

## Article

# The Effect of Temperature and Treatment Regime on the Physical, Chemical, and Biological Properties of Poultry Litter Biochar <sup>†</sup>

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<sup>†</sup> In the memory of our mothers.

**Abstract:** Poultry litter was converted to biochar by torrefaction and to hydrochar by hydrothermal carbonisation. Many parameters were measured for the resulting chars, to investigate the effects of the production method and production temperature. SEM showed the presence of large quantities of crystalline material on the surface of the biochars. The elemental composition of some crystals was determined as 35% K and 31% Cl. This was confirmed as sylvite (KCl) crystals, which explains the high levels of water-extractable potassium in the biochar and may also be important in germination inhibition. Biochars almost totally inhibited germination, whilst hydrochars decreased germination. Although germination occurred on hydrochar, root growth was severely inhibited. Consequently, the germination index may be better to determine total phytotoxicity as it measures both effects and could be used as a bioassay for chars used as soil amendments. Washing removed germination inhibition in a low-temperature char (350 °C), possibly by removing KCl; however, root toxicity remained. There were very low levels of heavy metals, suggesting they are not the source of toxicity. In biochars, pore mean size decreased with temperature from 350 °C to 600 °C, due to changes in pore size distribution. The mean pore size was measured directly using SEM. The merits of this method are discussed. Low-temperature biochars seem best suited for fuel as they have a high calorific value, high hydrophobicity, a low ash content and a high yield. Higher temperatures are better for soil amendment and sequestration applications with a smaller mean pore size, higher surface area, and higher pH.

**Keywords:** poultry litter; torrefaction; hydrothermal carbonisation; biochar; hydrochar; germination; sylvite; morphology



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## 1. Introduction

In the 21st century, there are several major environmental challenges that need to be solved; these include the reduction of greenhouse gas (GHG) emissions, maintenance of soil fertility and disposal of organic wastes. The discovery of the *Terra Preta* dark earth soils in Amazonia caused a surge of research interest in the applications of a special form of charcoal/biochar. The Amazonian soils appear to have retained their fertility for thousands of years because of the presence of biochar [1]. Therefore, converting organic wastes into biochar instead of combusting or landfilling them and then applying the biochar to soil has the potential to solve all three challenges simultaneously. Biochar is thus one of the most important emergent technologies for environmental management. Initial studies focused on biochar obtained from plant material [2], but there are now a number of studies on biochar from animal manure feedstocks, poultry litter (PL), and sewage sludge [3–6]. What is more, recent research studies have proved that the biochar produced from poultry litter releases more water-soluble dissolved P and K concentrations and increases soil plant-available P and K concentrations as compared to lignocellulosic-based biochars [7].

Biochar has complex physio-chemical and biological properties; it is also extremely heterogeneous due both to differences in the feedstock and variations in manufacturing. Heating organic matter to low temperatures causes a decrease in mass—due to loss of water and volatile organics. The loss of hydrogen and oxygen increases the proportion of carbon. At 250–350 °C, cellulose is depolymerised, and organic material is converted into a carbon honeycomb structure, where macropores are formed with their size and distribution determined by the cellular structure of the original biomass. There is further mass loss due to volatilisation and carbon is converted into aromatic structures. These aromatic structures are not easily broken down, giving biochar resistance to decay [8].

Above 600 °C, under an inert environment (Ar), the main process occurring is carbonisation, where most non-carbon atoms are lost. Very high temperatures cause a loss of structural complexity due to the melting of cell structure—which is why lower temperatures produce a more effective biochar [2]. As well as this change in structure, the surface of the biochar changes chemically in a very varied manner. During thermal degradation, potassium and chloride ions are highly mobile and can produce crystals on the surface of the biochar, as can other inorganics, producing a range of minerals. Silicon and calcium are present in cell walls and are released at higher temperatures, whilst nitrogen can be released at low temperatures. Potassium and phosphorus can start to vaporise above 760 °C and, above 800 °C, other elements follow [9]. As these are plant nutrients, biochar preparation temperatures must be kept below these values to retain them. In general, the biochar surface can be hydrophilic or hydrophobic, acidic or basic, and may be coated with bio-oil containing aromatic compounds. Biochar's basicity is due to the presence of alkali metals (K, Ca, Na, and Mg) present as carbonates and hydroxides [10].

The removal of water and volatiles creates a better fuel by increasing energy density—making fuel transport and storage easier. For example, woody biomass torrefied material contains 90% of the energy in 70% of the mass. The 30% mass loss is caused by the production of gas that only contains 10% of the energy but can still be used as a fuel. Torrefaction gives a less heterogeneous product than the original biomass [11]. Ash content varies depending on the biomass used—this is important commercially as ash produced needs disposal. Woody feedstock only produces very low amounts of ash (<1%), grasses, due to their high silica content, give higher values (up to 24%), whilst manures can produce up to 45% [2]. Apart from ash production, non-lignocellulose feedstocks, such as manure and sewage sludge, behave in much the same way as woody feedstock [5]. One problem with the manufacture of biochar is the possible production of toxic polyaromatic hydrocarbons (PAHs), which could be released in the gases from the furnace or remain in the biochar and be passed into the soil.

The disposal of poultry litter is a growing problem in industrialised countries, as is the disposal of other organic wastes, including human sewage. Broiler chicken numbers are increasing rapidly in the UK, so the amount of litter to be disposed of will also increase. Presently, for example, Northern Ireland produces 260 ktpa of poultry litter waste that needs disposal. This may rise to 400 ktpa within 5–10 years, assuming a 50% increase in capacity. Previously, poultry litter, which consists of a mixture of poultry manure and bedding material from poultry farms, was used on soil as manure. However, land application of manure without any treatment threatens the environment due to high concentrations of organics, ammonia, and other toxic pollutants [12].

Animal manure biochars are different in many ways to plant waste biochars in having a less regular and more heterogeneous nature. They also contain large quantities of plant nutrients, N, P, K, and may contain heavy metals. N, P, and K can be seen as a problem as they can be leached out and cause eutrophication, but they should be viewed as a source of nutrients to replace fertilisers instead. If these nutrients are to be used as a soil amendment, then it is important to know the chemical form of the element in the biochar. For example, K can be present as potassium chloride (KCl), or potassium carbonate ( $K_2CO_3$ ), or some other complex form. The form of K present is important as some minerals are far more water soluble than others. Sylvite (KCl in natural mineral form) is very soluble in water

but struvite ( $\text{KMgPO}_4 \cdot 6\text{H}_2\text{O}$ ), which may be present, has a far lower solubility so will be less available in soil [13]. The same is true of heavy metals; they may be present but not bioavailable and, thus, not harmful to living organisms. It has been found that the speciation, bioavailability, and environmental risk of heavy metals in chicken manure biochar produced by pyrolysis decrease by increasing the temperature. The residual portion of heavy metals is regarded as non-bioavailable and nontoxic because the residual solids, which mainly contain primary and secondary solids, contain metals in their crystal structures [14].

The most used technology to produce biochar from poultry litter is slow pyrolysis, as it maximises the amount of char and minimises the amount of bio-oil produced. Slow pyrolysis typically takes place at temperatures between 350 and 800 °C, with residence times from minutes to hours and a heating rate from 0.5 to 60 °C min<sup>−1</sup>. The characteristics of biochars will depend on the type of feedstock [15], particle size of the biochar [16], temperature [4,17], heating rate, and residence time [18]. Higher pyrolysis temperature will lead to more alkaline biochars [19] and to products with a higher surface area and a lower H/C ratio, indicating the formation of more aromatic structures [20]. One of the most important characteristics is the biochar's content of N, P and K, referred to as the primary macronutrients, because without them plants could not survive.

Despite the recent interest in biochars produced from animal origins as soil amendments for improving soil quality and increasing soil carbon sequestration, there is inadequate knowledge about the amounts, speciation, and availability to plants of the nutrients. There is also limited knowledge about how the different pyrolysis regimes and operating conditions affect the soil amendment properties of these materials produced from different feedstocks [21].

When waste is converted to biochar, this can then be utilised for three main applications: as fuel, as a soil amendment, or as an adsorbent. The use of chars as soil amendments is still controversial. The importance of biochar to reduce GHG and dispose of organic waste hinges on its complementary role in soil improvement. Without this, burying biochar is similar to landfilling; consequently, soil improvement is the application considered here. Many experiments show that biochar increases soil fertility and, therefore, crop yields [2], but the actual mechanism is still unknown.

Pot trials of biochar showed large increases in lettuce yield, particularly in unfertilised soil [22]. The addition of the biochar increased soil pH, Cation Exchange Capacity (CEC), and nutrient supply. Their experiment was on acidic soils, and the basic nature of biochar increased the pH, thus improving fertility. Although these seem to be positive results, the application rates were very high, and yields dropped after three years, contradicting the evidence from Amazonian soils, where fertility was retained for thousands of years. The study of lettuce yield did at least show positive effects. Another recent study applied PL biochar produced at 550 °C to field-grown corn, along with fertiliser. PL char alone gave the highest yield, but when urea was added with PL char, the yield was reduced [23].

In germination trials, most biochars show no inhibition. Germination testing is an example of a bioassay used to test phytotoxicity. These tests have been used for some years, initially to test new agricultural chemicals and then to assess ecological risks in contaminated soils [24]. As such, they should now be used to assess chars that are added to soils. A bioassay is a test that determines levels of toxic chemicals by their direct effects on living organisms, in this case by the effect on germination. In germination, seeds take up water by imbibition, which bursts the testa (seed coat). This allows more water uptake by osmosis. Enzymes are activated and convert insoluble stored food to soluble components, which move to the growing points of the radicle (root) and plumule (shoot). Respiration of stored food occurs to release energy. Most seeds do not need light for germination, only water, oxygen, and a suitable temperature. Germination inhibition is usually caused by factors that interfere with water uptake or direct toxic effects on enzymes at a later stage when enzymes become active.

This paper aims at making a significant contribution to the knowledge of the physical, chemical, and biological properties of poultry litter biochar for agricultural use, including morphology, structure, mineralogy, chemical composition, hydrophobicity, and germination, and at identifying the way the method and conditions of manufacture affect these properties. The emphasis is on the characterisation of the different char samples' morphology, mainly of the macropore structure and size, surface composition, and speciation. To the best of our knowledge, this is the first time such a detailed morphology characterisation has been performed, not only on PL char but on any other biochar.

## 2. Procedure

### 2.1. Poultry Litter Samples Collection

Poultry litter was supplied from a broiler farm in Northern Ireland. It was split into 20 samples and placed in a freezer. An equal amount was taken from each of ten subsamples and then mixed together to produce the working sample. This was homogenised in a food processor and then oven-dried at 105 °C for at least 12 h. The moisture content was about 30%.

### 2.2. Poultry Litter Samples Preparation

#### 2.2.1. Biochar Samples Obtained by Torrefaction

Biochar samples were generated by Carbogen Ltd., through the project funded by the Technology Strategy Board via the Small Business Research Initiative and Invest Northern Ireland. Biochar was produced by torrefaction (mild pyrolysis), in bulk (about 1 kg), under nitrogen flow, at specific temperatures. The residence time for all samples was 30 min. The samples were labelled TOR followed by the torrefaction temperature, e.g., TOR 350. Subsequently, some of these samples were "stripped", that is, boiled with 0.3 M nitric acid for 30 min. They were labelled TOR 350 stripped. Ashes were also produced from both stripped and non-stripped char samples using the ashing furnace as described below.

