites, fast electron transfer, and promoted proton transport in a natural OEC have been rarely reported.

In this work, we report holistic functional biomimetic inorganic–organic composites with clearly defined structure–function relationships to display the significance of the synergy of the catalytically active site, electron transfer, and proton transport in the design of water-oxidation electrocatalysts (Scheme 1). Coordinatively unsaturated edge, corner, and defect sites on two-dimensional (2D) manganese oxide nanosheets ( $\text{MnO}_x\text{-NS}$ ), which can be obtained from structural exfoliation of  $\delta\text{-MnO}_2$ , serve as biomimetic active sites. To achieve fast electron transfer between local active sites, material bulk, and the electrode interface,  $\text{MnO}_x\text{-NS}$  was assembled with graphene to form a nanocomposite structure. Proton transport is essential for the formation of water-oxidation key intermediates, such as  $\text{Mn}^{\text{VII}}\text{-oxo}$ ,<sup>14</sup>  $\text{Mn}^{\text{IV}}\text{O}$ ,<sup>15</sup> and  $\text{Mn}^{\text{V}}\text{-oxo}$ .<sup>16</sup> Inspired by Nakamura and co-

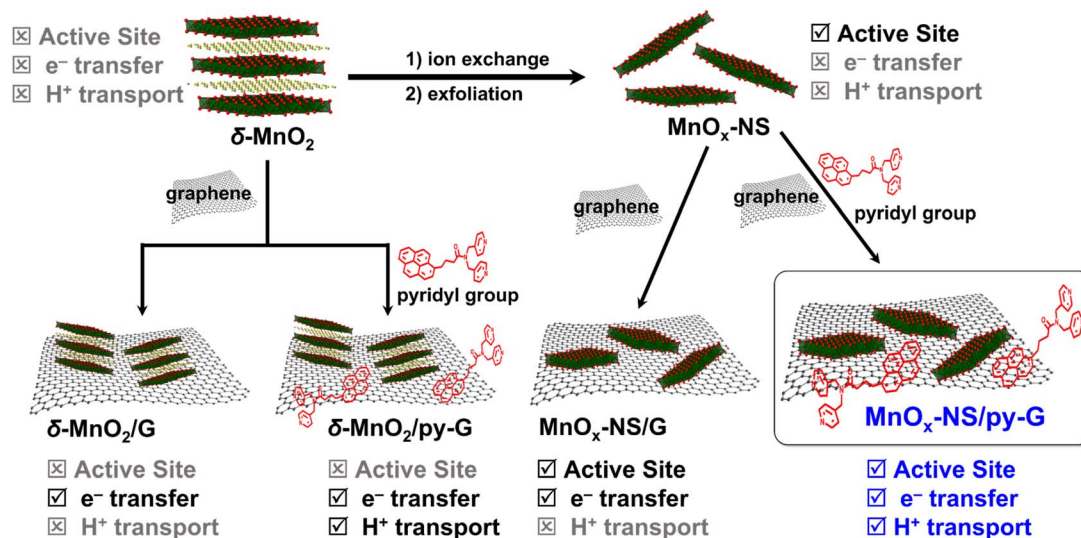
workers' study,<sup>17</sup> we further modified the graphene surface with pyridyl groups with a  $\text{pK}_a$  value of around 5 to accelerate proton transfer.

The fabricated holistic functional biomimetic composites of  $\text{MnO}_x\text{-NS}$  and pyridyl-modified graphene ( $\text{MnO}_x\text{-NS/py-G}$ ) showed 600-fold higher catalytic activity than  $\delta\text{-MnO}_2$ , being ranked on top in highly active manganese-based water-oxidation catalysts. Further in-depth studies revealed the structure–activity relationship of the  $\text{MnO}_x\text{-NS/py-G}$  composites. The activity improvement results from the synergy of sufficient exposure of catalytically active sites, fast electron transport, and efficient proton transport. Each of the factors contributes 30 times, 10 times, and 2 times to the overall activity increment, respectively. These results proved the effectiveness of the holistic functional biomimetic for electrocatalyst design.

