Co$_2$O$_3$/Co$_2$N$_{0.67}$ nanoparticles encased in honeycomb-like N, P, O-codoped carbon framework derived from corncob as efficient ORR electrocatalysts

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It is essential to develop cost-effective rechargeable metal–air batteries, with high activity, stability, and efficiency, that use non-precious metals (NPMs)-based cathodic oxygen reduction reaction (ORR) catalysts. Here, by using earth-abundant corncob (CC) as the carbon source, Co(OH)$_2$, NaH$_2$PO$_4$, and melamine as the precursors, and KOH as the chemical activator, CoNP@bio-C-a is obtained and comparative studies are carried out with three other types of CC-derived carbon-based catalytic materials, namely, bio-C, CoP@bio-C, and CoNP@bio-C. Depending mainly on the formation of Co$_2$O$_3$/Co$_2$N$_{0.67}$ active sites (as p–n heterojunctions) and N, P, O-containing functional groups, the resultant CoNP@bio-C-a catalyst exhibits best electrocatalytic activity among the four types of catalysts; via a 4-electron pathway, it has good stability and good methanol tolerance. In addition, its unique honeycomb-like porous structure, high graphitization degree, and abundant oxygen-containing groups contribute to its excellent ORR activity. This study provides insights for exploring the application of heteroatom-doped biomass-derived carbon catalysts.

1. Introduction

In the field of fuel cells and rechargeable metal–air batteries, it is essential to develop high-efficiency oxygen reduction reaction (ORR) electrocatalysts; however, their applications are affected by the rate-limiting processes in cathodic ORR.\textsuperscript{1,2} Pt-based catalysts are contemporary commonly used commercial catalysts. However, they have disadvantages, such as inactivation after poisoning, poor stability, high cost, and limited availability.\textsuperscript{3} Hence, exploring efficient and stable non-precious metal (NPM)-based ORR catalysts would be beneficial for energy conversion systems.

Biomass has attracted considerable attention because it is abundant and inexpensive. Biomass-derived carbon-based materials also have good electrical conductivities and high specific surface areas and porosities, along with three-dimensional morphologies. However, biomass-based carbon materials are not ideal ORR electrocatalysts. Heteroatom-doped biomass-based porous carbon materials have drawn considerable attention owing to their promising ORR activities.\textsuperscript{4} Introducing heteroatoms (such as N, P, S, B, and O) into the carbon framework causes carbon defects by modifying heteroatoms (such as P, S, B, and O) into the carbon framework, which enhance their hydrophilicities, and thus, accelerate the ORR activity.\textsuperscript{8,9} Hence, phosphorus has better capacity to supply electrons. The oxygen atom has high electronegativity, and so, O-doping can not only adjust the electronic structure of carbon materials, but also enhance their hydrophilicities, and thus, accelerate the ORR activity.\textsuperscript{11} In recent years, ternary heteratomic doping has attracted considerable attention because of its synergistic effect.
on improving the ORR activity of carbon materials. However, there are relatively few reports on oxygen ternary doping in carbon materials.

Herein, a new honeycomb-like porous carbon hybrid with Co, N, P, O-codoping (denoted as CoNP@bio-C-a) is prepared from waste biomass (corn cob, CC) mixed with cobalt hydroxide, sodium hypophosphite, and melamine (precursors) and KOH (activating agent) through a simple one-step pyrolysis process. The resultant CoNP@bio-C-a catalyst exhibits a 4-electron process, contains active sites from Co2O3/Co2N0.67 hetero-junctions and N, P, O-codoped carbon, and shows excellent electroactivity, good stability, and methanol resistance in 0.1 M KOH ($E_{\text{onset}} = 0.92 $ V, $E_{1/2} = 0.85 $ V, $J_L = -5.5 \text{ mA cm}^{-2}$).

2. Experimental section

2.1. Samples preparation for doped and undoped bio-C materials

Cleaned and dried CC was milled at 300 rpm. CC powder (5.7 g), Co(OH)$_2$ (0.05 g), and NaH$_2$PO$_2$·H$_2$O (0.25 g) were mixed in deionized water and subjected to ultrasound. The product was dried and then mixed with melamine (precursor) with a mass ratio of 2 : 1 and grinded till it was mixed uniformly. Then, KOH powder (0.2 g) was added to this mixture and grinded till it was mixed uniformly. The obtained CoNP@bio-C-a was first carbonized in an inert atmosphere of 10 $^\circ$C min$^{-1}$ at 500 $^\circ$C for 2 h and then activated using CO$_2$ gas at 800 $^\circ$C for 3 h. The obtained mixture was cooled, cleaned, and dried.

