Abstract: High-performance non-precious metal catalysts exhibit high electrocatalytic activity for the oxygen-reduction reaction (ORR), which is indispensable for facilitating the development of multifarious renewable energy systems. In this work, N-doped carbon-encapsulated Fe₃C nanosphere ORR catalysts were prepared through simple carbonization of iron precursors loaded with polyacrylamide microspheres. The effect of iron precursors loading on the electrocatalytic activity for ORR was investigated in detail. The electrochemical measurements revealed that the N-doped carbon-encapsulated Fe₃C nanospheres exhibited outstanding electrocatalytic activity for ORR in alkaline solutions. The optimized catalyst possessed more positive onset potential (0.94 V vs. reversible hydrogen electrode (RHE)), higher diffusion limiting current (5.78 mA cm⁻²), better selectivity (the transferred electron number n > 3.98 at 0.19 V vs. RHE) and higher durability towards ORR than a commercial Pt/C catalyst. The efficient electrocatalytic performance towards ORR can be attributed to the synergistic effect between N-doped carbon and Fe₃C as catalytic active sites; and the excellent stability results from the core-shell structure of the catalysts.

Keywords: polyacrylamide microspheres; N-doped carbon; Fe₃C; core-shell structure; oxygen reduction reaction

1. Introduction

With the limited supply of fossil fuels and growing concern about environmental problems, the development of renewable energy and efficient energy conversion systems has become more important than ever. Fuel cells are regarded as one of the most promising energy conversion devices accessible nowadays [1–6]. Among various fuel cells, proton exchange membrane fuel cells (PEMFCs) and polymer electrolyte fuel cells (PEFCs) are considered as advanced electrochemical energy conversion systems due to their advantages of fast electrode reaction kinetics, high conversion efficiency and environmental friendliness [7,8]. In fuel cells, small molecule fuels are oxidized at the anode and concurrently oxygen is reduced at the cathode [9]. Efficient catalysts are necessary due to the sluggish kinetics of the oxygen-reduction reaction (ORR) at the cathode which is of great limit of the energy
conversion efficiency of fuel cells. Pt and Pt-based materials have been used extensively as effective catalysts for ORR, which have extremely low overpotential as well as high current density in the ORR process [9–12]. However, the high price as well as the relatively low stability has been the main bottlenecks that hinder their broader commercialization [13,14]. Therefore, the development of low-cost and high-performance alternatives, especially non-precious metal catalysts, have aroused extensive research interest. Among these potential alternatives, N-doped carbon and transition metal species have attracted much attention due to the relatively low cost and promising electrocatalytic activity to ORR [15–19]. Furthermore, specific surface area and structure of the catalysts were found to determine the accessibility of the active sites and can largely improve the ultimate catalytic performance [20–24].

Recently, non-precious metal catalysts based on metal carbides, such as Fe3C, coated with N-doped carbon nanostructures, have been proved to be very efficient for ORR. The N-doped carbon-encapsulated structures offer a specific synergistic effect of Fe3C active sites and N-doped carbon matrix [25–28]. In addition, this kind of structure partly prevents metal composite from corrosion in electrolytes, which is necessary for long-term performance [29]. Typically, the N-doped carbon encapsulated Fe3C composite was prepared by Fe-MOF or a molecular precursor containing Fe, nitrogen and carbon precursor [30–32]. G. Ren [33] and R. Zhong [34] reported that porous core-shell Fe3C embedded N-doped carbon nanofibers were synthesized by the electrospinning method. The resulting nanofibers catalysts showed excellent ORR activities and high stabilities, which were attributed to well-dispersed nanocrystalline Fe3C active sites, high N-doping level, large surface areas, and a one-dimensional carbon nanostructure. Therefore, the development of a facile strategy for designing and synthesizing core-shell structured catalysts with high surface areas and abundant active sites is highly desirable for the further development of the N-doped carbon encapsulated metal composite ORR catalysts.

Herein, we report a facile and controllable fabrication of N-doped microporous carbon encapsulated Fe3C nanospheres derived from iron precursor-loaded polyacrylamide microspheres. Polyacrylamide microspheres act as the carbon and nitrogen precursor and possess excellent hydroscopicity, which can adsorb iron salts and ensure their uniform distribution. The amounts of the active sites are regulated by adsorbing different amount of iron precursors before carbonization of polyacrylamide microspheres and the porous structures are obtained by activation of KOH. The obtained catalysts are highly microporous with well-dispersed Fe3C nanospheres in the N-doped graphite carbon matrix. As expected, Fe3C@N-doped microporous carbon nanospheres exhibited better ORR electrocatalytic activity in alkaline media compared to the commercial Pt/C catalyst.

2. Experimental Section

2.1. Materials and Reagents

Tween-60 (C32H62O10), N,N'-Methylenebisacrylamide (C6H10N2O2, 98.0%), ammonium persulfate ((NH4)2S2O8, 98.5%), acrylamide (C3H5NO, 98.0%), Potassium hydroxide (KOH, >96.0%) and isopropanol (C3H8O, >99.7%) were obtained from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China). Ferroporphyrin (C34H32ClFeN4O4, 98.0%), Span-80 (C24H44O6) and mineral oil were purchased from Shanghai Yuanye Biological Co., Ltd, Shanghai, China). 20 wt % Pt/C catalyst and 5 wt % Nafion solution were obtained from Shanghai Macklin Biochemical Co., Ltd, Shanghai, China). All the chemicals were used as received without further purification.

