Understanding the Surface Characteristics of Biochar and Its Catalytic Activity for the Hydrodeoxygenation of Guaiacol

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Abstract: Biochar (BCR) was obtained from the pyrolysis of a palm-oil-empty fruit bunch at 773 K for 2 h and used as a catalyst for the hydrodeoxygenation (HDO) of guaiacol (GUA) as a bio-oil model compound. Brunauer–Emmet–Teller surface area analysis, NH3 and CO2-temperature-programmed desorption, scanning electron microscope–dispersive X-ray spectroscopy, CHN analysis and X-ray fluorescence spectroscopy suggested that macroporous and mesoporous structures were formed in BCR with a co-presence of hydrophilic and hydrophobic sites and acid–base behavior. A combination of infrared, Raman and inelastic neutron scattering (INS) was carried out to achieve a complete vibrational assignment of BCR. The CH–OH ratio in BCR is ~5, showing that the hydroxyl functional groups are a minority species. There was no evidence for any aromatic C–H stretch modes in the infrared, but they are clearly seen in the INS and are the majority species, with a ratio of sp3–CH:sp2–CH of 1:1.3. The hydrogen bound to sp2–C is largely present as isolated C–H bonds, rather than adjacent C–H bonds. The Raman spectrum shows the characteristic G band (ideal graphitic lattice) and three D bands (disordered graphitic lattice, amorphous carbon, and defective graphitic lattice) of sp2 carbons. Adsorbed water in BCR is present as disordered layers on the surface rather than trapped in voids in the material and could be removed easily by drying prior to catalysis. Catalytic testing demonstrated that BCR was able to catalyze the HDO of GUA, yielding phenol and cresols as the major products. Phenol was produced both from the direct demethylation of GUA, as well as through the demethylation pathway via the formation of catechol as the intermediate followed by deoxygenation.

Keywords: biochar; palm oil empty fruit bunch; hydrodeoxygenation; guaiacol

1. Introduction

The world has gradually shifted to renewable energy sources in an effort to reduce dependency on fossil fuels, as well as to address climate change and the global warming caused by excessive greenhouse gas emissions [1]. Indonesia, for example, has rolled out B30, a fuel blend of 30% biofuel from fatty acid methyl esters (FAMEs) and 70% conventional diesel, for nationwide commercial transportation in order to meet these targets by 2030 [2]. However, FAMEs are categorized as first-generation biofuels which rely heavily on food commodities and are, therefore, considered to be unsustainable, prompting...
massive deforestation as a result of industrial plantation expansion [3,4]. Second-generation biofuel, on the other hand, uses abundant and non-competitive biomass waste sources as their precursor. Among them, palm oil empty fruit bunches (POEFBs) are a potential source, since they are rich in lignocellulose feedstock [5].

The production of second-generation biofuels involves the pyrolysis of lignocellulose into bio-oil followed by upgrading via a hydrodeoxygenation (HDO) process to remove oxygen [3,6]. Bimetallic metal sulfides (e.g., NiMoS, CoMoS) have been one of the most widely studied catalysts for HDO [7–9], but the search for a suitable support material to increase the stability of the active metal centers, or act as the catalyst itself, remains a challenge. Activated carbons have been reported to offer higher stability in HDO systems compared to alumina and silica due to the weak acidity [10,11], but the extensive microporosity is a drawback which inhibits reactions involving large molecules [12]. Macroporous and mesoporous materials have proven favorable as the relatively larger pores can improve molecular diffusion for the large molecules [11,13,14].

Pyrolysis is considered to be a sustainable way to generate energy from bio-oil and syngas. However, biochar (BCR)—a solid carbon byproduct—is inevitably created in the production line [6,15,16]. BCR is a carbon-rich solid with easily tuned surface functionality and porosity [17–20]. The oxygen-containing functionalities of BCR, together with other functionalities upon modification, make the surface of this material both hydrophilic and hydrophobic, affecting the adsorption of the reactants for further activation [11,21,22]. It is known that the –OH, C–O, and C=O functional groups in BCR interact with phenol groups through H bonding [23,24]. This is beneficial for HDO, since the interaction of phenols via H bonding, such as that observed on silica and clay, is preferable compared to the chemisorption observed in alumina. This is because the formation of a strongly held carbonaceous species in the form of phenates was observed, leading to catalyst deactivation [9,25]. While BCR can be returned to farm fields as a fertilizer for carbon sequestration or other functional materials [26], it would be ideal to utilize the BCR as a catalyst for the HDO of bio-oil, creating a net-zero pyrolysis process.