## Results and discussion

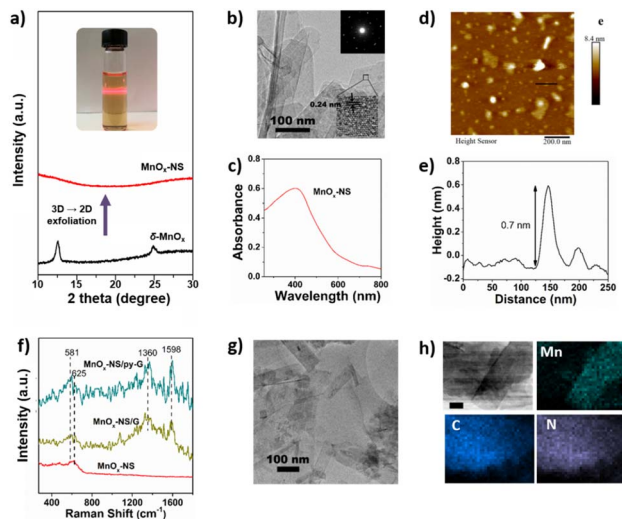
### Preparation and characterization

General synthetic procedures of  $\text{MnO}_x\text{-NS}$ ,  $\text{MnO}_x\text{-NS/G}$ , and  $\text{MnO}_x\text{-NS/py-G}$  are demonstrated in Scheme 2. Control samples  $\delta\text{-MnO}_2/\text{G}$  and  $\delta\text{-MnO}_2/\text{py-G}$  were also prepared by similar



**Scheme 2** Schematic illustration of the synthetic procedure of  $\text{MnO}_x\text{-NS/G}$ ,  $\text{MnO}_x\text{-NS/py-G}$ ,  $\delta\text{-MnO}_2/\text{G}$ , and  $\delta\text{-MnO}_2/\text{py-G}$ .





**Fig. 1** (a) Optical image of the colloidal solution of exfoliated  $\text{MnO}_x$ -NS and XRD pattern of  $\delta\text{-MnO}_2$  and  $\text{MnO}_x$ -NS. (b) The TEM image of  $\text{MnO}_x$ -NS. (c) The UV-vis absorption spectra of the colloidal suspension of  $\text{MnO}_x$ -NS. (d) The AFM image and (e) the height profile of  $\text{MnO}_x$ -NS. (f) The Raman spectrum of  $\text{MnO}_x$ -NS,  $\text{MnO}_x$ -NS/G, and  $\text{MnO}_x$ -NS/py-G. (g) The TEM image of  $\text{MnO}_x$ -NS/py-G. (h) STEM-EDX elemental mapping of Mn, C, and N of  $\text{MnO}_x$ -NS/py-G (scale bar is 20 nm).

methods. The graphene used is commercial, and the pyridyl molecule for proton transport is from laboratory synthesis (Fig. S1 and S2†). A colloidal solution of 2D  $\text{MnO}_x$ -NS was first obtained from the exfoliation of  $\delta\text{-MnO}_2$  by an ion-exchange approach.<sup>18–20</sup> A typical Tyndall effect is observed in the brown and transparent solution under laser beam illumination, demonstrating its good dispersion and colloidal properties (inset of Fig. 1a). Fig. 1a shows the powder X-ray diffraction (XRD) spectrum of  $\delta\text{-MnO}_2$  and  $\text{MnO}_x$ -NS. In contrast to the XRD pattern of  $\delta\text{-MnO}_2$  with typical sharp peaks at  $2\theta = 12.4^\circ$ , which is related to the layered structure with an interlayer spacing of 0.7 nm, the XRD spectrum of  $\text{MnO}_x$ -NS presents no diffraction peak. The disappearance of the peak at  $12.4^\circ$  illustrates the delamination of the long-range ordered layered structure to individual irregularly oriented nanosheets after exfoliation.<sup>18,20</sup>

$\text{MnO}_x$ -NS was characterized in detail by transmission electron microscopy (TEM), ultraviolet-visible (UV-vis) spectroscopy, and atomic force microscopy (AFM). The TEM image of  $\text{MnO}_x$ -NS displays a clear single-layer nanosheet morphology (Fig. 1b). The selected area electron diffraction (SAED) pattern of  $\text{MnO}_x$ -NS exhibits hexagonally arranged diffraction spots (inset of Fig. 1b). All these results are in sharp contrast to  $\delta\text{-MnO}_2$ , which exhibits a multi-layered morphology and polycrystal SAED image with diffraction rings, showing the exfoliation of a multi-layered structure to monolayer nanosheets (Fig. S3†).<sup>20</sup> Notably, the SAED results of  $\delta\text{-MnO}_2$  and  $\text{MnO}_x$ -NS present hexagonal reciprocal lattices, indicating that  $\text{MnO}_x$ -NS retains a similar crystalline structure to the  $\delta\text{-MnO}_2$  crystal. The HRTEM image of  $\text{MnO}_x$ -NS presents a lattice fringe with an interlayer spacing of 0.24 nm, which can be assigned to the  $\{100\}$  plane of  $\delta\text{-MnO}_2$ .

