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## CoP electrocatalysts embedded in nitrogen-doped carbon as a host toward fast iodine conversions†

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Herein, well-dispersed cobalt phosphide (CoP) electrocatalysts embedded in nitrogen-doped carbon (CoP@NC) were developed as an iodine host for zinc iodine batteries. Benefiting from the high electrical conductivity of the carbon matrix and the strong interaction as well as the efficient electrocatalytic activity of CoP with iodine species, the host achieved rapid iodine conversion while effectively suppressing the formation of polyiodides and zinc dendrites.

In recent years, lithium-ion batteries have faced challenges in meeting the growing demand for energy storage due to their high cost and safety risks, which has triggered the rapid development of alternative rechargeable batteries with high energy density, enhanced safety, and environmental sustainability. Among these, aqueous zinc-iodine batteries (AZIBs) have garnered significant attention from researchers due to their low cost, high safety and high capacity.<sup>1,2</sup> However, the inherent poor conductivity of iodine ( $\sim 10^{-6}$  S cm<sup>-1</sup>) severely hinders the iodine redox kinetics. Moreover, the shuttle effect, where iodine forms polyiodide intermediates (I<sub>3</sub><sup>-</sup> and I<sub>5</sub><sup>-</sup>) during cycling, results in irreversible loss of active mass, severe self-discharge, and corrosion of the zinc anode.<sup>2</sup>

To address these challenges, researchers have explored porous carbon-based materials as iodine hosts, such as carbon fibers,<sup>3</sup> nanosheets,<sup>4</sup> and porous carbon<sup>5</sup> to inhibit the dissolution and shuttling of polyiodides. However, the weak van der Waals forces of non-polar carbon materials are insufficient to

provide stable polyiodide inhibition during the cycling process. Therefore, introducing electrocatalysts to catalyze iodine conversion reactions and enhance the stable chemical adsorption of iodine species presents a promising strategy for improving the iodine conversion kinetics.

Metal phosphides have the characteristics of low cost, high conductivity and the ability to accelerate redox reactions, making them widely studied in electrocatalysis and lithium-sulfur batteries.<sup>6,7</sup> For example, Sun *et al.*<sup>6</sup> synthesized CoP nanoparticles/carbon nanocages, achieving a high capacity of 622 mA h g<sup>-1</sup> at 5C in a Li-S battery by enhanced S redox kinetics. Similarly, Ji *et al.*<sup>7</sup> developed CoP nanoframes that exhibited impressive bifunctional catalytic activities for both the HER and OER. Recently, a Ni<sub>2</sub>P@C host has been developed and showed a promoting effect on the iodine's redox in AZIBs.<sup>8</sup> These applications have greatly inspired us to explore the potential of metal phosphide hosts in catalyzing iodine's redox kinetics. However, highly surface-active CoP particles are prone to agglomerate, significantly reducing their catalytic sites. Here, integrating carbon materials with CoP nanoparticles offers a viable strategy to balance catalytic sites and improve iodine conductivity. First, the carbon matrix can effectively prevent the aggregation of CoP nanoparticles.<sup>9</sup> Second, nitrogen-doped carbon materials can greatly enhance the conductivity of the composite, accelerating the redox kinetics.<sup>2</sup> Notably, CoP has not been reported in AZIBs, and the fundamental understanding of its intrinsic mechanism for accelerating the redox of iodine remains to be fully elucidated.

