A pyrolysis-free path toward superiorly catalytic nitrogen-coordinated single atom

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Nitrogen-coordinated single-atom catalysts (SACs) have emerged as a frontier for electrocatalysis (such as oxygen reduction) with maximized atom utilization and highly catalytic activity. The precise design and operable synthesis of SACs are vital for practical applications but remain challenging because the commonly used high-temperature treatments always result in unpredictable structural changes and randomly created single atoms. Here, we develop a pyrolysis-free synthetic approach to prepare SACs with a high electrocatalytic activity using a fully π-conjugated iron phthalocyanine (FePc)–rich covalent organic framework (COF). Instead of randomly creating Fe-nitrogen moieties on a carbon matrix (Fe-N-C) through pyrolysis, we rivet the atomically well-designed Fe-N-C centers via intermolecular interactions between the COF network and the graphene matrix. The as-synthesized catalysts demonstrate exceptional kinetic current density in oxygen reduction catalysis (four times higher than the benchmark Pt/C) and superior power density and cycling stability in Zn-air batteries compared with Pt/C as air electrodes.

INTRODUCTION

Carbon-based catalysts with nitrogen-coordinated single transition metal atoms as active sites have emerged as promising systems for catalytic processes because of their maximal atom utilization and highly catalytic activity (1–8). As revealed both experimentally and theoretically, the nitrogen-coordinated single transition metal moieties supported in carbon provide versatile active sites, while the carbon matrix ensures a large specific area and efficient mass transport, provides a stable matrix for the metal atoms, and affects the electronic density owing to strong intermolecular interactions (9–16). Currently, reported strategies to prepare the N-coordinated single-atom catalysts (SACs) dominantly depend on high-temperature pyrolysis, which highly demands the rigorous manipulation of the carbonization process and precise elemental ratio of precursors (17–21). For instance, during the pyrolysis of zeolitic imidazolate frameworks to prepare catalysts with single Fe-N-C sites, slight changes in pyrolysis temperature will lead to notable differences in catalytic performance (22–25). Meanwhile, because the isolated metal atoms are thermodynamically unstable, for many situations, the metal atoms tend to agglomerate as their ratio in precursors increases but hardly form single-atom centers when the ratio decreases (14, 26). By this means, randomly creating Fe-N-C sites on the carbon matrix through pyrolysis is unpredictable and difficult to repeat in an atomic manner. There are many researches working on precisely tuning the nitrogen-coordinated SAC (25–28). However, developing precisely controllable synthesis of nitrogen-coordinated SACs in the long term for practical applications by means of high-temperature pyrolysis remains a challenge. To this end, we developed a pyrolysis-free synthetic approach SACs with a high electrocatalytic activity using a fully closed π-conjugated iron phthalocyanine (FePc)–rich covalent organic framework (COF).

RESULTS

Instead of randomly creating Fe-nitrogen moieties on the carbon matrix (Fe-N-C) through pyrolysis, we rivet the atomically well-designed Fe-N-C centers via intermolecular interactions between the Fe-N-C–rich pyrolysis-free COF network and the graphene matrix. Typically, as shown in Fig. 1A, we assembled benzene-1,2,4,5-tetra carbonitrile with Fe centers into a fully closed π-conjugated COF (termed COF_{BTC}) according to our previous method (29) to fabricate atomically dispersed Fe-N-C moieties (for additional details, see Materials and Methods), which were subsequently assembled with the graphene matrix via intermolecular interactions [see the extended x-ray absorption fine structure (EXAFS) and resistivity measurements discussed below]. Different from unpredictable pyrolysis, our single Fe-N-C centers were pre-assembled and directly riveted onto the graphene matrix; thus, the structures of the pyrolysis-free synthesized SACs (marked as pfSAC-Fe-X, where X represents the added mass ratio of COF_{BTC}) were well atomically arranged in the intact COF network. As expected, the graphene provided a stable matrix for the pre-assembled single Fe-N-C centers of COF_{BTC}. On the basis of the transmission electron microscopy (TEM) images (associated with elemental mapping), the as-obtained pfSAC-Fe demonstrated homogeneous elemental distribution (figs. S1 and S2), which was also confirmed by x-ray photoelectron spectroscopy (XPS) elemental analyses (fig. S3). We further performed atomic-resolution high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) and identified that the coordinated Fe atoms, as represented by bright dots, were exclusively in the single-atom format and uniformly anchored throughout the graphene matrix (Fig. 1B and fig. S4).

