Biomass-derived activated carbon as high-performance non-precious electrocatalyst for oxygen reduction

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Abstract
A new type of Fe and N doped carbon material is synthesized by pyrolyzing ferric chloride doped egg white (EW) and the proposed synthetic route is easy, green, and low-cost. In addition, the as-prepared sample exhibits a feasible magnetism and comparable oxygen reduction reaction (ORR) activity to commercial Pt/C.

As the cathodic ORR plays an important role in the performance of a fuel cell,1,2 efficient ORR electrocatalysts are highly desirable for practical applications. So far, Pt and its alloys have long been regarded as the most effective catalysts for the ORR in fuel cells.3,4 However, their large-scale commercial applications have been hindered by high costs.5 Besides, they still suffer from serious intermediate tolerance, anode crossover, sluggish kinetics, and poor stability in an electrochemical environment.6–8 For these sakes, extensive research has been expected to solve the problems that involve (i) replacing the noble metal by alloy to reduce the consumption of novel metals and to lower the cost;9,10 (ii) assuming certain novel carbon materials with particular shapes like carbon nanotubes (CNTs) and graphene to promote the conductivity of the catalyst or the intermediate product transport;11,12 (iii) the introduction of transition metals (such as Fe, Co)13,14 as well as metal oxides (such as Fe₃O₄ and CoO)15,16 to boost the ORR; (iv) doping some N, B, P, and S atoms17–20 to modify the electrocatalysts to enhance the methanol tolerance. Among these explorations, it is confirmed that nitrogen-doped carbon materials not only exhibit high catalytic activity, long-term stability, and excellent methanol tolerance in alkaline media, but also possess the advantages of low cost and environmental friendliness.

According to the theory of Dai et al.,21 the high catalytic activity by doping N may be attributed to the larger electronegativity of N compared to C atoms. And, N atoms can create positive charge density on the adjacent C atoms, which results in the very favorable adsorption of O₂. Consequently, intense research on the ways to modify the carbon materials with nitrogen atom has been greatly inspired. To our knowledge, the source of nitrogen is always derived from expensive organic monomer containing nitrogen element22 or NH₃.12 However, these
nitrogen-containing compounds are either expensive or harmful to human health. In addition, it is complicated to prepare the nitrogen-doped and non-precious electrocatalysts. Thus, a simple and convenient method to synthesize a new kind of ORR electrocatalysts with the features of low price, green origin as well as excellent performance can be expected.

Considering that several species of natural biomass were successfully given to be carbon materials and these materials showed potential applications in a variety of fields, particularly in fuel cells, egg white (EW) could be a promising candidate. Since EW abounds with carbon, nitrogen and oxygen elements owing to the existence of proteins, it is expected to provide an alternative precursor to synthesize advantageous nanomaterials. Herein, we demonstrated an easy and green strategy to synthesize a new type of N, Fe doped non-precious electrocatalyst by using EW as the sources of carbon and nitrogen sources. This new type of electrocatalyst (Fe-CEW) not only showed favorable magnetic properties but also exhibited surprising electrocatalytic activity towards ORR. Verified by the electrochemical experiments, the as-prepared samples had ORR catalytic activity comparable to the commercial carbon-supported Pt catalyst and had superior stability, thus leading to a new high performance catalyst for ORR. First of all, the practical composition of CEW and Fe-CEW were evaluated by both elemental analysis. The analytical results indicate that CEW consists of N (6.9 wt%), C (62.7 wt%), and the reminder inorganic salt residue like K, Na, Ca etc.; Fe-CEW consists of N (4.4 wt%) and C (48.4 wt%). Based on the ratio of each element, metallic Fe is calculated to be 16.8 wt%. The XRD patterns of CEW and Fe-CEW (Fig. S1, ESI) show two peaks located at 2h # 25u and 44u, which can be attributed to (002) and (101) diffraction peaks from different forms of carbon, referred to turbostratic carbon (carbon black) and graphite carbon (graphitic structure), respectively. The other diffraction peaks at ca. 28u, 31u, 41u, 45u in the two curves are assigned to certain inorganic salt residue of K, Na, Ca etc. originated from CEW. Besides, a diffraction peak at 2h = 35u can be observed for Fe-CEW sample, which is associated with the iron/carbon-black composite nanoparticle, manifesting that the doped Fe is bonded with EW after heat treatment. In the Raman spectrum of XC-72, CEW and Fe-CEW in Fig. S2, ESI, the peaks at y1310 and y1590 cm^{-1} assigned to the D and G band, respectively, can be observed. The peak intensity ratio of the D to G band (ID/IG) for the XC-72 and CEW is 1.03 and 1.14, respectively, suggesting that the CEW shows more defect sites (edge plans) and poorer electric conductivity than the XC-72 does. In terms of the raman spectrum of CEW and Fe-CEW, both the D band and G band do not significantly change in ether width or shape apart from the ID/IG. The ID/IG ratio of Fe-CEW is 1.07, closed to the value of XC-72 but smaller than that of CEW (1.14), indicating that (i) the Fe-CEW exhibits less defects sites than the CEW, which may be caused from the original defect sites that is covered by the doped Fe atoms, as reported in literature for the 1-aminopyrene-functionalized multi-walled carbon nanotubes; (ii) the Fe-CEW possesses much better electric conductivity compared to CEW, which means a higher current density and excellent performance for ORR. Fig. S3, ESI shows the surface functional groups of the carbon nitride materials are
characterized by FT-IR. Fig. S4, ESI3 displays TGA and DTG profiles of the decomposition process of EW and Fe doped egg white (Fe-EW), from which we can obtain information on residual masses at different temperatures. As shown in Fig. S4a, ESI3 two stages are included in the pyrolysis process. A slight weight loss occurs during the first stage ranging from the starting temperature to y200 uC, which can be related to the elimination of water (dehydration). The second stage, from y200 uC to the end, is characterized by a major loss, which corresponds to the main pyrolysis process (devolatilization). It proceeds with a high rate in the initial period but keep a comparatively steady rate in the later stage, demonstrating that prior to forming the nitrogen- doped carbon materials, the carbohydrate in EW is pyrolyzed firstly. With respect to the decomposition process of Fe-EW (as shown in Fig. S4b, ESI3), it can be also divided into two stages: (i) dehydration (from room temperature to y200 uC) due to the removal of adsorbed and bound water; (ii) devolatilization under the participation of FeCl3. However, we notice that the second stage in curve b is absolutely distinct from that in curve a. The curve b displays a broader platform region than curve a. We infer that during the pyrolysis process of EW, the reaction of forming the composite of Fe–N/C occurs simultaneously.

