Synthesis of Iron Oxide/Partly Graphitized Carbon Composites as a High-Efficiency and Low-Cost Cathode Catalyst for Microbial Fuel Cells

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Supporting Information

ABSTRACT: Waste cornstalks and pomelo skins are used as carbon resources for preparing nanocomposites of iron oxide and partly graphitized carbon ($\text{Fe}_3\text{O}_4$/PGC-CS and $\text{Fe}_3\text{O}_4$/PGC-PS). The results showed that $\text{Fe}_3\text{O}_4$ with a face-centered cubic structure is uniformly dispersed on the skeleton of $\text{Fe}_3\text{O}_4$/GC, and the highest $S_{\text{BET}}$ values of $\text{Fe}_3\text{O}_4$/PGC-CS (476.5 m$^2$ g$^{-1}$) and $\text{Fe}_3\text{O}_4$/PGC-PS (547.7 m$^2$ g$^{-1}$) are obtained at 1000 °C. The electrical conductivity and density of catalytic active sites are correspondingly improved by the introduction of Fe species. Microbial fuel cells (MFCs) with a mixed composite ($\text{Fe}_3\text{O}_4$/PGC-CS:$\text{Fe}_3\text{O}_4$/PGC-PS = 1:1) cathode (three-dimensional structures) generate the highest power density of $1502 \pm 30$ mW m$^{-2}$, which is 26.01% higher than that of Pt/C ($1192 \pm 33$ mW m$^{-2}$) and only declines by 7.12% after 18 cycles. The $\text{Fe}_3\text{O}_4$/PGC-CS cathode has the highest Coulombic efficiency (24.3 ± 0.7%). The $\text{Fe}_3\text{O}_4$/PGC composites exhibit high oxygen reduction reactivity, low charge transfer resistances, and long-term stability and can be used as a low-cost and high-efficiency catalyst for MFCs.

KEYWORDS: graphitized carbon, microbial fuel cells, oxygen reduction reactivity, stability, waste biomass

INTRODUCTION

Recently, a lot of attention has been paid to microbial fuel cells (MFCs) due to their potential use of electrochemically active microorganisms for clean, sustainable, renewable energy production associated with wastewater treatment. In addition, new chemical production (hydrogen, methanol, ethanol, methane, and formate), CO$_2$ fixation, decolorization, and energy utilization in miniized MFCs for low-power-consumption niches (marine wireless sensors and implantable medical devices) are also appealing microbial electrochemical technologies. However, the high power production cost has been a great challenge affecting the industrialization and commercial applications of MFC technology. The oxygen reduction reaction (ORR) as a kinetically slow process is undoubtedly the pivot among power density generation and investment of materials. Platinum-based materials are the most widely used precious-metal catalysts for their superior ORR activity. However, their susceptibility to CO poisoning and exorbitant price limit their large-scale application. Therefore, a novel, durable, highly active, and cheap catalyst has been desired in recent years.

New carbon materials with a unique structure and redox activity such as graphene and carbon nanotubes have been recognized as low-cost efficient catalysts. Feng et al. used chemical vapor deposition (CVD) to synthesize nitrogen-doped carbon nanotubes (NCNTs) as MFC ORR cathode catalysts. The vertical arrangement of the nanocluster structure of the CNTs diminished the internal resistance, then increased the cathode potential, and proceeded on a $4e^-$ pathway for efficient electrocatalysis in a neutral phosphate-buffered solution (PBS). The maximum power density (1600 ± 50 mW m$^{-2}$) and the stability were better than those of Pt/C (1393 ± 35 mW m$^{-2}$) after 25 cycles. Despite the complicated synthesis process, newly structured carbon has already become one of the most valuable substitutes in practical MFC applications due to its good ORR activity and high stability.

Recent studies on ORR catalysts have shown that graphitized carbon nanostructured composites exhibit a relatively high
electrocatalytic activity, electrical conductivity, and stability. Transition metals were initially reported to show comparable catalytic activity for electroreduction of O₂ to Pt. For example, Su et al. synthesized a cobalt oxide/nitrogen-doped graphene (Co₃O₄/NG) nanocomposite by the hydrothermal method which showed high ORR and oxygen evolution reaction (OER) activities. The power density was up to 1340 ± 10 mW m⁻², only a bit lower than that of Pt/C (1470 ± 10 mW m⁻²). Hao Yu et al. prepared a series of Fe-based and Co-containing ORR catalysts that could be considered as a potential cathode choice for biological electricity generation. Recently, Xia et al. used iron ethylenediaminetetraacetic acid solution to modify the activated carbon (MAC). The electron donor capacity of this MAC had been enhanced by weakening the O−O bond and was further improved by the transition metal Fe, which could stabilize the tetravalent nitrogen and promote the disproportionation of H₂O₂. It is generally recognized that the synergistic effect between the graphite carbon support and transition metal played a crucial role in the formation of active sites electrically conductive for ORR. However, considerable research is still required to solve the inherent problems associated with the complexity of the synthesis process, fallacious architecture, and poor interaction of graphite carbon and the active constituents.