Herein, well-dispersed CoP electrocatalysts embedded in a nitrogen-doped carbon matrix (CoP@NC) were customized *via* a simple pyrolysis and vapor-phase phosphating method. The CoP@NC host not only demonstrates strong chemical affinity to inhibit the polyiodide's penetration, but also offers abundant catalysis sites to promote iodine redox kinetics (I<sub>2</sub> to I<sup>-</sup>). At the same time, the n-doped carbon matrix can further enhance the conductivity of the composite and provide sufficient interfaces to load iodine. Consequently, the CoP@NC/I<sub>2</sub> exhibits a superior high-rate capacity of 191.1 mA h g<sup>-1</sup> at 0.1 A g<sup>-1</sup> and

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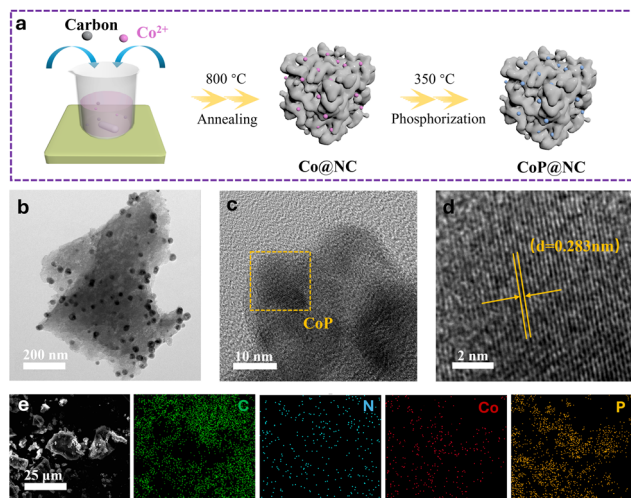
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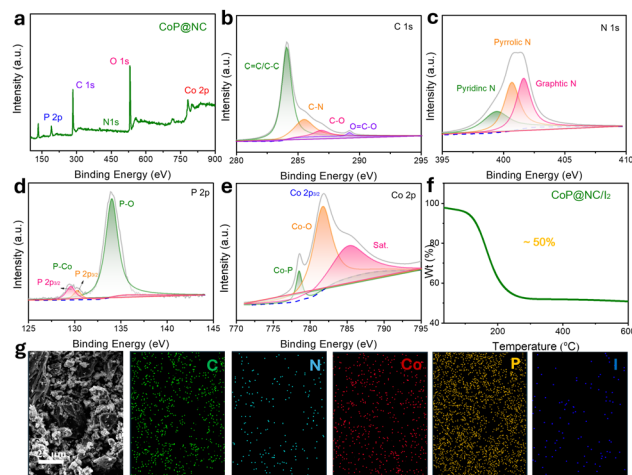


**Fig. 1** (a) Schematic illustration of the synthesis process for the CoP@NC host, (b) TEM images, (c) and (d) high-resolution TEM images, and (e) SEM image with element mappings of CoP@NC.

superior cycling stability of  $101.4 \text{ mA h g}^{-1}$  with around 100% coulomb efficiency (CE) after 1000 cycles at  $1.0 \text{ A g}^{-1}$ . These results highlight the outstanding performance of CoP@NC as an iodine host, establishing it as a competitive candidate among metal phosphide-based materials for AZIBs. Our work provides new insights into developing high-performance metal phosphide-based host materials for AZIBs.

The CoP@NC was prepared using a two-step strategy involving pyrolysis and phosphidation (Fig. 1a). The structure of CoP@NC was identified by X-ray diffraction (XRD) first. Fig. S1 (ESI<sup>†</sup>) displays obvious diffraction peaks at  $\sim 31.5, 35.2, 36.3, 46.1, 48.0, 52.2, 56.0$  and  $56.7$ , which are indexed to the CoP phase (PDF# 65-2593), without an impurity phase of cobalt oxide.<sup>10</sup> Transmission electron microscopy (TEM) in Fig. 1b showed that the CoP nanoparticles are evenly dispersed on the n-doped carbon with an average size of  $\sim 22 \text{ nm}$  (Fig. S2, ESI<sup>†</sup>). The high-resolution TEM (HRTEM) image further demonstrated the crystalline nature of the CoP (Fig. 1c), and the regular crystalline fringes with a  $d$ -spacing value of  $0.283 \text{ nm}$  can be observed in Fig. 1d, correspond to the (011) lattice plane of metallic CoP.<sup>11</sup> This result is consistent with the XRD's conclusion. The scanning electron microscope (SEM) and elemental distribution mapping (EDS) images further confirmed the uniform distribution of C, N, Co, and P elements throughout the CoP@NC (Fig. 1e), validating the successful integration of CoP within the nitrogen-doped carbon matrix.