3. Results and discussion

3.1. Morphological and structural characterization

Morphologies of bio-C, CoP@bio-C, CoNP@bio-C, and CoNP@bio-C-a were visualized using scanning electron microscopy (SEM, Fig. 2). The SEM images revealed the improvement of the pore structures from undoped bio-C, CoP@bio-C, and CoNP@bio-C to CoNP@bio-C-a. With the introduction of Co and P into the bio-C framework (Fig. 2a and b), the pore size of the fibrous network structure in bio-C decreased from approximately 40 $\mu$m for bio-C to 10 $\mu$m for CoP@bio-C. After incorporating Co, N, and P into the bio-C framework, an ordered honeycomb structure (diameter = 10–20 $\mu$m, wall thickness = 600 nm) was obtained (Fig. 2c). Fig. 2d shows no obvious morphological changes during the conversion of CoNP@bio-C to CoNP@bio-C-a, but it shows that uniform pores (size = 2–4 $\mu$m) appeared on the relatively thick walls of the honeycomb-like structure of CoNP@bio-C-a.

The microstructure of CoNP@bio-C-a was examined using high-resolution transmission electron microscopy (HRTEM) (Fig. 3). Some small inorganic spots (diameters = 5–11 nm) were encased in the carbon layers (Fig. 3a), indicating the presence of the active species of Co$_2$O$_3$ and Co$_2$N$_{0.67}$ in the carbon materials (Fig. 3b and c). The values of d-spacing were 0.39, 0.18, and 0.20 nm, assigned to the (002) plane of graphite carbon, (100) plane of Co$_2$O$_3$, and (101) plane of Co$_2$N$_{0.67}$ in CoNP@bio-C-a, respectively. EDX spectra reveal (Fig. S1†) the atomic percentage composition of Co : P as approximately 3 : 8 = 0.375 (0.24 : 0.83). The inductively coupled plasma optical emission spectrometry (ICP-OES) results (Table S1†) show that the atomic composition of CoNP@bio-C-a in an aqua regia solution is 7 : 18 = 0.389 (0.73 : 1.81), which matched well with the findings of EDX analysis. This confirms that these two elements
were successfully incorporated into the material.\textsuperscript{14,15} The elemental mapping images (Fig. S2†) further show that Co, C, P, N, and O were evenly distributed in CoNP@bio-C-a.

The crystal structures of bio-C, CoP@bio-C, CoNP@bio-C, and CoNP@bio-C-a, were evaluated using PXRD (Fig. 4a). The prepared samples had a wide (002) dihedral that the graphite structure was disordered and mainly an sp\(^2\) hybrid structure was observed, which is in good agreement with the HRTEM and Raman results.\textsuperscript{16} Closer inspection revealed that the diffraction peak of the (002) plane from graphitic carbon (GC) of bio-C, CoP@bio-C, and CoNP@bio-C shifted only slightly at 2\(\theta\) values of 22.07° (0.402 nm), 22.09° (0.402 nm), and 21.91° (0.405 nm), respectively. After KOH activation, the GC (002) diffraction peak of CoNP@bio-C-a moved to a higher angle (2\(\theta\) = 22.88°, \(d_{002} = 0.388\) nm) with a smaller \(d\)-spacing compared to that of CoNP@bio-C, implying a stronger layer-to-layer (i.e., π-π) interaction in the GC framework in CoNP@bio-C-a.\textsuperscript{17-19} The weak diffraction peaks in the PXRD patterns indicate that nanoparticles of Co\(_2\)N\(_{0.67}\) (JCPDS no. 06-0691) and Co\(_2\)O\(_3\) from the HRTEM image of CoNP@bio-C-a. The crystal structures of bio-C, CoP@bio-C, CoNP@bio-C, and CoNP@bio-C-a were used to characterize the surface compositions and chemical valence states (Fig. 5). The survey spectra revealed distinct peaks of C, Co, N, O, and P, with elemental contents of 71.28%, 9.02%, 1.09%, 17.58%, and 1.03%, respectively (Fig. 5a), showing that Co, P, N, and O were successfully doped in CoNP@bio-C-a. Particularly, the C 1s spectrum can be divided into four peaks centered at 284.8 (C\(-\text{C}\)), 285.6 (C\(-\text{C}–\text{O}\)), 286.5 eV (C\(-\text{C}–\text{N}\)), and 289.4 eV (O–C\(-=\text{O}\)) (Fig. 5b). The high content of sp\(^2\)-carbon (C\(-=\text{C}\) at 284.8 eV) further indicates the existence of GC, which enhances the electron conductivity of carbon materials, improving ORR activities and electrocatalytic performance.

