Hydrothermally Carbonized Waste Biomass as Electrocatalyst Support for $\alpha$-MnO$_2$ in Oxygen Reduction Reaction

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Abstract: Sluggish kinetics in oxygen reduction reaction (ORR) requires low-cost and highly durable electrocatalysts ideally produced from facile methods. In this work, we explored the conversion and utilization of waste biomass as potential carbon support for $\alpha$-MnO$_2$ catalyst in enhancing its ORR performance. Carbon supports were derived from different waste biomass via hydrothermal carbonization (HTC) at different temperature and duration, followed by KOH activation and subsequent heat treatment. Scanning electron microscopy (SEM), Fourier transform infrared spectroscopy (FTIR), energy dispersive X-ray spectroscopy (EDX) and X-Ray diffraction (XRD) were used for morphological, chemical, and structural characterization, which revealed porous and amorphous carbon supports for $\alpha$-MnO$_2$. Electrochemical studies on ORR activity suggest that carbon-supported $\alpha$-MnO$_2$ derived from HTC of corncobs at 250 $^\circ$C for 12 h (CCAC + MnO$_2$ 250-12) gives the highest limiting current density and lowest overpotential among the synthesized carbon-supported catalysts. Moreover, CCAC + MnO$_2$ 250-12 facilitates ORR through a 4-e$^-$ pathway, and exhibits higher stability compared to VC + MnO$_2$ (Vulcan XC-72) and 20% Pt/C. The synthesis conditions preserve oxygen functional groups and form porous structures in corncobs, which resulted in a highly stable catalyst. Thus, this work provides a new and cost-effective method of deriving carbon support from biomass that can enhance the activity of $\alpha$-MnO$_2$ towards ORR.

Keywords: hydrothermal carbonization; biomass; hydrochar; activated carbon; MnO$_2$/C; oxygen reduction reaction

1. Introduction

The use of electrochemical energy technologies, such as batteries and fuel cells, are effective and practical ways of dealing with stability issues brought by intermittent renewable energy sources. At present, lithium-ion chemistries dominate among energy storage technologies and cover a wide array of applications. However, metal-air batteries (MABs) are also seen to address the challenges for renewable energy applications and other niche applications [1,2].

The utilization of ambient air in energy storage applications is constrained by the sluggish kinetics of the oxygen reduction reaction (ORR) [3]. As such, previous studies have focused on the development of electrocatalysts to enhance ORR performance [4–6]. ORR electrocatalysts should provide low ORR
overpotential, high durability, and selectivity towards the ideal 4-electron (e⁻) pathway. Moreover, the material should be abundantly available and can be easily synthesized [7]. Electrocatalysts that were previously investigated include precious metals and their alloys, transition metals, metal oxides, and carbon-based materials [1]. Transition metal oxides have also been formed into a matrix with carbon-based structures to improve ORR activity [8–10]. Among the transition metal oxides, manganese oxides (MnO₂) are widely recognized as suitable ORR catalysts due to their high catalytic activity, low cost, and environmental friendliness [11–13].

MnO₂ comes in various polymorphs. Two of which, α-MnO₂ and δ-MnO₂, are reported to have the relatively high ORR catalytic activities [12,13]. The former is considered to have better performance due to higher Mn³⁺ surface concentration, which allows for more electroactive sites for catalysis [14]. The drawbacks of α-MnO₂, however, include poor conductivity and low stability. Thus, catalyst supports such as carbon black, graphene, and carbon nanotubes are typically used to improve MnO₂’s catalytic performance [10,15].

Catalyst supports assist in metal dispersion and morphological formation of the electrocatalyst. As a result, heat and mass transfer properties are enhanced, thereby improving catalyst efficiency [16]. Moreover, interactions of the carbon support and the metal species result in a faster electron transfer and improved absorption/desorption characteristics for oxygen intermediates [17]. Conversion of low-cost carbon-based materials to catalyst supports, however, involves complex methods [18]. Therefore, facile methods for carbon-based support synthesis are investigated in several studies [19–24].

For an efficient, low-cost production, carbon can be derived from different sources, including waste biomass [19]. Natural structures in biomass such as periodic patterns, hierarchical organization, and nanoarchitectures allow for special properties such as super hydrophobicity, structural coloration, and biological self-assembly. These structures can also provide networks for electrochemically active sites, which are beneficial for catalysis [20,21]. Mass producible biomass such as jackfruit seeds [19], eggplant [23], and silk [24] have been explored for electrocatalyst materials and showed promising results as carbon-based supports and air electrodes.