The surface chemical states of the CoP@NC were studied using X-ray photoelectron spectroscopy (XPS). As shown in Fig. 2a, the survey XPS spectra confirmed the presence of C, N, Co, and P elements in CoP@NC. Specifically, the high-resolution XPS spectrum of C 1s in Fig. 2b revealed four types of carbon species, namely C=C/C-C in  $284.0 \text{ eV}$ , C-N in  $285.4 \text{ eV}$ , C-O in  $286.8 \text{ eV}$  and O=C-O in  $289.1 \text{ eV}$ , respectively.<sup>12,13</sup> For the N 1s spectrum, three characteristic N types – pyridine N, pyrrolic N and graphitic N can be observed at binding energies of  $399.4, 400.6$  and  $401.6 \text{ eV}$ , respectively

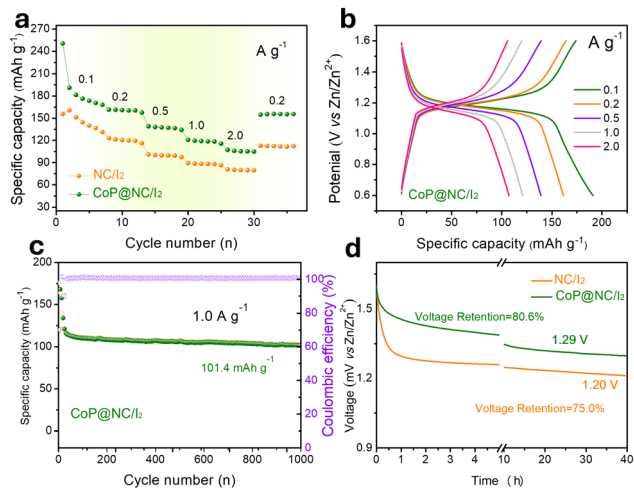


**Fig. 2** (a) XPS survey spectrum of CoP@NC. (b)–(e) High-resolution XPS spectra of C 1s, N 1s, P 2p and Co 2p of CoP@NC. (f) TGA curves of CoP@NC/I<sub>2</sub>. (g) SEM image with EDS images of CoP@NC/I<sub>2</sub>.

(Fig. 2c). These nitrogen dopants can increase the conductivity of the composites and facilitate the redox reaction kinetics.<sup>14,15</sup> Moreover, the P 2p spectra in Fig. 2d exhibited peaks at  $134.0, 130.3$  and  $129.5 \text{ eV}$ . The former peak belongs to the P-O bonds, while the latter two weak peaks are assigned to the P-Co bond (Co  $2p_{3/2}$  and Co  $2p_{1/2}$ ).<sup>10</sup> The Co 2p spectrum (Fig. 2e) further corroborated the presence of CoP, with peaks assigned to Co-P ( $778.5 \text{ eV}$ ), a satellite peak ( $785.1 \text{ eV}$ ) and Co-O ( $781.6 \text{ eV}$ ).<sup>16,17</sup> This Co-O is caused by the surface oxidation of the CoP@NC.<sup>7</sup> Notably, the Co  $2p_{3/2}$  in CoP@NC shifted positively ( $778.5 \text{ eV}$ ), compared with the Co ( $778.1 \text{ eV}$ ) in pure CoP, indicating a strong interfacial coupling between CoP and the N-doped carbon substrate.<sup>18,19</sup> These results collectively confirmed the successful incorporation of CoP electrocatalyst into the n-doped carbon matrix, with well-defined bonding configurations supporting its catalytic properties.