$\text{MnO}_2$ .<sup>21,22</sup> The UV-vis spectrum of  $\text{MnO}_x$ -NS shows a broad absorption peak at around 400 nm, which can be attributed to the d–d transition of Mn ions in  $\text{MnO}_6$  octahedra (Fig. 1c).<sup>18,20</sup> At last, the successful exfoliation of  $\delta\text{-MnO}_2$  to 2D  $\text{MnO}_x$  monolayer nanosheets was confirmed by AFM imaging of  $\text{MnO}_x$ -NS (Fig. 1d and e). A thickness of 0.7 nm was determined for  $\text{MnO}_x$ -NS, which is consistent with the thickness of the  $\text{MnO}_x$  monolayer.<sup>18,20</sup>

An apparent change of the electronic structure in  $\text{MnO}_x$ -NS compared to  $\delta\text{-MnO}_2$  is the increased ratio of  $\text{Mn}^{3+}/\text{Mn}^{4+}$  oxidation states, which was determined by X-ray photoelectron spectroscopy (XPS).  $\delta\text{-MnO}_2$  shows Mn peaks at 643.0, 642.1 eV, and 640.7 eV, which can be assigned to  $\text{Mn}^{4+}$ ,  $\text{Mn}^{3+}$ , and a minimal amount of  $\text{Mn}^{2+}$ , respectively (Fig. S4a†).<sup>23–26</sup> Based on the peak integration, its  $\text{Mn}^{3+}/\text{Mn}^{4+}$  content ratio was calculated to be 0.44. Similar Mn 2p XPS peaks were presented for  $\text{MnO}_x$ -NS. However, the ratio of  $\text{Mn}^{3+}/\text{Mn}^{4+}$  oxidation states increased to 0.67, much larger than the 0.44 of  $\delta\text{-MnO}_2$ . Moreover, the ratio of Mn–OH (531.0 eV)/Mn–O–Mn (529.7 eV) is calculated to be 0.46 for  $\text{MnO}_x$ -NS, much higher than 0.34 for  $\delta\text{-MnO}_2$  (Fig. S4b†).<sup>23–25</sup> In general, Mn–OH is a marker of active sites since the Mn ions with several coordinating –OH groups in the  $\text{Mn}_4\text{CaO}_5$  cluster, usually labeled as  $\text{Mn}_4$ , are considered to be the active sites for O–O bond formation.<sup>27</sup> These XPS results demonstrate that abundant  $\text{Mn}^{3+}$  species and coordinatively unsaturated sites are exposed on  $\text{MnO}_x$ -NS upon 3D  $\rightarrow$  2D structural exfoliation, serving as biomimetic active sites.

Then, single-layer  $\text{MnO}_x$ -NS were assembled with graphene and pyridyl-modified graphene (py-G) to give the  $\text{MnO}_x$ -NS/G and  $\text{MnO}_x$ -NS/py-G samples. The Raman spectra of  $\text{MnO}_x$ -NS/G and  $\text{MnO}_x$ -NS/py-G are presented in Fig. 1f. Both the samples show additional scattering bands at 1360 and 1598  $\text{cm}^{-1}$  compared with  $\text{MnO}_x$ -NS, which are the typical Raman signals of graphene.<sup>28,29</sup> The TEM images of  $\text{MnO}_x$ -NS/G and  $\text{MnO}_x$ -NS/py-G (Fig. 1g and S5†) and the energy dispersive spectroscopy (EDS) mapping (Fig. 1h) of  $\text{MnO}_x$ -NS/py-G display randomly distributed  $\text{MnO}_x$ -NS on the surface of graphene without apparent aggregation. A nitrogen fluorescence signal was presented in the EDS mapping of  $\text{MnO}_x$ -NS/py-G, confirming the homogeneous modification by the pyridyl molecule. The presence of the pyridyl molecule in  $\text{MnO}_x$ -NS/py-G can be further depicted by the electrochemical cyclic voltammetry (CV) study. A broad reduction wave was distinctly presented between  $-1.0$  and  $-1.35$  V (vs. Ag/AgCl) for  $\text{MnO}_x$ -NS/py-G, which is the typical redox peak of the pyridine moiety (Fig. S6†).<sup>30,31</sup> Overall, these above characterization studies thoroughly demonstrate the successful assembly of  $\text{MnO}_x$ -NS/py-G composites.