In this paper, we report a method (one step) for preparation of Fe₃O₄/partially graphitized carbon (Fe₃O₄/PGC) by using two recycled biomasses (cornstalks and pomelo skins) which have a low cost and are easily accessible as carbon resources. Especially the carbonized fluffy spongelike skins of pomelo and straw could offer a high specific surface area, a large number of active sites, and a large pore volume for effective mass transport. The conductivity, catalytic activity, and stability of Fe₃O₄/PGC could be greatly enhanced by the introduction of a nano-Fe species, which was expected to have a higher ORR capacity in MFCs. The performance of the synthesized Fe₃O₄/PGC as the cathode catalyst in MFCs was compared with that of the commonly used Pt/C catalyst. The electrocatalytic activity of the Fe₃O₄/PGC for ORR in MFCs, which was still unknown, was also investigated in detail.

**EXPERIMENTAL SECTION**

**Materials.** Wasted cornstalks from farmland and residual pomelo skins were used as carbon resources for synthesis of carbon composites. After the outside crust of the cornstalks and pomelo skins (yellow skins) were stripped down, the inside cores were kept in desiccators for further use. Inocula used in this study were obtained from the effluent of the previous reactors that had been running for 4 months as previously described. All chemicals used in the experiment were of analytical grade and were used without further purification.

**Synthesis of Fe₃O₄/PGC Composites.** The preparation of Fe₃O₄/PGC was conducted by using a simple in situ simultaneous synthesis method. Briefly, the spongelike parts of the cornstalks and pomelo skins were impregnated with FeCl₃ solution to form a carbon−Fe⁺ complex with a mass ratio of 1:1. The resulting products were dried at 60 °C for 12 h in an oven and carbonized at 1000 °C for 1.5 h with a heating rate of 5 °C min⁻¹ under a highly pure N₂ flow (50−60 ml min⁻¹). Finally, the black samples were ground into powder and washed with deionized water three times. The samples made from cornstalks and pomelo skins are designated as Fe₃O₄/PGC-PS and Fe₃O₄/PGC-CS, respectively. As a comparison, the above two products were washed with HCl (2 mol L⁻¹) for 12 h to completely remove the Fe species, and the resulting products are respectively designated as PGCC-PS and PGCC-CS.

**Cathode Preparation for MFCs.** The membrane-free single-chamber air-cathode MFCs were constructed by using a carbon fiber brush as the anode and stainless steel mesh as the cathode base material. The gas diffusion layer (GDL) of the air cathode was prepared by rolling carbon black and poly(tetrafluoroethylene) (PTFE) (60 wt %) with a mass ratio of 7:3 and then sintering the resulting material at 340 °C for 25 min. The catalyst layer was rolled onto the opposite side of the GDL with a mass ratio of 6:1 and the resulting material dried at 80 °C according to the previously mentioned method. Five kinds of carbon-based cathodes were prepared, including Fe₃O₄/PGC-PS and Fe₃O₄/PGC-CS and their physical mixture (at a mass ratio of 1:1; Figure S1, Supporting Information), PGC-PS, and PGC-CS. For comparison, commercially available Pt/C (10 wt %) powder was also used as the cathode catalyst (5 mg cm⁻²).

**MFC Configuration and Operation.** MFCs were conducted with single-chamber reactors 4 cm long and 3 cm in diameter previously described in the literature. The graphite fiber brush anode was pretreated with acetone and heated in the air. All the reactors were inoculated by using the effluent from MFCs operated for 4 months, and operated at a fixed external circuit resistance (1000 Ω). The substrate solution contained glucose (1 g L⁻¹) and PBS, which contained NH₄Cl (0.31 g L⁻¹), NaH₂PO₄·2H₂O (3.321 g L⁻¹), Na₂HPO₄·12H₂O (10.317 g L⁻¹), KCl (0.13 g L⁻¹), trace minerals (12.5 mL L⁻¹), and vitamins (5 mL L⁻¹). The MFC operating temperature was maintained at 30 °C, and the medium was replaced with the fuel when the voltage decreased to less than 50 mV, which formed one complete cycle. To achieve statistical soundness, at least two replicates were carried out.

**Materials Characterization.** X-ray diffraction (XRD) patterns were collected with a Rigaku D/max-IIIB diffractometer using Cu Kα radiation (θ = 1.5406 Å) at step scan of 0.02° from 10° to 80°. The voltage and the applied current were 40 kV and 20 mA, respectively. The resulting powder diffraction patterns were analyzed according to the Joint Committee on Powder Diffraction Standard data. Raman spectra were recorded with an HR 800 micro Raman spectrometer (Jobin-Yvon, France) at 457.9 nm. The laser beam was focused on the sample with a 50× objective. The laser intensity excited by an argon ion laser was kept below the threshold for any laser-induced changes in the Raman spectra or electrical transport. Transmission electron microscopy (TEM) images were taken by using a JEM-2100 electron microscope (JEOL) with an acceleration voltage of 200 kV.