#### 2.2.2. Biochar Samples Obtained by Using a Tube Furnace

These samples were produced at Teesside University with a Carbolite Tube Furnace (model CTF 12/75/700) following standard operating practice. All samples used oven-dried PL (between 0.4 and 0.5 g), which was heated in a crucible at a rate of 30 °C min<sup>-1</sup> to the specified temperature under nitrogen flow at 500 mL min<sup>-1</sup>. Residence time was 30 min, but for high temperatures it often took over 12 h to cool down; thus, residence time at high temperatures was effectively longer than the stated time. These samples were labelled TF followed by the temperature used, e.g., TF 200.

#### 2.2.3. Biochar Samples Obtained by Hydrothermal Carbonisation

Hydrothermal carbonisation biochar samples were produced at Teesside University in a Buchi AG miniclave with a Julabo 4 20–250 thermal heat system. In all cases, 40 mL 0.1 M citric acid was used with 2 g of poultry litter. The set point temperature was noted; however, a sensor gave the actual internal temperature. Pressure was up to 20 bar, unless otherwise specified. Residence time was usually 120 min. The samples were labelled HTC followed by temperature used, e.g., HTC 80. Samples were also produced in larger quantities by Carbogen in a pressure cooker at a temperature of about 120 °C and a pressure of 2 bar. These samples were washed and dried. Subsequently, they were used in a second HTC run at Teesside. These samples were labelled HTC-C followed by the temperature used; C1 had a residence time of 5 min, C2 of 30 min, e.g., HTC-C1-210.

### 2.3. Characterisation of Biochar Samples

#### 2.3.1. Total Digestion Method

Samples were prepared for total K determination by the total digestion method. Samples for water-extractable K were prepared by water extraction, as for pH. Due to time



constraints, only one digestion was performed for most of the char samples. Just for the TOR 350, 500, 550, and 650 samples, both methods were used.

A weighed sample of 0.1 g was mixed with 20 mL concentrated nitric acid (70% analar) in a conical flask with anti-bumping granules. This was boiled on a hot plate under air reflux for 3 h. Narrow-mouth reflux tubes were used. Samples were cooled and diluted to 100 mL with distilled water in a volumetric flask. The flask and reflux tube were washed three times into a volumetric flask, leaving anti-bumping granules in the flask. This gives 1/100 dilution, which can be diluted further for AAS and ICP-MS.

### 2.3.2. pH

One gram of sample was weighed and added to a 250 mL HDPE (high-density polyethylene) bottle. A total of 20 mL of distilled water was added and the bottle was shaken at 100 rpm for 30 min on a Stuart Orbital Shaker. The solutions were filtered using Whatmans no.1 paper, and the pH was read using a calibrated Hanna pH meter.

### 2.3.3. X-ray Diffraction (XRD)

The structure of the biochar samples was assessed by XRD measurements taken with a Siemens diffractometer (D500) at a voltage of 40 kV, using Cu K $\alpha$  radiation. Both low- and wide-angle runs were performed. Low-angle used a 0.018° slit, wide-angle used 0.15°. Wide-angle originally used 2 $\theta$  of 5–90° but a later run used 5–70° with a step size of 0.02°. Identification of compounds was performed by comparing peak values to the RRUFF database.

### 2.3.4. Scanning Electron Microscopy (SEM)

A Hitachi 3400N scanning electron microscope having the capability of energy dispersive X-ray spectroscopy (EDX) was used to assess the morphology and elemental composition of the biochar samples.

The pulverised samples were placed on 14 mm aluminium stubs with 12 mm double-sided carbon tabs. Samples were blown with compressed air to ensure there were no loose particles. The samples were viewed at  $\times 50$  magnification and an area that was well covered with sample was chosen. Then a transect was performed to give a minimum of three sites for the elemental analysis, from which a mean value was calculated. The whole area of the view was used for this analysis. The samples were then viewed at a higher magnification ( $\times 250$  or  $\times 500$ ) to find pore structures. These pores were measured in their largest dimension using the SEM software—at least ten pores per site if available, with a minimum of three sites per sample for analysis, from which a mean value was calculated. The distribution of pore size was also plotted as a histogram in bands of 10  $\mu\text{m}$  as it would be expected that there would be different groups of pore sizes. Any crystalline or other unusual structures were viewed at higher magnification; up to  $\times 2000$ , depending on size. Elemental analysis was performed on the structure using EDX with the point-and-ID facility.

### 2.3.5. Flame Photometry (FP)

A Sherwood Flame Photometer 410 was used to determine the potassium levels. After flame ignition, deionised water was aspirated for 10 min. Then, zero was set using deionised water and a 10 ppm standard was set at 100 (to give greater accuracy). Blanks and standard were rechecked until steady and then a set of standards was read at 2.5, 5, 7.5, and 10 ppm to plot a calibration curve. These standards were made up by diluting a bought 1000 ppm standard with distilled water (for comparison, the standards were diluted with deionised water in the same way as the samples, although the standard is in nitric acid). Then, a sample was aspirated, and the reading noted. Samples were prepared by the total digestion method and diluted 1:10. Other samples were prepared by water extraction as for pH. In some cases, the reading was too high for the machine to read; in these cases, the sample was diluted.

### 2.3.6. Atomic Absorption Spectroscopy (AAS)

AAS was performed on a Thermo-scientific iCE 3000 AA with an air acetylene flame, with acetylene at a flow rate of  $0.9 \text{ dm}^3 \text{ min}^{-1}$ . A distilled water blank with a small amount of nitric acid and three standards with varied concentrations for the different metals were used. As the lamp must be changed for each element, it is more time consuming than ICP-MS.

### 2.3.7. Inductively Coupled Plasma–Mass Spectrometry (ICP-MS)

Samples from total digestion were diluted 1:100 by serial dilution. They were stored in HDPE containers in the fridge (important as metals can stick to glass). Samples were measured using an Agilent 7500 series ICP-MS machine with an octapole detector. A blank of nitric acid was used to calibrate the machine, then standards of 10 and 100 ppb were run for 28 elements (details in results). The samples were then aspirated and read. A two-minute wash of nitric acid was performed between each sample. To check the accuracy of the machine, three replicates of each standard were read to give an idea of machine error. A total digestion blank of a standard was also performed to quantify method errors. A comparison was also performed between AAS and ICP-MS by calculating the  $\text{mg g}^{-1}$  of metals. This was also used to compare the results with other studies.

### 2.3.8. Germination

The method was based on the International Biochar Initiative technical paper [25]. However, that method suggested mixing with soil, which would add more variables. So, germination was performed on biochar only. A few grams of biochar were used to cover the bottom of a petri dish. Fifty seeds of cress (*Lepidium sativum*) were placed on the char [26]. The seeds were not surface-sterilised as germination occurs within about 48 h, so sterilisation was deemed unnecessary. An amount of 2 mL of water was added over the seeds. The dish was closed, and germinated seeds were counted roughly each day to gain an idea of germination rate. It is not possible to do a perfect count without untangling the seeds, which could damage them, so final germination was counted accurately after 7 days. Germination was considered to have occurred when the cotyledon became visible.

Observation showed that root growth was affected in some germination, so a sample of 25 roots from each germination was measured after 7 days. This allowed calculation of a germination index (GI) [27].

$$\text{Relative germination} = \frac{\text{Number of germinated seeds in on substrate}}{\text{Number of germinated seeds in control}} \times 100$$

$$\text{Relative root growth} = \frac{\text{Mean root length in substrate}}{\text{Mean root length in control}} \times 100$$

$$\text{Germination index} = \frac{(\text{Relative seed germination \%}) \times (\text{relative root growth \%})}{100}$$

### 2.3.9. Thermogravimetric Analysis (TGA)

This was performed on pulverised samples using Perkin Elmer STA-1500 (Simultaneous Thermal Analyser) with Infinity Pro thermal analysis software. A small quantity of sample (between 12–28 mg depending on density) was placed in the crucible. The furnace was run at 25–800 °C at a heating rate of  $10 \text{ °C min}^{-1}$  and a residence time of 60 min. The heating was performed under argon at  $50 \text{ mL min}^{-1}$ .

### 2.3.10. Statistics and Calculations

All calculations and graph plots were performed using Excel 2010. The add-on data analysis package was used to perform univariate analysis of variance (ANOVA).

### 3. Results and Discussion

#### 3.1. Prepared Samples and Characterisation Methods

All the prepared samples and their respective characterisation methods are shown in Table 1.

**Table 1.** Prepared samples and the characterisation methods used for each of them. Number of ticks indicates number of replicates.

Sample	pH	ICP-MS	AAS	TGA	Pore Size	EDX	XRD	LOI * and Ash	FP	Yield	Germination
PL	✓✓	✓									✓
TOR 350	✓✓	✓	✓	✓	✓✓✓	✓✓✓	✓	✓✓	✓		✓
TOR 400	✓✓✓	✓			✓✓✓	✓✓✓					
TOR 450	✓✓	✓		✓	✓✓✓	✓✓✓	✓	✓✓✓✓			
TOR 500		✓			✓✓✓	✓✓✓			✓		
TOR 550	✓✓	✓			✓✓✓	✓✓✓		✓✓	✓		
TOR 600	✓✓	✓	✓	✓	✓✓✓	✓✓✓	✓	✓✓	✓		✓
TF 135					✓✓✓	✓✓✓				✓	
TF 200	✓✓	✓			✓✓✓	✓✓✓		✓	✓	✓	
TF 275	✓✓	✓		✓	✓✓✓	✓✓✓		✓	✓	✓	
TF 350	✓	✓		✓	✓✓✓	✓✓✓			✓	✓	
TF 400										✓	
TF 450	✓									✓	
TF 500	✓									✓	
HTC 80	✓	✓			✓✓✓	✓			✓		
HTC 95		✓		✓	✓✓✓	✓			✓		
HTC 120	✓	✓			✓✓✓	✓		✓	✓	✓	✓
HTC-C1-210								✓		✓	✓
HTC 250		✓			✓	✓			✓		

\* LOI stands for Loss on ignition.