2.2. Synthesis of Fe3C@N-Doped Carbon Nanospheres

Polyacrylamide microspheres were firstly synthesized by inverse emulsion polymerization. Briefly, 50 mL mineral oil and 6.2 g of Span-80 were mixed for 10 min to act as oil phase. 20.0 g of acrylamide monomer, 1.8 g of Tween-60 and 1 mL N,N'-Methylenebisacrylamide solution (0.04 wt %) were dissolved into 25 mL deionized water and then 1 mL ammonium persulfate solution (0.4 wt %) was dripped after stirring for 5 minutes, the mixture solution was served as water phase. The water phase was added dropwise into the oil phase with stirring at the speed of 600 r/min. After 30 min of vigorous
stirring, the mixed solution was put into a 55 °C water bath and the N₂ was aerated continuously to remove the air for about 40 min. Then the mixture was kept in a 55 °C water bath with stirring at the speed of 300 r/min in nitrogen atmosphere for 3 h to obtain crosslink-polyacrylamide reverse-phase microemulsion. The resulting w/o microemulsion was poured into excess acetone for demulsification. After washing with deionized water and absolute ethyl alcohol, the collected solids were dried at 70 °C overnight to obtain polyacrylamide microspheres. In order to obtain carbonization yield of polyacrylamide microspheres, in preliminary experiment, 2 g of dried polyacrylamide microspheres was pyrolyzed under flowing N₂ for 1 h at 900 °C with a heating rate of 5 °C min⁻¹ and 0.4 g of carbon was collected. This indicates that the carbonization yield of polyacrylamide microspheres was about 20%.

Fe₃C@N-doped carbon nanospheres were prepared by direct carbonization of iron precursors-loaded polyacrylamide microspheres. In a typical preparation process, a certain amount of ferroporphyrin and 1.6 g of KOH solid were dissolved in 15 mL deionized water, the mixture solution was completely absorbed by 4 g dried polyacrylamide microspheres. The KOH was added to dissolve the Ferroporphyrin and then activate the carbon. The sample was dried at 70 °C for 24 h and calcinated at 900 °C for 1 h filled with N₂ atmosphere and the ramping rate was 5 °C min⁻¹. The resulting dark powder was washed with deionized water for several times to remove alkali metal K which was produced during the activation and calcination process, followed by drying at 70 °C overnight. The final as-synthesized product has been obtained and labeled as Fe₃C@N/C-x, and x represents the mass ratio of ferroporphyrin and carbon according to the theoretical dosage. For comparison, a series of Fe₃C@N/C-x composites were also prepared via the same procedure except for the difference of ferroporphyrin additive amounts. These as-synthesized catalysts with various ferroporphyrin additive amounts were labeled as Fe₃C@N/C-0, Fe₃C@N/C-0.5, Fe₃C@N/C-1, Fe₃C@N/C-2, respectively.

2.3. Characterizations

Transmission electron microscope (TEM) images were collected using a JEM-2100F TEM (Tokyo, Japan), XRD measurements were measured with a D8 Advance X-ray diffractometer from Bruker AXS Company (Karlsruhe, Germany). X-ray photoelectron spectroscopy (XPS) analysis was recorded on a VG Multilab 2000 X-ray photoelectron spectrometer (VG Scientific, Waltham, MA, USA). The pore structure analysis was performed through nitrogen adsorption and desorption isotherm measurements (BET, ASAP 2020, Micromeritics). The dynamic function theory (DFT) method was applied to calculate the distribution of the micropore size.

2.4. Electrochemical Measurements

To evaluate ORR activity, the electrochemical performance of Fe₃C@N-doped microporous carbon nanospheres catalysts were performed in a standard three-electrode glass cell with N₂ or O₂-saturated 0.1M KOH solution at room temperature on the electrochemical workstation (CHI660E, CHI instrument) using a glassy carbon electrode (GCE, of 5.00 mm in diameter) as working electrode, a platinum wire as counter electrode, and Hg/HgO electrode as reference electrode, respectively. The electrochemical properties of as-synthesized catalysts were referred to the Hg/HgO electrode. The Hg/HgO reference electrodes were calibrated with respect to the reversible hydrogen electrode (RHE) before measurement [2]. The calibration values were $E_{\text{RHE}} = E_{\text{Hg/HgO}} + 0.89 \text{ V}$ in 0.1M KOH. 5.0 mg of as-synthesized catalysts power, 0.1 mL of deionized water, 0.9 mL of isopropanol and 0.02 mL of 5 wt % nafion solution was ultrasonically mixed to form the ink of the catalysts. The GCE was modified with 20 µL of the ink to serve as the working electrode. Cyclic voltammetry (CV) and linear sweep voltammetry (LSV) were carried out to evaluate the ORR performances. CV curves were obtained in N₂ or O₂-saturated 0.1M KOH electrolyte solutions without any rotation. The CV measurement data were carried out in the potential range from -0.6 V to 0.4 V at a sweep rate of 50 mV s⁻¹, and LSV measurements were carried out at a scan rate of 5 mV s⁻¹ in the potential range from -0.8 V to 0.2 V under various electrode rotation rates (400, 800, 1200, 1600, 2000 rpm).
in O$_2$-saturated 0.1M KOH electrolyte solutions. All samples were also tested in N$_2$-saturated for comparison. Chronoamperometric measurements for each as-synthesized catalyst were investigated in O$_2$-saturated 0.1M KOH (−0.3 V vs. Hg/HgO) at room temperature. The ORR process kinetics was analyzed using the Koutecky–Levich (K–L) equation.