**Electrochemical Analysis.** Electrochemical measurements were conducted on a computer-controlled electrochemical workstation (BAS100B electrochemical workstation, Germany) with a typical three-electrode cell equipped with gas flow systems. Linear sweep voltammetry (LSV) was performed at 1 mV s⁻¹ on the cathodes, and the reactor was filled with 28 mL of 50 mM PBS solution and equipped with a 2.25 cm² platinum square counter electrode and a Ag/AgCl reference electrode (±495 V vs the standard hydrogen electrode, 3.0 M KCl). The cyclic voltammetry (CV) scan was performed in the potential range of −0.6 to +0.4 V at a scan rate of 10 mV s⁻¹.

A rotating disk electrode (RDE) can be used to explore the mechanism of the complex electrode reaction and obtain more information on the catalyst properties. Therefore, RDE tests were conducted to evaluate the catalyst activity of the Fe₃O₄/PGC composites by comparison with that of Pt/C. The RDE type used for ORR measurements was a glassy carbon rotating disk electrode, which was 0.4 cm in diameter (the effective disk area was 0.126 cm²). Catalyst ink was prepared by adding a 10 mg powdered sample to 10 mL of isopropyl alcohol, and the resulting solution was then treated ultrasonically for 15 min. The ink suspension (10 μL) was dropped onto a 0.4 cm diameter glassy carbon electrode and dried in the air.
until formation of a uniform film (sample loading of 0.079 mg cm\(^{-2}\)). Nafion (5 wt % solution, 5 \(\mu\)L) was then added to the coated catalyst for kinetic analysis. The RDE tests were carried out by using a modulated speed rotator (BAS Inc. Japan) immersed in a medium of 50 mM PBS at rotation rates of 400–3600 rpm with a pure oxygen flow. The solutions should be sparged with \(O_2\) for at least 30 min. The average number of transferred electrons (\(n\)) and kinetic current (\(i_k\)) for the oxygen reduction reaction were calculated on the basis of the Koutecky–Levich (K–L) equation:\(^{25}\)

\[
\frac{1}{i} = \frac{1}{k_c} + \left(\frac{1}{0.62nFAD^{2/3}v^{1/6}C}\right)\omega^{-1/2}
\]

where \(i\) is the measured current (A), \(F\) is the Faraday constant (96 485 C mol\(^{-1}\)), \(A\) is the effective projected area of the disk electrode (0.126 cm\(^2\), 0.4 cm in diameter), \(D\) is the diffusion coefficient of oxygen (2.7 \(\times\) 10\(^{-5}\) cm\(^2\) s\(^{-1}\)), \(v\) is the kinematic viscosity (8.08 \(\times\) 10\(^{-3}\) cm\(^2\) s\(^{-1}\)), \(C\) is the concentration of oxygen in the solution (2.3 \(\times\) 10\(^{-5}\) mol cm\(^{-3}\)), and \(\omega\) is the rotation rate of the electrode.

Electrochemical impedance spectroscopy (EIS) was conducted at the open circuit voltage (OCV) over a frequency range of 10\(^3\) to 10\(^{−2}\) Hz with a sinusoidal perturbation of 10 mV amplitude. EIS data were analyzed by fitting the spectra to an equivalent circuit model (Figure S2, Supporting Information). The produced voltage (\(E_{\text{cell}}\)) and electrode potentials were measured at a fixed external circuit resistance (1000 \(\Omega\)) by using a multimeter and Ag/AgCl reference electrode. After the reactors were operated with fresh substrate for about 2 h, polarization and power density curves were obtained by measuring the stable voltage generated at various external resistances (50–5000 \(\Omega\)). All the other parameters, including current density, power density, and Coulombic efficiency (CE), were calculated from the ratio of the total electrical charge produced within every cycle at different external resistances to the theoretical amount of electrons available from the oxidation of glucose to carbon dioxide. All the above electrochemical experiments were carried out at 30 °C. The chemical oxygen demand (COD) in MFCs was determined by the potassium dichromate oxidation method.