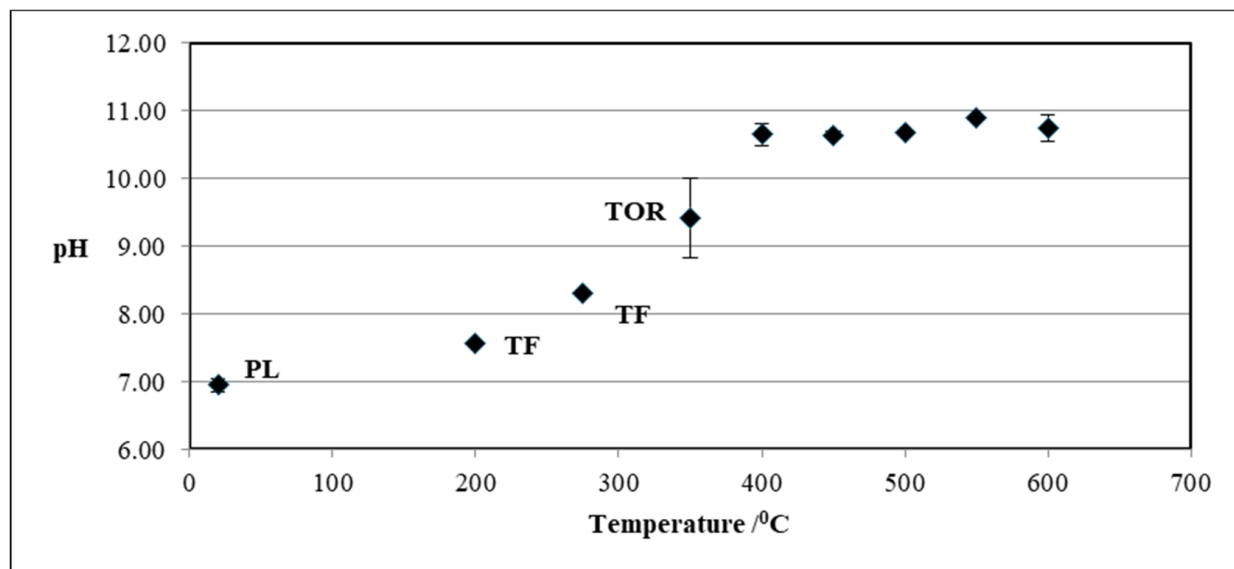
#### 3.2. Effect of Production Method and Temperature on PL Biochars' pH

Figure 1 below shows the influence of temperature on biochar's pH in the case of tube furnace and torrefied samples, merged together to give changes over a greater temperature range, as compared with the poultry litter's pH. Poultry litter is neutral, but as temperature increases, so does pH, levelling out at about 10.7 from 400 °C onwards, demonstrating that the most acidic groups were lost during the pyrolysis process and, as all the volatile matters were leached from the pyrolytic structure, the pH became constant. Other studies have found similar pH values of 9.5 at 300 °C to 11.5 at 600 °C [9,10], and reported that biochars generated from PL, peanut hulls, and pine chips through 400 °C pyrolysis had pHs of 10.1, 10.5, and 7.6, respectively. A higher pH at the higher pyrolysis temperatures is expected because of the increased relative concentration of non-pyrolyzed inorganic elements in the feedstocks and the formation of hydroxides and basic surface oxides [28]. It is reported that inorganic carbonates were the major alkaline components of the biochar generated at high temperature, and that organic anions contributed to the alkalinity of biochar generated at low temperatures [29].

There is a consensus in the literature that pH increases with pyrolysis temperature, but the magnitude of this increment depends on the raw material characteristics [30].

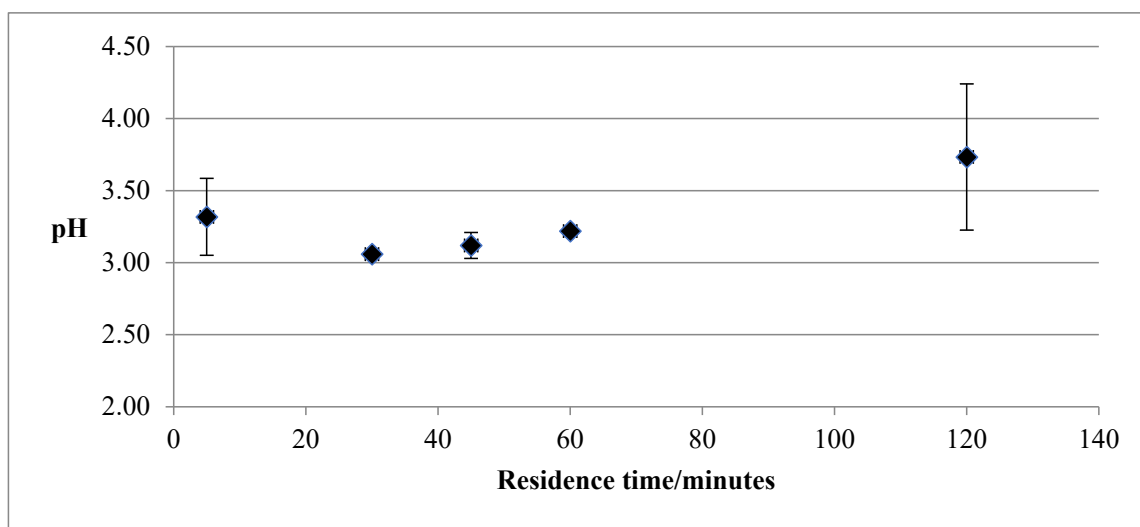
Hydrothermal carbonisation produces also a liquor (process water) along with the solid biochar samples. Therefore, the pH of the HTC samples was measured for both, as a function of temperature and residence time. Although the HTC was run at increasing temperatures, from 80 °C to 237 °C, there was insufficient solid formed at temperatures higher than 120 °C to do the pH tests. Therefore, there are only results for two temperatures,

i.e., 80 °C and 120 °C, respectively. For each temperature, the residence time to run the HTC varied from 5–120 min. The pH of the solid samples was 3.8 at 80 °C and 5.6 at 120 °C, respectively.



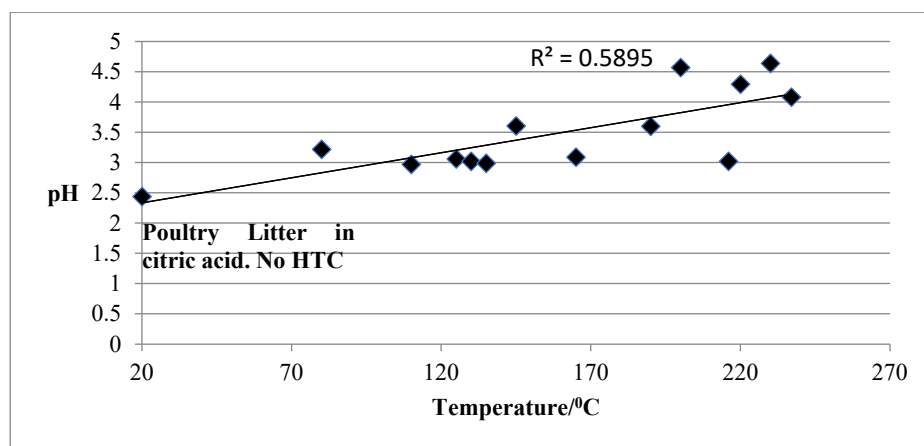
**Figure 1.** Effect of temperature on mean pH of char.

Figure 2 shows the pH variation of the liquor as a function of the residence time for the HTC temperature of 120 °C. The primary determinants of the pH of the HTC liquor are the raw material, PL in this case, and the liquid added, which was citric acid. Therefore, as expected, the pH was acidic, varying slightly with the residence time. This is another expected result, as the pH of the liquor was measured when the system was cooled down; therefore, the actual residence time was much longer than the HTC's running time.



**Figure 2.** Effect of HTC residence time on pH.

Figure 3 shows the pH of the process water as a function of temperature. For all temperatures, the pH of the liquor was acidic, with values from 2.97, at lower temperatures, to 4.57 at higher temperatures. The poultry litter's pH in the HTC medium (citric acid) at 20 °C was 2.44.



**Figure 3.** Variation in pH of HTC liquor with temperature.

Acidic pH was observed for the process water samples even when the HTC was performed in pure water. The pH of the spent liquor from hydrothermal carbonisation of PL, at the PL-to-water ratio of 1:5, 225 °C, and 15 min residence time, was 5.5 and unchanged by recirculation [31]. This finding supports the approach of using the spent liquor as an HTC medium for fresh HTC treatments, although, at one point in time, the process water should be disposed of, which will be a challenge due to its acidic nature.

However, the initial pH of the HTC medium is important as it impacts the yields and properties of the hydrochar (HC). It has been found that undertaking HTC in the presence of acids ( $\text{CH}_3\text{COOH}$ ,  $\text{H}_2\text{SO}_4$ ) significantly affects the yields and properties of HC. The C content and HHV of the HC increased with a decreasing initial pH. In the presence of  $\text{H}_2\text{SO}_4$ , the hydrochar yield increased while the ash content was significantly reduced. The lowest ash content and the highest hydrochar yield were measured in the HC produced from the suspension with an initial pH of 2 using  $\text{H}_2\text{SO}_4$ .

The acidic pH is due to the formation of organic acids that dissolve in the process water.

As is well known, HTC is an exothermic process that lowers both the oxygen and hydrogen content of feed (described by the molecular O/C and H/C ratio) by five main reaction mechanisms, which include hydrolysis, dehydration, decarboxylation, polymerization, and aromatization [32,33].

Although the catalytic effect of the acidic pH on the hydrolysis and dehydration of biomass is acknowledged, the effects of acidic conditions on other reaction mechanisms, such as decarboxylation and condensation polymerization, is largely unknown. It has been reported, however, that weakly acidic conditions improve the overall rate of reaction of hydrothermal carbonisation [34].

It is important that the pH of different biochars is accurately determined because changes in pH have great impacts on many soil processes, such as nitrogen mineralization, mineral precipitation, ion exchange, and greenhouse gas emissions [35].

### 3.3. Effect of Production Method and Temperature on Samples' Morphology, Composition, and Structure

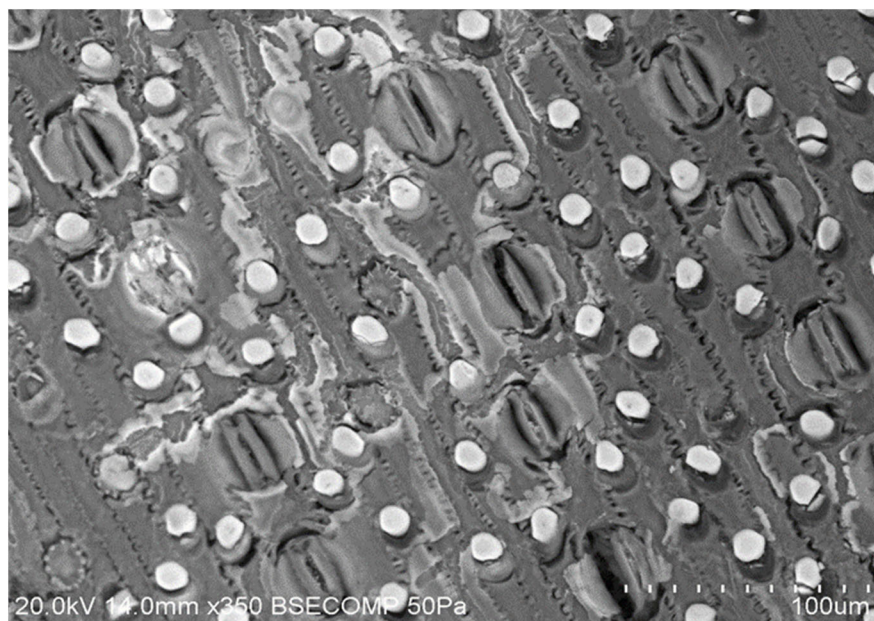
SEM-EDX measurements were performed over most of the samples (Table 2). The SEM micrographs showed the presence of different structures; some crystalline ones, coating the biochar particles or grown on the surface of a complex 3D lattice; some amorphous ones, while the morphology of the char samples was mainly a macroporous one. The composition of the different surface structures was determined, and the identification of their type and provenience was done based on the EDX results alone or by comparing them with those obtained on barley, feathers, or wheat straws. An attempt was made to determine the pore sizes and their distribution.

