

Cite this: *Chem. Sci.*, 2022, 13, 2385

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# A carbonyl-rich covalent organic framework as a high-performance cathode material for aqueous rechargeable zinc-ion batteries†

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Aqueous rechargeable zinc-ion batteries (ZIBs) provide high theoretical capacity, operational safety, low-cost and environmental friendliness for large-scale energy storage and wearable electronic devices, but their future development is plagued by low capacity and poor cycle life due to the lack of suitable cathode materials. In this work, a covalent organic framework (Tp-PTO-COF) with multiple carbonyl active sites is synthesized and successfully introduced in aqueous rechargeable ZIBs for the first time. Tp-PTO-COF delivers high specific capacities of 301.4 and 192.8 mA h g<sup>-1</sup> at current densities of 0.2 and 5 A g<sup>-1</sup>, respectively, along with long-term durability and flat charge–discharge plateaus. The remarkable electrochemical performance is attributed to the abundance of nucleophilic carbonyl active sites, well defined porous structure and inherent chemical stability of Tp-PTO-COF. Moreover, the structural evolution and Zn<sup>2+</sup> ion intercalation mechanism are discussed and revealed by the experimental analysis and density functional theory calculations. These results highlight a new avenue to develop organic cathode materials for high performance and sustainable aqueous rechargeable ZIBs.

Received 18th November 2021

Accepted 31st January 2022

DOI: 10.1039/d1sc06412f

rsc.li/chemical-science

## Introduction

To overcome the worldwide energy crisis, the utilization of green renewable energy sources such as solar, wind and tidal energy is highly desirable. Therefore, efficient energy storage devices play a crucial role in realizing the storage and release of renewable energy. Although widespread success has been achieved for rechargeable lithium-ion batteries (LIBs),<sup>1,2</sup> increasing concerns about safety issues, high cost and constrained lithium resources strongly limit their applications.<sup>3</sup> Compared to expensive and flammable LIBs, aqueous rechargeable zinc-ion batteries (ZIBs) are becoming an attractive alternative due to the natural abundance and high operational safety of zinc anodes.<sup>4–6</sup> Furthermore, aqueous rechargeable ZIBs can theoretically achieve higher specific capacity and energy density, because of Zn<sup>2+</sup> ions as multivalent charge carriers involved in

the redox reaction and aqueous electrolytes offering much higher ionic conductivities compared to organic electrolytes. Overall, these economic and technical superiorities of aqueous ZIBs hold particular promise for their application in large-scale energy storage systems and even in wearable electronic devices. Nevertheless, one of the primary obstacles hindering the development of aqueous rechargeable ZIBs is the lack of suitable cathode materials which can reversibly host Zn<sup>2+</sup> ions and maintain structural stability during the process of charge–discharge.<sup>7,8</sup> As a result, great efforts have been devoted to building high-performance ZIBs using inorganic compound cathodes such as manganese-based oxides,<sup>9–12</sup> vanadium-based oxides,<sup>13–15</sup> spinel-type materials<sup>16–18</sup> and Prussian blue analogues.<sup>19–21</sup> However, these inorganic electrodes were plagued by structural and chemical instabilities, unsatisfactory capacity, significant capacity fading, and toxic elements in their components. Recently, various organic materials have been recognized as the next promising green electrodes because they offer the advantages of light weight, low-toxicity, inexpensiveness and sustainability.<sup>22–25</sup> Organic compounds with carbonyl groups can store Zn<sup>2+</sup> ions by coordination reactions, therefore, some carbonyl compounds have been explored as cathode materials in aqueous rechargeable ZIBs, and exhibited high specific capacity and electrochemical reversibility.<sup>26–30</sup> Although these carbonyl compounds are almost insoluble in aqueous electrolytes, their discharge products always display high solubility, which inevitably limits their cycle stability and creates an urgent need for organic polymer electrode systems owing to the

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/d1sc06412f



stability and structural tunability of organic polymer. However, the simple polymerization of carbonyl compounds often leads to reduced capacity and poor rate performance.<sup>31</sup> Hence, both cycle stability and high capacity have to be further considered in the design of novel organic polymer cathode materials.