**RESULTS AND DISCUSSION**

**Characteristics Analyses of Fe\(_3\)O\(_4\)/PGC Composites.** As shown in Figure 1, the XRD patterns of Fe\(_3\)O\(_4\)/PGC-CS and Fe\(_3\)O\(_4\)/PGC-PS carbonized at 900–1050 °C were almost the same. The typical crystalline phases of the two composites carbonized at 900 °C were Fe\(_3\)O\(_4\) nanoparticles (JCPDS card no. 75-1609) with a face-centered cubic structure (30°, 35.4°, 43°, 53.4°, 56.9°, and 62.5°). At 950, 1000, and 1050 °C, the peaks at 44.5° corresponded to the typical diffraction peaks of \(\alpha\)-Fe with a body-centered cubic structure, which was the main difference from those at 900 °C. This implied that, at higher temperatures, part of the Fe\(_3\)O\(_4\) particles in Fe\(_3\)O\(_4\)/PGC composites were reduced to \(\alpha\)-Fe by in situ carbothermal reduction. At 1050 °C, the graphitized carbon (26.2°), which originated from the carbon graphitization catalyzed by iron species, was obviously detected in the two Fe\(_3\)O\(_4\)/PGC composites. The graphitization of carbon was a gradual and slow process, which means that the PGC was already present in the Fe\(_3\)O\(_4\)/PGC composites at 950 or 1000 °C.

Figure 2 shows the Raman spectra of Fe\(_3\)O\(_4\)/PGC-CS and Fe\(_3\)O\(_4\)/PGC-PS carbonized at 900–1050 °C. As the temperature increased, the \(I_d/I_g\) (the D and G bands usually correspond to the sp\(^3\) defect sites and the bond stretching of sp\(^2\)-bonded pairs, respectively) value increased gradually, which means that the graphitization degree of Fe\(_3\)O\(_4\)/PGC-CS or Fe\(_3\)O\(_4\)/PGC-PS was increased.\(^{26}\) Meanwhile, the peak for the G bands of Fe\(_3\)O\(_4\)/PGC-CS (Figure 2A) was transferred from 1588 cm\(^{-1}\) (900 °C) to 1576 cm\(^{-1}\) (1050 °C); for Fe\(_3\)O\(_4\)/PGC-PS (Figure 2B), the peak was transferred from 1582 cm\(^{-1}\) (900 °C) to 1570 cm\(^{-1}\) (1050 °C). The peak shift for the G bands of Fe\(_3\)O\(_4\)/PGC-PS (10 cm\(^{-1}\)) was higher than that of Fe\(_3\)O\(_4\)/PGC-CS (4 cm\(^{-1}\)), taking the highly oriented pyrolytic graphite (1580 cm\(^{-1}\)) as the standard. This implies that the greater the
shift of the G bands to lower wavenumber, the higher the degree of carbon curling.\textsuperscript{27} 

\(\) 

\(\text{N}_2\) adsorption−desorption isotherms and pore size distribution curves of Fe\(\text{O}_4\)/PGC-CS and Fe\(\text{O}_4\)/PGC-PS carbonized at 900−1050 °C are shown in Figures S3 and S4 (Supporting Information), respectively. It can be seen that both Fe\(\text{O}_4\)/PGC-CS and Fe\(\text{O}_4\)/PGC-PS showed typical type IV isotherms with a sharp capillary condensation step and H2-type hysteresis loops at relative pressures of 0.5−0.8. Their pore size distributions were very wide and mainly consisted of mesopores, and the average pore size decreased gradually as the temperature increased (Figures S3 and S4). The highest \(S_{BET}\) values of Fe\(\text{O}_4\)/PGC-CS (476.5 m\(^2\) g\(^{-1}\)) and Fe\(\text{O}_4\)/PGC-PS (547.7 m\(^2\) g\(^{-1}\)) were both obtained at 1000 °C, which showed that the development of the porous structure of the composite made from pomelo skins was better than that from cornstalks (Table 1). One reason was that the structure of the spongelike part of pomelo skins was quite different from that of cornstalks, which resulted in the difference in \(S_{BET}\).

According to the above analyses, 1000 °C was selected as the optimal temperature for Fe\(\text{O}_4\)/PGC composite preparation. The morphological features of Fe\(\text{O}_4\)/PGC-CS and Fe\(\text{O}_4\)/PGC-PS carbonized at 1000 °C were observed by TEM (Figure 3). Fe\(\text{O}_4\) with particle sizes of 10−20 nm was uniformly distributed in Fe\(\text{O}_4\)/PGC-CS, and no agglomeration phenomena are observed in Figure 3A. Figure 3B shows that the GC was already formed in the body of Fe\(\text{O}_4\)/PGC-CS and tightly surrounded the Fe\(\text{O}_4\) particles. As observed in Figure 3D,E, some of the aggregated Fe\(\text{O}_4\) particles were connected tightly to each other and inlaid in the nanocapsule structure of Fe\(\text{O}_4\)/PGC-PS. The conductivity and electron transfer of the Fe\(\text{O}_4\)/PGC composites were inevitably enhanced by the high degree of carbon graphitization.\textsuperscript{15,16} As shown in Figure 3C,F, the lattice spacing marked in the pattern between the (311) planes of Fe\(\text{O}_4\) was 0.25 nm (HRTEM measurements). The cathodes made with the stable Fe\(\text{O}_4\)/PGC composites were expected to have multiple catalytic active sites and high electrocatalytic activity.