**Table 2.** Biochar samples characterised by SEM only. Number of ticks indicates number of replicates.

Sample	Pore Size	EDX	Structures and Crystals
TOR 350 strip	✓✓✓	✓✓✓	✓✓✓
TOR 600 strip	✓✓✓	✓✓✓	✓✓✓
TOR 350 ash	✓✓✓	✓✓✓	✓✓✓
TOR 400 ash	✓✓✓	✓✓✓	✓✓✓
TOR 450 ash	✓✓✓	✓✓✓	✓✓✓
TOR 500 ash	✓✓✓	✓✓✓	✓✓✓
TOR 550 ash	✓✓✓	✓✓✓	✓✓✓
TOR 600 ash	✓✓✓	✓✓✓	✓✓✓
TOR 550 ash strip	✓✓✓	✓✓✓	✓✓✓
TOR 600 ash strip	✓✓✓	✓✓✓	✓✓✓
Feather		✓✓✓	✓✓✓
Wheat		✓✓✓	✓✓✓
Barley		✓✓✓	✓✓✓
Egg shell	✓✓✓	✓✓✓	✓✓✓
KCl crystals		✓✓✓	✓✓✓

### 3.3.1. Amorphous Structures

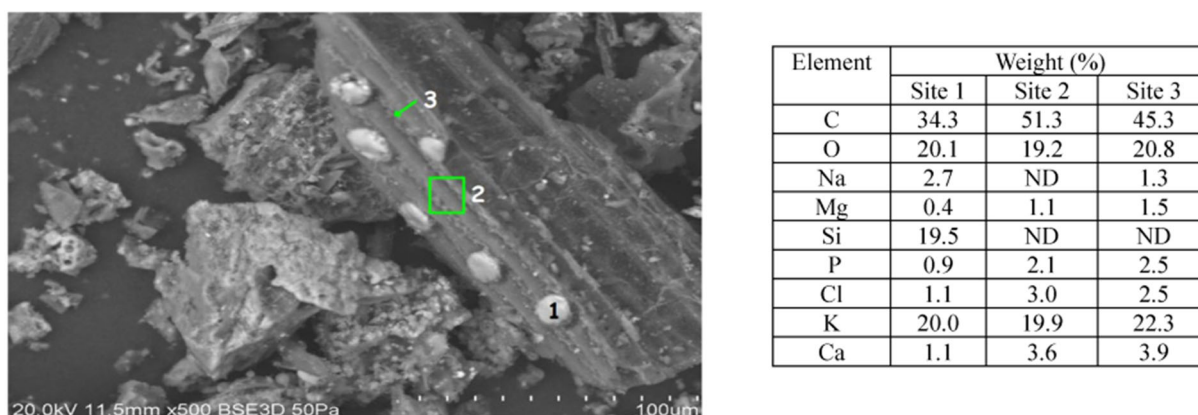
As shown in Figure 4, two more obvious structures are present in the mixture of the HTC chars, one white and round, and another one that looks like a grass epidermis, showing stomata and phytoliths. The elemental analysis proved that the round structures are siliceous and about 10–20  $\mu\text{m}$  in size. These siliceous structures were found in many of the chars and appear to be loosely attached to the surface.

**Figure 4.** SEM micrograph of a mixture of chars obtained by HTC at 120 °C, but with different residence times.

In order to identify the provenience of these siliceous structures, EDX measurements were performed on them and on the background material, and the results were compared with those obtained on siliceous structures in wheat and barley. The stems did not show the siliceous structures, but the leaves did, showing that these structures are not caused by torrefaction. Elemental composition in wheat and barley is just C, O, and Si. However, these



non-torrefied structures contain more oxygen, which would be expected as combustion reduces oxygen content. The other elements on the char samples may be material deposited on the surface during torrefaction (Figure 5, Figures S1 and S2).



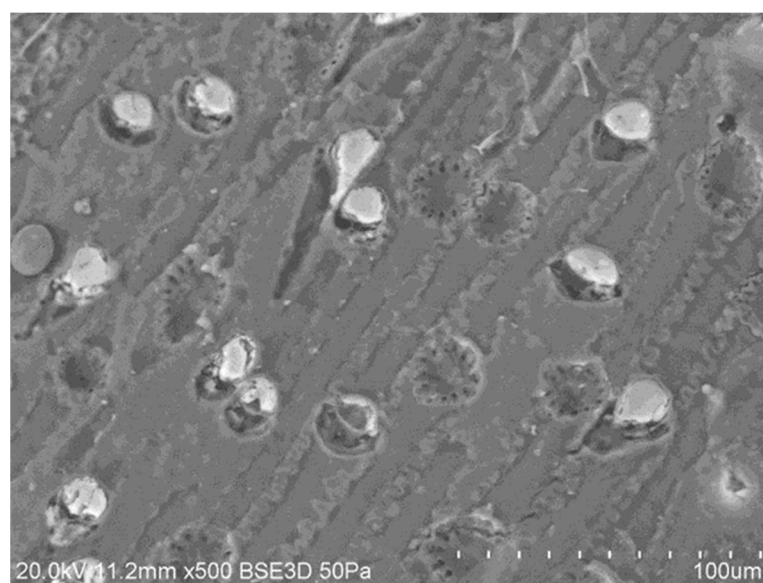
**Figure 5.** SEM/EDX results from TOR 600 sample.

Figure 5 shows the results obtained from the TOR 600 sample, while Figures S1 and S2 present the SEM-EDX results obtained on the oven-dried wheat straw and barley leaves, respectively.

As shown in Figure S1, the wheat sample presents siliceous structures and stomata. Barley (see Figure S2) has similar structures, although the elemental composition is more varied, containing K, Cl, and Na. Stomata are also visible.

The size of the siliceous structures was measured for torrefied samples as well and compared with the size of the siliceous structures in PL, wheat, and barley. The average size decreases as the temperature increases, i.e., for TOR 400 it is 20.4  $\mu\text{m}$ , while for TOR 500 the value is 15  $\mu\text{m}$ . The size of the siliceous structures in PL is, on average, 15.6  $\mu\text{m}$ ; for wheat it is 17.6  $\mu\text{m}$ , while for barley it is 13.6  $\mu\text{m}$ , respectively. The size comparison and appearance suggest that the siliceous structures are probably wheat. However, the sample is too small to be certain—it is possible that both are present.

What is more, SEM images of PL (Figure 6) reveal the presence of other siliceous structures that do not look like those found in either oven-dried wheat straw or barley leaves.



**Figure 6.** SEM micrograph of PL sample.

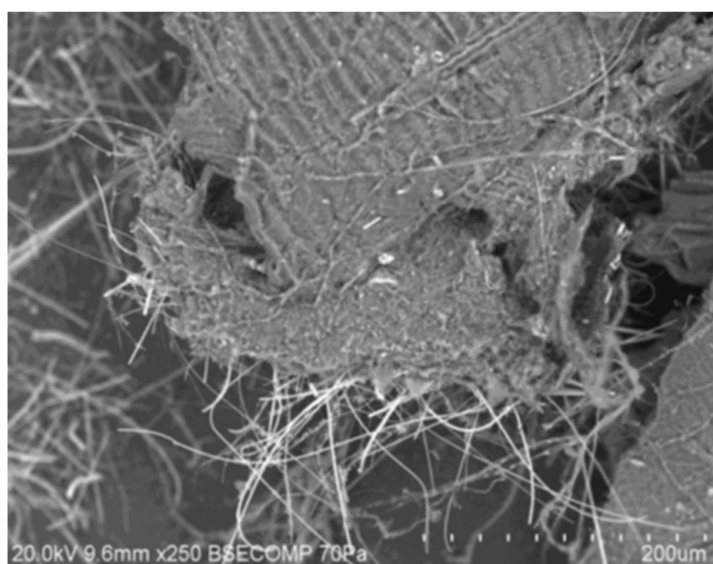
For PL, the structures seem to be in hollows. Also visible are what appear as wiggly lines between cells. These are probably phytoliths and can be seen in greater detail on the barley leaf (Figure S3). They are about 124  $\mu\text{m}$  in length and are composed mainly of silicon. Phytoliths are used in archaeology to identify species as they are resistant to decay, so they could be used to identify the plant material a char is derived from [36]. Phytoliths are primarily silica and can be a problem if feedstock is used as a fuel [37].

These phytolith structures remained and were visible in ash samples—not surprising as they are separated from plant material by ashing [36].

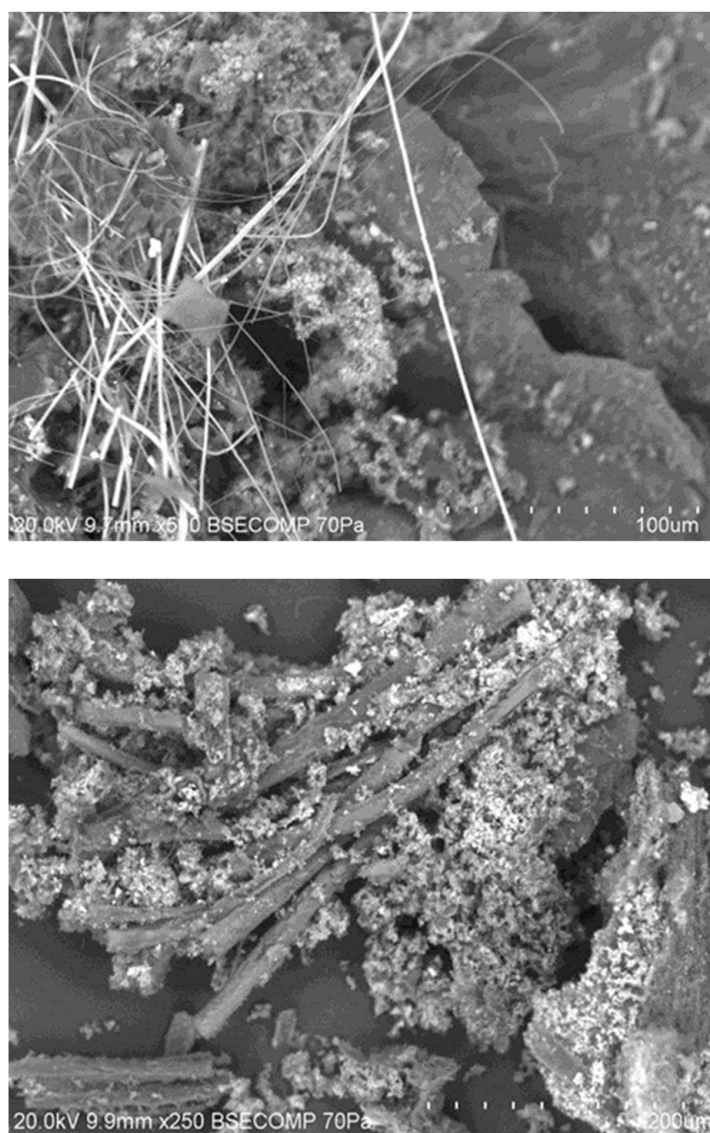
There were other amorphous structures found in torrefied material that looked like feathers (see Figure S4) and had a similar chemical composition. To test their provenience, feathers and torrefied feathers were analysed.