The BET specific surface areas of the four types of samples were compared and revealed a clear trend toward higher specific surface areas, following the order CoP@bio-C (291.82 m\(^2\) g\(^{-1}\)) < CoNP@bio-C-a (365.40 m\(^2\) g\(^{-1}\)) < bio-C (453.74 m\(^2\) g\(^{-1}\)) < CoNP@bio-C (475.55 m\(^2\) g\(^{-1}\)). The large accessible surface area of CoNP@bio-C-a is beneficial for electrolyte penetration and oxygen adsorption, which can positively impact the electrocatalytic activity, as discussed below.\textsuperscript{24} In contrast, CoP@bio-C and CoNP@bio-C exhibited slightly low BET surface areas, which may be caused by the excessive corrosion and distortion due to Co, N, P-codoping into the bio-C structure.\textsuperscript{25}

X-ray photoelectron spectroscopy (XPS) measurements of CoNP@bio-C-a were used to characterize the surface compositions and chemical valence states (Fig. 5). The survey spectra revealed distinct peaks of C, Co, N, O, and P, with elemental contents of 71.28%, 9.02%, 1.09%, 17.58%, and 1.03%, respectively (Fig. 5a), showing that Co, P, N, and O were successfully doped in CoNP@bio-C-a. Particularly, the C 1s spectrum can be divided into four peaks centered at 284.8 (C\(-\text{C}\)), 285.6 (C\(-\text{C}–\text{O}\)), 286.5 eV (C\(-\text{C}–\text{N}\)), and 289.4 eV (O–C\(-=\text{O}\)) (Fig. 5b). The high content of sp\(^2\)-carbon (C\(-=\text{C}\) at 284.8 eV) further indicates the existence of GC, which enhances the electron conductivity of carbon materials, improving ORR activities and electrocatalytic performance.

**Fig. 3** (a) TEM image of CoNP@bio-C-a (inset: particle size distribution of CoNP@bio-C-a); lattice fringe images of (b) graphitic carbon and (c) Co\(_2\)N\(_{0.67}\) and Co\(_2\)O\(_3\) from the HRTEM image of CoNP@bio-C-a.

**Fig. 4** (a) PXRD patterns; (b) Raman spectra; (c) N\(_2\) adsorption/desorption isotherms of bio-C, CoP@bio-C, CoNP@bio-C, and CoNP@bio-C-a.
activities. The other three peaks of C 1s at 285.6, 286.5, and 289.4 eV are attributed to the defects and functional groups due to nitrogen and phosphorus heteroatom doping in the carbon matrix. In the Co 2p spectra for CoNP@bio-C-a (Fig. 5c), the peaks at approximately 782.1 and 786.8 eV belong to Co–N species, while the peaks at 798.3 and 803.6 eV originate from Co–O species. Both Co–N and Co–O species also can improve the catalytic activity of ORR under alkaline conditions. Moreover, some groups indicate that the N-rich porous carbon shell surrounding exposed Co3O4 particles improves the conductivity of the particles, which in turn increases the ORR activity. The fitted N 1s spectra for CoNP@bio-C-a centered at 398.3, 399.0, 401.6, and 400.7 eV can be assigned to pyridinic, pyrrolic, and graphitic nitrogen and Co–N, respectively (Fig. 5d). Studies have demonstrated that the incorporation of nitrogen and a metal as active centers in a carbon matrix can reduce the local work function on the carbon surface. The deconvolution of the O 1s region of CoNP@bio-C-a (Fig. 5e) comprises peaks at 530.9 (O1), 532.2 (O2), and 533.5 eV (O3). The sharp peak at 530.9 eV is caused by the lattice oxygen (O2-) of Co3O4. The peaks at 532.2 and 533.5 eV belong to the oxygen-containing functional groups on the carbon surface. To qualitatively characterize the chemical functional groups, FTIR spectra of CoNP@bio-C-a were collected (Fig. S4, ESI†). The peaks near 1697, 1460, and 1133 (wide peak) cm−1 are attributed to C=O stretching vibration, C=C and/or C=N bands, and the C–O bond, respectively. Three absorption bands are observed at 896, 1050, and 1280 cm−1, which can be assigned to P–O–C or P=O–C stretching. The high-resolution P 2p peak for CoNP@bio-C-a was further verified using XPS analysis (Fig. 5f). The P 2p peak centered at 133.6 eV belongs to P–C bonds, indicating that P elements have been doped in the biomass-based carbon material. These results show that Co, N, P, and O were successfully incorporated in CoNP@bio-C-a. Doping of nitrogen and oxygen in carbon materials can improve the catalytic activity of ORR under alkaline conditions. The authors also explain the role of nitrogen and phosphorus in enhancing the conductivity and activity of the catalyst.