Among the different methods used to carbonize biomass, hydrothermal carbonization (HTC) is considered as the most cost-effective since it allows for the conversion of high moisture materials [25,26]. HTC requires no prior drying, utilizes water as media, and produces a lignite-like product [27]. This process is carried out at 180–250 °C under autogenous pressure. Carbon and hydrogen in the feedstock undergo decomposition by dehydration and decarboxylation [28]. The properties of the HTC product (called hydrochar) can be controlled by varying the synthesis conditions such as temperature, pH, and residence time [28–32]. Activation using potassium hydroxide (KOH) can also be employed to further enhance the porosity, surface area, and total pore volume of the hydrochar [33].

In this work, we investigate the derivation of carbon supports from different waste biomass materials including corncobs, coffee waste grounds, rice hulls, and coconut lumber sawdust. Their potential viability as carbon supports are then evaluated based on their morphology, and their ORR activity when used as support for α-MnO₂. Carbon supports were prepared via HTC at different temperatures and durations, followed by KOH activation. Treatment of waste biomass at 250 °C for 12 h produced carbon-supported MnO₂ with the highest ORR catalytic activity. This work provides a new and cost-effective method of deriving carbon supports from biomass through HTC for effective and durable ORR electrocatalysts.

2. Results and Discussion

2.1. Hydrothermal Carbonization

The presence of oxygen functional groups from cellulose and aromatic carbon bonds in lignin were investigated using Fourier transform infrared (FTIR) spectroscopy (Figure 1). The summary of peaks and the corresponding functional groups are summarized in Table 1.
Figure 1. FTIR spectra of raw and hydrochars of (a) corncobs (CC), (b) coffee waste grounds (CWG), (c) rice hull (RH) and (d) coconut sawdust (SD) at different temperatures (200 and 250 °C) and residence time (6 and 12 h).

Table 1. Peaks and associated functional groups of produced hydrochars.

<table>
<thead>
<tr>
<th>Peaks (cm⁻¹)</th>
<th>Functional Group</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>1655–1700</td>
<td>C=O and C=C vibration from ketones and aromatic ring structures</td>
<td>[34]</td>
</tr>
<tr>
<td>1450–1510</td>
<td>C=C–C bonds vibration in an aromatic ring</td>
<td>[35]</td>
</tr>
<tr>
<td>1310–1410</td>
<td>O–H bends and aliphatic CH₃ and CH₂ deformation</td>
<td>[35]</td>
</tr>
<tr>
<td>1050–1270</td>
<td>Aryl–O and C–O stretch from ether compounds</td>
<td>[36]</td>
</tr>
<tr>
<td>1000–1055</td>
<td>C–O bonds in glucose</td>
<td>[36]</td>
</tr>
<tr>
<td>900</td>
<td>C–H groups with aromatic out-of-plane bends</td>
<td>[37,38]</td>
</tr>
</tbody>
</table>

The HTC process reduced the volatile oxygen functional groups from cellulose (1655, 1310, 1270 and 1050 cm⁻¹). The high thermal stability of lignin [39], however, resulted in the preservation of functional groups in aromatic rings (1600, 1500 and 1450 cm⁻¹). Increasing the temperature from 200 to 250 °C resulted to a higher degree of degradation compared to increasing the HTC duration from 6 to 12 h.

The changes in the morphological structures were examined using scanning electron microscopy (SEM). SEM images of raw and hydrothermally carbonized samples are shown in Figure 2. After HTC, the porous structures are preserved with lignin serving as the backbone [39]. Increasing the HTC temperature resulted in size reduction of globular masses found in the hydrochars. This is due to nucleation and polymerization reactions, which led to the breakage of the fibrous structure and the creation of micrometric cellulose fragments [40]. Similarly, an increase in the residence time resulted in the formation of a different hydrochar with polyaromatic structure, which consists of the polymerized fragments [41]. From the SEM images, rice hull (RH) was observed to have the least surface variation before and after treatment compared to the other samples. Similar observations were made for the HTC of RH in other studies, which were attributed to partial carbonization at the given HTC conditions. Therefore, the hydrochar from RH is expected to have low surface area and pore volumes [42].
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Figure 2. SEM images at 2500× magnification of the (a–d) corncobs (CC), (e–h) coffee waste grounds (CWGs), (i–l) rice hull (RH) and (m–p) coconut sawdust (SD) at varying HTC temperatures (200 and 250 °C) and duration (6 and 12 h). A measurement scale is located at the lower-right of the figure.