3. Results and Discussion

The morphology of polyacrylamide microspheres and the N-doped carbon encapsulated Fe$_3$C nanospheres catalysts were characterized by TEM. Figure 1a shows the surface morphology of polyacrylamide microspheres. The results showed that the polyacrylamide microspheres are aggregates of spherical particles with an average diameter of about 30 nm. After adsorbing iron precursors and carbonization, the spherical morphology of samples can be maintained. The TEM images of Fe$_3$C@N/C-1 were shown in Figure 1b,c. It can be observed that the core-shell structured catalysts were successfully synthesized. The high-resolution TEM (HRTEM) results in Figure 1d showed that the spacing between adjacent lattice fringe in nanospheres core was 0.21 nm, corresponding to the (211) crystal planes of Fe$_3$C. The lattice fringe in outer shell was 0.348 nm, corresponding to the graphite (002) plane. It indicated that the Fe$_3$C nanospheres were obtained and wrapped within graphite carbon layer during the carbonization process.

![Figure 1](https://via.placeholder.com/150)

**Figure 1.** (a) Transmission electron microscope (TEM) image of polyacrylamide microspheres; (b,c) TEM and (d) high-resolution TEM (HRTEM) image of Fe$_3$C@N/C-1.

In order to further identify the crystal structure of samples, X-ray diffraction (XRD) patterns of these samples were recorded and shown in Figure 2. The major peaks located at 37.8°, 43.1°, 43.1°, 43.9°,
44.9°, 46.0°, 48.7° and 49.3° correspond to the Fe₃C phase [35]. In addition, the peaks at 26.4° and 44.0° were assigned to the (002) and (004) reflection of graphitic carbon [26]. There are only broad graphitic carbon diffraction peaks in Fe₃C@N/C-0 sample. With the introduction of ferroporphyrin, the graphitic carbon and Fe₃C crystalline phases coexisted in the other samples. The relative intensity of Fe₃C diffraction peaks increased with the iron content further enhanced in the synthesis process, indicating an increased ratio of Fe₃C. The Fe₃C nanocrystalline should originate from the reduction of iron precursor by carbon during the pyrolysis process. In the synthesis process of catalysts, ferroporphyrin was firstly dissolved in the KOH solution and homogeneously adsorbed in the polyacrylamide microspheres. Iron compound could be reduced by carbon to form metallic iron at high temperature, and these metallic iron atoms contribute to the catalytic graphitization of carbon [36]. When the iron atoms concentration is high enough, it would aggregate to form crystalline iron which react with the carbon atom to form a graphite layer-encapsulated Fe₃C crystal during the process of pyrolysis and cooling. With the growth of Fe₃C crystal, it is difficult for the large-size Fe₃C crystal to make the inside carbon atoms diffuse onto the surface. Therefore, the relative intensity of graphitic carbon diffraction peaks weakens with the increase of iron content in the catalysts [35–39].

XPS was conducted to illustrate the surface elements of catalysts. XPS spectra of Fe₃C@N/C catalysts are exhibited in Figure 3a, which revealed the existence of C, O, Fe and N in catalysts. There is only a weak Fe peak was detected in catalysts, which could be resulted from the coverage of graphitic layers on the Fe₃C surface [33,39]. The high-resolution C 1s spectrum of Fe₃C@N/C-1 shown in Figure 3b can be deconvoluted into four individual peaks that are assigned to C–C (284.6 eV), C–O (285.2 eV), C=O (286.5 eV), C–N (288.2 eV), respectively. The N 1s spectrum of Fe₃C@N/C-1 shown in Figure 3c can be divided into three peaks, assigned to pyridinic N (398.4 eV), pyrrolic N (400.1 eV), graphitic N (401.2 eV), suggesting that nitrogen has been doped into the carbon authentically. It is generally believed that pyridinic N and graphitic N can serve as the efficient active sites for ORR. Pyridinic N increases the spin density and the density of π states of the carbon atoms near the Fermi level which could enhance the reduction of O₂, while graphitic N can increase the conductivity of the catalysts [35,40,41]. In the case of the high-resolution Fe 2p spectrum (Figure 3d), the peaks at 711.0 eV and 724.4 eV are consistent with Fe 2p₃/2 and Fe 2p₁/2 states, respectively [42–44]. Fe, C and N content, N/C weight ratios and the relative atomic amount of N species in all of Fe₃C@N/C-x samples are shown in Table 1. The amount of Fe incorporated into the final catalyst increases with the enhancement of ferroporphyrin loading during the synthesis process, while the amount of N declines slightly at the same time. In all cases, graphitic N and pyridinic N are dominant species. In general, with the
increase of the amount of Fe$_3$C catalytic active sites, the amount of N-doped carbon catalytic active sites declines.

![Figure 3.](image)

**Figure 3.** (a) X-ray photoelectron spectroscopy (XPS) survey spectra of Fe$_3$C@N/C-x samples. (b) C 1s, (c) N 1s and (d) Fe 2p spectra of Fe$_3$C@N/C-1.