### Electrocatalytic water oxidation

Electrocatalytic oxygen evolution reaction (OER) performances of  $\delta\text{-MnO}_2$ ,  $\text{MnO}_x$ -NS,  $\text{MnO}_x$ -NS/G, and  $\text{MnO}_x$ -NS/py-G were evaluated by loading on glass carbon (GC) electrodes under alkaline conditions (1 M KOH).  $\delta\text{-MnO}_2$  shows negligible activity within a wide potential range, which is consistent with the generally low OER activity characteristics of  $\delta\text{-MnO}_2$  (Fig. 2a).<sup>32–34</sup>  $\text{MnO}_x$ -NS displays significantly enhanced OER





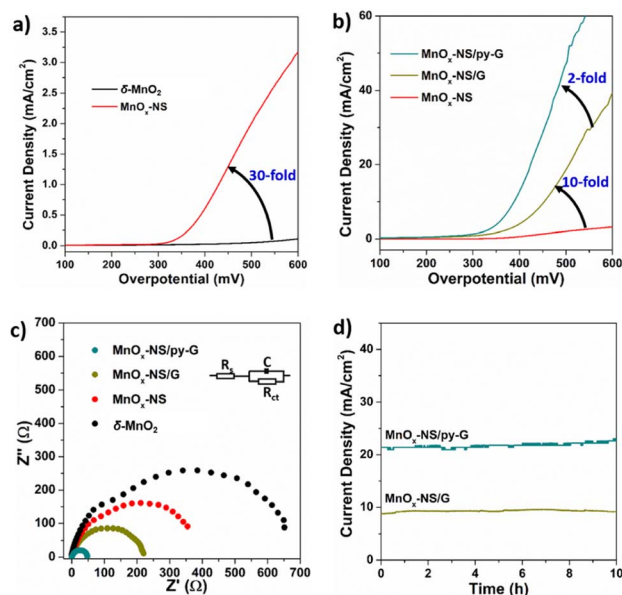


Fig. 2 (a) Polarization curves of  $\delta$ -MnO<sub>2</sub> and MnO<sub>x</sub>-NS. (b) Polarization curves of MnO<sub>x</sub>-NS, MnO<sub>x</sub>-NS/G, and MnO<sub>x</sub>-NS/py-G. (c) Nyquist diagrams of  $\delta$ -MnO<sub>2</sub>, MnO<sub>x</sub>-NS, MnO<sub>x</sub>-NS/G, and MnO<sub>x</sub>-NS/py-G. (d) Chronoamperometry curve of MnO<sub>x</sub>-NS/G and MnO<sub>x</sub>-NS/py-G at a 450 mV overpotential.

activity, which provides approximately 30 times higher current densities compared with  $\delta$ -MnO<sub>2</sub> (Fig. 2a). Notably, after assembling with graphene, the catalytic current density of MnO<sub>x</sub>-NS/G increases further by 10-fold compared to that of MnO<sub>x</sub>-NS (Fig. 2b). Then, with the effect of the pyridyl molecule modification, 2 times improvement in OER performance was additionally achieved for the final MnO<sub>x</sub>-NS/py-G catalyst compared to that of MnO<sub>x</sub>-NS/G. In contrast to the initial  $\delta$ -MnO<sub>2</sub>, MnO<sub>x</sub>-NS/py-G eventually displays 600 times higher OER activity due to the synergy of active site exposure, nano-composite structure, and molecular modification, being ranked as a highly active manganese-based water oxidation catalyst (Fig. S7†).

The electrochemical impedance spectroscopy (EIS) measurements under water oxidation conditions can further reflect the improved catalytic rate. MnO<sub>x</sub>-NS/py-G exhibits a charge transfer resistance ( $R_{ct}$ ) of 41.09  $\Omega$ , which is 20 times lower compared with 584.8  $\Omega$  of  $\delta$ -MnO<sub>2</sub> (Fig. 2c). A low Tafel slope of 86 mV dec<sup>-1</sup> was determined for MnO<sub>x</sub>-NS/py-G on the MnO<sub>x</sub>-NS/py-G catalyst to assess the catalytic stability (Fig. S8†). A stable current density of 21 mA cm<sup>-2</sup> was maintained during 10 h electrolysis at a 450 mV overpotential, indicating the excellent electrocatalytic durability of MnO<sub>x</sub>-NS/py-G (Fig. 2d). However, due to the bubble removal, mass transfer, and mechanical stability problems, long-term electrolytic assessment at a large current density (e.g. 200 mA cm<sup>-2</sup>) cannot be carried out using a GC working electrode in this study.<sup>42</sup> For a powder catalyst, a flow electrochemical cell and anion exchange membrane electrolyzer are better and important evaluation methods to achieve the performance of the catalyst in practical application.<sup>42,43</sup>

TEM, EDS-mapping, and electrochemical cyclic voltammetry of the MnO<sub>x</sub>-NS/py-G catalyst after reaction shows comparable results with the as-synthesized catalyst, which confirm its good catalytic stability (Fig. S9 and S10†). In a separate experiment, the OER faradaic efficiency was determined to be 98% for MnO<sub>x</sub>-NS/py-G under 400 mV overpotential electrolysis, verifying that a vast majority of charges was contributed to water oxidation (Fig. S11†).