2.2. Activation

The hydrochars from the different biomass samples were then subjected to KOH activation followed by heat treatment at 600 °C in an N₂ environment. The effect of activation on the functional groups of hydrochars was then examined using FTIR, as shown in Figure 3. Results showed reduction of peak strength of the functional groups in all waste biomass samples due to volatilization and degradation [30,43]. Coffee waste grounds (CWG) and RH showed a relatively higher degree of functional group degradation compared to corn cobs (CC) and coconut sawdust (SD). Among the samples subjected to HTC at 250 °C for 12 h prior to activation, activated CWG (CWGAC) and activated RH (RHAC) exhibited the lowest peaks while activated CC (CCAC) and activated SD (SDAC) retained aromatic groups at peaks located at 1655, 1310, 1270 and 1050 cm⁻¹.

The morphological examination of hydrochars after KOH activation and heat treatment are shown in Figure 4. Under the same activation conditions, samples that were pre-carbonized through HTC at 250 °C and 12 h showed higher degrees of formation of porous structures compared to those at lower temperature and residence time. Activation allowed for the opening of more pores, which were not achieved through HTC alone. Increased pore formation translates to formation of networks that can provide active sites for ORR [44]. Further, pre-carbonization reduces the formation of carbonaceous intermediate defects while it increases the micropore formation [45]. The pre-carbonization step (HTC) before activation was therefore essential in achieving stable and effective carbon support structures.
Figure 3. FTIR spectra of activated carbon from (a) corncobs (CCAC), (b) coffee waste grounds (CWGAC), (c) rice hull (RHAC) and (d) coconut sawdust (SDAC).

Figure 4. SEM images at 2500× magnification after activation of (a–d) corncobs (CCAC), (e–h) coffee waste grounds (CWGAC), (i–l) rice hull (RHAC) and (m–p) coconut sawdust (SDAC) that were pre-carbonized in HTC temperatures (200 and 250 °C) and residence times (6 and 12 h).
The elemental composition of the raw biomass and activated carbons that were analyzed using energy dispersive X-ray (EDX) spectroscopy are summarized in Table 2. The results show an increase in the carbon concentrations and the reduction of other elements after HTC and KOH activation. Of the samples that were synthesized, CCAC 250-12 showed the highest carbon concentration. Analysis of RHAC 250-12 showed that silicon (Si) was still present after carbonization. EDX data also indicate that there were no metals or other impurities in the synthesized carbon supports other than silicon (in the form of silica), which is inherently present in rice hulls [42]. This result also supports the FTIR results of the samples that were subjected to HTC at 250 °C for 12 h since no peaks were found at 1342–1260 cm\(^{-1}\) and 710–570 cm\(^{-1}\). Such peaks are used to gauge the presence of C–N and C–S bonds respectively [36–38].

**Table 2. Elemental composition of raw biomass and activated carbons.**

<table>
<thead>
<tr>
<th>Sample</th>
<th>%Weight Concentration</th>
<th>C</th>
<th>O</th>
<th>N</th>
<th>K</th>
<th>Si</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>CC Raw</td>
<td>42.36</td>
<td>52.66</td>
<td>3.95</td>
<td>1.02</td>
<td>100.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CCAC 250-12</td>
<td>83.40</td>
<td>16.60</td>
<td></td>
<td></td>
<td></td>
<td>100.00</td>
<td></td>
</tr>
<tr>
<td>CWG Raw</td>
<td>39.01</td>
<td>47.90</td>
<td>12.02</td>
<td>1.08</td>
<td>100.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CWGAC 250-12</td>
<td>76.79</td>
<td>23.21</td>
<td></td>
<td></td>
<td></td>
<td>100.00</td>
<td></td>
</tr>
<tr>
<td>RH Raw</td>
<td>46.06</td>
<td>22.70</td>
<td>3.56</td>
<td>0.00</td>
<td>27.68</td>
<td>100.00</td>
<td></td>
</tr>
<tr>
<td>RHAC 250-12</td>
<td>61.92</td>
<td>31.71</td>
<td></td>
<td></td>
<td>6.38</td>
<td>100.00</td>
<td></td>
</tr>
<tr>
<td>SD Raw</td>
<td>41.33</td>
<td>49.38</td>
<td>8.41</td>
<td>0.89</td>
<td>0.00</td>
<td>100.00</td>
<td></td>
</tr>
<tr>
<td>SDAC 250-12</td>
<td>73.66</td>
<td>26.34</td>
<td></td>
<td></td>
<td></td>
<td>100.00</td>
<td></td>
</tr>
</tbody>
</table>

The crystal phase of activated carbon was determined using XRD (Figure 5). In the analysis, a broad peak was identified at 15–30 degrees, which indicates the presence of amorphous carbon [46,47]. The broad diffraction peaks in 20.8 and 21.6 in XRD pattern refers to the (002) facet [46]. Among the samples, higher intensities were observed for CCAC 250-12 and SDAC 250-12. Sharp peaks that were generated were attributed to the gold (Au) metal holder used in the XRD analysis.