Simultaneous TG−DSC analyses of Fe\(\text{O}_4\)/PGC-CS and Fe\(\text{O}_4\)/PGC-PS are shown in Figure 4. The exothermic process with a corresponding weight loss of 65% appeared in the range of 400−720 °C, which should be attributed to the burning loss of carbon (in the form of CO\(_2\) or CO) from Fe\(\text{O}_4\)/PGC-PS (Figure 4A). The corresponding weight loss of 80% for Fe\(\text{O}_4\)/PGC-CS occurred in the range of 420−660 °C (Figure 4B). These results suggest that the content of Fe species in Fe\(\text{O}_4\)/PGC-PS (about 35%) was higher than that of Fe\(\text{O}_4\)/PGC-CS (about 20%), which could be attributed to the different complexing capacities of Fe\(^{3+}\) and the carbon resource.

**MFC Performance of Fe\(\text{O}_4\)/PGC Composite Cathodes.** LSV scan tests for the cathode materials (Figure S5, Supporting Information) showed that the mixture cathode of Fe\(\text{O}_4\)/PGC-CS and Fe\(\text{O}_4\)/PGC-PS (mass ratio of 1:1) had the highest current density in the potential range of −1.5 to +0.4 V vs the Ag/AgCl reference electrode. The unique highly porous stack structures of the mixture made it more favorable for chemisorption of oxygen onto the catalyst surface, which inevitably enhanced the ORR electrocatalytic activity.\textsuperscript{28} Fe\(\text{O}_4\)/PGC-CS cathodes produced a slightly higher current density than Fe\(\text{O}_4\)/GC-PS cathodes. To further evaluate the ORR

\begin{table}[h]
\centering
\caption{Textural Properties of Fe\(\text{O}_4\)/PGC-CS and Fe\(\text{O}_4\)/PGC-PS Obtained at 900−1050 °C}
\begin{tabular}{|c|c|c|c|}
\hline
sample & surface area/m\(^2\) g\(^{-1}\) & pore vol/cm\(^3\) g\(^{-1}\) & av pore width/nm \\
\hline
Fe\(\text{O}_4\)/PGC-CS-900 & 391.22 & 0.22 & 12.59 \\
Fe\(\text{O}_4\)/PGC-CS-950 & 400.86 & 0.19 & 12.13 \\
Fe\(\text{O}_4\)/PGC-CS-1000 & 476.55 & 0.27 & 11.31 \\
Fe\(\text{O}_4\)/PGC-CS-1050 & 382.01 & 0.26 & 10.75 \\
Fe\(\text{O}_4\)/PGC-PS-900 & 371.78 & 0.25 & 10.63 \\
Fe\(\text{O}_4\)/PGC-PS-950 & 482.68 & 0.29 & 10.69 \\
Fe\(\text{O}_4\)/PGC-PS-1000 & 545.95 & 0.34 & 9.85 \\
Fe\(\text{O}_4\)/PGC-PS-1050 & 265.92 & 0.14 & 8.77 \\
\hline
\end{tabular}
\end{table}

Figure 3.

Figure 4. Thermogravimetry and differential scanning calorimetry (TG−DSC) analyses of Fe\(\text{O}_4\)/PGC-PS (A) and Fe\(\text{O}_4\)/PGC-CS (B).

Figure 3. TEM image of Fe\(\text{O}_4\)/PGC-CS (A), HRTEM images of PGC-CS (B), and HRTEM images of Fe\(\text{O}_4\) (C) for Fe\(\text{O}_4\)/PGC-CS. TEM image of Fe\(\text{O}_4\)/PGC-PS (D), HRTEM images of PGC-CS (E), and HRTEM images of Fe\(\text{O}_4\) (F) for Fe\(\text{O}_4\)/PGC-CS.
electrocatalytic activity, CV tests of different cathodes in 50 mM PBS saturated with O₂ and N₂ were conducted. As shown in Figure S6 (Supporting Information), PGC-PS and PGC-CS exhibited a weaker current response and also showed poor catalytic activity for ORR, and PGC-PS had a minimum current of 7.43 mA in O₂-saturated solution (Figure S7A, Supporting Information). The current of Fe₃O₄/PGC-CS (13.84 mA) was higher than that of Fe₃O₄/PGC-PS (11.96 mA), which might be due to the large surface area, high conductivity, and excellent ORR activity of the Fe₃O₄/PGC-CS composite. 15,16 The current values for Fe₃O₄/PGC composites were higher than that of Pt/C (10.82 mA). A significant decrease in current generation happened in the N₂-saturated solution test (Figure S7B). It should be noted that the redox peaks were unobvious in this study, consistent with the previously reported results,29 which might be attributed to the confines of CV testing methods for MFCs.