**Table 1.** Fe, C and N content, N/C weight ratios and the relative atomic amount of N species in all of the Figure 3 samples derived from XPS analyses.

| Sample         | Weight Content (%) | N/C Weight Ratio | Relative Atomic Amount of N Species |
|----------------|--------------------|------------------|-------------------------------------|
|                | C  | N  | Fe  | Pyridinic N | Pyrrolic N | Graphitic N |
| Fe$_3$C@N/C-0  | 87.03 | 8.26 | 0.00 | 0.094 | 0.32 | 0.27 | 0.41 |
| Fe$_3$C@N/C-0.5| 87.96 | 7.18 | 0.78 | 0.082 | 0.35 | 0.26 | 0.39 |
| Fe$_3$C@N/C-1  | 88.68 | 6.82 | 1.02 | 0.080 | 0.36 | 0.25 | 0.39 |
| Fe$_3$C@N/C-2  | 89.72 | 6.13 | 1.25 | 0.068 | 0.37 | 0.22 | 0.41 |

The nitrogen adsorption and desorption isotherms were conducted to investigate the pore structure of these catalyst samples and the results are shown in Figure 4a. The nitrogen adsorption and desorption isotherm curves of all catalysts reveal a Type-I sorption isotherm with no hysteresis, manifesting the microporous structure of the Fe$_3$C@N/C-x samples. In the synthesis process, KOH was used not only as an alkaline medium to dissolve ferroporphyrin, but also as an activator to form micropores. The abundant micropores and high specific surface area can improve the accessibility of electrolyte and oxygen to the active sites. The specific surface area and pore volume of all samples are shown in Table 2. With the increase of iron content, the total surface area of Fe$_3$C@N/C-x catalysts reduce from 2484 to 1687 m$^2$ g$^{-1}$. Figure 4b illustrates the pore size distribution (PSD) curves of Fe$_3$C@N/C-x catalysts. The results of PSDs are greatly distributed in the microporous region which is less than 2.0 nm and the peaks are mostly 0.78 nm and 0.59 nm.
which is essential for ORR. Therefore, the enhanced electrocatalytic performance of Fe \(_3\)C@N/C-1 catalyst is compared to some other Fe \(_3\)C-based electrocatalysts reported in literature. Overall, the Fe \(_3\)C@N/C-1 catalyst possessed comparative onset potential with other Fe \(_3\)C based electrocatalysts. Moreover, the diffusion limiting current of the Fe \(_3\)C@N/C-1 catalyst was higher than that of most catalysts, indicating the excellent electrochemical activity towards ORR.

CV and LSV measurements were performed to study the electrochemical activity of these catalysts. Figure 5 shows the CV curves of Fe\(_3\)C@N/C-1 catalyst and 20 wt % Pt/C in N\(_2\) and O\(_2\)-saturated 0.1 M KOH solution. Well-defined oxygen reduction peak for Fe\(_3\)C@N/C-1 catalyst and 20 wt % Pt/C could be seen in the CV curves when saturating the alkaline solution with O\(_2\), illustrating the pronounced ORR activity of Fe\(_3\)C@N/C-1 catalyst. LSV curves were carried out on a rotating disk electrode (RDE) for the further evaluation of the electrocatalytic activity of catalyst samples for ORR. The LSV measurement was tested in O\(_2\)-saturated 0.1M KOH solution at a rotation rate of 1600 rpm with a potential scan rate of 5 mV s\(^{-1}\) (the solid line). All samples were also tested in N\(_2\)-saturated for comparison (the same colour scheme as the dotted line). As shown in Figure 6, the Fe\(_3\)C@N/C-1 catalyst with onset potential of 0.94 V (vs. RHE) and diffusion-limiting current of 5.78 mA cm\(^{-2}\) reveals the remarkable catalytic performance towards ORR compared to the benchmark Pt/C (0.91 V and 5.55 mA cm\(^{-2}\) correspondingly). By contrast, the Fe\(_3\)C@N/C-0 catalyst shows the lowest catalytic activity to ORR. It is reported that the encapsulated metal compound’s nanoparticles could produce host–guest electronic interaction and change the local work function of the carbon shell, creating additional active sites to ORR [33]. Doping of nitrogen into the carbon shell could further improve the catalytic activity of these core-shell structured catalysts by modifying the electronic properties of carbon surface. In addition, there is a general agreement that N-doped graphitic carbon is playing an important role in enhancing the conductivity of the catalysts which is essential for ORR. Therefore, the enhanced electrocatalytic performance of Fe\(_3\)C@N/C-1 can be attributed to the synergistic effect between N-doped carbon and Fe\(_3\)C as catalytic active sites. It should be noted that Fe\(_3\)C@N/C-2 showed obviously lower electrocatalytic activity than that of Fe\(_3\)C@N/C-1, due to the reduced specific surface area and depressed graphitization degree of the carbon shell in the Fe\(_3\)C@N/C-2 catalyst. In Table 3 below, the performance of the Fe\(_3\)C@N/C-1 catalyst is compared to some other Fe\(_3\)C-based electrocatalysts reported in literature. Overall, the Fe\(_3\)C@N/C-1 catalyst possessed comparative onset potential with other Fe\(_3\)C based electrocatalysts. Moreover, the diffusion limiting current of the Fe\(_3\)C@N/C-1 catalyst was higher than that of most catalysts, indicating the excellent electrochemical activity towards ORR.

**Figure 4.** (a) Nitrogen adsorption and desorption curves. (b) Pore size distribution of Fe\(_3\)C@N/C-x samples.

**Table 2.** The specific surface area and pore volume of Fe\(_3\)C@N/C-x samples.

| Sample      | \(S_{BET}\) (m\(^2\) g\(^{-1}\)) | \(V_{micro}\) (cm\(^3\) g\(^{-1}\)) |
|-------------|-----------------------------------|-------------------------------------|
| Fe\(_3\)C@N/C-0 | 2484.37                           | 0.76                                |
| Fe\(_3\)C@N/C-0.5 | 2121.87                           | 0.71                                |
| Fe\(_3\)C@N/C-1 | 1967.83                           | 0.68                                |
| Fe\(_3\)C@N/C-2 | 1687.45                           | 0.58                                |
Fe/C HN-700C-60M 0.1 M KOH 1600 0.98 5.95 [45]
Fe@C-NG/NCNTs 0.1 M KOH 1600 0.93 5.11 [46]
Fe@C-NG@C-1 catalyst count from the slope of K–L plots is 3.85–4.00, which is close to 4, suggesting the Fe@C-NG@C-1 catalyst catalyze ORR process through a quasi-four-electron process. 