![Figure 5. XRD data for (a) CCAC 250-12, (b) CWGAC 250-12, (c) RHAC 250-12 and (d) SDAC 250-12.](attachment:image.png)
2.3. Incorporation of MnO₂

The hydrochars and activated carbons from the different biomass were embedded with an MnO₂ catalyst via the hydrothermal method. Figure 6 shows SEM images of activated samples after MnO₂ embedding. Based on the images, the change in the surface roughness of the samples indicate that MnO₂ was successfully embedded into the carbon supports. In Figure 7, the presence of the α-MnO₂ was confirmed by the XRD peaks.

![Figure 6. SEM images at 2500× magnification for (a) α–MnO₂ and after embedding MnO₂ to activated hydrochar (b) CCAC + MnO₂, (c) CWGAC + MnO₂, (d) RHAC + MnO₂ and (e) SDAC + MnO₂ from HTC at 250 °C at 12 h.](image)

![Figure 7. XRD pattern of the synthesized MnO₂.](image)

2.4. ORR Activity

In Figure 8, the cyclic voltammetry (CV) curves of activated carbon (AC) derived from hydrochars with embedded α-MnO₂ were obtained after three cycles from 0.035 to −0.096 V vs. Ag/AgCl at a scan rate of 50 mV s⁻¹ in 0.1 M KOH electrolyte saturated with N₂ or O₂. The CV curves in the N₂-saturated solutions showed capacitive behavior in all cases. The appearance of peaks in the O₂-saturated solution is thus attributed to ORR. Well-defined cathodic peaks corresponding to ORR were obtained from CCAC + MnO₂ and CWGAC + MnO₂ in contrast with RHAC + MnO₂ and SDAC + MnO₂. In the same figure, CCAC + MnO₂ 250-12 shows the highest area enclosed in the CV curve, which is indicative of the power per active surface area of the electrocatalyst (W cm⁻²) resulting to a better ORR performance.
The ORR performance of pure AC were then compared using linear sweep voltammetry (LSV) along with hydrochars and AC embedded with α-MnO₂ (Figure 9). LSV was carried out at 1600 rpm in 0.1 M KOH solution. The results show that the activation process consistently improved the ORR performance of their unactivated counterparts. The activation process could have potentially increased the open edge sites and curvatures, which assisted oxygen molecules to reach active sites for ORR catalysis [42]. Furthermore, the figures also suggest that the incorporation of α-MnO₂ in the activated carbon improved the ORR performance. This is observed also when doped carbon materials were added to the active electrocatalyst in other studies [10,15,18]. Among the samples that were analyzed, hydrochar and activated carbon produced from HTC at 250 °C for 12 hours showed the highest performance, which is consistent with the CV results. Increasing the temperature and duration of HTC leads to the formation of micrometric structures [29,35], thereby improving ORR catalytic activity.

The activity of α-MnO₂ supported by biomass-derived carbons were also compared with Vulcan XC-72 (VC) supported α-MnO₂, and 20% Pt/C (Figure 10). Comparison of the limiting current density ($i_{\text{lim}}$), onset potential ($E_{\text{onset}}$), and overpotential of samples are shown in Table 3. Results indicate that among the biomass derived samples, CCAC + MnO₂ 250-12 exhibited the highest limiting current density (2.85 mA cm⁻²) and lowest overpotential (0.50 V) at the onset of ORR. However, the values were still lower compared to VC + MnO₂ and 20% Pt/C. The large difference between the limiting current density of 20% Pt/C with the different carbon-supported α-MnO₂ could be attributed to the loading ratio of α-MnO₂ to the carbon support. In the study, α-MnO₂ was estimated to constitute 33% of the catalyst loading. The synthesis method, however, involved the direct incorporation of the carbon during α-MnO₂ synthesis, which may have resulted in a lower catalyst loading. Investigation of an appropriate α-MnO₂ to carbon ratio should be pursued in future works to optimize the performance of the developed carbon-supported electrocatalyst. Onset potentials of hydrothermally-prepared catalysts follow the behavior of biomass derived carbon for ORR since carbonaceous compounds have inherently low ORR activity in alkaline media [48–50]. However, if carbon is used as a support for

![Figure 8. Cyclic voltammetry (CV) results for (a) CCAC + MnO₂ 250-12, (b) CWGAC + MnO₂ 250-12, (c) RHAC + MnO₂ 250-12 and (d) SDAC + MnO₂ 250-12 in N₂ and O₂ saturated 0.1 M KOH electrolyte.](image-url)
the metal oxide, rather than as a pure electrocatalyst, the conductivity of the composite enhances and facilitates the electron transfer to the electroactive sites of the metal oxide. [51]. The support also provides a conducting network that enables charge transfer on the metal oxide surface during electrochemical reactions such as ORR [52].