Subsequently, the CoP@NC's ability to serve as an iodine host was further evaluated, and the CoP@NC/I<sub>2</sub> cathode was obtained through a melt-diffusion strategy (Fig. S3, ESI<sup>†</sup>). The mass content of I<sub>2</sub> in the CoP@NC/I<sub>2</sub> was determined to be  $\sim 50 \text{ wt\%}$ , as calculated from the thermogravimetric analysis (TGA) curve (Fig. 2f). The stable iodine-carrying properties can be attributed to the synergistic effect of the physical confinement provided by the carbon matrix and the strong chemical adsorption of iodine by the CoP, which laid the foundation for the subsequent rapid and efficient iodine redox. Furthermore, SEM and EDS images of CoP@NC/I<sub>2</sub> confirmed the uniform distribution of I<sub>2</sub> in CoP@NC (Fig. 2g).

Leveraging the great potential of CoP@NC as a host material, the electrochemical performance of the CoP@NC/I<sub>2</sub> cathode was systematically studied in AZIBs. As shown in Fig. 3a, the CoP@NC/I<sub>2</sub> cathode delivered high reversible rate capacities of  $191.1, 161.5, 138.9, 120.8,$  and  $107.1 \text{ mA h g}^{-1}$  at  $0.1, 0.2, 0.5, 1.0,$  and  $2.0 \text{ A g}^{-1}$ , respectively, which are higher than that of the NC/I<sub>2</sub> cathode across all tested conditions. When the current density is recovered to  $0.2 \text{ A g}^{-1}$ , the CoP@NC/I<sub>2</sub>

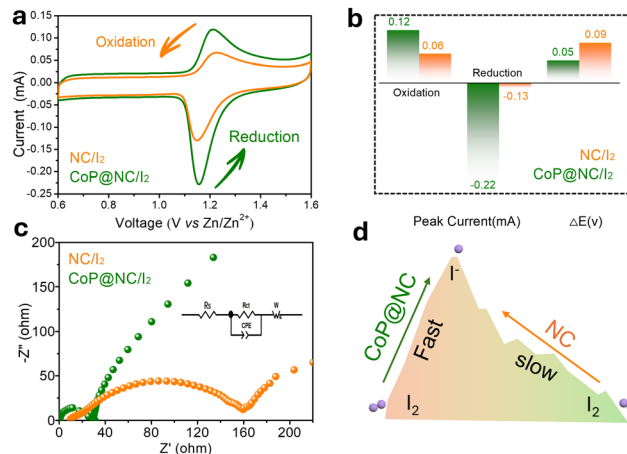


**Fig. 3** (a) Rate capacity of CoP@NC/I<sub>2</sub> and NC/I<sub>2</sub>. (b) Discharge/charge curves and (c) cycling performance of CoP@NC/I<sub>2</sub>. (d) Self-discharge performance of CoP@NC/I<sub>2</sub> and NC/I<sub>2</sub>.

cathode achieved a high specific capacity of 155.6 mA h g<sup>-1</sup>, demonstrating its remarkable reversibility. In contrast, the NC/I<sub>2</sub> cathode only showed a poor performance of 113.4 mA h g<sup>-1</sup> at 0.2 A g<sup>-1</sup>. Besides, as the current density increases, the discharge/charge (GCD) curves of the CoP@NC/I<sub>2</sub> electrode maintained distinct plateaus, while the NC/I<sub>2</sub> electrode showed a rapid tightening trend, especially at high current density (Fig. 3b and Fig. S4, ESI<sup>†</sup>). These results indicated that the CoP@NC host can greatly improve iodine utilization. Notably, the rate performance of CoP@NC/I<sub>2</sub> was highly competitive compared to other reported Zn/I<sub>2</sub> battery hosts (Fig. S5, ESI<sup>†</sup>), underscoring its potential for advanced AZIBs.