Fig. 1 displays the N 1s spectra for CEW, Fe-CEW and Fe 2p spectra for Fe-CEW. As reported in the literature,12 the N 1s spectrum can be further deconvoluted into three different signals with binding energies of 398.0, 400.0, and 401.3 eV, each of which correspond to pyridinic-N, pyrrolic-N and graphitic-N, respectively. Of these three types of N species, pyridinic-N and pyrrolic-N, particularly for pyrrolic-N, have a lone electron pair in the plane of the carbon matrix, resulting in an increase of electron-donor property for the catalysts.21 The O–O bond would be weaken by the bonding between oxygen and nitrogen (and/or the adjacent carbon atom), and thus the reduction of oxygen is facilitated.27 The Fe 2p spectrum for Fe-CEW shows two peaks at 726.0 and 711.5 eV (Fig. 1c), corresponding to Fe 2p1/2 and Fe 2p3/2, respectively. The value is similar to that of a certain kind of Fe–N/C compound reported previously.28 It could be inferred that Fe atoms bond with nitrogen and/or carbon. Considering the introduction of Fe, the as-prepared samples may exhibit the properties of magnetism after heat treatment. As shown in Fig. 2a and b, a reasonable amount of samples is dispersed into a standard bottle containing ethanol, and then a magnet is kept closed to the bottle. After a few times, the sample of Fe-CEW is separated from the ethanol solution, revealing that the Fe-CEW sample exhibits good magnetic features and re-disperse properties. This suggests a potential application for targeting and separation.
Fig. 1  N 1s peaks in the XPS spectrum for (a) CEW, (b) Fe-CEW and Fe 2p peaks in the XPS spectrum for (c) Fe-CEW.

Fig. 2  The separation process of the as-prepared (a) BW, CEW and (b) Fe-EW, Fe-CEW with magnet; SEM images with different magnification for the as-prepared (c, d) Fe-CEW.
Fig. 2c and d show scanning electron microscopy (SEM) images of the as-prepared Fe-CEW. It can be observed that Fe-CEW has formed coral-like structure with some anomalous pores, which possibly provides large specific surface area for the interface reaction, stores liquid electrolyte, and promotes the mass transfer and separation.