Several authors have demonstrated the use of BCR as a catalyst for the esterification and transesterification of vegetable oils to produce FAME in high yields [27–29]. However, its utilization as a catalyst for bio-oil upgrading in second-generation biofuel production has not been fully explored [30–32], despite the fact that BCR is also produced during the biofuel production. The physicochemical properties of BCR have been demonstrated to vary significantly based on the type of raw biomass materials, the temperature and retention time used in pyrolysis and activation, or functionalization techniques employed [33–35]; hence, it is imperative to understand the surface characteristics of BCR, as tailoring the catalytic activity of BCR would be challenging without such knowledge.

Herein, we investigate the surface characteristics of BCR obtained from the pyrolysis of POEFB and apply it as a catalyst for the HDO of guaiacol (GUA) as a bio-oil model compound. The physicochemical properties of BCR were studied using Brunauer–Emmet–Teller (BET) surface area analysis, NH₃- and CO₂-temperature programmed desorption (TPD–NH₃, TPD–CO₂), scanning electron microscope–dispersive X-ray spectroscopy (SEM-EDS), CHN analysis, X-ray fluorescence spectroscopy (XRF) and powder X-ray diffraction (XRD). A complete vibrational assignment of BCR was carried out by a combination of infrared, Raman and inelastic neutron scattering (INS) techniques to understand in detail the functional groups on the BCR surface, which greatly influence the surface characteristics and its catalytic activity towards HDO.

2. Results and Discussion
2.1. Characterization of BCR
2.1.1. Morphology and Elemental Analyses

The morphology of BCR examined by SEM (Figure 1) was observed to have irregular texturing with convoluted smooth and porous surfaces. Parts of the structure had a highly complex network of pores, channels and fibrous ridged surfaces. The release of volatiles
during pyrolysis leads to the formation of bubbles, open and closed pores in BCR, similar to that observed for biochar obtained from palm kernel shells at pyrolysis temperatures higher than 773 K [36].

![Figure 1](image1.png)

**Figure 1.** SEM analysis results of BCR: (a) ×500 and (b) ×1000 magnification.

The elemental analyses of BCR measured by CHN analysis, SEM-EDS and XRF are summarized in Table 1. The H/C and O/C molar ratios are informative to evaluate the hydrophilicity and hydrophobicity of BCR and as an indication of the development of aromatic structures [34,36,37]. Chen [34] evaluated the pyrolysis of various types of biomasses and found that the H/C of the biomass precursor ranged from 1.4 to 1.8 and the O/C ratio was found to be approximately 0.55–0.75, indicating that the starting biomass has low aromaticity and a high aliphatic content. After pyrolysis, a significant decrease in the H/C and O/C atomic ratios was observed that decreased linearly as the pyrolysis temperature increased. In the case of BCR, the H/C and O/C ratio was recorded as 0.80 and 0.34, respectively, confirming that the surface of BCR was more hydrophobic than the reported biomass precursors. The loss of H and O during the pyrolysis process is a consequence of the dehydration and decarboxylation reactions of the weak linkages within the BCR structure, making the surface less hydrophilic with doped hydrophobic sites. The co-presence of both hydrophobic and hydrophilic sites has proven to be beneficial in heterogeneous catalysis involving substrates with varied polar functionalities [22,38]. The mineral content in BCR was dominated by Si, Ca, K and Fe, with minor amounts of other minerals detected.