Figure 9. Linear sweep voltammetry (LSV) curves for hydrochar with MnO₂, pure activated carbon (AC) and AC embedded with MnO₂ derived from (a–c) corncobs, (d–f) coffee waste grounds, (g–i) rice hull and (j–l) coconut sawdust in 0.1 M KOH at 0.244 mg cm⁻² loading, scan rate of 5 mV s⁻¹ and rotation speed of 1600 rpm.
The results showed that carbon supported MnO$_2$ generally follow a 4-e$^-$ pathway for CCAC + MnO$_2$ 250-12. The number of electrons transferred to oxygen bond cleaves more easily which benefits the ORR performance [53]. The FTIR results included C=O [34], aryl–O, and C–O [36]. The presence of oxygen-containing groups were found to improve ORR activity in an alkaline environment [53]. It was postulated that the oxygen adsorbed to the active carbon site has a synergistic effect on the oxygen of the carboxyl group. As a result, the carbon to oxygen bond cleaves more easily which benefits the ORR performance [53]. The FTIR results for SDAC 250-12 from Table 2 also indicated the presence of the said functional groups. However, its low activity can be attributed to insufficient development of a porous structure, which hindered the progress of electrochemical reactions on the catalyst surface. In addition, CCAC showed the highest carbon concentration in the EDX analysis which can allow for a greater area for the electrocatalyst.

The higher performance of the support from corncobs compared to the other biomass can be attributed to the structures left after HTC and KOH activation as discussed in the FTIR results. The regions 1600–1000 cm$^{-1}$ wherein peaks still remain for CCAC 250-12 while others have been greatly reduced, are regions with oxygen-containing functional groups. These functional groups include C=O [34], aryl–O, and C–O [36]. The presence of oxygen-containing groups were found to improve ORR activity in an alkaline environment [53]. It was postulated that the oxygen adsorbed to the active carbon site has a synergistic effect on the oxygen of the carboxyl group. As a result, the carbon to oxygen bond cleaves more easily which benefits the ORR performance [53]. The FTIR results for SDAC 250-12 from Table 2 also indicated the presence of the said functional groups. However, its low activity can be attributed to insufficient development of a porous structure, which hindered the progress of electrochemical reactions on the catalyst surface. In addition, CCAC showed the highest carbon concentration in the EDX analysis which can allow for a greater area for the electrocatalyst. Therefore, it is important to consider both the functional groups and the pore structure resulting from HTC and activation to produce the most suitable carbon support.

To further elucidate the electrochemical properties of activated-carbon-supported MnO$_2$ samples, the electron transfer number was determined through LSV at different rotation speeds (Figure 11). Koutecky–Levich (KL) plots were then generated from $-0.96$ to $-0.75$ V vs. Ag/AgCl (Figure 12). The results showed that carbon supported MnO$_2$ generally follow a 4-e$^-$ pathway for CCAC + MnO$_2$ 250-12 and CWGAC + MnO$_2$ 250-12. On the other hand, RHAC + MnO$_2$ and SDAC + MnO$_2$ 250-12

**Figure 10.** LSV curves for carbon supported MnO$_2$ of different activated HTC-treated biomass, (Vulcan) supported MnO$_2$ and 20% Pt/C in 0.1 M KOH at 0.244 mg cm$^{-2}$ loading, scan rate of 5 mV s$^{-1}$ and at 1600 rpm rotation speed.

**Table 3.** Electrocatalytic activity of carbon supported electrocatalysts and 20% Pt/C.

<table>
<thead>
<tr>
<th>Sample</th>
<th>$j_L$ (mA cm$^{-2}$)</th>
<th>$E_{ons}$ (V vs. Ag/AgCl)</th>
<th>Overpotential (V)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CCAC + MnO$_2$ 250-12</td>
<td>-2.85</td>
<td>-0.23 (0.73)$^1$</td>
<td>0.50</td>
</tr>
<tr>
<td>CWGAC + MnO$_2$ 250-12</td>
<td>-2.62</td>
<td>-0.25 (0.71)$^1$</td>
<td>0.52</td>
</tr>
<tr>
<td>RHAC + MnO$_2$ 250-12</td>
<td>-2.72</td>
<td>-0.35 (0.61)$^1$</td>
<td>0.62</td>
</tr>
<tr>
<td>SDAC + MnO$_2$ 250-12</td>
<td>-2.58</td>
<td>-0.36 (0.6)$^1$</td>
<td>0.63</td>
</tr>
<tr>
<td>VC + MnO$_2$</td>
<td>-2.67</td>
<td>-0.17 (0.79)$^1$</td>
<td>0.44</td>
</tr>
<tr>
<td>20% Pt/C</td>
<td>-4.63</td>
<td>-0.098 (0.87)$^1$</td>
<td>0.36</td>
</tr>
</tbody>
</table>

$^1$ 1 V vs. RHE.
showed a combination of a 2-e\textsuperscript{−} and 4-e\textsuperscript{−} pathway. The number of electrons transferred is greatest in CCAC + MnO\textsubscript{2} 250-12 (Figure 12a) among tested samples, which further supports that carbon support derived from CC has also better ORR performance.