Fig. 5 (a) XPS survey spectra of CoNP@bio-C-a. High-resolution XPS spectra of (b) C 1s, (c) Co 2p, (d) N 1s, (e) O 1s, and (f) P 2p core levels for CoNP@bio-C-a.
electrical conductivity and surface wettability, which is helpful for improving the electrochemical catalytic performance.\(^4\)

### 3.2. Electrochemical properties

The ORR activity of the as-prepared carbon catalysts (bio-C, CoP@bio-C, CoNP@bio-C, and CoNP@bio-C-a) was assessed using cyclic voltammetry (CV) in a 0.1 M KOH solution (O\(_2\)-saturated: solid lines, N\(_2\)-saturated: dotted lines; Fig. 6a and Table S3, ESI\(^\dagger\)). Comparison of the CV profiles showed that CoP@bio-C and CoNP@bio-C demonstrated slightly improved ORR activity than bio-C. As expected, upon adding KOH activator in CoNP@bio-C, the resultant CoNP@bio-C-a exhibited enhanced catalytic ORR activity due to shifting of the peak potential to a more positive value (\(E_{\text{onset}} = 0.85\) V), further confirming the effectiveness of the chemical activations of carbon materials in improving the catalytic activity of ORR.\(^4\)

The ORR activities of the four types of carbon-based catalytic materials were further investigated using LSV polarization curves (Fig. 6b). For comparison, ORR activity of 20 wt% Pt/C for industrial use was studied. The ORR activity of all the doped biomass-based carbon materials was better than that of undoped bio-C (Fig. 6b, S5–S7, and Table S3, ESI\(^\dagger\)), and the order of ORR activities was bio-C (\(E_{\text{onset}} = 0.85\) V, \(E_{1/2} = 0.69\) V, \(J_L = -2.78\) mA cm\(^{-2}\)) < CoP@bio-C (\(E_{\text{onset}} = 0.85\) V, \(E_{1/2} = 0.79\) V, \(J_L = -3.47\) mA cm\(^{-2}\)) < CoNP@bio-C (\(E_{\text{onset}} = 0.89\) V, \(E_{1/2} = 0.82\) V, \(J_L = -3.84\) mA cm\(^{-2}\)) < CoNP@bio-C-a (\(E_{\text{onset}} = 0.92\) V, \(E_{1/2} = 0.85\) V, \(J_L = -5.49\) mA cm\(^{-2}\)), indicating that the treatment for the bio-C materials proceeded stepwise. However, the BET surface areas of CoP@bio-C and CoNP@bio-C were lower than that of bio-C-800 (Table S2, ESI\(^\dagger\)). Furthermore, to assess the contribution of the Co compound and carbon doping carbon on ORR, CoP@bio-C-a, N@bio-C-a, and NP@bio-C-a were prepared from CC powder by adding only auxiliary precursors of Co(OH)\(_2\)-NaH\(_2\)PO\(_4\)-H\(_2\)O-KOH, melamine-KOH, and NaH\(_2\)PO\(_4\)-H\(_2\)O-melamine-KOH, respectively. CV and LSV profiles were employed to study the ORR catalytic activity of CoP@bio-C-a (\(E_{\text{onset}} = 0.89\) V, \(E_{1/2} = 0.83\) V, \(J_L = -3.77\) mA cm\(^{-2}\)), N@bio-C-a (\(E_{\text{onset}} = 0.90\) V, \(E_{1/2} = 0.71\) V, \(J_L = -3.06\) mA cm\(^{-2}\)), and NP@bio-C-a (\(E_{\text{onset}} = 0.85\) V, \(E_{1/2} = 0.71\) V, \(J_L = -3.06\) mA cm\(^{-2}\)) in an O\(_2\)-saturated 0.1 M KOH electrolyte (Fig. S8–S10 and Table S3, ESI\(^\dagger\)), and they both displayed lower catalytic activity than CoNP@bio-C-a. For comparison, the catalytic activities of CoNP@bio-C-a generated via pyrolysis at 700 °C and 900 °C were also examined using CV and LSV profiles with the RDE configuration (Fig. S11, S12 and Table S3, ESI\(^\dagger\)). Pyrolysis temperatures lower or higher than 800 °C resulted in poor catalytic activity of ORR compared with CoNP@bio-C-a. This phenomenon was further analyzed using BET and SEM characterization. At 700 °C, it had a three-dimensional bowl-shaped structure; however, it had a small specific surface area due to the insufficient temperature of activated pore creation. At 900 °C, it had a large specific surface area; however, it did not maintain a good honeycomb structure during the activation process, resulting in fewer attachment points for the active sites (Fig. S13, S14 and Table S2, ESI\(^\dagger\)). In addition, the ORR measurements of CoNP@bio-C-a-N\(_2\) in the N\(_2\) process were compared with the corresponding results (summarized in Fig. S15 and Table S3, ESI\(^\dagger\)). Compared with the CO\(_2\)-activated CoNP@bio-C-a catalyst, the CoNP@bio-C-a-N\(_2\) catalyst showed poor ORR catalytic properties (\(E_{\text{onset}} = 0.86\) V, \(E_{1/2} = 0.65\) V) in