**Table 1.** Elemental analyses of BCR measured by CHN analyzer, SEM-EDS and XRF.

| Element | CHN/wt % | SEM-EDS/wt % | XRF/wt % |
|---------|----------|--------------|----------|
| C       | 40.59    | 66.71        | -        |
| H       | 2.72     | -            | -        |
| N       | 3.53     | -            | -        |
| O       | 53.16*   | 30.45        | -        |
| Mg      | -        | -            | 4.98     |
| Al      | -        | -            | 3.76     |
| Si      | -        | 1.18         | 37.63    |
| P       | -        | -            | 4.62     |
| S       | -        | -            | 3.88     |
| Cl      | -        | -            | 1.27     |
| K       | -        | 0.52         | 14.16    |
| Ca      | -        | 0.56         | 18.45    |
| Ti      | -        | -            | 0.48     |
| Mn      | -        | -            | 0.56     |
| Fe      | -        | -            | 9.51     |
Table 1. Cont.

| Element          | CHN/wt % | SEM-EDS/wt % | XRF/wt % |
|------------------|----------|--------------|----------|
| H/C molar ratio  | 0.80     | -            | -        |
| O/C molar ratio  | -        | 0.34         | -        |

*Calculated by difference (100%—CHN content).

2.1.2. BET Surface Area, Pore Size, and Total Acidity–Basicity

The N\textsubscript{2} isotherm of BCR is shown in Figure 2. This shows that the N\textsubscript{2} adsorption–desorption isotherms for BCR are classified as type II, which is a typical isotherm of macroporous materials (containing pores > 50 nm width) [14].

![Figure 2. Adsorption–desorption isotherm curve of BCR.](image)

The hysteresis loop of BCR is seen as an H3 type, as found in layered aggregates with gaps in mesoporous or macroporous materials [39]. It is a typical feature of non-rigid aggregates of plate-like particles, analogous to a Ni-biochar catalyst from rice husks [22,40]. The specific surface area, pore volume and pore size of the catalyst are listed in Table 2. The surface area was recorded as 2.14 m\textsuperscript{2}g\textsuperscript{−1}. This low surface area of BCR is affected by the incomplete removal of volatile matter from biomass and has been reported for several untreated biochars from different feedstocks [23]. The mean pore size of BCR is 27.31 nm, indicating the formation of a mesoporous structure in the material.

Table 2. BET surface area, total acidity, and total basicity of BCR measured by TPD–NH\textsubscript{3} and TPD–CO\textsubscript{2}.

| Properties                  | Value  |
|-----------------------------|--------|
| BET surface area/m\textsuperscript{2} g\textsuperscript{−1} | 2.14   |
| Pore size/nm                | 27.31  |
| Pore volume/cm\textsuperscript{3} g\textsuperscript{−1} | 0.01   |
| Total acidity/mmol g\textsuperscript{−1} | 0.15   |
| Total basicity/mmol g\textsuperscript{−1} | 0.50   |

The total acidity and basicity of the catalysts were analyzed via TPD–NH\textsubscript{3} and TPD–CO\textsubscript{2}. The TPD profiles are presented in Figure 3 and the calculated acidity/basicity values are listed in Table 2. It has been reported that the TPD–NH\textsubscript{3} for rice husk char are generally divided into three peaks: the peaks in the 100–230 °C range represent weak acidity; the peaks in the range of 230–380 °C represent medium-strong acidity; and strong acidity is
observed for peaks in the 380–500 °C range [41]. Our TPD–NH₃ results show that the acidity peak of BCR was found in the range of 150–300 °C, demonstrating that BCR mostly contains weak acid sites with a total acidity value of 0.15 mmol g⁻¹. Meanwhile, the TPD–CO₂ shows a basicity peak in the range of 100–250 °C, with a total basicity value of 0.50 mmol g⁻¹. An increase in basic functional groups may be the result of charring and liming induced by the decreasing amount of acidic functional groups, as well as alkali salts separating from the organic compounds [42]. Hence, TPD measurements show that BCR has both weak acid and base sites present on its surface. Yakovlev [43] suggested that the acidic–basic properties play an important role during GUA HDO because of the possible additional activation of oxy-compounds on the support surface, as found for TiO₂ and ZrO₂ [12].