To examine the ORR kinetic current and the transferred electron number, RDE was tested in 50 mM O₂-saturated PBS at rotation rates of 400−3600 rpm. At the highest rotation rate (3600 rpm), Fe₃O₄/PGC-CS generated the highest limiting current of 0.186 mA, which was slightly higher than that of Fe₃O₄/PGC-PS (0.177 mA) and 10.06% higher than that of Pt/C (0.169 mA) (Figure 5A−C). The lowest limiting current (0.065 mA) was generated by PGC-PS, which indicated that Fe₃O₄ could be the important active functional site for ORR (Figure 5D,E). The kinetic current of different catalysts was calculated by the K−L equation (Figure 5F) and was 0.23, 0.22, 0.21, 0.11, and 0.07 mA for Fe₃O₄/PGC-CS, Fe₃O₄/PGC-PS, Pt/C, PGC-CS, and PGC-PS, respectively. The average number (n) of transferred electrons on Fe₃O₄/PGC-CS and Fe₃O₄/
PGC-PS was 3.70 and 3.61, respectively, which indicates the dominant pathway for ORR was the four-electron oxygen reduction, including Pt/C (n = 3.56). Additionally, Fe₃O₄-free PGC-CS (n = 2.5) and PGC-PS (n = 2.0) exhibited much poorer ORR activity, suggesting the presence of a two-electron ORR pathway on PGC. The Fe₃O₄/PGC composites afforded higher ORR activity than PGC composites due to the comparable synergistic catalytic activities of nano-Fe₃O₄ particles and PGC. The RDE results were consistent with those observed in CV scans, which could confirm the better ORR activity of the prepared composites.

Generally, the ORR activity was inevitably related to the unique structure and properties of the catalysts. The superior electrocatalytic activity obtained here was due to the synergistic effect of PGC and Fe₃O₄ nanoparticles. The size of the Fe₃O₄ nanoparticles (10–20 nm in diameter) obtained here was smaller than the previously reported 110 and 150 nm, which could improve the migration of oxygen atoms between PGC and Fe₃O₄, and provide a large surface area and various active sites for ORR. Furthermore, the carbon resources were derived from natural biomass, which was favorable for generating abundant oxygen-containing functional groups in PGC. All these factors made the complete oxygen reduction occur more easily in the four-electron pathway on Fe₃O₄/PGC composites. Two-electron ORR occurred on PGC in which oxygen was mainly reduced to H₂O₂ when catalyzed by pure carbon without an active metal component.

As shown in Figure 6, EIS was used to investigate the kinetic activities on the cathodes, and the diameters in the semicircles developed with Nyquist plots indicated that the charge transfer resistances (Rₑ) of the Fe₃O₄/PGC-PS (15.3 ± 0.2 Ω) and Fe₃O₄/PGC-CS (10.8 ± 0.3 Ω) electrodes were lower than that of Pt/C (18.1 ± 0.5 Ω) and 52.79% and 62.45% lower than those of PGC-PS (32.2 ± 0.2 Ω) and PGC-CS (28.76 ± 0.3 Ω), respectively (Figure 6, Table 2). The decrease in charge transfer resistance verified the hypothesis that the introduction of Fe species could enhance the conductivity, electron transportation, and entire ORR performance of the composites. Furthermore, the density of catalytic active sites could be effectively improved by the uniform distribution of Fe species in the PGC skeleton. The Rₑ value of the mixture cathode (7.2 ± 0.2 Ω) was the lowest one, which was probably caused by the parallel arrangement of two resistors (Fe₃O₄/PGC-PS and Fe₃O₄/PGC-CS).

As shown in Figure S7 (Supporting Information), the voltage output of MFCs with the mixed cathode was higher than those of the other five cathodes. The voltage output of all MFCs at a fixed resistance (1000 Ω) gradually decreased after 800 h due to the formation of a biofilm on the cathode surface. It can be seen from Figure 7A that the average maximum voltage output of MFCs with the mixed cathode (0.64 ± 0.03 V) over the 21 feeding cycles was the highest. The voltage outputs of Fe₃O₄/PGC-PS (0.61 ± 0.04 V) and Fe₃O₄/PGC-CS (0.61 ± 0.02 V) were almost the same and were slightly higher than that of Pt/C (0.58 ± 0.05 V) and 52.13% and 37.54% higher than those of PGC-PS (0.292 ± 0.02 V) and PGC-CS (0.381 ± 0.04 V), respectively. By using the same anodes, the properties of the cathodes were dramatically influenced by the oxygen permeability and biofilm grown on the solution side of the cathodes. Three-dimensional structures of the mixed cathode were constructed by mixing 50% Fe₃O₄/PGC-CS and 50% Fe₃O₄/PGC-PS, which was favorable for ORR activity and antibiotic carbon. The Coulombic efficiencies (CEs) and COD removal rates of MFCs with Fe₃O₄/PGC-CS were slightly higher (about 5%) than those of Pt/C and almost the same as those of the other two cathodes (Figure 7B). Although the voltage output of MFCs with the mixture cathode was the highest, this did not mean that the CEs related to the electronic recycling efficiency must be high.