The long-term cycling performance of both electrodes was evaluated in Fig. 3c. Specifically, the CoP@NC/I<sub>2</sub> cathode maintained a high capacity of 101.4 mA h g<sup>-1</sup> with ~100% CE after 1000 cycles at 1.0 A g<sup>-1</sup>, indicating the exceptional efficiency of its electrochemical redox reactions. Importantly, the CoP@NC/I<sub>2</sub> electrode also maintained a stable high voltage of 1.29 V over 40 h, while the NC/I<sub>2</sub> electrode just kept 1.20 V, indicating the excellent self-discharge performance of CoP@NC/I<sub>2</sub>. This excellent performance may be due to the rapid adsorption-catalytic ability of the CoP@NC to iodine species, which greatly reduces the formation of polyiodide and improves the utilization efficiency of iodine.

To elucidate the excellent electrochemical performance of the CoP@NC/I<sub>2</sub> cathode, cyclic voltammetry (CV) and electrochemical impedance spectroscopy (EIS) tests were conducted to investigate the iodine conversion kinetics. As shown in Fig. 4a, the CV curves of both electrodes showed a pair of redox peaks at ~1.15/1.20 V at 0.2 mV s<sup>-1</sup>, corresponding to the conversion from I<sub>2</sub> to I<sup>-</sup>.<sup>5</sup> Notably, the CoP@NC/I<sub>2</sub> electrode exhibited a higher current (0.12/−0.22 mA) and a lower voltage polarization (0.05 V) compared to the NC/I<sub>2</sub> electrode, indicating significantly enhanced redox kinetics for the CoP@NC/I<sub>2</sub> electrode (Fig. 4b). Besides, Tafel plots derived from the CV data further support this finding.<sup>16,20</sup> As illustrated in Fig. S6 (ESI<sup>†</sup>), the CoP@NC/I<sub>2</sub> electrode exhibited a smaller Tafel slope for both



**Fig. 4** (a) CV profiles, (b) the peak current and polarization voltage, and (c) EIS curves of CoP@NC/I<sub>2</sub> and NC/I<sub>2</sub>. (d) Mechanism diagram illustrating the I<sub>2</sub> redox reaction.

oxidation (133 mV dec<sup>-1</sup>) and reduction (350 mV dec<sup>-1</sup>), compared to the NC/I<sub>2</sub>'s Tafel slope (208 and 402 mV dec<sup>-1</sup>). These results demonstrated that CoP@NC rapidly catalyzes bidirectional iodine redox processes. EIS curves in Fig. 4c showed the Nyquist plots of both electrodes between 100 kHz and 0.01 Hz before cycling. Specifically, the R<sub>ct</sub> value of the CoP@NC/I<sub>2</sub> electrode is 29.0 Ω, fitted by the equivalent electrical circuit (Fig. 4c, inset),<sup>17</sup> which is smaller than that of the NC/I<sub>2</sub> electrode (160.0 Ω), suggesting the fastest charge-transfer kinetics in the CoP@NC/I<sub>2</sub> electrode. Fig. 4d summarizes the mechanism of the CoP@NC host promoting the iodine conversion process. Unlike the NC, which only physically restricts iodine species and results in slow iodine conversion, the CoP@NC host has highly exposed active sites, enhances the binding ability with intermediates, and thereby improves the catalytic iodine conversion kinetics.<sup>21–23</sup> Besides, the good contact interface between CoP and the carbon substrate can promote electron transfer.<sup>7,21</sup> This synergistic effect achieves fast iodine conversion while effectively suppressing shuttling, ultimately improving the overall battery performance.