As can be seen, the actual feather structure is a main rib with side branches emerging, similar to the structure shown in TF135. These branches appear to be strap-like and narrow as they proceed further from the main rib. Torrefaction causes feathers to lose most of their fine structure. The size is also similar. Elemental analysis also suggests that this is a feather. The C and O percentages are similar for the actual feathers and the structures in TF135. Feathers are made of the protein keratin, which has high levels of S, as seen in the elemental analysis. The TF135 feather has a low S value, but this could be a result of natural variation. Another problem is the difference between the feather heated to 450  $^{\circ}\text{C}$  and one heated to 105  $^{\circ}\text{C}$  (oven-dried). Heating seems to cause a loss of Na and Ca; however, the difference may be because the feather was not of the same type; it was taken from a local bird rather than from the PL. Therefore, the long fibres in the HTC chars could have been keratin protein fibres from feathers, which have “an intricate network of connective and hollow fibrous structure [38]”. However, it was found that HTC would hydrolyse keratin from chicken feathers into small protein [39].

As for the hydrothermal carbonisation samples, some more and different amorphous structures were shown (see Figure 7). Many fine white structures, 700 nm–2  $\mu\text{m}$  long, are visible. These could be cellulose fibrils, as the literature suggests that HTC treatment breaks down hemicellulose, which causes individual cellulose fibres to separate. Debris from the cell wall is then deposited on the surface of the hydrochars [40]. However, microfibrils are 28 nm in diameter, so these fibres are too large—they may be aggregations of microfibrils.



**Figure 7.** *Cont.*



**Figure 7.** SEM images of HTC 250 sample.

Structures like roots were also observed. The diameter of these structures is about 20 µm for the larger ones, going down to 2 µm for finer branches. These are about the correct size of and similar structure to roots.

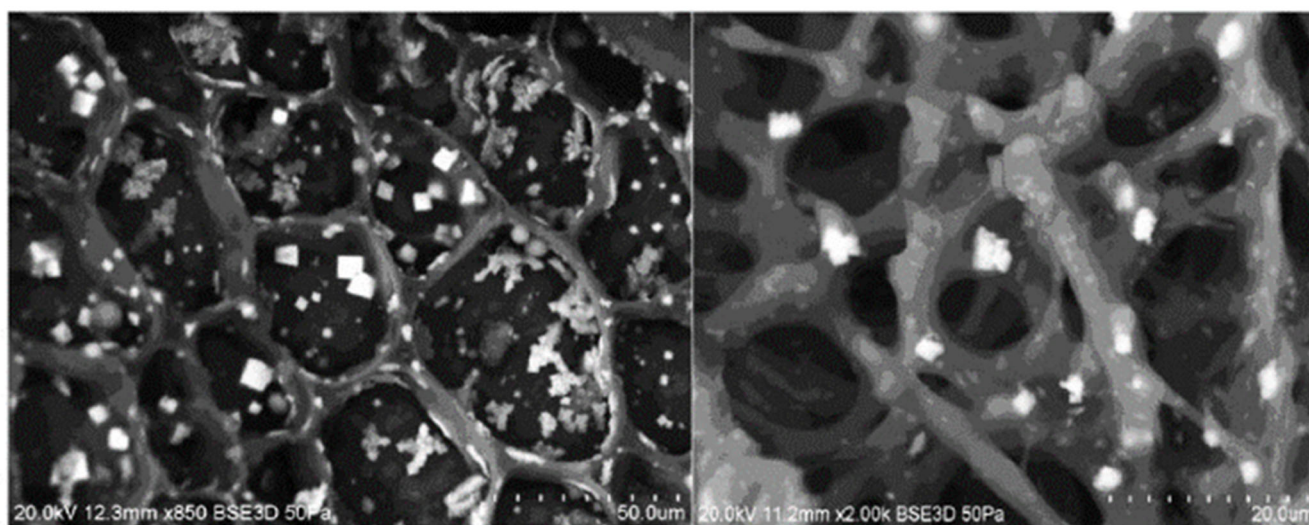
### 3.3.2. Crystalline Structures

Many structures appear crystalline with very regular shapes and distinct angles. These structures, which are found on the surface, seem primarily inorganic and may, therefore, be very important when PL is used as a soil amendment.

The crystalline structures found on the TOR 350, TOR 450, and TOR 550 samples, respectively, along with those found on the TF 350 and TF 275 samples, were studied in detail. Results are as follows.

Figure 8 presents the SEM images of the crystalline structures observed on the complex 3D network of the TOR 350 char sample along with their composition. These crystals are about 2 µm on one face and appear cuboid. The elemental analysis shows high amounts of K and Cl, so these crystals could be potassium chloride.





Element	Weight%
C	38.9
O	18.7
Na	0.9
Mg	0.2
Al	5.8
Si	0.2
S	1.1
Cl	14.0
K	19.7
Cu	0.5

**Figure 8.** SEM/EDX results from TOR 350 char sample.

In order to support this statement, pure KCl crystals were characterised by SEM/EDX (Figure S5). The K:Cl ratio is 1.1 for the pure KCl crystal, very close ( $\pm 1\%$ ) to the published value [41], while for the TOR 350 char sample, this ratio is 1.4. Therefore, even all the Cl is in the form of KCl, that still leaves some K and, with high amounts of carbon and oxygen present, also suggests a possible presence of potassium carbonate,  $K_2CO_3$ , which is deliquescent. KCl crystals are also hygroscopic.

Crystalline structures were found in other samples as well, in some cases in very large quantities, as for the TOR 450 char sample. There are two types of crystalline structures observed, some bright white ones and some grey ones, with very similar composition, but with far lower Cl and higher Ca percentages, respectively, for the grey ones. The K:Cl ratio for the bright white ones is 1.5, which supports the assumption that they are most probably KCl crystals. These crystals are about  $2\ \mu\text{m}$  on one face and appear cuboid (see Figure 9).

As for the TOR 550 sample, the crystalline structure measured has a high percentage of P, K, and Mg, confirmed as pyrochroite ( $K_2MgO_7P_2$ ) [42] via XRD. Also, although it has a very high concentration of K but only a little Cl, it means that K is present in forms other than KCl [3].

Wide-angle XRD measurements were performed in order to identify the nature of all crystals present on the torrefied samples.

Figure 10 shows the diffractograms for the three torrefaction samples, namely TOR 350, TOR 450, and TOR 550, respectively.

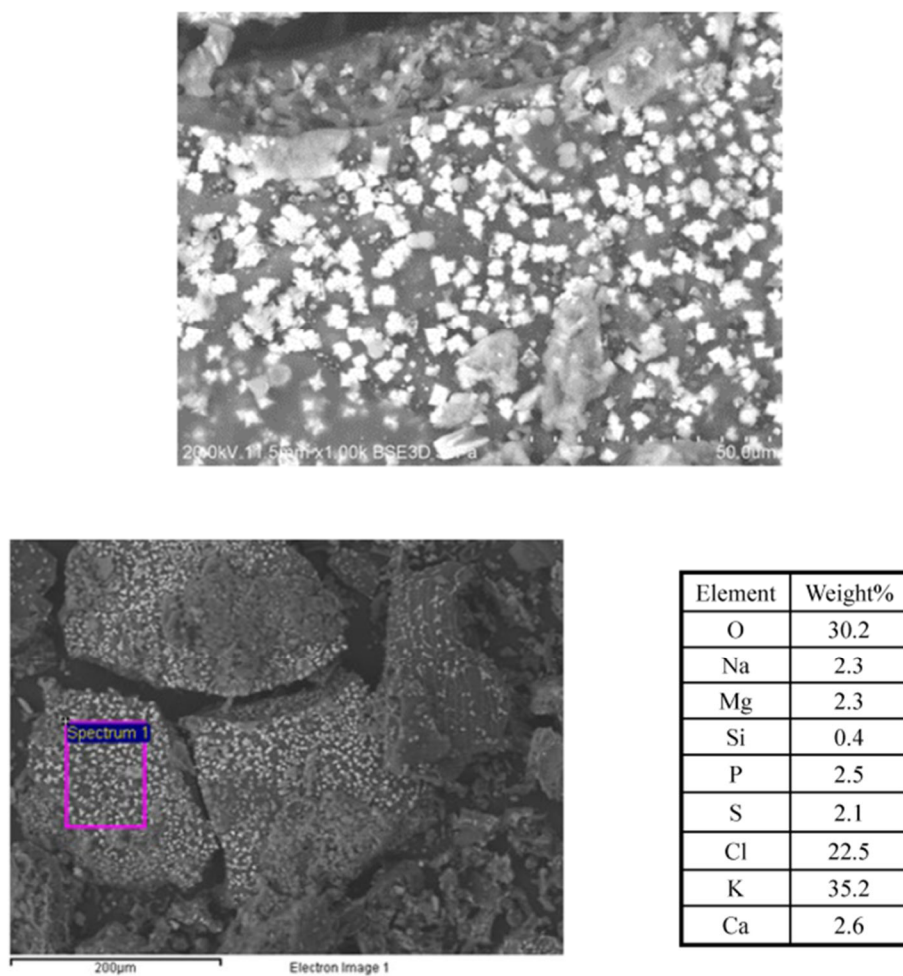


Figure 9. SEM/EDX results from TOR 450 char sample.

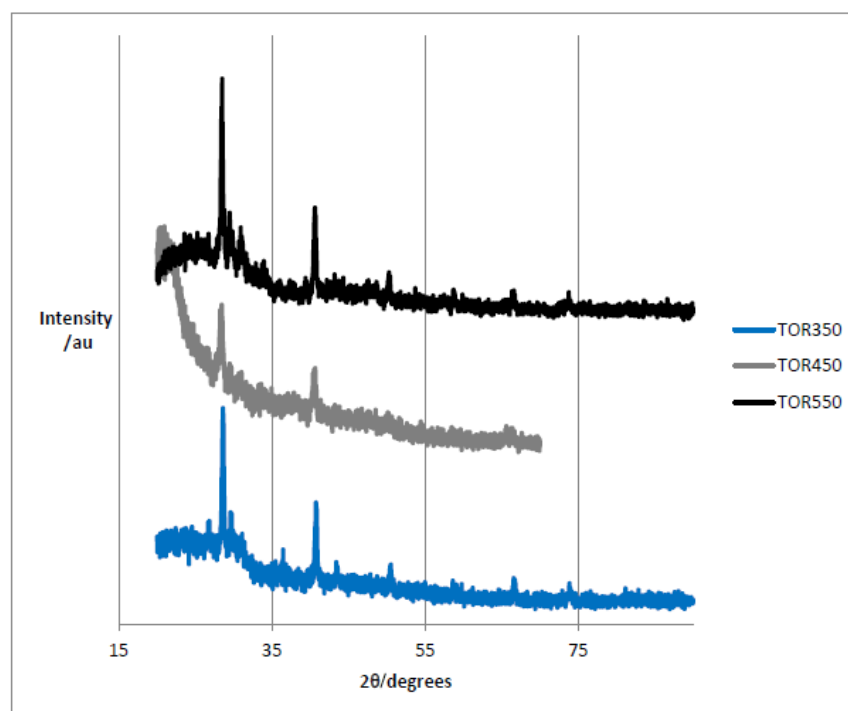


Figure 10. Wide-angle XRD for the TOR 350, TOR 450, and TOR 550 char samples, respectively.

The samples are broadly similar, with several peaks at the same positions for all chars. These peaks are considered more reliable and will be analysed below.