Following our design strategy in Scheme 1, the OER activities of the  $\delta$ -MnO<sub>2</sub>, MnO<sub>x</sub>-NS, MnO<sub>x</sub>-NS/G, and MnO<sub>x</sub>-NS/py-G catalysts have been successively advanced owing to the directed improvement in each sample, achieving a remarkable OER performance for the final MnO<sub>x</sub>-NS/py-G. To understand the activity enhancement from the starting inactive  $\delta$ -MnO<sub>2</sub> to the final highly active MnO<sub>x</sub>-NS/py-G, we then thoroughly examined the specific effects of structural exfoliation, graphene substrate assembly, and pyridyl molecule modification on electrocatalytic water oxidation.

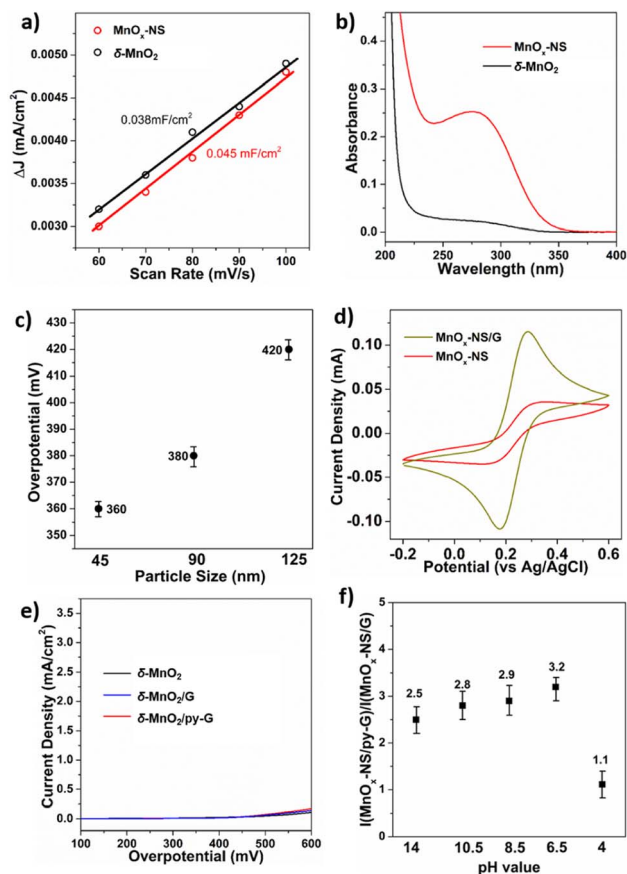
### Effects of the exposure of the active site

Compared to  $\delta$ -MnO<sub>2</sub>, the activity of MnO<sub>x</sub>-NS increased by 30 times, indicating the significant effects of the structural exfoliation. Two possible ways may induce this dramatic improvement: (i) enlarged electrochemically active surface area (ECSA) caused by a morphology change; (ii) increased numbers of catalytic sites due to 3D  $\rightarrow$  2D structural changes. To identify the effects of structural exfoliation, we first estimated the ECSA of  $\delta$ -MnO<sub>2</sub> and MnO<sub>x</sub>-NS from electrochemical double-layer capacitance ( $C_{dl}$ , Fig. 3a and S12†). The linear slope of MnO<sub>x</sub>-NS was calculated to be 0.045 mF cm<sup>-2</sup>, which is only 18% higher than the 0.038 mF cm<sup>-2</sup> of  $\delta$ -MnO<sub>2</sub>. The similar ECSA of  $\delta$ -MnO<sub>2</sub> and MnO<sub>x</sub>-NS indicates that the layered structure of  $\delta$ -MnO<sub>2</sub> is already widely accessible for electrochemical charging, which has also been reported on double layer hydroxide materials.<sup>44</sup> This slight increase in ECSA contributes negligibly to the 30 times increase in activity upon exfoliation of  $\delta$ -MnO<sub>2</sub> to MnO<sub>x</sub>-NS.