**Figure 11.** LSV curves for different rotation speeds for (a) CCAC + MnO\textsubscript{2} 250-12, (b) CWGAC + MnO\textsubscript{2} 250-12, (c) RHAC + MnO\textsubscript{2} 250-12 and (d) SDAC + MnO\textsubscript{2} 250-12.

**Figure 12.** Koutecky–Levich (KL) plots at different potential for (a) CCAC + MnO\textsubscript{2} 250-12, (b) CWGAC + MnO\textsubscript{2} 250-12, (c) RHAC + MnO\textsubscript{2} 250-12 and (d) SDAC + MnO\textsubscript{2} 250-12.
The ORR mechanism for the electron pathway that an electrocatalyst follows depends on the interaction of oxygen molecules with the catalyst surface. Adsorption of O$_2$ through a horizontal mode on $\alpha$-MnO$_2$ proceeds via a bridge-type connection, which is consistent with the 4-e$^-$ pathway [54,55]. While, a vertical mode adsorption follows a two electron 2-e$^-$ reduction pathway [55,56]. Most carbonaceous materials follow a 2-e$^-$ pathway [57] while MnO$_2$ follows a 4-e$^-$ in an alkaline environment [12,51]. Although the synthesized supports are essentially carbonaceous in nature, differences in the structure of each biomass potentially resulted in different adsorption mechanisms. This implies that CCAC and CWGAC have structures, which supports the electrocatalytic activity of $\alpha$-MnO$_2$ towards a horizontal mode of O$_2$ adsorption and thus favoring the 4-e$^-$ pathway. On the other hand, it can be concluded that RHAC- and SDAC-supported $\alpha$-MnO$_2$ exhibits ORR that proceeds both in 2-e$^-$ and 4-e$^-$ pathways, which resulted in an electron transfer number approaching three (3). In other studies [10,19,58], the electron pathway is influenced both by the individual ORR mechanism of the carbon support and the electrocatalyst. For such materials, ORR proceeds both in the composites, hence a mixed behavior can be expected. In this work, the $\alpha$-MnO$_2$-mediated pathway for ORR provided a greater contribution compared to the carbon-mediated pathway for CCAC + MnO$_2$ 250-12 and CWGAC + MnO$_2$ 250-12. Whereas, ORR in RHAC + MnO$_2$ 250-12 and SDAC + MnO$_2$ 250-12 proceeded to undergo ORR equally in the carbon support and in $\alpha$-MnO$_2$. Therefore, the carbon from CC and CWG were more effective supports for enhancing ORR activity of $\alpha$-MnO$_2$ compared to those derived from RH and SD since the former group were able to retain the activity of the electrocatalyst.

Material stability is also a concern for carbonaceous compounds and noble metals in ORR electrocatalysis. Accelerated durability tests (ADT) were conducted for CCAC + MnO$_2$ 250-12, VC + MnO$_2$ and 20% Pt/C in 0.1 M KOH at 50 mV s$^{-1}$ for 5000 cycles. A comparison of limiting current density ($j_L$) and half wave potential ($E_{1/2}$) is summarized in Table 4 while the voltammetric curves are illustrated in Figure 13. Figure 13e,f show significant reduction of performance for 20% Pt/C. On the other hand, CCAC-MnO$_2$ 250-12 showed less reduction in activity compared to VC + MnO$_2$. The $E_{1/2}$ for 20% Pt/C showed no changes while significant reduction was observed for VC + MnO$_2$. A reduction of the limiting current density value was significant for both 20% Pt/C and VC + MnO$_2$ which amounts to 140% and 98% reduction, respectively. The activity decrease implies the degradation of surface-active sites over time. Meanwhile, CCAC + MnO$_2$ 250-12 showed minor changes in activity compared to the 20% Pt/C and VC + MnO$_2$, which suggests higher durability.