Low-angle XRD was measured as well to assess the broad peak at  $2\theta$  between  $17^\circ$  and  $27^\circ$ . The broad peak showed a complete amorphous structure. There were no diffraction peaks observed except for a broad band centred at  $2\theta$  decreasing from  $22.5^\circ$  for TOR 350 to  $20.5^\circ$  for TOR 550, with the peak maxima at  $2\theta = 21^\circ$  for TOR 450, which is a well-known feature for amorphous silica material. The results were in good agreement with those on the JCPDS file for  $\text{SiO}_2$ .

Amorphous carbon also shows a peak in this area [13,43].

Table 3 presents the peaks' identification for the three torrefaction samples. The conclusion from the XRD measurements is that sylvite is present in all samples; calcite and quartz are also present. There are some diffraction peaks in TOR 550, which are difficult to identify. The complex mixture of minerals that might be present causes problems with identification. In rice husk char, for example, as well as sylvite and calcite, a number of complex potassium minerals were present: archerite ( $\text{KH}_2\text{PO}_4$ ), chlorocalcite ( $\text{KCaCl}_3$ ), kalicinite ( $\text{KHCO}_3$ ), pyrocoprite ( $\text{K}_2\text{MgO}_7\text{P}_2$ ), struvite ( $\text{KMgPO}_4 \cdot 6\text{H}_2\text{O}$ ) ([13]). All of these are possibly present given the elemental composition found by EDX but may be present in only small quantities and be undetectable.

**Table 3.** XRD peaks' identification for TOR 350, TOR 450, and TOR 550 biochar samples, respectively.

TOR 350		TOR 450		TOR 550		Identification
$2\theta$	Intensity (Counts)	$2\theta$	Intensity (Counts)	$2\theta$	Intensity (Counts)	
28.6	514	28.44	390	28.44	504	Sylvite
29.6	266			29.48	278	Calcite
30.46	178			30.92	244	Pyrocoprite ( $\text{K}_2\text{MgP}_2\text{O}_7$ )
36.46	178					Quartz ( $\text{SiO}_2$ )
40.74	290	40.64	239	40.62	291	Sylvite
43.4	148					Quartz ( $\text{SiO}_2$ )
50.52	142			50.28	137	Sylvite
				58.7	96	Sylvite
60.56	110			59.86	84	Quartz ( $\text{SiO}_2$ )
				66.34	93	Sylvite
73.84	98			73.74	90	Sylvite (Anon, 2008)
				83.56	70	Unidentified
				86.62	74	Unidentified

Another XRD study of chicken manure identified calcite, hydroxyapatite, struvite, dolomite, quartz, and magnesium phosphate [44]. This study found calcite and quartz—again, the others may be present in low quantities in the studied PL, but the fact that sylvite is not present is surprising. This implies that the sylvite possibly originates from the plant material, which is not present in manure. This is supported by another study that found sylvite in a  $300^\circ\text{C}$  biochar from straw (*Brassica campestris*). In this case, the higher-temperature biochars had no sylvite but did show calcite and dolomite [45].

The size of the sylvite crystallites was determined by using the Scherrer equation. For  $2\theta$  around  $28^\circ$ , the size of the sylvite crystal was  $28.5$  at  $350^\circ\text{C}$ ,  $21.4$  at  $450^\circ\text{C}$ , and  $21.4$  at  $550^\circ\text{C}$ , respectively. As for  $2\theta$  around  $40$ , the size of the sylvite crystal was  $24.6$  at  $350^\circ\text{C}$ ,  $22.1$  at  $450^\circ\text{C}$ , and  $22.1$  at  $550^\circ\text{C}$ , respectively. There is no obvious trend for crystal size with temperature, with all values being similar for a given peak.

One problem with XRD is that it is best suited to the analysis of homogenous material rather than these heterogeneous chars. It has a detection limit of 2% for components in mixtures, so substances present in low concentrations may not be detected [46].

A variety of crystalline structures were observed as well on the surface of the biochar samples prepared in the tube furnace. Figure 11 presents the SEM micrograph along with EDX composition for the TF 275 sample, while Figure 12 shows SEM/EDX results from the TF 350 biochar sample.



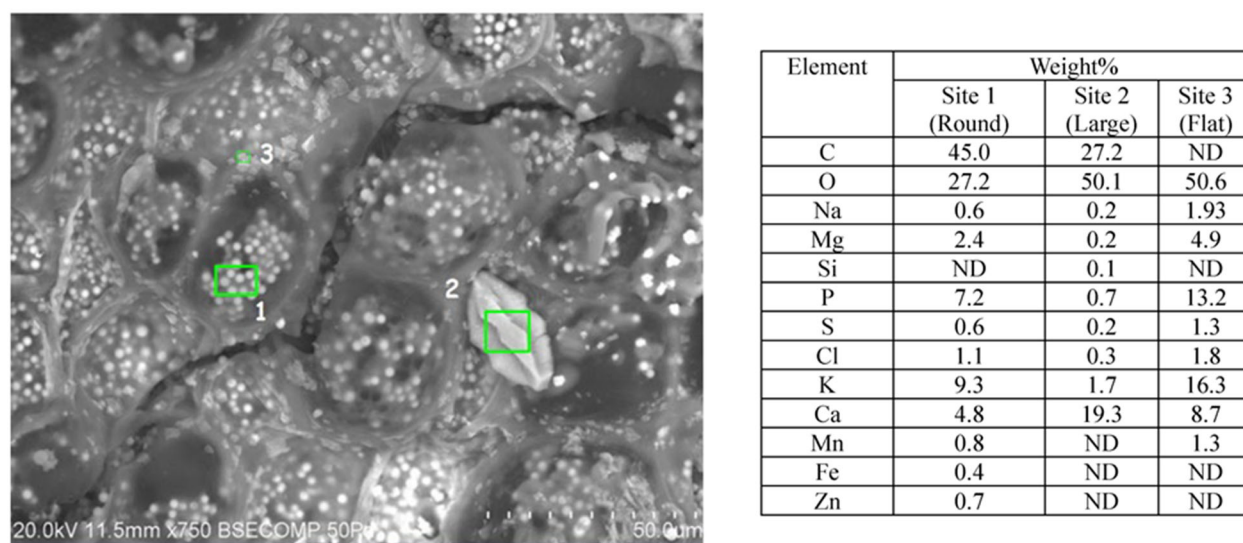


Figure 11. SEM/ EDX results from TF 275 char sample.

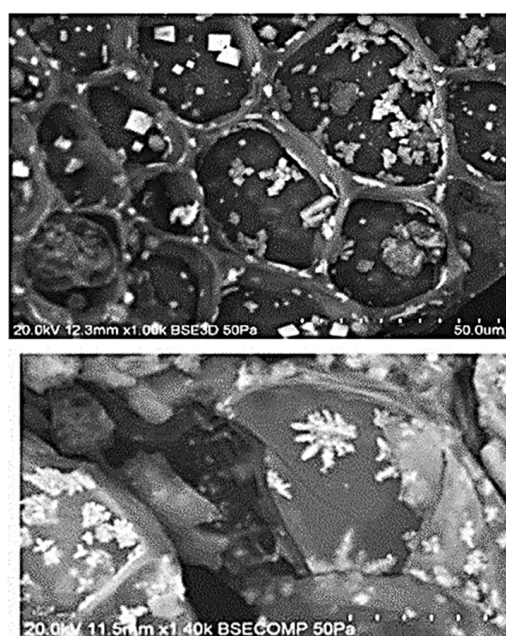


Figure 12. SEM/EDX results from TF 350 char sample.

The large crystal observed on the TF 275 sample is probably calcium carbonate (calcite), whilst the flat crystal is possibly some type of phosphate. The round crystal could be potassium carbonate ( $K_2CO_3$ ) or possibly pyrocoproite ( $K_2MgP_2O_7$ ), which was found to be present in the XRD results for the TOR samples.

The crystalline structure shown on the TF 350 sample is most probably aluminosilicate.

To the best of our knowledge, this is the first time that the crystalline structures on the surface of the poultry litter biochar were identified, and their composition determined.

Some of the feedstock nutrients, namely P, K, Ca, and Mg, were concentrated on the surface, and their content increases with increasing the pyrolysis temperature. It is important to know the type and composition of the crystalline structures on the surfaces of the biochars as these crystals are the first to be released into the soil, to cycle nutrients back into agricultural fields. As a measure of the direct nutrient value of biochars, it is not the total content but, rather, the availability of the nutrient that is an important consideration.

Hydrochar samples showed no crystals on the surface as they, presumably, dissolved in the process water.

### 3.3.3. Macroporosity and Elemental Composition (Surface, Subsurface, and Bulk) of Biochar Samples

The SEM/EDX technique was also employed to visualise the porous structure and to determine the pore size and the elemental composition of the biochar samples, and to assess their changes with the biochar production conditions. As seen in Figures 13 and S6–S10, distinct morphologies of increasing porous structure are present.