Cyclic voltammogram (CV) curves on the CEW and Fe-CEW electrodes in N2-saturated and O2-saturated 0.1 mol L⁻¹ KOH solutions are shown in Fig. 3a and b. As can be seen, an oxidation peak appears at around 20.47 V and 20.42 V in the CV curve of Fe-CEW in N2 and O2 atmosphere, arising from the Fe(II) is oxidized to Fe(III), whereas no peaks have response to the curve of CEW. Beyond that, it can also be found that the CV of the Fe-CEW shows a current density comparable with that of CEW in N2 atmosphere, but a larger electric double layer than that of CEW in O2 atmosphere. This result implies that the doped Fe is involve into the electrochemical process, including the ORR and provides a larger electrochemically accessible area, an area where the
To further confirm whether the N and Fe work in the process of ORR, we conduct the linear sweep voltammograms (LSV) experiments as shown in Fig. 3c. First of all, the LSV curve of CEW is compared with that of XC-72 to verify the function of N contained in EW. As illustrated in the LSV curves, the half-wave potential of the CEW is 2212 mV, which is a shift of 56 mV more positive to that of the XC-72 (2268 mV). Subsequently, the similar method is used to confirm the role of doped Fe. It is found that, once the Fe is introduced, the half-wave potential of Fe-CEW (2130 mV) shifted positively to around 82 mV compared to CEW, but a little lower (30 mV) than that of Pt/C (2100 mV). In terms of the onset potential, it decreases in the order of Pt/C and Fe-CEW (220 mV) > CEW (181 mV) > XC-72 (40 mV) (as shown in Fig. S5, ESI3). Hence, based on the above facts, it has clearly demonstrated a significant enhancement of electro-catalytic activity for Fe-CEW towards the ORR as a result of the introduction of N and Fe in respect to CEW and XC-72. Subsequently, to investigate the ORR kinetic character of Fe-CEW and Pt/C, Tafel plots (Fig. 3d) are carried out. In Tafel plots, two well-defined linear regions similar with Tafel slope values are displayed at high and low potentials, respectively. The Tafel slope values of 60 mV dec21 at high potentials suggest that ORR rate may be determined by migration of adsorbed oxygen intermediates.30–32 While at low potentials, the Tafel slope of 120 mV dec21 could be ascribed to the transfer of the first electron as a rate-determining step.31,32 The above results suggest that the ORR mechanism of Fe-CEW may be as similar as that of Pt/C.

Then, rotating disk electrode (RDE) current-potential curves of Fe-CEW at various rotating speeds are plotted in Fig. 4a. The transferred electron numbers per O2 involved in the oxygen reduction at the Fe-CEW electrode is determined by the Koutecky–Levich (K–L) equation as given below

\[ \frac{1}{j} = \frac{1}{j_k} + \frac{1}{B_0^{0.5}} \]  

where \( j \) is the measured current density, \( j_k \) is the kinetic current density, \( v \) is the rotation speed, \( B \) is the slope of K–L plots which can be obtained from Fig. 4b based on the Levich equation as follows:33,34

\[ B = 0.2 nF(D_0)^{2/3}v^{-1/6}C_{O_2} \]  

where \( n \) is the number of electrons transferred in the reduction of one O2 molecule in ORR, \( F \) is the Faraday constant (\( F = 96485 \text{ C mol}^{-1} \)), \( D_0 \) is the diffusion coefficient of O2 in 0.1 mol L21 KOH (\( D_0 = 1.9 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1} \)), \( C_{O_2} \) is the bulk concentration of O2 in the electrolyte (\( C_{O_2} = 1.2 \times 10^{-6} \text{ mol cm}^{-3} \)), and \( n \) is the kinematic viscosity of the electrolyte (\( n = 0.01 \text{ cm}^2 \text{ s}^{-1} \)).35 The constant 0.2 is adopted when the rotation speed is expressed in rpm. Calculated
from the slope of the K–L plots, the number of the average number of the electrons transferred per O₂ molecule for the sample of Fe-CEW in the process of ORR is 3.4 at a potential ranging from 20.6 to 20.3 V for Fe-CEW. Typically, the above results demonstrate that the process of ORR on Fe-CEW proceeds close to 4e² pathway.

![Fig. 5 CVs for ORR at (a) CEW and (b) Fe-CEW electrodes before and after a continuous potentiodynamic swept for 500 cycles in 0.1 mol L⁻¹ KOH under air atmosphere. Rotation rate: 1600 rpm, scan rate: 50 mV s⁻¹ for CV, temperature: 30 °C.](http://repository.uwc.ac.za)

Finally, the stability of CEW and Fe-CEW is evaluated by CV method. As indicated in Fig. 5a and b, 500 cycles have been measured so that the catalysts can trend to be a steady state, and then the following 500 cycles can be used to assess the stability of the obtained Fe-CEW catalyst. According to the method that reported in literature, the CV shows a reliable stability of Fe-CEW less than 3.1% current density loss at 0.6 V, in contrast, the deterioration of the CEW catalyst reaches 4.5%. Based on the above result, the Fe-CEW catalyst exhibits a superior durability in alkaline medium, which may be explained as the reported principle that the relative strong strength of covalent N–C bond and coordination linkage between Fe and N.

In summary, a novel approach to prepare Fe and/or N doped carbon materials is developed by pyrolyzing EW and Fe-EW, which can be used as non-precious electrocatalysts for ORR. The synthetic route is green and low-cost, thus providing an original perspective for the development of new catalytic carbon materials in fuel cells. Interestingly, the Fe-CEW exhibits fine magnetism, which suggests a potential application for targeting and separation. Among the as-prepared samples, Fe-CEW exhibits excellent performance for the ORR. It is proved that N (including pyridinic-N and pyrrolic-N) and Fe immensely promote the ORR. In addition, compared to commercial Pt/C, Fe-CEW displays a comparable ORR activity, which suggests that it could be a promising candidate for ORR catalysts in the future.

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Notes and references