Figure 3. TPD profiles of BCR: (a) TPD–NH₃ and (b) TPD–CO₂.

2.1.3. Powder X-ray Diffraction Analysis

The diffraction pattern of BCR is shown in Figure 4. The most prominent peak is seen at 2θ = 26.7° which is attributed to the stacking of the graphitic basal planes of the char crystallites [36]. The relatively high intensity and symmetry are a result of graphitization. The small features at 2θ = 20.9 and 29.5° possibly show the presence of amorphous silica and fluorite [44], which was confirmed by XRF to be one of the major minerals contained in BCR. The broad band at 2θ = 20–30° indicates that the majority of the BCR remains amorphous [45].

2.2. Vibrational Studies

The infrared, Raman and INS spectra of BCR are shown in Figure 5 and selected regions are also presented in Figure 6. The infrared spectrum is very similar to that found for rice straw-derived biochar after pyrolysis at 773 K [46].

It is striking that the infrared, Raman and INS spectra are so different. This is a consequence of the different strengths of the techniques, emphasizing the need to have all three types of spectra available for a complete analysis. Thus, the infrared spectra are dominated by modes relating to the oxygen functionalities present, the Raman by the aromatic structure and the INS by modes involving displacement of hydrogen.

In the C–H and O–H stretch region (2800–3700 cm⁻¹), the infrared spectrum (Figure 5a) is dominated by the extremely intense and very broad (~800 cm⁻¹ FWHM) O–H stretch mode of hydroxyls. The intensity would suggest that there is a large number of these present; however, the MAPS INS spectrum (Figure 5d) shows that this is not the case. The intensity of an INS mode depends on the amplitude of motion, which in the harmonic approximation is inversely proportional to the transition energy, as described in the litera-
Crucially, this does not depend on the electronic structure, unlike the infrared and Raman spectra, where the intensity derives from the dipole moment derivative and the polarizability tensor, respectively. This means that, to a reasonable degree of accuracy, in the INS, the ratio of the integrated intensity of the C–H and O–H stretch modes is a direct measure of their relative population. Figure 6 (top) shows a curve fit to the spectrum in this region. From this we find that the CH:OH ratio is ~5, showing that the hydroxyls are a minority species. Their intensity in the infrared spectrum is misleading and occurs due to the large increase in the dipole moment derivative that is characteristic of hydrogen-bonded systems [48].

The C–H stretch modes are seen at 2924 and 2855 cm$^{-1}$ in the infrared spectrum, which is characteristic of aliphatic methylene groups [49]. The associated scissors mode is clearly seen in both the infrared and INS spectra at 1450 cm$^{-1}$. There is no evidence for any aromatic C–H stretch modes in the infrared spectrum, but they are clearly seen in the INS spectra and are the majority species. Figure 6 (top) shows that the ratio of sp$^3$–CH:sp$^2$–CH is 1:1.3.
Figure 6. Top: Curve fit to the MAPS INS spectrum in the C–H and O–H stretch region. Middle: Curve fit to the Raman spectrum in the range of the G and D bands. Bottom: comparison of the TOSCA INS spectrum of biochar (a) and that of glassy carbon (b) [50] in the region of the aromatic C–H-bending modes. The inset shows the definition of H1 (an isolated C–H bond) and H2 (two adjacent C–H bonds) termination.
In the 1000–1700 cm$^{-1}$ range, the Raman spectrum shows the characteristic D and G bands of sp$^2$ carbons. These are seen in a wide range of materials including soot [51], coal [52], industrial carbons [53] and carbon nanotubes [54]. Curve fitting the 1100–1600 cm$^{-1}$ shows the presence of four bands—see Figure 6 (middle). Following Sadezky et al. [51], these are labeled as G-band (1578 cm$^{-1}$, ideal graphitic lattice), D1 (1350 cm$^{-1}$, disordered graphitic lattice, e.g., edges, defects, surface layers), D3 (1472 cm$^{-1}$, amorphous carbon) and D4 (1180 cm$^{-1}$ defective graphitic lattice). The G-band is also seen in the infrared spectrum. Carbons typically show a 2D band at 2700 cm$^{-1}$ [54]. However, there is no evidence for this in our materials. The INS spectrum has a strong broad peak at 1370 cm$^{-1}$, close to the D1-band. However, its intensity shows that it must originate from a hydrogenous mode; we assign it to the methylene wag. The infrared band at 1378 cm$^{-1}$ could be either (or both) of the D1-band or the methylene wag.