![Fig. 6](image-url) (a) CVs of bio-C, CoP@bio-C, CoNP@bio-C, and CoNP@bio-C-a. (b) LSV curves of as-prepared catalysts and commercial 20 wt% Pt/C catalyst at 1600 rpm. (c) LSV curves of CoNP@bio-C-a at various values of revolutions per minute (inset: K-L plots of CoNP@bio-C-a). (d) Nyquist plots of bio-C, CoP@bio-C, CoNP@bio-C, and CoNP@bio-C-a catalysts-modified electrodes.

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an O₂-saturated 0.1 M KOH solution. These results demonstrated that the CO₂ activation process has a positive influence on biomass-based carbon materials.²⁴ Consistent with the CV study, CoNP@bio-C-a displayed the best performance of ORR activity among the heteroatom-doped bio-C and undoped bio-C catalysts, that is, with E_{onset} = 0.92 V and E_{1/2} = 0.85 V (O₂-saturated 0.1 M KOH), which are comparable to the corresponding values of the commercial Pt/C catalyst (E_{onset} = 0.96 V, E_{1/2} = 0.86 V; J_f = −5.61 mA cm⁻²).

To further understand the electron transfer ability of CoNP@bio-C-a, LSV tests were carried out at 400–2025 rpm; the results are shown in Fig. 6c. Typically, current density is enhanced upon increase in the rotation rate. The Koutecky–Levich (K–L) plots of CoNP@bio-C-a catalyst showed good linearity, which indicates a first-order reaction kinetics process under different potentials. The electron transfer number (n) for CoNP@bio-C-a at a potential range of 0.40–0.60 V was calculated to be approximately 4 (inset of Fig. 6c), with an average value of 3.94, indicating that the oxygen reduction process followed the 4-electron reduction process, which was similar to that of the Pt/C catalyst. Furthermore, the electrochemical impedance spectroscopy (EIS) images of the four types of carbon-based catalytic materials were assessed. As shown in Fig. 6d, the smallest charge transfer resistance (Rt) of CoNP@bio-C-a obtained from the Nyquist plots indicates the beneficial role of the honeycomb-like structure and the Co₂O₃/Co₂N₀.₆₇ nanoparticles encased in the N, P, O-codoped carbon framework. We also performed Mott–Schottky (M–S) analysis at an AC frequency of 10 Hz to investigate the conductivity of CoNP@bio-C-a (Fig. S16, ESIF). A typical inverted “V-shape” plot was obtained in the M–S plot of CoNP@bio-C-a, confirming the formation of p–n junctions.⁸ To evaluate the impact of p–n junctions on ORR performance, the generated Co₂O₃ nanoparticles of the CoNP@bio-C-a were leached using an HCl solution. A positive slope (n-type) in the M–S plots appeared instead of the typical inverted “V-shape” plot of CoNP@bio-C-a (Fig. S17, ESIF). The LSV tests revealed that the ORR activity of the acid-treated samples decreased considerably (E_{onset} = 0.90 V, E_{1/2} = 0.82 V; J_f = −3.88 mA cm⁻²) (Fig. S18, ESIF), which confirms the crucial role of the p–n junctions in ORR. Thus, the existence of p–n heterojunctions in CoNP@bio-C-a created sufficient space charge layers to improve the current generation, which facilitated interfacial electron transfer in the ORR.⁴⁵–⁴⁶ This, combined with the change of the surface electronic structure of the carbon matrix because of the formation of C–P, C–N, and C–O active sites, led to the high ORR performance of CoNP@bio-C-a. Meanwhile, Co₂O₃/Co₂N₀.₆₇ nanoparticles, as active sites encased within a honeycomb-like N, P, O-codoped carbon framework, further enhanced the ORR activity.