Fe3C@NCNF-900 0.1 M KOH 1600 0.93 4.51 [33]

Fe3C/b-NCNT 0.1 M KOH 1600 0.96 6.25 [35]
Fe3C/b-NCNF 0.1 M KOH 1600 1.012 4.81 [34]
Fe3C/C-700 0.1 M KOH 1600 0.89 4.21 [31]
PMF-800 0.1 M KOH 1600 0.95 5.78 [29]

RDE measurement of Fe3C@N/C-1 catalyst was evaluated under different rotation speeds from 400 rpm to 2000 rpm and the results are shown in Figure 7a. The Koutecky–Levich plots (Figure 7b) were calculated by the recorded reaction currents at ~0.3 V on the LSV curves of the Fe3C@N/C-1 catalyst under various rotating speeds to study the reaction kinetics of this catalyst. The electron-transfer numbers of the Fe3C@N/C-1 catalyst were calculated by the K–L equation [47]. The n value for Fe3C@N/C-1 catalyst count from the slope of K–L plots is 3.85–4.00, which is close to 4, suggesting the Fe3C@N/C-1 catalyze ORR process through a quasi-four-electron process. H2O2 was released during 2e− process which degrades the membrane electrolyte; therefore, the 4e− process is desired for a fuel cell.
The stability of catalyst is necessary for the practical application of fuel cells. The chronoamperometric measurements were investigated to evaluate the durability of the Fe₃C@N/C-1 catalyst. The glassy-carbon electrode modified with Fe₃C@N/C-1 and commercial Pt/C were tested at constant voltage of −0.3 V in an O₂-saturated 0.1 M KOH aqueous solution with rotation rate of 1600 rpm for 20,000 s. The current-time response is shown in Figure 8b. As we can see, the current-time chronoamperometric response for commercial Pt/C exhibited a rapid current decrease and the current loss is about 25% after 20,000 s. In comparison, the current retention for the Fe₃C@N/C-1 catalyst is 94% after 20,000 s, implying the excellent stability under working conditions. The excellent stability of Fe₃C@N/C-1 catalyst may in virtue of the appearance of the sufficient outer graphene layer. Thus, it can be concluded that the Fe₃C@N-doped microporous carbon nanosphere catalyst is a promising alternative for costly Pt-based electrocatalysts to apply in fuel cells.

![Figure 7](image1.png)

**Figure 7.** (a) LSV curves of Fe₃C@N/C-1 catalyst with different rotating speed from 400 to 2000 rpm. (b) Koutecky–Levich (K–L) plots of Fe₃C@N/C-1 catalyst calculated from Figure 7a.

![Figure 8](image2.png)

**Figure 8.** (a) Tafel plots of Fe₃C@N/C-1 catalyst and Pt/C obtained from the rotating disk electrode (RDE) measurements. (b) current-time chronoamperometric response of Fe₃C@N/C-1 catalyst and Pt/C in O₂-saturated 0.1 M KOH at −0.3 V for 20000s.
4. Conclusions

In summary, Fe$_3$C@N-doped microporous carbon nanosphere catalysts were facilely and successfully synthesized by using polyacrylamide microspheres as a carbon and nitrogen resource and ferroporphyrin as an iron resource. The composite and nanostructure of catalysts were characterized by TEM, XRD, XPS and BET analysis and the electrochemical activity of these catalysts were investigated by CV and LSV measurements. The Fe$_3$C@N/C-1 catalyst showed excellent ORR activity and long-term stability in alkaline medium, with an onset potential and diffusion limiting current of 0.94 V vs. RHE and 5.78 mA cm$^{-2}$, which is superior to that of commercial Pt/C catalyst. The outstanding performance is attributed to the synergistic effect between N-doped carbon and Fe$_3$C as catalytic active sites, and the abundant microporous structure can improve the accessibility of electrolyte and oxygen to active sites. In addition, the carbon shell prevents Fe$_3$C active sites from leaching out. These results show that this kind of core-shell catalyst could be a promising alternative to Pt-based catalysts for the further development of durable and efficient ORR catalysts.