The INS bands in the region 750–1200 cm$^{-1}$ are characteristic of the in-plane (~1200 cm$^{-1}$) and out-of-plane (700–900 cm$^{-1}$) C–H-bending modes of hydrogen that decorate the edges of the graphene sheets [30,55]. Figure 6 (bottom) shows a comparison of this region with that of a non-graphitizing glassy carbon [50]. While the peak at 955 cm$^{-1}$ is consistent with it being one of the H2 peaks (two adjacent C–H bonds, see inset of Figure 6 (bottom) for an illustration of H1 and H2 hydrogen), the corresponding peak at 795 cm$^{-1}$ is absent. This suggests that the hydrogen is largely present as H1 (isolated C–H bonds). The presence of the D bands in the Raman spectrum shows that the material is highly disordered. Hence, we assign the other modes in this region at 955, 781 and 752 cm$^{-1}$ to out-of-plane C–H-bending modes of isolated hydrogen perturbed by the presence of defects or oxygen functionalities. The very strong infrared band at 1099 cm$^{-1}$ is assigned to the C–O–C-stretching modes of the β–O–4 linkages of lignin. There are a number of weak bands in the 200–700 cm$^{-1}$ region in all three spectra. These are assigned to torsional modes of the graphene planes of the biochar.

The features at 1700–2500 cm$^{-1}$ in the Raman spectrum are very difficult to assign, as we can find no precedent for them. The elemental analysis shows the presence of nitrogen. Therefore, the possibility of the presence of nitriles –C≡N must be considered. These show strong bands in both the infrared and Raman spectra in the range 2200–2500 cm$^{-1}$, depending on whether the nitrile is attached to sp$^2$–C, sp$^3$–C or is conjugated [49]. While a nitrile would account for the 2230 cm$^{-1}$ band, it would not explain the other bands in this region. Additionally, the absence of a corresponding band in the infrared spectrum militates against this assignment. We believe that the only plausible assignments are to resonance-enhanced overtones and combinations, because there are no functionalities present that could produce fundamental modes in this region. The observed modes and their vibrational assignments are summarized in Table 3.

Table 3. Transition energies (cm$^{-1}$) of BCR and vibrational assignments.

| FTIR | Raman | INS | Assignment $^1$ |
|------|-------|-----|----------------|
| 3300 | -     | 3300| ν OH           |
| -    | -     | 3060| ν sp$^2$-CH    |
| 2924 | -     | 2940| ν sp$^3$-CH$_2$|
| 2855 | -     |     |                |
| -    | 2475 | -   | G + 915        |
| -    | 2230 | -   | D1 + 915       |
| -    | 2070 | -   | D1 + 790       |
| -    | 1835 | -   | 2 × 915        |
| 1585 | 1578 | -   | aromatic ring stretch (G-band) |
BCR readily absorbs water, which is removable by heating in a vacuum at ~393 K. The amount of water is significant. Drying resulted in an 11% weight loss. Figure 7 shows the infrared and INS spectra of the adsorbed water, generated from the difference spectra: as received–dried. In the INS spectrum shown in Figure 7a, it can be seen that the adsorbed water does not resemble that of ice, as shown in Figure 7b. The slope to the leading edge of the librational modes at 4000–1000 cm\(^{-1}\) is typical of disordered forms of water, as seen on other types of carbon [50] and on metal oxides, e.g., those shown in [56]. Similarly, the infrared spectrum resembles that of liquid water much more closely than that of ice. This suggests that the water is present as disordered layers on the surface of the BCR rather than trapped in voids in the material.