Because the stability of long-term operation was one of the most significant indicators of the performance of MFCs, the power density and corresponding electrode potentials of four catalysts were tested in the initial and final cycles (Figure 8). In the initial cycle (Figure 8A), the OCV for the mixture cathode was about 0.85 ± 0.05 V, and the mixture cathode showed the highest power density of 1502 ± 2,0.77 mW m⁻², which was about 40.09% and 12.26% higher than those of Fe₃O₄/PGC-CS (1443 ± 63 mW m⁻², 0.80 ± 0.02 V) and Fe₃O₄/PGC-PS (1338 ± 86 mW m⁻², 0.79 ± 0.02 V), respectively. The electricity production capacities for the mixture, Fe₃O₄/PGC-CS, and Fe₃O₄/PGC-PS were all better than that of Pt/C (1192 ± 33 mW m⁻², 0.77 ± 0.02 V) and extremely superior to those of PGC-PS (292 ± 21 mW m⁻², 0.51 ± 0.02 V) and PGC-CS (381 ± 23 mW m⁻², 0.54 ± 0.05 V), consistent with the voltage output (Figure 7A). Figure 8B shows the unobvious change of the anode potential and distinctive variations of the cathode potential as the current density increased, which suggests that the difference in power generation is attributed to the different ORR properties of the cathode. The maximum power density obtained by this mixture cathode was higher than that of the
Table 2. Electrochemical Impedance Fitting Results of Different Cathodes

<table>
<thead>
<tr>
<th>param</th>
<th>Pt/C</th>
<th>Fe3O4/PGC-PS</th>
<th>Fe3O4/PGC-CS</th>
<th>mixture</th>
<th>PGC-PS</th>
<th>PGC-CS</th>
</tr>
</thead>
<tbody>
<tr>
<td>$R_s$ (Ω)</td>
<td>9.86</td>
<td>6.42</td>
<td>7.81</td>
<td>5.15</td>
<td>16.87</td>
<td>13.86</td>
</tr>
<tr>
<td>$R_i$ (Ω)</td>
<td>18.1 ± 0.5</td>
<td>15.3 ± 0.2</td>
<td>10.8 ± 0.3</td>
<td>7.2 ± 0.2</td>
<td>32.2 ± 0.2</td>
<td>28.76 ± 0.3</td>
</tr>
<tr>
<td>$Z_m$ (Ω)$^{-1/2}$</td>
<td>$3.86 \times 10^{-13}$</td>
<td>$4.23 \times 10^{-13}$</td>
<td>$6.89 \times 10^{-13}$</td>
<td>$3.25 \times 10^{-13}$</td>
<td>$7.79 \times 10^{-14}$</td>
<td>$8.64 \times 10^{-14}$</td>
</tr>
<tr>
<td>$Q$ (F s$^{-1}$)</td>
<td>$7.77 \times 10^{-3}$</td>
<td>$7.81 \times 10^{-3}$</td>
<td>$4.68 \times 10^{-3}$</td>
<td>$6.57 \times 10^{-3}$</td>
<td>$8.83 \times 10^{-3}$</td>
<td>$8.93 \times 10^{-3}$</td>
</tr>
<tr>
<td>$n$</td>
<td>0.78</td>
<td>0.56</td>
<td>0.71</td>
<td>0.52</td>
<td>0.98</td>
<td>0.88</td>
</tr>
</tbody>
</table>

Fe3O4/PGC-CS samples were much looser than the Fe3O4/PGC-PS samples, which led to a similar increase of thickness of the Fe3O4/PGC-CS cathode. Therefore, the power density of Fe3O4/PGC-CS was slightly better than that of Fe3O4/PGC-PS in the initial operation stage, but declined rapidly in the next operations, consistent with the change of electricity production (Figure 7A) and electronics utilization (Figure 7B). Generally, good oxygen permeability was favorable for ORR and unfavorable for inhibiting the growth of bacteria.37 A balance of these two effects could be achieved by the mixture cathode, which had the highest power density and stability.