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References
1. Steele, B.C.H.; Heinzel, A. Materials for fuel-cell technologies. *Nat. Cell Boil.* 2001, 414, 345–352. [CrossRef] [PubMed]
2. Sa, Y.J.; Seo, D.J.; Woo, J.; Lim, J.T.; Cheon, J.Y.; Yang, S.Y.; Lee, J.M.; Kang, D.; Shin, T.J.; Shin, H.S.; et al. A General Approach to Preferential Formation of Active Fe-Nx Sites in Fe-N/C Electro catalysts for Efficient Oxygen Reduction Reaction. *J. Am. Chem. Soc.* 2016, 138, 15046–15056. [CrossRef]
3. Gasteiger, H.A.; Kocha, S.S.; Sompalli, B.; Wagner, F.T. Activity benchmarks and requirements for Pt, Pt-alloy, and non-Pt oxygen reduction catalysts for PEMFCs. *Appl. Catal. B: Environ.* 2005, 56, 9–35. [CrossRef]
4. Debe, M.K. Electrocatalyst approaches and challenges for automotive fuel cells. *Nat. Cell Boil.* 2012, 486, 43–51. [CrossRef] [PubMed]
5. Shao, M.; Chang, Q.; Dodelet, J.-P.; Chenitz, R. Recent Advances in Electro catalysts for Oxygen Reduction Reaction. *Chem. Rev.* 2016, 116, 3594–3657. [CrossRef]
6. Mistry, H.; Varela, A.S.; Kühl, S.; Strasser, P.; Cuema, B.R. Nanostructured electro catalysts with tunable activity and selectivity. *Nat. Rev. Mater.* 2016, 1, 16009. [CrossRef]
7. Vinothkannan, M.; Kim, A.R.; Kumar, G.G.; Yoo, D.J. Sulfonated graphene oxide/Nafion composite membranes for high temperature and low humidity proton exchange membrane fuel cells. *RSC Adv.* 2018, 8, 7494–7508. [CrossRef]
8. Vinothkannan, M.; Kim, A.R.; Kumar, G.G.; Yoon, J.-M.; Yoo, D.J. Toward improved mechanical strength, oxidative stability and proton conductivity of an aligned quadratic hybrid (SPEEK/FPAPB/Fe 3 O 4 -FGO) membrane for application in high temperature and low humidity fuel cells. *RSC Adv.* 2017, 7, 39034–39048. [CrossRef]
9. Niu, W.; Li, L.; Liu, X.; Wang, N.; Liu, J.; Zhou, W.; Tang, Z.; Chen, S. Mesoporous N-Doped Carbons Prepared with Thermally Removable Nanoparticle Templates: An Efficient Electrocatalyst for Oxygen Reduction Reaction. *J. Am. Chem. Soc.* 2015, 137, 5555–5562. [CrossRef]
10. Ramakrishnan, S.; Karuppannan, M.; Vinothkannan, M.; Ramachandran, K.; Kwon, O.J.; Yoo, D.J. Ultrathin Pt Nanoparticles Stabilized by MoS2/N-Doped Reduced Graphene Oxide as a Durable Electrocatalyst for Alcohol Oxidation and Oxygen Reduction Reactions. *ACS Appl. Mater. Interfaces* 2019, 11, 12504–12515. [CrossRef] [PubMed]
11. Arukula, R.; Vinothkannan, M.; Kim, A.R.; Yoo, D.J. Cumulative effect of bimetallic alloy, conductive polymer and graphene toward electrooxidation of methanol: An efficient anode catalyst for direct methanol fuel cells. *J. Alloy. Compd.* **2019**, *771*, 477–488. [CrossRef]

12. Kang, Y.J.; Ye, X.C.; Chen, J.; Cai, Y.; Diaz, R.E.; Adzic, R.R.; Stach, E.A.; Murzyn, C.B. Design of Pt-Pd Binary Superlattices Exploiting Shape Effects and Synergistic Effects for Oxygen Reduction Reactions. *J. Am. Chem. Soc.* **2013**, *135*, 42–45. [CrossRef]

13. Proch, S.; Wirth, M.; White, H.S.; Anderson, S.L. Strong Effects of Cluster Size and Air Exposure on Oxygen Reduction and Carbon Oxidation Electrocatalysis by Size-Selected Pt-n (n ≤ 11) on glassy carbon electrodes. *J. Am. Chem. Soc.* **2013**, *135*, 3073–3086. [CrossRef]

14. Yang, H.; Geng, L.; Zhang, Y.; Chang, G.; Zhang, Z.; Liu, X.; Lei, M.; He, Y. Graphene-templated synthesis of palladium nanoplates as novel electrocatalyst for direct methanol fuel cell. *Appl. Surf. Sci.* **2019**, *466*, 385–392. [CrossRef]

15. Liang, H.-W.; Wei, W.; Wu, Z.-S.; Meng, K.; Müllen, K. Mesoporous Metal–Nitrogen-Doped Carbon Electro catalysts for Highly Efficient Oxygen Reduction Reaction. *J. Am. Chem. Soc.* **2013**, *135*, 16002–16005. [CrossRef]

16. Yin, H.; Zhang, C.; Liu, F.; Hou, Y. Hybrid of Iron Nitride and Nitrogen-Doped Graphene Aerogel as Synergistic Catalyst for Oxygen Reduction Reaction. *Adv. Funct. Mater.* **2014**, *24*, 2930–2937. [CrossRef]

17. Serov, A.; Artyushkova, K.; Atanassov, P. Fe-N-C Oxygen Reduction Fuel Cell Catalyst Derived from Car bendazim: Synthesis, Structure, and Reactivity. *Adv. Nat. Mater.* **2014**, *4*, 1301735. [CrossRef]

18. Lin, L.; Zhu, Q.; Xu, A.-W. Noble-Metal-Free Fe–N/C Catalyst for Highly Efficient Oxygen Reduction Reaction under Both Alkaline and Acidic Conditions. *J. Am. Chem. Soc.* **2014**, *136*, 11027–11033. [CrossRef] [PubMed]

19. Chung, H.T.; Won, J.H.; Zelenay, F. Active and stable carbon nanotube/nanoparticle composite electrocatalyst for oxygen reduction. *Nat. Commun.* **2013**, *4*, 1922. [CrossRef]

20. Wang, H.; Liu, R.; Li, Y.; Lü, X.; Wang, Q.; Zhao, S.; Yuan, K.; Cui, Z.; Li, X.; Xin, S.; et al. Durable and Efficient Hollow Porous Oxide Spinel Microspheres for Oxygen Reduction. * Joule* **2018**, *2*, 337–348. [CrossRef]