A

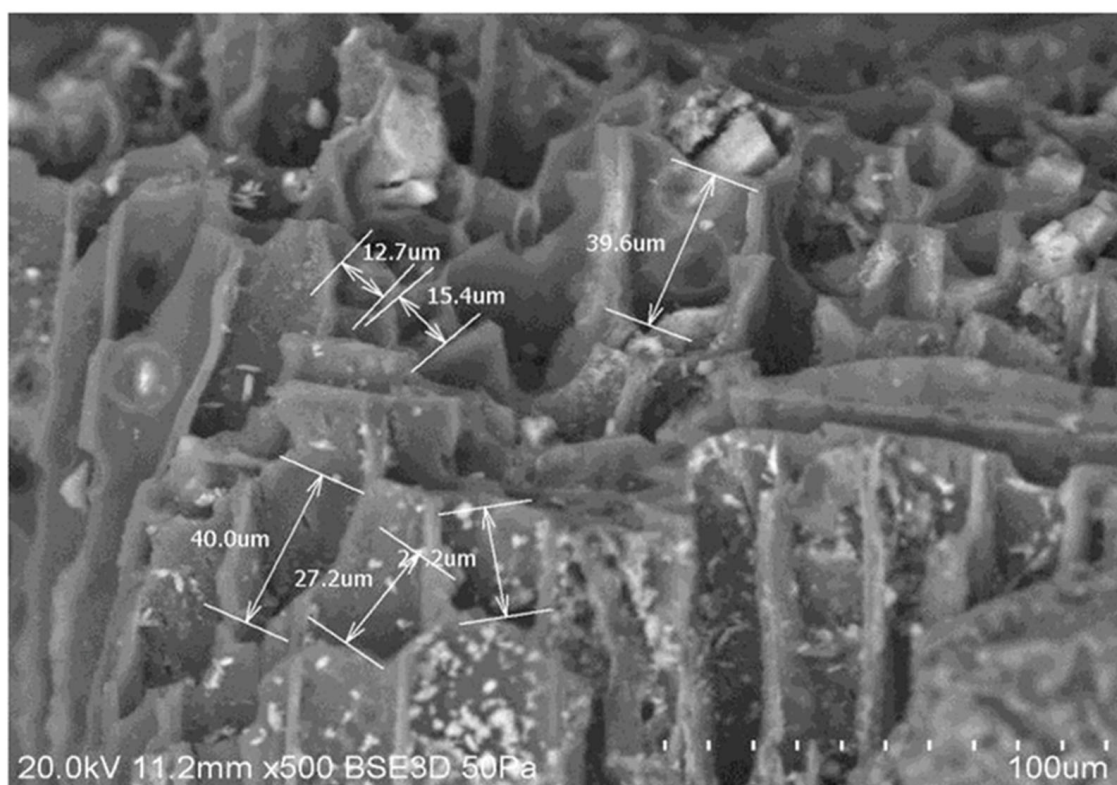
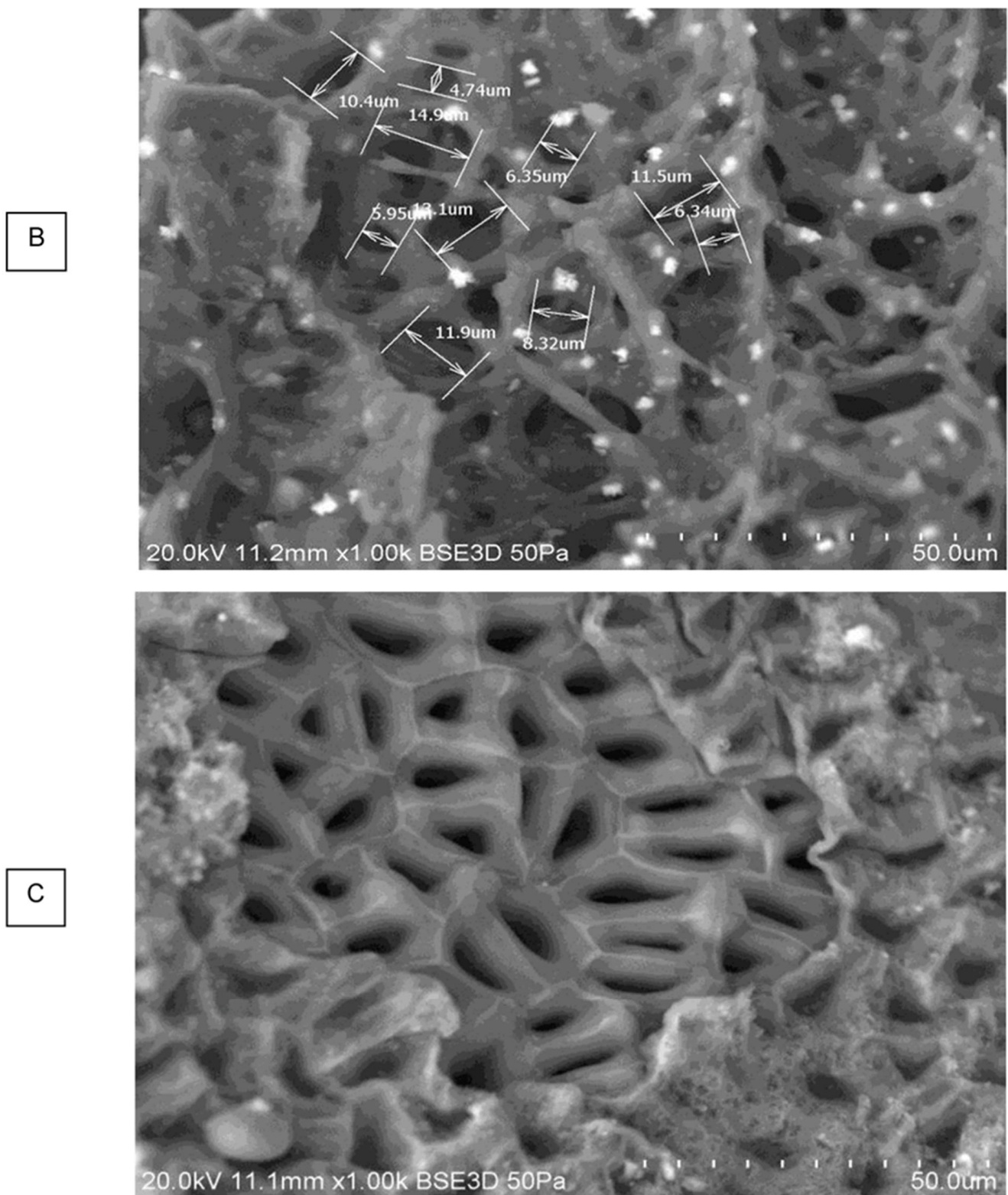


Figure 13. Cont.



**Figure 13.** SEM images of TOR 350 sample showing complex macroporosity: (A) pores formed by the open ends of cells; (B) complex 3D lattice; (C) true honeycomb lattice.

The represented microscopic honeycomb-like structures, typical of fibrous plant materials, are present in the PL biochar from the wood shavings used as bedding material. These microstructures evolve in shape and complexity as the torrefaction temperature increases.

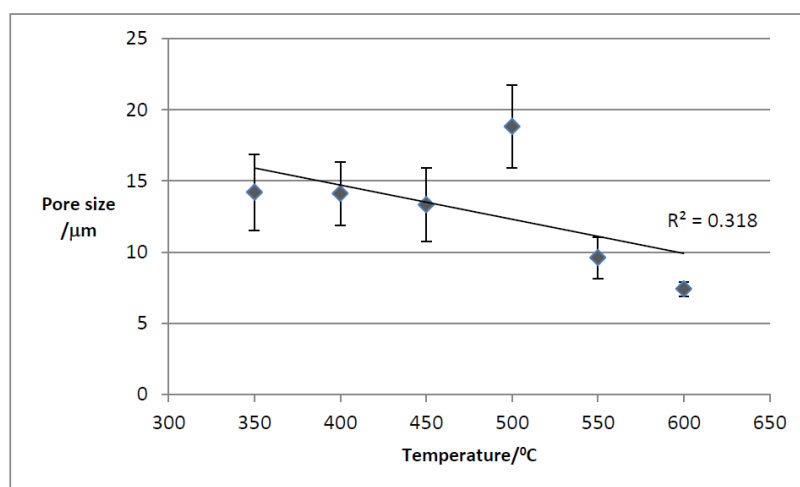


The TOR 600 sample, for example, shows a very regular pore structure, but still has a wide range of pore sizes, from 4.2  $\mu\text{m}$  to 12.3  $\mu\text{m}$  (Figure S10A).

Along with the external macroporosity between the biochar's particles and the residual macroporosity based on the plant cellular structure shown in the SEM images, a pyrogenic nanoporosity develops within the solid biochar volume and increases with production temperature but constitutes only a small portion of total porosity, even in higher-production-temperature biochars [47]. What is more, the pyrogenic nanopores are formed because of chemical changes at higher pyrolysis temperatures, higher than 600  $^{\circ}\text{C}$ , which was the maximum torrefaction temperature used to prepare the samples in this study.

Due to this, our study focused mainly on the macroporosity of the biochar samples, which is a key parameter influencing their water uptake.

Using the SEM software, the pore size was directly measured. Figure 14 presents the variation in the mean pore size with temperature.

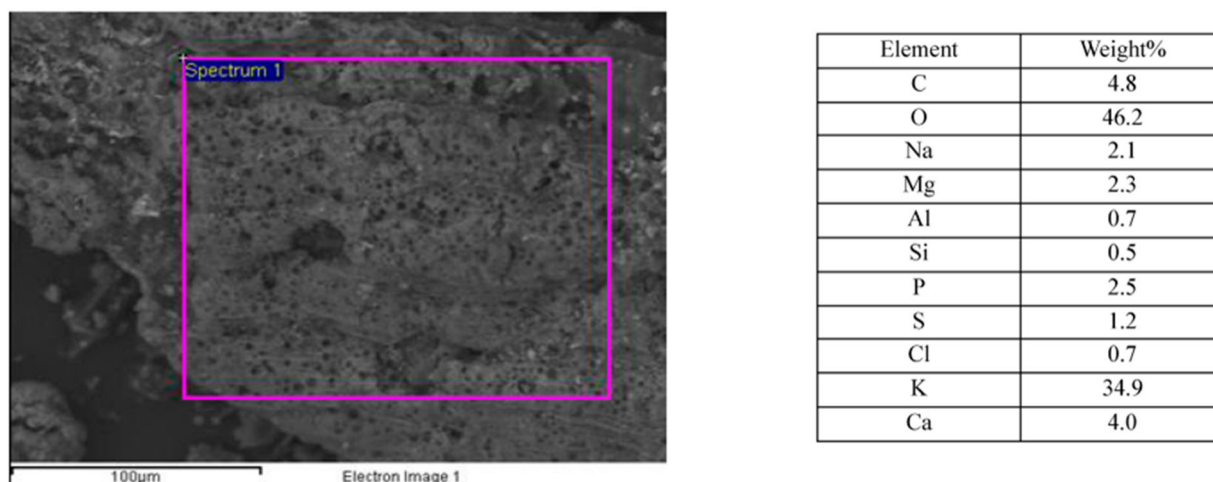


**Figure 14.** Variation in the mean pore size with temperature for the TOR biochar samples.

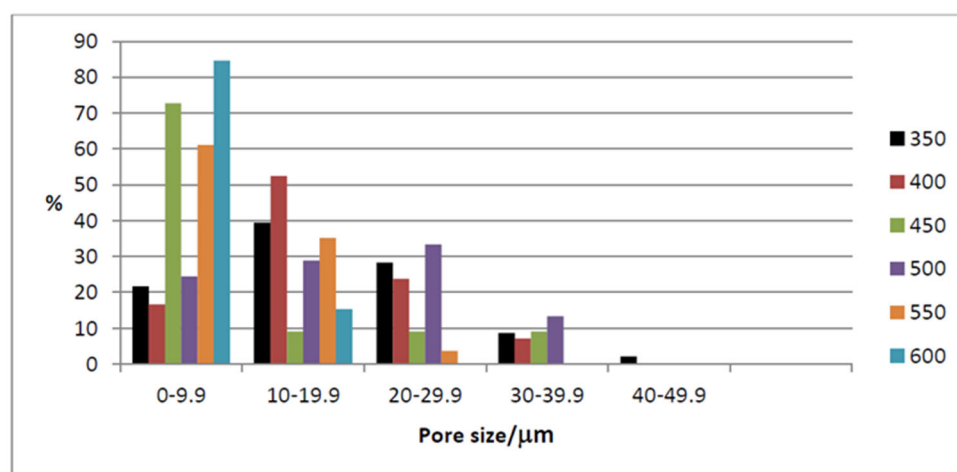
Mean pore size decreases with torrefaction temperature as expected but TOR 500 seems anomalous. In fact, there are many problems with this method of measuring pore size. It is often difficult to decide where the maximum diameter is, as this involves subjective judgement. Also, some pores are seen in oblique view, so that the measurement is larger. Most studies that find a decrease in pore size with temperature are considering a single feedstock [47]. One of the main features of poultry litter is its heterogeneous nature. Pore size is linked to the cell size of the feedstock—here, there are many different plant species present, including grasses and wood shavings as well as inorganic material. The SEM images and EDX composition of such an inorganic structure from TOR 350 are presented in Figure 15. Its elemental analysis shows very little carbon, suggesting its inorganic nature. It looks mineral and contains both calcium and potassium at high levels, with magnesium and sodium also present.

The heterogeneity of the PL means that not all SEM measurements are from the same material; thus, the observed variation in morphology. However, there are some advantages of the SEM direct measurement of the pore size method. The pores are seen, so the wide variation in size, in even a single site, can be seen and measured.