### 2.3. Catalytic HDO of GUA

The catalytic HDO reaction of GUA was tested using BCR as the catalyst under 20 bar H\(_2\) pressure for 6 h without the use of a solvent. A control reaction without the presence of the catalyst was carried out and the results were compared to our previous study using pillared clay as the catalyst (PILC) [9]. Five products were identified in the liquid product mixture: phenol (1), o-cresol (2), p-cresol (3), catechol (4) and veratrole (5). The unidentified product mixtures were labeled as others; the results are summarized in Table 4. The products (1)–(3) are HDO products with partial oxygen removal, whereas (4) and (5) are

| FTIR | Raman | INS | Assignment \(^1\) |
|------|-------|-----|-----------------|
| 1450 | -     | 1450| CH\(_2\) scissors |
| -    | 1472 | -   | aromatic ring stretch (D3-band) |
| 1378 | -     | 1370| CH\(_2\) wag |
| -    | 1350 | 1300| aromatic ring stretch (D1-band) |
| -    | 1180 | -   | aromatic ring stretch (D4-band) |
| -    | -     | 1160| \(\delta\) sp\(^2\)-CH |
| 1099 | -     | -   | \(\gamma\) C–O–C |
| 1028 | -     | 1032| \(\gamma\) CH |
| -    | 920   | -   | - |
| 874  | -     | 882 | \(\gamma\) CH (H1) |
| -    | 790   | -   | - |
| 781  | -     | -   | \(\gamma\) CH |
| -    | 712   | -   | \(\gamma\) CH |
| -    | 604   | 605 | - |
| -    | -     | 525 | - |
| 465  | 470   | 457 | \(\tau\) CC |
| 423  | -     | 431 | - |
| -    | -     | 232 | - |
| -    | -     | 148 | - |
| -    | -     | 99  | - |

\(^1\) \(\nu\) = stretch, \(\delta\) = in-plane bend, \(\gamma\) = out-of-plane bend, \(\tau\) = torsion.
the result of a bimolecular transalkylation reaction and generally occur in the presence of Bronsted acid sites [57].

![Figure 7. INS difference spectrum of (a) adsorbed water, (b) INS spectrum of ice Ih, (c) infrared difference spectrum of adsorbed water, (d) infrared spectrum of ice Ih and (e) infrared spectrum of liquid water.](image)

**Table 4.** HDO of GUA using BCR as the catalyst.

| Catalyst | Temperature /K | Pressure/bar | Time/h | Conversion/% | Product Distribution/% |
|----------|----------------|--------------|--------|--------------|-----------------------|
|          |                |              |        | (1) (2) (3) (4) (5) Others |
| BCR      | 523            | 20           | 6      | 100          | 60 10 19 0 0 11       |
| PILC [9] | 623            | 20           | 6      | 78           | 42 10 5 6 5 32       |
| No catalyst | 523        | 20           | 6      | 22           | 0 0 0 0 60 40       |

The reaction testing data show that the BCR was able to convert all the GUA (100%) yielding 60% (1), 10% (2) and 19% (3) as the major HDO products with no detection of (4) and (5). The selectivity towards the desired products (1)–(3), in particular for (1), was higher when BCR was used as the catalyst, as compared to PILC. Notably, no benzene ring hydrogenation products were formed in either case. The higher selectivity of BCR most likely arises from the presence of both weak acid and base sites, similar to ZrO₂ [12]. In addition, the contribution of both hydrophobic and hydrophilic sites in BCR allows the adsorption stability of GUA on the surface to be increased. As reported by Wang [22], the multiple absorption modes in a Ni–biochar catalyst were able to facilitate the adsorption/activation of the C–O functionality in vanillin and the aliphatic C–C in eugenol, enhancing its hydrogenation activity.