The riveting of single Fe-C-N centers was successfully realized via the intermolecular interactions between the COF_{BTC} and the graphene matrix. As investigated by x-ray absorption fine structure (XAFS) spectroscopy, the Fe K-edge of x-ray absorption near-edge structure (XANES) of these specimens suggested that the pfSAC-Fe contained near-edge structures similar to those of the original FePc but totally different from those of the Fe foil (Fig. 1C), indicating a N-coordinated chemical state of single Fe atoms. In the Fourier transform analyses, the curve of pfSAC-Fe demonstrated a main peak belonging to the Fe-N scattering paths, while the Fe-Fe peak at about 2.2 Å

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was not detected (Fig. 1D). We further simulated the Fe K-edge EXAFS, finding the signals of the two-body backscattering path of Fe-N in both pfSAC-Fe and COF BTC, while Fe-C was uniquely formed in the pfSAC-Fe (Fig. 1E and figs. S5 to S7). Accordingly, only van der Waals interaction was observed during the electron localization function (ELF) analysis (Fig. 2A) with the density functional theory (DFT) calculations. As confirmed by the comparison of charge density differences, the electrons of graphene were attracted to the N-coordinated Fe sites, forming an Fe-C electron pathway (Fig. 2B). Moreover, once oxygen was introduced, we observed that it was only absorbed onto the single Fe atoms (Fig. 2C), which offered superiorly catalytic sites for the oxygen reduction reaction (ORR) (vide infra).

Similar to randomly creating Fe-N-C centers on the carbon matrix through pyrolysis, our approach to rivet the pre-assembled Fe-N-C onto graphene also generated a facile electron transfer environment. In the powder electrical resistivity measurement, we observed that once the intermolecular interactions were constructed, the resistivity of as-prepared pfSAC-Fe and COF BTC decreased markedly, becoming very close to that of the raw graphene (Fig. 3A). By constructing the equivalent circuit based on the experimental data, we noted that the resistivity of the pfSAC-Fe-0.2 was in between the parallel connection and the series-parallel connection at low pressure (<12 MPa) while climbing higher than either the parallel connection or the series-parallel connection as the pressure increases (Fig. 3B). Because of the high pressing force leading to a much tighter connection and better conductivity, this phenomenon suggested that the intermolecular interactions generated an electrically conducting bridge between the Fe-N-C moieties and the graphene matrix, which was not as efficient as the conjugated systems indwelling the graphene, but still greatly benefitted electron transport. As observed from the density of state (DOS), the COF BTC was a semiconductor with a small direct bandgap. With the intermolecular interactions with graphene, the pfSAC-Fe exhibited excellent electrical conductivity similar to metal, which prominently enhanced the charge transfer (fig. S12). Furthermore, the work function of pfSAC-Fe-0.2 obtained from the ultraviolet photoelectron spectroscopy (UPS) spectra in the valence band emission region was as small as 4.60 eV (Fig. 3C), suggesting a high driving force for donating electrons from the catalytic surface and facilitating the catalytic performance during ORR (30, 31).

The electrocatalytic performance of pfSAC-Fe catalysts with different added ratios of COF BTC was evaluated with a traditional three-electrode system. In cyclic voltammetry (CV) and linear sweep voltammetry (LSV) measurements with a rotating disk electrode, all of the pfSAC-Fe catalysts exhibited excellent catalytic performance for ORR compared to the benchmark Pt/C (Fig. 4A). In addition, the performance of pfSAC-Fe was enhanced with the increase in the added ratio of COF BTC, demonstrating considerable high half-wave ($E_{1/2}$) potentials from 0.88 to 0.910 V versus reversible hydrogen electrode (RHE) (Fig. 4B). Meanwhile, the $E_{1/2}$ of the benchmark Pt/C was 0.86 V versus RHE, demonstrating the same activity as that in state-of-the-art literature (32–34). Consistent with the DFT
calculations, the single Fe atoms offered superior catalytic sites for the absorption and electron transfer of oxygen molecules during ORR. The optimized catalyst, pfSAC-Fe-0.2, exhibited the best ORR activity with a high kinetic current density ($J_k$) of 25.86 mA cm$^{-2}$ at 0.85 V (versus RHE), which was about four times higher than that of Pt/C (6.49 mA cm$^{-2}$), and with a low Tafel slope of 31.7 mV decade$^{-1}$ (Fig. 4, B and C). The efficient kinetics were also confirmed by the Koutecky-Levich (K-L) plots and rotating ring disk electrode (RRDE) test, revealing a highly efficient four-electron pathway (3.85 to 4.0 per O$_2$) and a low H$_2$O$_2$ yield (below 7%) over 0.5 to 0.9 V (Fig. 4, D and E) (35). We also noted that the electrocatalytic performance, such as half-wave potential and kinetic current density,
was enhanced with increased catalytic loading (fig. S13), which was consistent with pioneering work (36, 37). Superior to commercial Pt/C, the pfSAC-Fe catalysts exhibited better methanol tolerance during the methanol crossover test (Fig. 4F) and much longer durability during the continuous cycles (fig. S14A). The XAFS measurement of the pfSAC-Fe-0.2 test demonstrated similar curves after ORR tests, indicating that the structure of the catalysts was quite stable during the reaction (fig. S15). In addition, TEM and HAADF-STEM analyses confirmed that the single Fe atoms were still uniformly anchored throughout the graphene matrix after the ORR test (figs. S16 and S17).