<table>
<thead>
<tr>
<th>Sample</th>
<th>$E_{1/2}$ (V vs. Ag/AgCl)</th>
<th>$j_L$ (mA cm$^{-2}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Initial</td>
<td>Final</td>
</tr>
<tr>
<td>CCAC + MnO$_2$ 250-12</td>
<td>-0.32</td>
<td>-0.39</td>
</tr>
<tr>
<td>VC + MnO$_2$</td>
<td>-0.29</td>
<td>-0.38</td>
</tr>
<tr>
<td>20% Pt/C</td>
<td>-0.11</td>
<td>-0.11</td>
</tr>
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In this work, we developed a new facile method for the conversion of waste biomass to carbon-based supports for ORR electrocatalysts. The MnO$_2$-embedded carbon-based catalysts were also successfully synthesized, and their ORR performance analyzed. The use of the carbon-supported MnO$_2$ catalysts developed in this study could be further tested for metal-air battery systems, which have promising applications as compact power sources and energy storage devices [59]. Successful commercialization has already been reported for primary MAbs particularly in zinc-air battery systems [60]. Still, different studies have been made and are still ongoing to address constraints arising from high material costs, low catalytic efficiencies, and poor durability of materials [4–6,18–20]. The catalysts developed in this work are cheap and active alternative cathode materials that can potentially improve the performance of primary MAbs. As such, future works are intended to explore the characteristics of the developed electrocatalysts for metal-air battery applications.
In this work, we developed a new facile method for the conversion of waste biomass to carbon-based supports for ORR electrocatalysts. The MnO₂-embedded carbon-based catalysts were also successfully synthesized, and their ORR performance analyzed. The use of the carbon-supported MnO₂ catalysts developed in this study could be further tested for metal-air battery systems, which have promising applications as compact power sources and energy storage devices [59]. Successful commercialization has already been reported for primary MABs particularly in zinc-air battery systems [60]. Still, different studies have been made and are still ongoing to address constraints arising from high material costs, low catalytic efficiencies, and poor durability of materials [4–6,18–20]. The catalysts developed in this work are cheap and active alternative cathode materials that can potentially improve the performance of primary MABs. As such, future works are intended to explore the characteristics of the developed electrocatalysts for metal-air battery applications.

3. Materials and Methods

3.1. Hydrothermal Carbonization

Coffee waste grounds (CWG) were prepared by boiling at 100 °C using deionized water for three hours to remove volatile organic compounds. Corncobs (CC) were first milled into 1 mm product while rice hull (RH) and coconut sawdust (SD) were directly subjected to hydrothermal carbonization (HTC). A ratio of 32 g biomass to 100 mL deionized water was placed in a polytetrafluoroethylene (PTFE)-lined stainless-steel autoclave and was treated in a furnace. Residence time was varied at 6 and 12 h while temperatures were varied at 200 and 250 °C. The hydrochar produced was recovered using filtration and dried in a vacuum oven at 80 °C for 8 h. Samples were then cooled down to room temperature.
3.2. Activation

Hydrochar samples were mixed at a 1:3 ratio with KOH solution (RCI Labscan Ltd., Bangkok, Thailand) and heated to 90 °C for 30 minutes with constant stirring. KOH treated samples were then filtered and dried in a vacuum oven at 90 °C for 4 h. Dried samples were then heat treated in a tube furnace (LTF 12, Lenton Furnaces & Ovens, Hope Valley, United Kingdom) at 600 °C for one hour under N₂ atmosphere (200–300 mL min⁻¹) with a ramp rate of 5 °C min⁻¹. After cooling, the samples were washed with 1 M HCl (RCI Labscan Ltd., Bangkok, Thailand) followed by copious amounts of deionized water. Finally, the activated samples were dried at 90 °C for 4 h in a vacuum oven.

3.3. Incorporation of MnO₂

Equal amounts of 2 M manganese sulfate (MnSO₄; Loba Chemie Pvt. Ltd., Mumbai, India) and 2 M ammonium persulfate (NH₄)₂S₂O₈ (Loba Chemie Pvt. Ltd., Mumbai, India) were added to activated and hydrochar samples corresponding to the 1:2 w/w ratio of MnO₂ to carbon. The solution was then transferred to a PTFE lined stainless steel autoclave and hydrothermally treated for 3 h at 120 °C [13]. Samples were then taken out of the autoclave and allowed to cool to room temperature. Finally, the MnO₂ samples were filtered and dried for 6 h in a vacuum oven at 90 °C.

3.4. Material Characterization

Morphological analysis was performed using scanning electron microscopy (SEM; ZEISS Gemini SEM 500, Jena, Germany), determination of crystalline structures using X-ray diffraction (XRD; Maxima XRD-7000 Shimadzu Corporation, Kyoto, Japan), identification of functional groups on the samples surface using Fourier transform infrared (FT-IR) spectroscopy (Nicolet 6700, Wisconsin, USA) and elemental analysis using energy dispersive X-ray spectroscopy (EDX; Phenom XL SEM-EDX, Phenom-World B.V., Eindhoven, Netherlands).