It is noteworthy that, although no metals were immobilized on BCR, we suggest that the iron contained in BCR (9.51 wt%) played a role in the dissociation of H₂ during HDO, as our reaction conditions are not far from those generally used in an iron catalyzed Fischer-Tropsch process [58]. The activation of guaiacol on Fe-based catalysts has been reported to favour C–O bond breaking at the expense of C–C bond breaking [59,60]. The Fe catalysts show a low level of activity in the hydrogenation of aromatic rings, resulting in benzene,
toluene, xylenes (BTX) and phenol as the HDO products. The chemical mechanisms are catalyzed in the presence of the supported Fe particles, which supply the active H-species coming from the dissociation of H$_2$ on the metal phase. A different selectivity and mechanism are observed over supported Ni-based catalysts, which provides a higher level of selectivity for the hydrogenation of the benzene ring [61].

The presence of potassium in BCR (14.16 wt%) correlates with the reduction in the total acidity and changes the acid–base balance of BCR, which positively influenced the selectivity of the catalyst, as reported in the case of potassium-modified NiMo and CoMo–alumina catalysts [7]. Without the presence of a catalyst, a GUA conversion rate of only 20% was obtained, yielding 60% of (5) with no formation of the HDO products (1)–(3). At the current stage, we speculate that (5) was formed via a free radical mechanism rather than transalkylation in the absence of a catalyst, similar to that reported in the pyrolysis of guaiacol [62].

In order to evaluate the reaction profile, we examined the catalytic activity of BCR under different reaction times. As shown in Figure 8, product (1) was rapidly, and largely, formed by 0.5 h, resulting in a 50% yield. With increasing reaction time, (1) continuously increased to a 60% yield at 3 h, but no further increase in the yield was seen with a prolonged time of 6 h. Products (2) and (3) were also formed faster in the first 0.5 h, with no significant changes in their yields at 3 and 6 h. Interestingly, a 17% yield of (4) was observed at 0.5 h, which decreased with time. Only 3% of (4) was seen at 3 h and none was detected at 6 h. This indicates that (4) is an intermediate in the production of (1). It has been reported that the hydrogenolysis of the methyl–oxygen bond (demethylation) of the methoxy group of GUA to form catechol is the first stage of the reaction, followed by a second stage leading to the elimination of one of the hydroxyl groups (deoxygenation) to produce phenol [12,63,64]. Over the BCR catalyst, phenol was also produced via the direct demethoxylation of GUA. Fan [41] has reported that the impregnation of Ni on rice husk char was able to provide a 100% GUA conversion rate and had an 11.4% selectivity for hydrocarbons. Work from Santos [65] shows that complete HDO conversion of vanillin was achieved over Pd-biochar catalysts resulting in 92% selectivity of p-cresol. An MoWC bimetallic carbide catalyst has been reported to exhibit higher catalytic activity and oxygen-free aromatic hydrocarbon selectivity compared to the monometallic MoC in the HDO of GUA [66]. Our previous work has also shown that immobilization of bimetallic NiMoS on PILC further increases the yield and selectivity of HDO products [9]. In order to increase the catalytic activity and selectivity toward benzene ring hydrogenation products, we will investigate BCR as a catalyst support for bimetallic and metal sulfide catalysts in future HDO studies.

Figure 8. HDO of GUA over BCR catalyst at various reaction times.
3. Materials and Methods

3.1. Materials

BCR was produced through the pyrolysis of POEFB at \( T = 773 \) K for 2 h. The heating process started at room temperature and was raised to 773 K at a heating rate of \( 6 ^\circ C/\text{min} \) (75 min) followed by a holding period for 45 min at 773 K. GUA and reference products were purchased from Sigma-Aldrich (Gillingham, UK) and used without further purification.