Since the change of ECSA does not account for activity improvement, the remaining possibility is the increment of catalytic sites. Mn<sup>3+</sup> species have been commonly recognized as an indicator of active sites for Mn-based water oxidation catalysts.<sup>45,46</sup> To investigate the promotion effect of exfoliation on active site exposure, we introduce a pyrophosphate (PP) molecule to the solution of  $\delta$ -MnO<sub>2</sub> and MnO<sub>x</sub>-NS, which can extract Mn<sup>3+</sup> from the catalyst through coordination (Fig. 3b).<sup>46–48</sup> The resulting Mn<sup>3+</sup>-PP species was then monitored by using the UV-vis absorption spectrum. MnO<sub>x</sub>-NS exhibits a much higher Mn<sup>3+</sup>-PP absorption peak at around 258 nm compared with  $\delta$ -MnO<sub>2</sub>. Consistent with XPS, these results demonstrate the abundance of Mn<sup>3+</sup> upon exfoliation. These Mn<sup>3+</sup> species can be positively involved in water oxidation as highly reactive sites, improving the catalytic activity.

With further investigations, we could propose two aspects that effectively increase catalytically active sites by exfoliation. On the one hand, complete structural exfoliation can spontaneously promote active site exposure at the surface of the





**Fig. 3** (a) The ECSA measurement:  $\Delta J$  ( $J_a - J_c$ ) from the CV curves of  $\delta$ -MnO<sub>2</sub> and MnO<sub>x</sub>-NS plotted against scan rates. (b) UV-vis absorption spectra of a 20 mM sodium pyrophosphate solution after dipping  $\delta$ -MnO<sub>2</sub> and MnO<sub>x</sub>-NS. (c) Comparison of the required overpotential for 0.5 mA cm<sup>-2</sup> current density on MnO<sub>x</sub>-NS with a particle size of 45, 90, and 125 nm. (d) CV curves of MnO<sub>x</sub>-NS and MnO<sub>x</sub>-NS/G in 0.1 M KCl electrolyte containing 10 mM Fe(CN)<sub>6</sub><sup>3-/4-</sup> ions. (e) LSV curves of  $\delta$ -MnO<sub>2</sub>,  $\delta$ -MnO<sub>2</sub>/G, and  $\delta$ -MnO<sub>2</sub>/py-G. (f) Comparison of  $I(\text{MnO}_x\text{-NS/py-G})/I(\text{MnO}_x\text{-NS/G})$  under pH 14, pH 10.5, pH 8.5, pH 6, and pH 4 conditions.

monolayer nanosheets. To verify this, Na<sup>+</sup> was introduced into the MnO<sub>x</sub>-NS colloidal suspension to promote the self-reassembly of MnO<sub>x</sub>-NS, regenerating layered  $\delta$ -MnO<sub>2</sub>.<sup>20</sup> The XRD pattern of the re-assembled sample presents a typical diffraction peak of a layered structure with an interlayer space of 0.73 nm, indicating the reformation of the layered  $\delta$ -MnO<sub>2</sub> (Fig. S13a†). The electrocatalytic performance of this reformed  $\delta$ -MnO<sub>2</sub> dramatically decreased compared with that of MnO<sub>x</sub>-NS (Fig. S13b†). The remarkable activity difference between MnO<sub>x</sub>-NS and  $\delta$ -MnO<sub>2</sub> and our previous active “c-disordered”  $\delta$ -MnO<sub>x</sub> demonstrates the importance of the exposure of the MnO<sub>2</sub> nanosheet for efficient electrocatalytic water oxidation. However, we cannot exactly explain why  $\delta$ -MnO<sub>2</sub> has the same charge transfer properties as MnO<sub>x</sub>-NS, *i.e.*, the same ECSA, but the ordered MnO<sub>2</sub> nanosheet has deficient catalytic performance for water oxidation.

On the other hand, based on the TEM images, the average particle size of MnO<sub>x</sub>-NS was estimated to be around 90 nm

(Fig. S14†), which is much smaller than the 240 nm of  $\delta$ -MnO<sub>2</sub>. Without a well-ordered layered structure, the MnO<sub>x</sub> monolayer will collapse and break, leading to a decrease in particle size. The formed smaller nanosheets promoted the exposure of coordinatively unsaturated edge- and corner-sites, which are believed to be involved in water oxidation as highly reactive catalytic sites.<sup>44,49</sup> To further verify this, MnO<sub>x</sub>-NS with different lateral sizes were separated by controlled centrifugation, and denoted as MnO<sub>x</sub>-45 and MnO<sub>x</sub>-125 (Fig. S15†). A close relationship is presented between the lateral size and OER activity: where MnO<sub>x</sub>-45 with a smaller particle size shows higher activity than MnO<sub>x</sub>-90; whereas MnO<sub>x</sub>-125 with a larger particle size displays lower activity compared with MnO<sub>x</sub>-90 (Fig. 3c and S16†). These results demonstrate OER activity enhancement with particle size decrement. By the exfoliation process, plenty of coordinated unsaturated edge sites were generated on MnO<sub>x</sub>-NS (Fig. S17†), which are much easier to be oxidized to high-valent key intermediate states for water oxidation.<sup>14,47,48,50–53</sup>