Covalent organic frameworks (COFs),<sup>32–36</sup> an emerging class of crystalline and porous polymeric materials, often possess high chemical stability, well defined porosity, and ability to predictably organize redox-active groups. The unique features of COFs endow them with great advantages and potential for application as electrode materials for rechargeable batteries.<sup>37–41</sup> Unlike traditional amorphous organic polymers, COFs can achieve accurate assembly of the redox-active building blocks, which makes it reasonable to obtain COF electrodes with specific structure and redox properties. Besides, COFs provide continuous open channels due to various topological structures or the stacking of conjugated polymeric layers. Because of this porosity advantage, the Zn<sup>2+</sup> ions are allowed to rapidly move in the COF electrode and reach the active sites. More importantly, the highly crystalline  $\pi$ -conjugated structure of COFs can provide a stable physical and chemical reaction environment, which is particularly beneficial to achieve stable operation at variable cathodic and anodic polarizations. However, the relevant research on COF-based aqueous rechargeable ZIBs is just in the beginning stage. Recently, Banerjee *et al.* proposed for the first time that a classical two-dimensional (2D) COF (HqTp) could be used as a cathode for aqueous rechargeable ZIBs.<sup>42</sup> Unfortunately, HqTp delivered limited specific capacities of 276.0 mA h g<sup>-1</sup> at 0.125 A g<sup>-1</sup>, and carbon nanofibers must be used as an electrical conductivity amplifier in the cathode. Furthermore, HqTp exhibited slopping charge–discharge curves without apparent plateaus, which was detrimental to achieving a stable voltage output. Although the state-of-the-art capacity performance of COF-based aqueous rechargeable ZIBs is far from expectation, these studies hinted at the promise of COF redox chemistry and structural stability.<sup>43</sup>

To achieve better electrochemical performance for COFs as cathodes of aqueous ZIBs, we report herein the synthesis and research of a 2D COF (Tp-PTO-COF) with dual active sites of adjacent carbonyl groups and  $\beta$ -keto carbonyl groups as nucleophilic centers to provide reversible and efficient Zn<sup>2+</sup> ion storage during the charge–discharge process.<sup>44</sup> Benefiting from the abundance of nucleophilic active sites, well-defined porous structure and inherent chemical stability, Tp-PTO-COF shows a remarkably high experimental capacity of 301.4 mA h g<sup>-1</sup> at a current density of 0.2 A g<sup>-1</sup>. Impressively, even at a high current density of 5 A g<sup>-1</sup>, Tp-PTO-COF displays an impressive reversible capacity of 192.8 mA h g<sup>-1</sup>. Moreover, the structural evolution and Zn<sup>2+</sup> ion intercalation mechanism in Tp-PTO-COF have been demonstrated by experimental analysis and density functional theory (DFT) calculations.

## Results and discussion

### Synthesis and characterization

Tp-PTO-COF was synthesized by condensing 1,3,5-triformylphloroglucinol (Tp) with 2,7-diaminopyrene-4,5,9,10-tetraone

(DAPTO) *via* an acid-catalyzed solvothermal reaction (Fig. 1). The successful condensation and structural characterization of Tp-PTO-COF were confirmed by both Fourier transform infrared spectroscopy (FT-IR) and solid state <sup>13</sup>C NMR spectroscopy. The FT-IR spectrum (Fig. S6†) exhibits two strong peaks at 1580 cm<sup>-1</sup> and 1272 cm<sup>-1</sup> for Tp-PTO-COF, which are typical characteristics of the C=C bond and C–N bond, indicating the formation of a  $\beta$ -ketoenamine-linked framework structure. The concomitant disappearing characteristic peaks of aldehydic C=O (1640 cm<sup>-1</sup>) and diamine N–H (3349, 3436 cm<sup>-1</sup>) verify the complete consumption of the reactants. In addition, the characteristic peak of the  $\beta$ -keto C=O stretch appears at 1616 cm<sup>-1</sup>, and the peak at 1672 cm<sup>-1</sup> of Tp-PTO-COF is attributed to the carbonyl groups (C=O) of 1,2-diketone units. Solid-state <sup>13</sup>C NMR studies also indicate the local structures of Tp-PTO-COF (Fig. S7†). The appearance of the representative signal for the enamine carbon atom at 146.8 ppm validates the successful formation of the  $\beta$ -ketoenamine-linked framework. Furthermore, two well-resolved peaks are detected at 182.7 and 176.6 ppm indicating the two different carbonyl carbon atoms in Tp-PTO-COF. The structural stability of Tp-PTO-COF was evaluated by thermogravimetric analysis (TGA), which reveals its high thermal stability up to 400 °C (Fig. S8†). Furthermore, the permanent porosity of Tp-PTO-COF was assessed by N<sub>2</sub> sorption-desorption measurements at 77 K. As indicated in Fig. S9,† the typical type-II sorption isotherms of Tp-PTO-COF reveal its mesoporous properties, and the surface area of Tp-PTO-COF was calculated to be 601 m<sup>2</sup> g<sup>-1</sup> on the basis of the Brunauer–Emmet and Teller (BET) model.