To further evaluate the ability of CoP@NC to limit polyiodide diffusion, static adsorption experiments were performed by immersing CoP@NC and NC in zinc triiodide solutions. As shown in Fig. 5a and Fig. S7 (ESI<sup>†</sup>), after standing for 12 hours, the solution containing CoP@NC appeared much more transparent than the solution containing NC, indicating a stronger anchoring effect of CoP@NC on polyiodides. UV-visible absorption spectroscopy (Fig. 5b) further confirmed that the supernatant of the CoP@NC-impregnated solution exhibited the lowest adsorption intensity, demonstrating the superior chemical adsorption of CoP@NC for polyiodides than NC. This strong adsorption capacity was corroborated by XPS analysis of the CoP@NC after adsorption (Fig. 5c). First, the XPS survey spectrum of the CoP@NC after adsorption of I<sub>3</sub><sup>-</sup> solution in Fig. S8 (ESI<sup>†</sup>) showed peaks at ~630.6 and 618.0 eV, meaning the chemical interaction between I<sub>3</sub><sup>-</sup> and CoP@NC.<sup>2</sup> Compared with the Co spectrum of CoP@NC (Fig. 2e), all peaks of Co 2p in

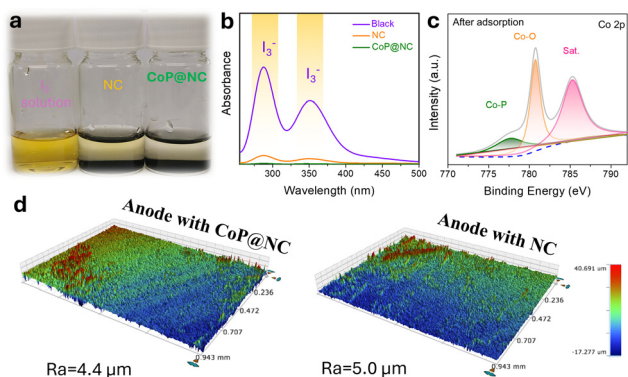


Fig. 5 (a) Visual comparison of adsorption capacity between CoP@NC and NC. (b) UV-vis spectra. (c) High-resolution XPS spectrum of Co 2p of CoP@NC-I<sub>3</sub><sup>-</sup>. (d) 3D surface profiles of the Zn anode after cycling.

CoP@NC-I<sub>3</sub><sup>-</sup> shifted in the negative direction. Specifically, the Co-O/Co-P bonds shifted from 781.6/778.5 eV to 780.7/777.5 eV, respectively, confirming the electron transfer interactions between CoP@NC and I<sub>3</sub><sup>-</sup>. The above results substantiate the existence of a strong chemical interaction between CoP@NC and polyiodides rather than weak physical adsorption.<sup>2</sup>

To verify the role of CoP@NC in inhibiting polyiodide-related degradation, the zinc anode was carefully inspected after long-term cycling. SEM of the zinc anodes with CoP@NC and NC electrodes after 100 cycles at 1.0 A g<sup>-1</sup> are shown in Fig. S9 (ESI<sup>†</sup>). The NC-based battery exhibited disordered flake/flower-like dendrites on the Zn's surface, while the CoP@NC-based battery retained an intact and smooth anode morphology. The corresponding 3D images in Fig. 5d also confirmed less byproduct coverage on the Zn's surface with the CoP@NC/I<sub>2</sub> cathode (4.4 μm) compared with the NC/I<sub>2</sub> cathode (5.0 μm), as evidenced by the reduced surface height variation. These findings certify that CoP@NC can effectively inhibit the occurrence of side reactions on the surface of the Zn anode, providing enhanced stability during cycling.

L. Z.: methodology, data curation, writing – original draft; X. G., Z. Z., Y. F., Z. Y., C. Z., and Y. W.: data curation & formal analysis. H. F., H. X., X. L. and H. L.: visualization, B. J. and H. Y.: writing – review & editing. Y. S. and T. M.: writing – review & editing, supervision, funding acquisition.

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## Data availability

Data are contained within the article and ESI<sup>†</sup>.

## Conflicts of interest

There are no conflicts to declare.

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