As shown in Figure 9, the CE of the Fe3O4/PGC-PS cathode varied in a wide range of 25–61% as the current density increased from 0.82 to 5.4 A m$^{-2}$, which was higher than that of Pt/C (20–50%) over the same range of current densities and slightly higher than those of the mixture (23–55%, 0.88–6.1 A m$^{-2}$) and Fe3O4/GC-CS (23–55%, 0.89–4.6 A m$^{-2}$). It was interesting that the mixed cathode for MFCs had the highest power density output, while the Fe3O4/GC-CS and Fe3O4/PGC-CS cathodes had the highest CEs (vs cycles and vs current densities, respectively). The Fe3O4/GC composite cathode could efficiently decrease the ORR overpotential and improve the interfacial contact for proton transport.38 Besides, the gradually graphitized accumulation process provided smaller, highly dispersed, and sufficient surface active sites for ORR.39 However, they could also be attacked by peroxide radicals (H2O2) formed on the surface of the three-phase boundary via insufficient ORR progress, which could also have an adverse effect on mass transfer and stability.40 From this perspective, the actual work number of catalytic active sites of the nano-Fe3O4/GC composites was larger than that of the commercial Pt/C catalyst. Therefore, the Fe3O4/GC composite or mixture could be used as an alternative MFC cathode catalyst for its high-efficiency and low-cost electricity production without losing power efficiency and stability.

Co3O4/stainless steel mesh or Fe/Fe3C/C nanorod cathode catalyst reported previously.35,36 After the 18th cycle (Figure 8C,D), the highest power density (1395 ± 46 mW m$^{-2}$, 0.82 ± 0.06 V) was still obtained by the mixed composite cathode, and it only slightly declined by 7.12%, which was lower than that (17.0%) of Pt/C (989 ± 81 mW m$^{-2}$, 0.71 ± 0.02 V). The power density of the Fe3O4/GC-CS and Fe3O4/GC-CS-PS reactors declined by 16.9% (1182 ± 5 mW m$^{-2}$, 0.78 ± 0.02 V) and 10.2% (1202 ± 67 mW m$^{-2}$, 0.71 ± 0.01 V), respectively. The decline of the power density for Pt/C, Fe3O4/GC-CS, and Fe3O4/GC-CS-PS was higher than that of the mixed composites, which was attributed to the higher oxygen permeability and greater biofilm growth. For the mixed composite cathode, biofilm growth might be slower than that of the Pt/C cathode because it was thick enough for inhibiting oxygen permeability (~0.5 mm for the mixed composite cathode and ~0.15 mm for Pt/C). Besides, the Fe3O4/GC-CS mixture cathode had the highest CEs (vs cycles and vs current densities, respectively). The Fe3O4/GC composite cathode could efficiently decrease the ORR overpotential and improve the interfacial contact for proton transport.38 Besides, the gradually graphitized accumulation process provided smaller, highly dispersed, and sufficient surface active sites for ORR.39 However, they could also be attacked by peroxide radicals (H2O2) formed on the surface of the three-phase boundary via insufficient ORR progress, which could also have an adverse effect on mass transfer and stability.40 From this perspective, the actual work number of catalytic active sites of the nano-Fe3O4/GC composites was larger than that of the commercial Pt/C catalyst. Therefore, the Fe3O4/GC composite or mixture could be used as an alternative MFC cathode catalyst for its high-efficiency and low-cost electricity production without losing power efficiency and stability.

**CONCLUSIONS**

Nano-Fe3O4/GC composites including Fe3O4/GC-CS and Fe3O4/GC-CS-PS were synthesized from waste agricultural biomass (cornstalks and pomelo skins) via the in situ simultaneous reduction method. The electrochemical properties of Fe3O4/GC used in single air-cathode microbial fuel cells (MFCs) as cathode catalysts were examined. The results showed that the Fe3O4/GC composites exhibited high electrocatalysis activity for ORR. High-efficiency electricity generation could be obtained in MFCs with the Fe3O4/GC and mixture cathodes and was more economical and sustainable than that with the Pt/C cathode. The Fe3O4/GC composites, by utilization of low-cost carbon resources, could be promising catalysts for practical application and commercialization of MFCs.
ASSOCIATED CONTENT

Supporting Information
Schematic diagram of the structure of carbon-constructed cathodes (Figure S1), equivalent circuit (Figure S2), nitrogen adsorption/desorption isotherms and pore size distribution of Fe₃O₄/GC-CS and Fe₃O₄/GC-PS obtained at different temperatures (Figures S3 and S4), linear sweep voltammetry curves (Figure S5), cyclic voltammetry current response for ORR (Figure S6), and voltage output of MFCs with different cathodes (Figure S7). This material is available free of charge via the Internet at http://pubs.acs.org/.

Figure 8. Power density and the corresponding electrode potentials using different cathodes as a function of the current density at the 4th cycle (A, B; the cathodes include Pt/C, Fe₃O₄/GC-PS, Fe₃O₄/GC-CS, a mixture of the two composites, PGC-PS, and PGC-CS) and the 18th cycle (C, D; the cathodes include Pt/C, Fe₃O₄/GC-PS, Fe₃O₄/GC-CS, and a mixture of the two composites).

Figure 9. Relationship of Coulombic efficiency and current density for microbial fuel cells with different cathodes.

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Notes
The authors declare no competing financial interest.

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