### Effects of fast electron transfer

MnO<sub>x</sub>-NS/G shows a 10 times higher activity compared with MnO<sub>x</sub>-NS. LSV and long-term electrolysis were used to show that bare graphene displays minor water oxidation. This indicates that the graphene substrate promotes the catalytic system rather than introducing more active sites (Fig. S18†). Graphene serves as a loading substrate to promote the uniform distribution of MnO<sub>x</sub>-NS with less stacking, as displayed in the XRD and TEM images of MnO<sub>x</sub>-NS/G and MnO<sub>x</sub>-NS/py-G (Fig. 1g and S19†). More importantly, the graphene substrate with excellent charge transportation capacity can facilitate electron transfer during the OER process.<sup>54,55</sup> To probe the enhancement in electron transfer by graphene, CV of K<sub>4</sub>Fe(CN)<sub>6</sub> was performed using a MnO<sub>x</sub>-NS and MnO<sub>x</sub>-NS/G covered GC electrode as the working electrode. In contrast to the broad redox peak of Fe<sup>2+</sup>/Fe<sup>3+</sup> for MnO<sub>x</sub>-NS/GC, a sharp Fe<sup>2+</sup>/Fe<sup>3+</sup> peak was observed in the CV curve measured with MnO<sub>x</sub>-NS/G/GC (Fig. 3d). The promoted Fe<sup>2+</sup>/Fe<sup>3+</sup> redox couple on MnO<sub>x</sub>-NS/G/GC illustrates that the combination of MnO<sub>x</sub>-NS with graphene effectively solves the slow charge transfer problems. The lowered Tafel slopes can also demonstrate the acceleration of electron transfer after graphene introduction (Fig. S8†). The Tafel slopes were calculated to be 89 mV dec<sup>-1</sup> and 86 mV dec<sup>-1</sup> (approaching 60 mV dec<sup>-1</sup>) for MnO<sub>x</sub>-NS/G and MnO<sub>x</sub>-NS/py-G, which are much smaller than the 134 mV dec<sup>-1</sup> (~120 mV dec<sup>-1</sup>) of pristine MnO<sub>x</sub>-NS, indicating that the rate-determining step has been changed from the first electron transfer to the chemical reaction that takes place after the one-electron transfer,<sup>48,56,57</sup> demonstrating the improvement in catalytic kinetics.<sup>44,47,58</sup>

The synergy of active site exposure and fast electron transfer can be displayed by the catalytic performance of  $\delta$ -MnO<sub>2</sub>/G and  $\delta$ -MnO<sub>2</sub>/py-G. No noticeable activity enhancement was observed when  $\delta$ -MnO<sub>2</sub> was assembled with graphene or py-graphene, indicating that the promoting effect of graphene requires optimal assembly between graphene and an exfoliated 2D nanosheet structure (Fig. 3e). These results demonstrate that the further 10-fold enhancement of the OER activity of MnO<sub>x</sub>-



NS/G compared with that of  $\text{MnO}_x\text{-NS}$  arises from the synergy of 3D  $\rightarrow$  2D structural exfoliation and fast electron transfer of graphene.

### Effects of molecular pyridyl modification

It has been reported that introducing pyridine into electrolytes can prominently improve the catalytic performance of  $\text{MnO}_x$  electrocatalysts<sup>17</sup> and  $\alpha\text{-Fe}_2\text{O}_3$  photoanodes.<sup>59</sup> In these systems, pyridine can work as a proton transfer relay, facilitating the proton-coupled electron transfer in the catalytic process. For  $\text{MnO}_x\text{-NS/py-G}$ , we directly modified the graphene surface, *i.e.*, the catalytic environment, with a pyridyl molecule to accelerate proton transfer during water oxidation. The  $\text{MnO}_x\text{-NS}$  loading on graphene afforded a two-fold increase in activity. A pH dependence of OER activity was established to evaluate the proton transfer properties in the catalytic process (Fig. S20†). Fig. 3f shows the OER activities of  $\text{MnO}_x\text{-NS/G}$  and  $\text{MnO}_x\text{-NS/py-G}$  in electrolytes at different pH values (pH ranging from 4 to 14). In contrast to the significantly enhanced performance under  $\text{pH} \geq 6.5$  conditions, the pyridine modification shows minor enhancements in the OER at pH 4. The absence of increased activity at pH 4 can be attributed to the inhibited proton transfer relay effect of the pyridine group, which possesses a  $\text{pK}_a$  value around 5 and is thus mainly present as its conjugate acid. The deuterium kinetic isotope effect (KIE) was evaluated to study the role of proton transfer in the rate determining step (RDS). The KIE values of  $\text{MnO}_x\text{-NS/py-G}$  and  $\text{MnO}_x\text{-NS/G}$  were determined to be 3.4 and 4.3, respectively, which exhibit primary deuterium kinetic isotope effects, indicating that the rate-determining step (RDS) during water oxidation involves proton transfer. The lower KIE value of  $\text{MnO}_x\text{-NS/py-G}$  compared with that of  $\text{MnO}_x\text{-NS/G}$  illustrates the facilitated proton transfer by the introduced pyridyl functionality (Fig. S21†). These pH-dependence and KIE studies demonstrate the proton transfer relay effect of pyridine modification.<sup>60,61</sup>