3.2. Characterization of BCR

The morphology and elemental analyses were measured using SEM-EDS JSM-IT200 Jeol operated at 20 kV and by a Bruker XRF. The CHN analysis was carried out on a LECO CHN628 Series Elemental Determinator (LECO, St. Joseph, MI, USA). Surface areas and pore size were analyzed based on the BET method using Micromeritics TriStar II 3020 Version 2.00 (Micromeritics, Norcross, GA, USA). The samples were degassed at 573 K prior to the measurement. TPD–NH\(_3\) and TPD–CO\(_2\) were measured on a Micromeritics Chemisorb 2750 (Micromeritics, Norcross, GA, USA) and XRD was carried out on a Panalytical AERIS (Malvern Panalytical, Almelo, Netherlands) using CuK\(\alpha\) \((\lambda = 0.15406)\) at 40 kV and 15 mA. The infrared spectra were measured on a Bruker Diamond ATR accessory (Bruker UK Limited, Coventry, UK) using the Bruker Vertex 70 (Bruker UK Limited, Coventry, UK). Raman spectroscopy was performed using a Horiba Scientific LabSpec 6 spectrometer (Horiba, Villeneuve-d’Ascq, France), with 532 nm excitation. The inelastic neutron scattering (INS) of BCR was carried out at the ISIS Neutron and Muon Source, Oxfordshire, United Kingdom \([67]\) using the spectrometers TOSCA \([68]\) and MAPS \([69,70]\). As explained in more detail in the literature \([69]\), the two spectrometers are complementary. TOSCA provides good resolution in the 0–2000 cm\(^{-1}\) region, while MAPS enables access to the C–H and O–H stretch regions (2800–4000 cm\(^{-1}\)). Prior to the analyses, the sample was dried at 383 K in vacuo overnight and the INS spectra were recorded at 20 K for ~6 h.

3.3. Catalytic Testing

The conversion of GUA was performed following the HDO procedure previously reported in our work \([9]\). In brief, the BCR catalyst (0.5 g, 5 wt %) and GUA (10 g) were placed in a high-pressure autoclave and reacted at 523 K and 20 bar H\(_2\) for varying reaction times (0.5–6 h). The BCR was dried overnight at 383 K prior to catalyst testing. After the reaction, the liquid product was extracted using dichloromethane and analyzed on an Agilent GC/MS 7890B system (Agilent, Santa Clara, CA, USA) with an HP-5MS U1 column. Products with lower boiling points, such as cyclohexane and cyclohexanone, were measured using an Agilent 7890A GC/FID system (Agilent, Santa Clara, CA, USA) equipped with a DB-WAX column. However, they were only found in trace amounts and are, therefore, not included in the mass balance calculation.

4. Conclusions

BCR from POEFB was developed as a new HDO catalyst. An evaluation of the physicochemical properties shows that pyrolysis at 773 K for 2 h generated amorphous BCR with macroporous and mesoporous structures that have irregular texturing and convoluted smooth and porous surfaces. The co-presence of hydrophilic and hydrophobic sites was observed based on the H/C and O/C molar ratios of BCR as compared to reported biomass precursors. BCR shows both acid and base behavior. Si, Ca, K and Fe were found to be the major minerals present. The co-existence of aliphatic and aromatic C–H stretch modes suggests that the pyrolysis of POEFB produced incomplete carbonization, leading to a model of BCR having a slightly higher population of aromatic components than aliphatic compounds. The INS clearly shows that hydroxyls are only present as a minority species, despite the strength of the bands in the infrared spectrum. Raman spectroscopy shows that there are graphitic domains present but that there is also considerable disorder as shown by the presence of the D3 and D4 bands. Adsorbed water in BCR is present as disordered layers on the surface rather than trapped in voids in the material. The catalytic
HDO of GUA over the BCR yielded phenol and cresols as the major products. Phenol was produced either from a two-step demethoxylation–deoxygenation pathway via the intermediate compound catechol, or by a one-step demethoxylation of GUA directly to phenol. No benzene ring hydrogenation products were formed in the reaction conditions of this experiment. Although no metals were immobilized on BCR, we speculate that the iron contained in BCR played a role in H$_2$ dissociation during HDO, whereas potassium potentially correlates with the reduction in the total acidity and changes the acid-base balance of BCR, which positively influenced the catalyst selectivity. These results provide reference information for the use of BCR as a catalyst for HDO, demonstrating how further optimization of the catalytic activity may be achieved by using BCR as a catalyst support for the widely studied bimetallic and metal sulfide HDO catalysts.

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