Furthermore, in the homemade Zn-air battery with pfSAC-Fe-0.2 as the electrocatalysts for air cathode (Fig. 5A), we observed an open circuit potential as high as 1.41 V (Fig. 5B). Compared with the battery using conventional Pt/C as the electrocatalysts, the Zn-air battery driven by pfSAC-Fe-0.2 delivered similar discharge behaviors, higher power density (123.43 mW cm⁻², compared to 113.81 mW cm⁻²), the power density of the battery driven by Pt/C), and larger specific capacity (732 mAh g⁻¹ at 100 mA cm⁻² with a zinc utilization of 89.3%) at the same loading (0.2 mg cm⁻²) (Fig. 5C and D). Besides, the Zn-air battery based on pfSAC-Fe-0.2 exhibited a notably long life cycle over 300 hours with a less than 0.1% decrease, suggesting the considerable stability of the pfSAC-Fe and the practical potential of the as-designed batteries, while the discharge voltage of the Pt/C-based system markedly decreased within 20 hours under the same conditions (Fig. 5E). We further periodically changed the current density from 5 to 40 mA cm⁻² to evaluate the stability of cycle discharge performance and only observed a small voltage difference over 30 hours (Fig. 5F). Unexpectedly, even when the current density was as high as 100 mA cm⁻², the battery was still able to operate for more than 300 min with little potential drop (Fig. 5G). By increasing the loading of pfSAC-Fe catalysts to 0.5 mg cm⁻², the peak power density of the battery was raised up to 126.83 mW cm⁻² (fig. S23), demonstrating a promising growth potential to replace the commercial Pt/C catalysts for metal-air batteries with high power density.

**DISCUSSION**

Rather than randomly creating single atoms on carbon through pyrolysis, we developed a pyrolysis-free synthetic approach to prepare SACs by riveting the pre-assembled COF BTC on a conductive matrix (e.g., graphene) via intermolecular interactions. The as-synthesized pfSAC-Fe demonstrated a small work function and highly efficient catalytic performance for ORR. The Zn-air battery driven by the pfSAC-Fe exhibited superior power density and cycling stability for practical applications. The invented pyrolysis-free approach toward SACs not only paves a promising way to atomically optimize the SAC structures but also simplifies the accurate theoretical prediction calculations based on well-defined structures to develop more excellent catalysts for various energy conversions and storage devices (e.g., CO₂ reduction, N₂ fixation, and photoelectronic catalysis) besides oxygen reduction.
MATERIALS AND METHODS

Synthesis of pfSAC-Fe-X

To synthesize the pfSAC-Fe-X electrocatalysts, different mass ratios of as-synthesized COF<sub>RTC</sub> (1, 5, 10, and 20%) were dispersed in glycol solution, followed by grinding with corresponding amounts of graphene. After grinding the mixture for 1 hour, the as-obtained composite material was kept in argon at 250°C to remove the glycol solvent. The as-prepared composite electrocatalysts were marked as pfSAC-Fe-X (X = 0.01, 0.05, 0.1, and 0.2, which represents the added mass ratios of the COF<sub>RTC</sub>).
DFT calculation
We used the Vienna ab initio simulation package herein for all DFT calculations. To describe the interactions of the core electrons within different atoms, projected augmented wave was used. For exchange-correlation interactions, we adopted the generalized gradient approximation by Perdew-Burke-Ernzerhof. The plane wave basis sets had a cutoff energy of 500 eV to describe all atoms' valence electrons. The total energy of all the structures converged to $10^{-4}$ eV, and the Hellmann-Feynman force converged to 0.02 eV/Å. The Brillouin zone was set as the γ point. The graphene and COF-BTC were optimized separately before building the composite structure model, which was built by periodical single-layer graphene-based systems with a 15-Å vacuum to avoid the interactions between the layers. The lattice constants of these models were 19.7 Å by 19.7 Å by 15 Å. Van der Waals correction was applied in all composite structure calculations. ISPIN values were considered when calculating the models with magnets.

Assembly of the Zn-air battery
Zn-air battery was assembled with homemade frameworks. On the air electrode, the catalysts were coated along with a gas diffusion layer at an effective size of 1.0 cm by 1.0 cm. A polished zinc foil was assembled as the negative electrode with the same effective size. The electrolyte was composed of 8 M KOH and 0.5 M ZnO. The battery performance was tested using a constant current charging-discharging process by Land Instruments at room temperature.

SUPPLEMENTARY MATERIALS
Supplementary material for this article is available at http://advances.sciencemag.org/cgi/content/full/5/8/eaaw2322/DC1

Table S3. Inductively coupled plasma atomic emission spectroscopy results for Fe contents in COF-BTC and the as-obtained pSAC-Fe-0.2.

Table S4. Comparison of the performance of the zinc-air batteries based on pSAC-Fe-0.2 and other electrocatalysts.

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