21. Wu, Z.; Ly, Y.; Xia, Y.; Webley, P.A.; Zhao, D. Ordered Mesoporous Platinum@Graphitic Carbon Embedded Nanophase as a Highly Active, Stable, and Methanol-Tolerant Oxygen Reduction Electro catalyst. *J. Am. Chem. Soc.* **2012**, *134*, 2236–2245. [CrossRef]

22. Silva, R.; Voiry, D.; Chhowalla, M.; Asefa, T. Efficient Metal-Free Electrocatalysts for Oxygen Reduction: Polyaniline-Derived N- and O-Doped Mesoporous Carbons. *J. Am. Chem. Soc.* **2013**, *135*, 7823–7826. [CrossRef] [PubMed]

23. Liang, Y.; Wang, H.; Diao, P.; Chang, W.; Hong, G.; Li, Y.; Gong, M.; Xie, L.; Zhou, J.; Wang, J.; et al. Oxygen Reduction Electro catalyst Based on Strongly Coupled Cobalt Oxide Nanocrystals and Carbon Nanotubes. *J. Am. Chem. Soc.* **2012**, *134*, 15849–15857. [CrossRef]

24. Zhou, K.; Zhou, W.; Liu, X.; Wang, Y.; Yan, J.; Chen, S. Nitrogen Self-Doped Porous Carbon from Superpl Slugde as Metal-Free Electrocatalysts for Oxygen Reduction Reactions. *ACS Appl. Mater. Interfaces* **2014**, *6*, 14911–14918. [CrossRef] [PubMed]

25. Ferrero, G.A.; Preuss, K.; Marinovic, A.; Jorge, A.B.; Mansor, N.; Brett, D.J.L.; Fuertes, A.B.; Solis, M.S.; Titirici, M.-M. Fe–N-Doped Carbon Capsules with Outstanding Electrochemical Performance and Stability for the Oxygen Reduction Reaction in Both Acid and Alkaline Conditions. *ACS Nano* **2016**, *10*, 5922–5932. [CrossRef] [PubMed]

26. Jiang, W.J.; Gu, L.; Li, L.; Zhang, Y.; Zhang, X.; Zhang, L.J.; Wang, J.Q.; Hu, J.S.; Wei, Z.D.; Wan, L.J. Understanding the High Activity of Fe–NC Electro catalysts in Oxygen Reduction: Fe/Fe3C Nanoparticles Boost the Activity of Fe-Nx. *J. Am. Chem. Soc.* **2016**, *138*, 3570–3578. [CrossRef] [PubMed]

27. Wei, J.; Liang, Y.; Hu, Y.X.; Kong, B.A.; Simon, G.P.; Zhang, J.; Jiang, S.P.; Wang, H.T. A Versatile Iron-Tannin-Framework Ink Coating Strategy to Fabricate Biomass-Derived Iron Carbide/Fe–N-Carbon Catalysts for Efficient Oxygen Reduction. *Angew. Chem. Int. Ed.* **2016**, *55*, 1355–1359. [CrossRef] [PubMed]

28. Wu, Z.-Y.; Xu, X.-X.; Hu, B.-C.; Liang, H.-W.; Lin, Y.; Chen, L.-F.; Yu, S.-H. Iron Carbide Nanoparticles Encapsulated in Mesoporous Fe-N-Doped Carbon Nanofibers for Efficient Electrocatalysis. *Angew. Chem.* **2015**, *127*, 8297–8301. [CrossRef]

29. Yang, W.X.; Liu, X.J.; Yue, X.Y.; Jia, J.B.; Guo, S.J. Bamboo-like Carbon Nanotube/Fe3C Nanoparticle Hybrids and Their Highly Efficient Catalysis for Oxygen Reduction. *J. Am. Chem. Soc.* **2015**, *137*, 1436–1439. [CrossRef] [PubMed]
30. Galeano, C.; Meier, J.C.; Peinecke, V.; Bongard, H.; Katsounaros, I.; Topalov, A.A.; Lu, A.; Mayrhofer, K.J.J.; Schuth, F. Toward Highly Stable Electrocatalysts via Nanoparticle Pore Confinement. *J. Am. Chem. Soc.* 2012, 134, 20457–20465. [CrossRef]

31. Hu, Y.; Jensen, J.O.; Zhang, W.; Cleemann, L.N.; Xing, W.; Bjerrum, N.J.; Li, Q. Hollow Spheres of Iron Carbide Nanoparticles Encased in Graphitic Layers as Oxygen Reduction Catalysts. *Angew. Chem.* 2014, 126, 3749–3753. [CrossRef]

32. Xiao, M.; Zhu, J.; Feng, L.; Liu, C.; Xing, W. Meso/Macroporous Nitrogen-Doped Carbon Architectures with Iron Carbide Encapsulated in Graphitic Layers as an Efficient and Robust Catalyst for the Oxygen Reduction Reaction in Both Acidic and Alkaline Solutions. *Adv. Mater.* 2015, 27, 2521–2527. [CrossRef]

33. Ren, G.; Lu, X.; Li, Y.; Zhu, Y.; Dai, L.; Jiang, L. Porous Core–Shell Fe 3 C Embedded N-doped Carbon Nanofibers as an Effective Electrocatalysts for Oxygen Reduction Reaction. *ACS Appl. Mater. Interfaces* 2016, 8, 4118–4125. [CrossRef]