The crystalline structure feature of Tp-PTO-COF was analyzed by powder X-ray diffraction (PXRD) measurement in conjunction with structural simulation. As observed in Fig. 2a, Tp-PTO-COF exhibits well defined peaks, and the experimental pattern matches well with the simulated pattern of the eclipsed AA stacking model. The experimental PXRD pattern displays an apparent peak located at 3.4°, corresponding to the (100) reflection plane, and the hexagonal pore size of Tp-PTO-COF is calculated to be ~25.7 Å, which is in accordance with the simulated result (Fig. 2b). In addition, the presence of a broad peak around 27° can be ascribed to the  $\pi$ - $\pi$  stacking construction, revealing the existence of a multilayered COF structure. A full profile Pawley refinement was also performed for the experimentally obtained PXRD patterns based on the simulated AA stacking model, which has led to satisfactorily low

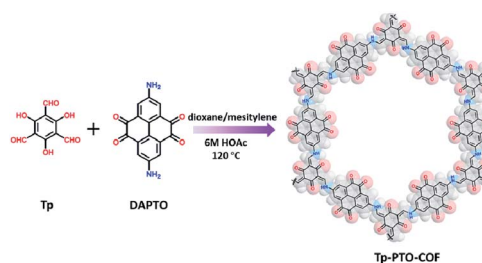


Fig. 1 Schematic presentation of the synthesis process for Tp-PTO-COF.



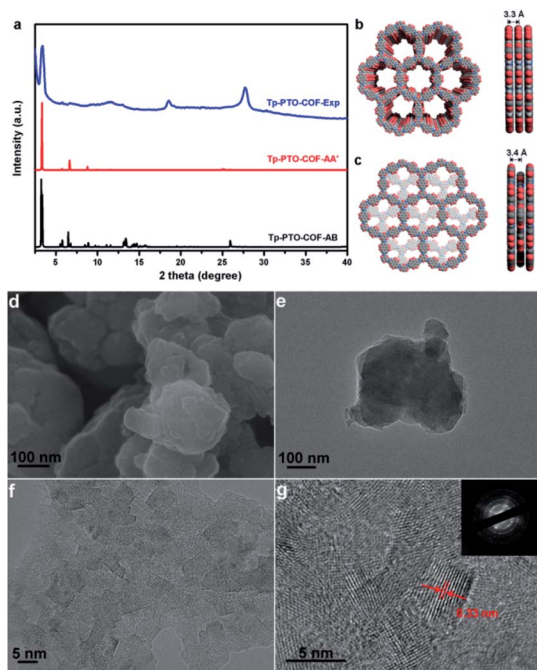


Fig. 2 (a) Comparison of the PXR D patterns among the as-synthesized structure (blue), the simulated eclipsed AA structure (red), and the simulated staggered AB structure (black); (b) simulated eclipsed AA structure of Tp-PTO-COF; (c) simulated staggered AB structure of Tp-PTO-COF; (d) SEM image of Tp-PTO-COF; (e) TEM image of Tp-PTO-COF; (f and g) HRTEM images of Tp-PTO-COF. The inset is the corresponding SAED image.

residual values and acceptable profile differences (Fig. S13<sup>†</sup>). These obvious PXR D characteristic peaks in conjunction with structural simulation indicate a high crystallinity and typical 2D layered hexagonal network structure of the as-prepared Tp-PTO-COF.

The morphology of Tp-PTO-COF was examined by scanning electron microscopy (SEM) and transmission electron microscopy (TEM). The as-synthesized COF exhibits a 2D lamellar stacking morphology, and the boundaries of the TEM image show the layered-sheet arrangement of Tp-PTO-COF (Fig. 2d and e). In addition, an interplanar distance of approximately 0.33 nm can be clearly observed from the high-resolution TEM (HRTEM) images of Tp-PTO-COF (Fig. 2f and g), which is in accordance with the  $\pi$ - $\pi$  stacking distance calculated based on the PXR D data and structural simulation. The HRTEM image and diffraction rings obtained from selected-area electron diffraction (SAED) analysis further confirm the intrinsic crystal structure of Tp-PTO-COF.