3.5. Electrochemical Measurements

Electrocatalytic activity was measured against a platinum on carbon standard using linear sweep voltammetry (LSV). Samples from hydrochar, and unactivated and activated hydrochar embedded with MnO₂ were analyzed for the difference in electrocatalytic activities. The catalyst ink was prepared by dispersing 6 mg of samples in 950 μL ethanol (RCI Labscan Ltd., Bangkok, Thailand), which was then subjected to ultrasonication for 30 minutes. Then, 50 μL of Nafion solution (5 wt%) (Fuel Cell Store, Texas, USA) was added into the dispersion and ultrasonicated for 20 minutes. Vulcan XC-72 (Fuel Cell Store, Texas, USA) and 20% Pt/C (HiSPEC 3000, John Matthey Fuel Cells, Fuel Cell Store, Texas, USA) catalyst ink were also prepared to serve as standards for ORR performance. Finally, 0.244 mg cm⁻² loading of the catalysts was dropped in a glassy carbon electrode then onto a rotating ring disk electrode (RRDE). Electrochemical measurements were performed using a potentiostat (PGSTAT204, Metrohm, Herisau, Switzerland). The set-up used for voltammetry consisted of RRDE working electrode, Ag/AgCl reference electrode, and a platinum sheet counter electrode. Cyclic voltammetry was performed under N₂- and O₂-saturated 0.1 M KOH solutions at 50 mV s⁻¹. Linear sweep voltammetry was performed at 5 mV s⁻¹ scan rate from 0.035 to −0.96 V vs. Ag/AgCl at 1600 rpm.

The number of electrons transferred (n) was estimated according to the Koutecky–Levich equation [60].

\[ \frac{1}{j} = \frac{1}{B \omega^{-\frac{1}{2}}} + \frac{1}{j_k} \]  
\[ B = 0.62nFCE\sigma \frac{2}{\pi} \nu^{-\frac{1}{6}} \]  

where \( j \) is the recorded current density, \( j_k \) is the kinetic current density, \( \omega \) is the electrode rotating rates (rad s⁻¹), \( n \) means the transferred electron number, and \( F \) is the Faraday constant (96,485 C mol⁻¹).
C₀ (1.2 × 10⁻³ mol L⁻¹) is the bulk concentration, and D₀ (1.9 × 10⁻⁵ cm² s⁻¹) corresponds to the diffusion coefficient of O₂ in 0.1 mol L⁻¹. The ν represents the kinematic viscosity of the electrolyte (0.01 cm² s⁻¹) [61–63].

Accelerated degradation test was then performed from −0.36 to 0.035 V vs. Ag/AgCl at 0.1 M KOH for 5000 cycles. Changes in LSV and CV data were obtained from 0.035 to −0.96 V vs. Ag/AgCl for 1000, 3000 and 5000 cycles.

4. Conclusions

In this work, we successfully synthesized and evaluated carbon support for α-MnO₂ through hydrothermal carbonization followed by KOH activation and heat treatment. High temperature (250 °C) and longer duration (12 h) provided biomass-derived carbon support with better ORR performance due to the formation of pores and micrometric fragments, and the preservation of the lignin backbone structure, which enhances the contact of oxygen to active sites of MnO₂. Further, it was found that activation is a necessary step in the conversion of biomass to carbon support as it functions as the main factor for cellulosic structure degradation and pore formation. Finally, CCAC + MnO₂ 250-12 showed the greatest ORR catalytic performance among prepared samples since the conditions were favorable in the formation of porous structures and the preservation of oxygen containing functional group, which facilitated ORR performance. The CCAC + MnO₂ 250-12 also showed higher stability compared to VC + MnO₂ and 20% Pt/C. This work provides a new and cost-effective method of deriving carbon support from biomass for enhanced electrocatalytic activity towards ORR. In addition, this synthesized carbon-supported electrocatalyst is seen as a potential cathode for fuel cells and metal-air battery application.

Author Contributions: H.O.P., J.D.A.P. and E.A.E.J. conceived, designed, and performed the synthesis experiments. H.O.P. and J.D.A.P. performed the electroanalytical experiments; J.D.A.P. and E.A.E.J. assisted in project administration; J.A.D.d.R. performed validation of results; H.O.P. prepared the draft manuscript; E.A.E.J., J.A.D.d.R. and J.D.O. reviewed and edited the draft manuscript; J.D.O. provided the overall supervision of the project. All authors have read and agreed to the published version of the manuscript.

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