34. Zhong, R.Q.; Wu, Y.X.; Liang, Z.B.; Guo, W.H.; Zhi, C.X.; Qu, C.; Gao, S.; Zhu, B.J.; Zhang, H.; Zou, R.Q. Fabricating hierarchically porous and Fe3C-embedded nitrogen-rich carbon nanofibers as exceptional electrocatalysts for oxygen reduction. *Carbon* 2018, 142, 115–122. [CrossRef]

35. Aijaz, A.; Masa, J.; Rössler, C.; Antoni, H.; Fischer, R.A.; Schuhmann, W.; Muhler, M. MOF- Templated Assembly Approach for Fe3C Nanoparticles Encapsulated in Bamboo-Like N-Doped CNFs: Highly Efficient Oxygen Reduction under Acidic and Basic Conditions. *Chem. A Eur. J.* 2017, 23, 12125–12130. [CrossRef]

36. He, Z.B.; Maurice, J.L.; Kohno, H.; Homma, Y. Atomic-Scale In-situ Observation of Carbon Nanotube Growth from Solid State Iron Carbide Nanoparticles. *Nano Lett.* 2008, 8, 2082–2086. [CrossRef] [PubMed]

37. Rodriguez-Manzo, J.A.; Terrones, M.; Terrones, H.; Kroto, H.W.; Sun, L.; Banhart, F. In situ nucleation of carbon nanotubes by the injection of carbon atoms into metal particles. *Nat. Nanotechnol.* 2007, 2, 307–311. [CrossRef] [PubMed]

38. Tang, H.; Zeng, Y.; Zeng, Y.; Wang, R.; Cai, S.; Liao, C.; Cai, H.; Lu, X.; Tsiakaras, P. Iron-embedded nitrogen doped carbon frameworks as robust catalyst for oxygen reduction reaction in microbial fuel cells. *Appl. Catal. B Environ.* 2017, 202, 550–556. [CrossRef]

39. Park, S.H.; Choi, C.H.; Woo, S.I. Heteroatom doped carbons prepared by the pyrolysis of bio-derived amino acids as highly active catalysts for oxygen electro-reduction reactions. *Green Chem.* 2011, 13, 406–412.

40. Wang, Y.; Liu, Y.; Wang, K.; Song, S.; Tsiakaras, P.; Liu, H. Preparation and characterization of a novel KOH activated graphite felt cathode for the electro-Fenton process. *Appl. Catal. B Environ.* 2015, 165, 360–368. [CrossRef]

41. Zhao, Y.; Watanabe, K.; Hashimoto, K. Self-Supporting Oxygen Reduction Electrocatalysts Made from a Nitrogen-Rich Network Polymer. *J. Am. Chem. Soc.* 2012, 134, 19528–19531. [CrossRef]

42. Liang, J.; Jiao, Y.; Jaromiec, M.; Qiao, S.Z. Sulfur and Nitrogen Dual-Doped Mesoporous Graphene Electrocatalyst for Oxygen Reduction with Synergistically Enhanced Performance. *Angew. Chem.* 2012, 124, 11664–11668. [CrossRef]

43. Wohlgemuth, S.-A.; Fellinger, T.-P.; Jäker, P.; Antonietti, M. Tunable nitrogen-doped carbon aerogels as sustainable electrocatalysts in the oxygen reduction reaction. *J. Mater. Chem. A* 2013, 1, 4002. [CrossRef]

44. Wang, R.; Li, J.; Cai, S.; Zeng, Y.; Zhang, H.; Cai, H.; Tang, H. Facile Synthesis of Fe3 C@Graphene Hybrid Nanorods as an Efficient and Robust Catalyst for Oxygen Reduction Reaction. *ChemPlusChem* 2016, 81, 464–465. [CrossRef]

45. Wang, Q.; Lei, Y.; Chen, Z.; Wu, N.; Wang, Y.; Wang, B.; Wang, Y. Fe/Fe 3 C@CNPs nanoparticles encapsulated in N-doped graphene–CNTs framework as an efficient bifunctional oxygen electrocatalyst for robust rechargeable Zn-air batteries. *J. Mater. Chem. A* 2018, 6, 516–526. [CrossRef]

46. Chakrapani, K.; Sampaum, S. The morphology dependent electrocatalytic activity of Ir nanostructures towards oxygen reduction. *Phys. Chem. Chem. Phys.* 2014, 16, 16815–16823. [CrossRef] [PubMed]

47. Wu, G.; More, K.; Johnston, C.M.; Zelenay, P. High-Performance Electrocatalysts for Oxygen Reduction Derived from Polyamine, Iron, and Cobalt. *Science* 2011, 332, 443–447. [CrossRef] [PubMed]
49. Proietti, E.; Larouche, N.; Jaouen, F.; Lefevre, M.; Tian, J.; Herranz, J.; Dodelet, J.-P. Iron-based cathode catalyst with enhanced power density in polymer electrolyte membrane fuel cells. Nat. Commun. 2011, 2, 416. [CrossRef] [PubMed]

50. Wu, Z.-S.; Yang, S.; Sun, Y.; Parvez, K.; Feng, X.; Müllen, K. 3D Nitrogen-Doped Graphene Aerogel-Supported Fe₃O₄ Nanoparticles as Efficient Electrocatalysts for the Oxygen Reduction Reaction. J. Am. Chem. Soc. 2012, 134, 9082–9085. [CrossRef] [PubMed]

51. Liang, C.; Tao, N.; Wang, B.; Lin, S.; Lei, M. Graphite carbon coated TiN nanoparticles as high durable oxygen reduction reaction catalyst in alkaline electrolyte. Micro Nano Lett. 2018, 13, 1260–1264. [CrossRef]