### Electrochemical characterization

Such an ordered porous structure and high-density carbonyl centers are particularly favorable in  $\text{Zn}^{2+}$  ion storage electrodes due to their ability to offer ion diffusion channels and facilitate  $\text{Zn}^{2+}$  ion storage *via* reversible coordination/uncoordination. In order to evaluate the potential of Tp-PTO-COF as a cathode material for aqueous rechargeable ZIBs, the electrochemical performance was initially evaluated on a typical coin-type cell

(Fig. 3a), which was assembled with Zn foil as the counter electrode and a glass fiber membrane adsorbed electrolyte (2 M  $\text{ZnSO}_4$ ) as the separator. Fig. 3b shows the cyclic voltammetry (CV) curve of Tp-PTO-COF at a scan rate of  $0.5 \text{ mV s}^{-1}$  in the potential range of 0.4–1.5 V vs.  $\text{Zn}^{2+}/\text{Zn}$ . Two pairs of redox peaks around 0.72/1.01 V and 0.86/1.13 V (denoted as R1/O1 and R2/O2) can be observed, which correspond to a multistep  $\text{Zn}^{2+}$  ion intercalation process followed by the electrochemical reduction of Tp-PTO-COF. Furthermore, CV curves at different scan rates have also been recorded (Fig. S14<sup>†</sup>). There is a linear relationship between its peak current and the square root of the scan rate (Fig. S15<sup>†</sup>), indicating that the redox-reaction kinetics are controlled by the  $\text{Zn}^{2+}$  ion diffusion process. Furthermore, the  $\text{Zn}^{2+}$  ion diffusion coefficient of the Tp-PTO-COF electrode was studied using the galvanostatic intermittent titration technique (GITT) method. As shown in Fig. S16<sup>†</sup>, the diffusion coefficient of  $\text{Zn}^{2+}$  ions in the Tp-PTO-COF cathode is around  $10^{-11} \text{ cm}^2 \text{ s}^{-1}$ . The fast kinetic performance is further proven by electrochemical impedance spectroscopy (EIS) measurements, which reflect a charge-transfer resistance ( $R_{\text{ct}}$ ) value of 29.08  $\Omega$  (Fig. S17 and Table S2<sup>†</sup>).

The energy-storage properties of Tp-PTO-COF can be observed in the galvanostatic charge–discharge (GCD) profiles at different current densities in Fig. 3c, where the achieved current densities and capacities are calculated based on the mass of Tp-PTO-COF in the cathode. The Tp-PTO-COF cathode exhibits a remarkably high specific capacity of  $301.4 \text{ mA h g}^{-1}$  at a current density of  $0.2 \text{ A g}^{-1}$  and maintains specific capacities of 281.3, 250.1, and  $228.6 \text{ mA h g}^{-1}$  at current densities of 0.5, 1, and  $2 \text{ A g}^{-1}$  respectively. Remarkably, even at extremely high current densities of 5 and  $10 \text{ A g}^{-1}$ , it also delivers capacities of 192.8 and  $140.07 \text{ mA h g}^{-1}$ , much higher than that of many reported COF cathodes and inorganic or organic electrode materials (Fig. S18 and Table S3<sup>†</sup>). In addition, we further evaluated the rate performance of batteries with different Tp-PTO-COF loading masses (Fig. S19<sup>†</sup>). By comparing the results, similar capacity performances were found. It is worth mentioning that Tp-PTO-COF not only exhibits significantly better specific capacity



Fig. 3 (a) Illustration of the aqueous Zn/Tp-PTO-COF battery; (b) CV curve at a scan rate of  $0.5 \text{ mV s}^{-1}$ ; (c) galvanostatic charge–discharge curves of the aqueous Zn/Tp-PTO-COF battery at various current densities; (d) cycle stability at a current density of  $2 \text{ A g}^{-1}$ .







Fig. 5 (a) Electronegativity simulation of Tp-PTO-COF; (b) simulation of binding energies with different numbers of  $\text{Zn}^{2+}$  ions in MO-Tp-PTO.

## Conclusions

In summary, Tp-PTO-COF with dual carbonyl active sites was proposed as a novel cathode material and successfully applied in aqueous rechargeable ZIBs. Tp-PTO-COF possesses abundant carbonyl nucleophilic centers, an ordered porous structure and inherent chemical stability, which are beneficial for  $\text{Zn}^{2+}$  ion storage and diffusion. The aqueous rechargeable ZIBs assembled with Tp-PTO-COF electrodes display superior capacity output, flat charge–discharge plateaus and excellent cycle stability. In addition, the  $\text{Zn}^{2+}$  ion intercalation mechanism has been studied through experimental analysis and theoretical simulations. The encouraging performance opens new possibilities for the design of advanced COF cathodes for high performance and sustainable aqueous rechargeable ZIBs.

## Data availability

All experimental and computational data are available in the ESI.†

## Author contributions

D. Ma and K. Liu designed the study. D. Ma, H. Zhao, H. Zhao and J. Li performed the experiments and analyzed the data. F. Cao and J. Li contributed to the theoretical calculations of the work. D. Ma, K. Liu and L. Wang prepared the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by the National Natural Science Foundation of China (No. 21601103 and 21701097) and the Natural Science Foundation of Shandong Province, China (No. ZR2020QB035).

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