Long-term durability of ORR catalysts is generally examined using chronoamperometric responses. The stability of CoNP@bio-C-a and Pt/C at 1600 rpm under a high constant voltage of 0.8 V was measured (Fig. S19a, ESIF). For CoNP@bio-C-a, 73% of the initial current density was retained after continuous operation for 10,000 s, while for the Pt/C catalyst, rapid activity loss occurred, and it could retain only approximately 60% of the initial current density. Therefore, abundant nitrogen species may provide stable anchoring sites for Co-based active components and improve the stability of catalysts.⁴⁷–⁴⁹ Furthermore, methanol crossover experiments revealed that when methanol was added to the solution, there was no significant influence on the current density of CoNP@bio-C-a. In contrast, the current of the Pt/C catalyst decreased significantly under methanol oxidation (Fig. S19b, ESIF). The internal composition and morphology of CoNP@bio-C-a were studied and analyzed after long-term stability tests. XRD patterns of CoNP@bio-C-a after durability tests showed no changes in its chemical composition, indicating that the catalyst composition was stable (Fig. S20, ESIF). In the XPS spectrum of Co 2p, the Co–O peak moved to a higher energy region, while the Co–N peak moved to a lower energy region. The initial sharp peaks of CoNP@bio-C-a for O 1s and P 2p at 532.4 eV and 133.6 eV moved toward higher energy regions, 532.5 eV and 134.2 eV, after the ORR durability test, and new peaks appeared at approximately 536.3 eV and 140.9 eV, likely attributed to the chemisorbed oxygen (Fig. S21, ESIF).⁵⁰ Samples of CoNP@bio-C-a were collected after the stability tests, and their SEM profiles showed that the honeycomb-like morphology of the catalyst was retained (Fig. S22, ESIF). These results indicate that CoNP@bio-C-a has good structural stability.

Surface wettability of electrocatalysts is an important factor.³¹–³³ Hence, we carried out surface wettability tests of the four materials. The contact angle decreased in the order bio-C (~72°) > CoNP@bio-C (~73°) > CoNP@bio-C (~51°) > CoNP@bio-C (~46°), indicating that the treatment of the bio-C materials was followed by further functionalization in the above order (Fig. S23, ESIF). This contributed to the best ORR catalytic activity of CoNP@bio-C-a. The heteroatoms of Co, N, O, and P were also beneficial in enhancing the contact between the electrolyte and the electrode surface compared to that in the pure bio-C (~72°) and N-doped N@bio-C (~69°) electrodes, respectively (Fig. S23e, ESIF). The resulting enhanced affinity to H₂O is favorable for enhancing the catalytic activity of CoNP@bio-C-a toward ORR.

4. Conclusions

We explored a high-performance ORR catalyst, CoNP@bio-C-a, which was obtained using earth-abundant CC as the carbon source, sodium hypophosphite, melamine, and cobalt hydroxide as the precursors, and KOH as the chemical activator. This catalyst exhibited excellent electroactivity in 0.1 M KOH (E_{onset} = 0.92 V, E_{1/2} = 0.85 V; J_f = −5.5 mA cm⁻²), which was superior to that of the commercial Pt/C catalyst. The high activities are attributed to the unique nanostructures of Co₂O₃/Co₂N₀.₆₇ (as p–n junctions), heteratomic doping (Co, N, P, and O), and large specific surface areas. This study showed that high-performance electrocatalysts can be prepared from biomass derivatives through rational design.

Author contributions

Writing—original draft and writing—review & editing, Y. -l. W.; investigation, Q. -g. H.; data curation, F. -y. Q.; supervision, C. -c.
S.; funding acquisition, M. -l. Q.; resources, Y. -l. C. All authors have read and agreed to the published version of the manuscript.

Conflicts of interest
There are no conflicts to declare.

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