The above analyses elucidate that the origin of the dramatically enhanced catalytic activity of  $x\text{-NS/py-G}$  is a combined

result of 3D  $\rightarrow$  2D structural exfoliation, the graphene substrate, and pyridyl molecule modification. The 2D structure of  $\text{MnO}_x\text{-NS}$  achieved by exfoliation significantly increased the number of catalytically active sites in the  $\text{MnO}_x\text{-NS/py-G}$  catalyst. The nanocomposite structure between 2D  $\text{MnO}_x\text{-NS}$  and the graphene substrate provided a fast electron transfer pathway. And the pyridyl group modification accelerated proton transport during water oxidation. The synergy of catalytically active site exposure, fast electron transport, and efficient proton transport accounts for the increased activity by 600 times (Scheme 3), respectively contributing 30 times, 10 times, and 2 times to the overall activity enhancement (Fig. 2a and b). The clear revelation of the structure–activity relationship of the  $\text{MnO}_x\text{-NS/py-G}$  composites demonstrates the significance of the holistic functional biomimetic in electrocatalyst design, as the impact of each factor on the overall catalytic performance is multiplied.

## Conclusions

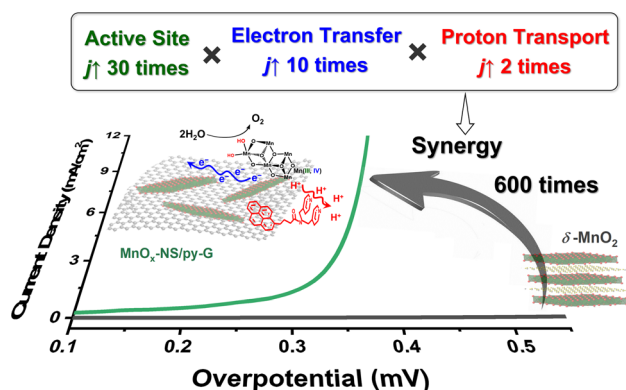
A natural OEC catalyzes water oxidation with a low overpotential and high turnover frequency because it is an integrated system that realizes the synergy of water molecule activation, O–O bond formation, electron transfer, and proton transport. The design of water oxidation electrocatalysts need to follow nature's course, not only simulating the catalytic site structure but also creating an overall synergistic system. Starting from typical  $\delta\text{-MnO}_2$ , we developed  $\text{MnO}_x\text{-NS/py-G}$  composites with a 600-fold activity improvement owing to the synergy of active site exposure, fast electron transport, and facilitated proton transport. Further studies indicate that the influence of each of the above factors on the overall catalytic performance is multiplied; therefore, outstanding activity can be achieved when the catalyst possesses efficient active sites, electron transfer, and proton transport simultaneously. In the development of water oxidation catalysts, in-depth analysis of factors leading to low catalytic activity can guide precise modification of the catalyst design for significantly improving the catalytic performance. This work demonstrates the success and significance of holistic functional biomimetics in electrocatalyst design, thereby providing a universal and *de novo* electrocatalyst design concept.

## Author contributions

L. S., B. Z. and L. F. conceived and designed the idea. B. Z. and L. F. designed the experiments. L. F. collected and analyzed the data. Y. S., B. J. J. T., and A. K. assisted with the experiments and characterization. F. Z. carried out the AFM measurements. L. F. and B. Z. discussed the results and wrote the manuscript. L. S. revised the manuscript. All the authors reviewed and contributed to this paper.

## Conflicts of interest

There are no conflicts to declare.



Scheme 3 Schematic illustration of the origin of the dramatic activity enhancement owing to the synergy of catalytically active site exposure, fast electron transport, and efficient proton transport.





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