In a heterogeneous feedstock like PL, it potentially allows the contributions of different materials to pore size to be identified. It is also possible to see how the pores are formed from cell walls. Perhaps a more representative measurement of pore size would use only one type of pore, such as what will be called a “true honeycomb”, as seen for TOR 600 (Figure S10B). The outer layer of plant material has been lost, exposing the cellular structure. The SEM pore size distribution measurement results are presented in Figure 16.



**Figure 15.** SEM images of an unusual structure on TOR 350 along with its EDX composition.

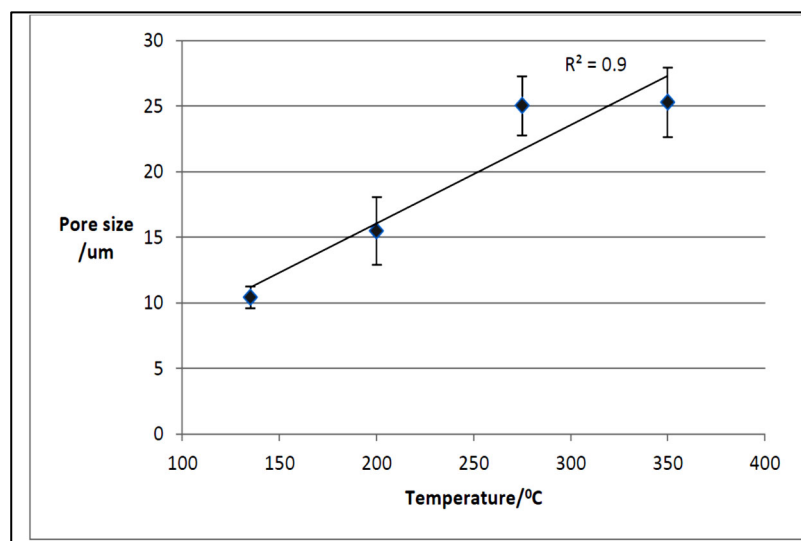


**Figure 16.** Frequency distribution of pore size with temperature for torrefied biochar samples.

TOR 350 has the widest range of pore sizes, and the most frequent size is between 10 and 19.9 μm. TOR 600 has a small range, and the most frequent size is 0–9.9 μm. TOR 500 seems anomalous with a peak frequency at 20–29 μm.

As for the TF biochar samples, SEM measurements showed that the mean pore size is increasing with temperature (see Figure 17), which was an expected result as well. Since the poultry litter feedstock was a mixture of wood, barley, wheat straws, feathers, chicken excreta, and spilled feed, etc., the microscopy images of TF samples show, as for the TOR samples, the presence of different types of biochar particles of varying size and morphology, which led to external pores, i.e., those pores between the biochar particles, of different sizes. The residual macroporosity is formed by the evolution of volatiles from the solid during the thermal degradation of the poultry litter. Knowing that the corresponding temperatures for maximum decomposition of hemicellulose and cellulose are 300 °C and 355 °C, respectively, while lignin decomposition begins at about 280 °C with a maximum rate occurring between 350 and 450 °C and the completion of the reaction at 450 and 500 °C, the increase in the pore size with temperature for the TF sample is most probably due to the volatilisation of hemicellulose and cellulose within the temperature range used in the tube furnace.

As the temperature of the conversion of poultry litter to biochar increases to produce the TOR samples, the pore size decreases since almost all the volatile compounds were lost at temperatures of around 350 °C.



**Figure 17.** Mean pore size variation with temperature for the TF biochar samples.

There is a slight difference between the mean pore size for TF 350 and TOR 350, 25  $\mu\text{m}$  compared to 14  $\mu\text{m}$ . It might be due to the different amount of poultry litter used to obtain the two samples: two grams were used for the TF samples' production, while for the TOR samples, 1 kg of poultry litter was used, with the heat transfer for carbonisation being faster for the surface samples (TF) than that for the bulk samples (TOR).

Figures S11–S14 show the SEM images and pore size measurements of the TF samples. There is still a wide range of pore sizes in each temperature as in TOR samples. For TF 135, the residual macropores are not very wide; some of them are not completely open, so the surface area and volume, respectively, of the pores is low (Figure S11). As the conversion temperature is increased at 200  $^{\circ}\text{C}$ , pores with different sizes and appearance (far more open), similar to those observed for the TOR samples, are more likely to be detected along with pores such as those on the TF 135 sample (Figure S12).

At higher temperatures, i.e., 275 and 350  $^{\circ}\text{C}$ , respectively, pores are now similar to those found in the TOR samples.

Figure 18 shows the frequency distribution of pore size with temperature for the tube furnace biochar samples. There is a smaller range of pore sizes at lower temperatures, with most pores in the smaller size range for TF 135 and TF 200.

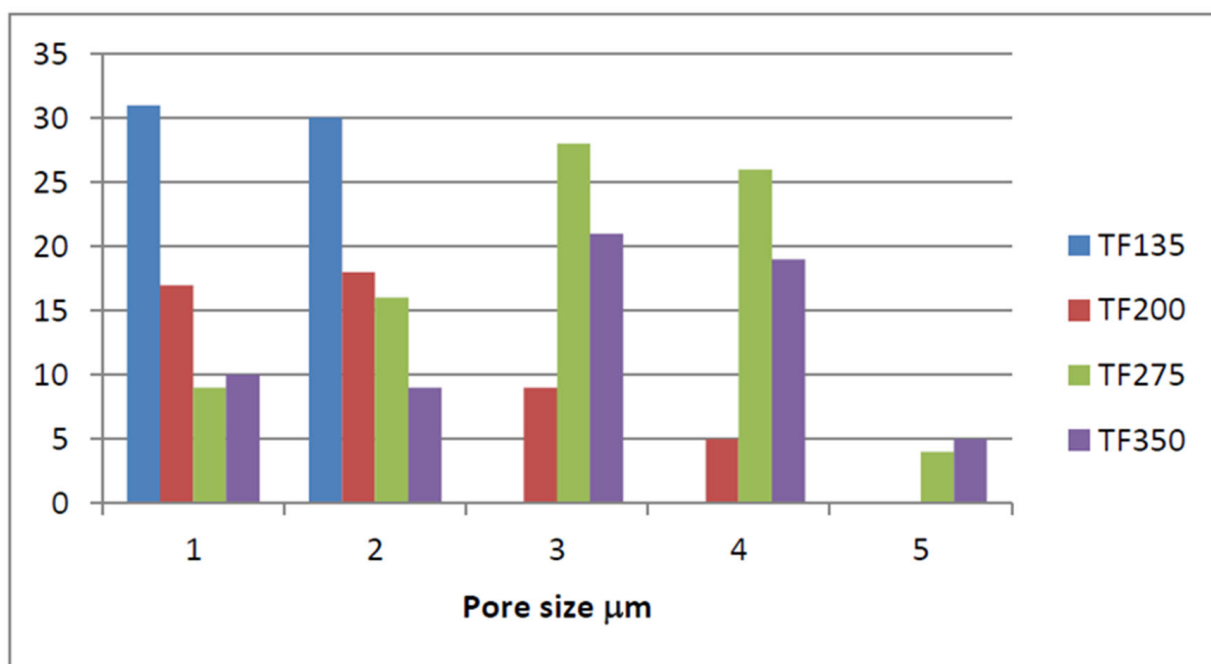
SEM images were taken for the HTC samples as well. Figures S15–S18 show the morphology and pore size measurements of the carbonised samples. As seen in Figure S15, for the HTC 80 sample, despite quite a low temperature, there is a large amount of honeycomb morphology with many pores in the process of formation.

True honeycomb morphology, but with no cavity in the centre, was observed for the HTC 95 sample (see Figure S16).

For the HTC 120 sample (Figure S17), the honeycomb has a different structure to previous samples; these are arguably not true pores, or they are pores in the process of forming. The pores may be formed from a different plant material. It is also possible that the differences in pore morphology in the HTC samples are due to the different decomposition of polymers in HTC, where hydrolysis is the most important mechanism [33]. The difference could also be due to pressure, although this is a low-pressure sample (2 bar).

If they are pores in the process of formation, a higher carbonisation temperature should be beneficial. Therefore, an extra HTC biochar sample was prepared at 221  $^{\circ}\text{C}$ , HTC 221. Figure S18 shows the SEM micrograph and pore size measurements for this sample. The microscopic pore morphology seems to be improved but there still are closed cavities. Similar observations were made about PL hydrochars obtained at temperatures of 150 and 175  $^{\circ}\text{C}$ , respectively; SEM micrographs showed incomplete decomposition and a corrugated surface with holes [48].

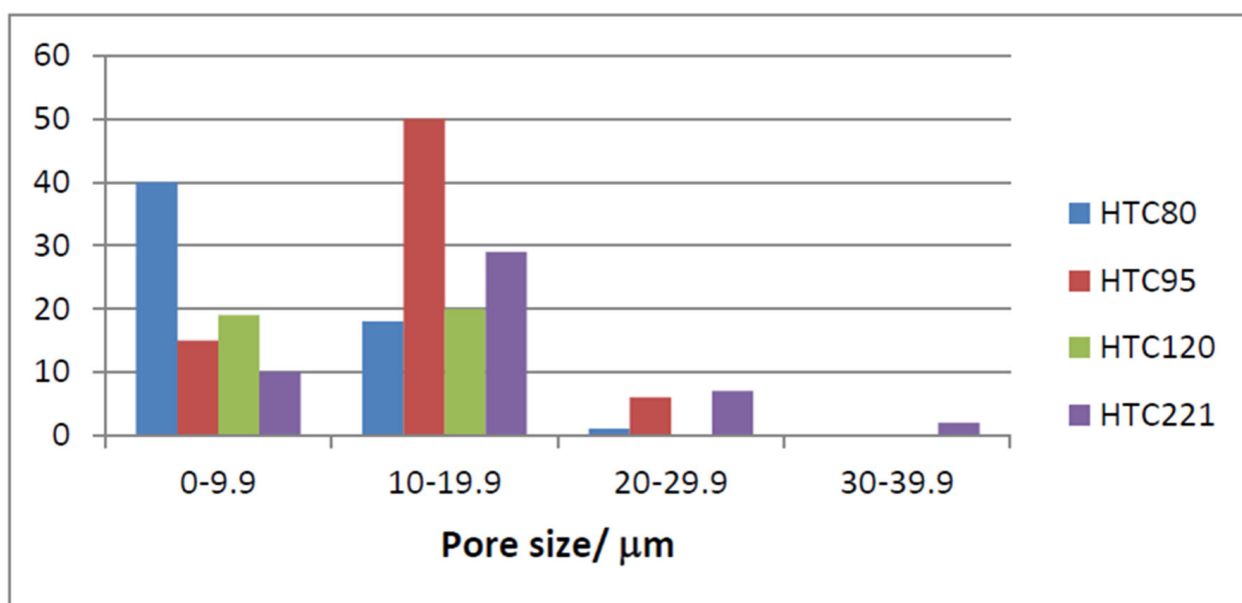




**Figure 18.** Frequency distribution of pore size with temperature for tube furnace biochar samples.

As expected, the mean pore size of the HTC samples (slightly) increases with temperature, from 10 µm at 80 °C to 15 µm at 221 °C.

Figure 19 presents the frequency distribution for the HTC samples' pore size against temperature showing a lower range than for the torrefied material with very few pores 20 µm or above. However, HTC 221 has the highest number of large pores. HTC 80 and HTC 95 are quite similar, but HTC 120 appears different. This could simply be sample variation; however, HTC 120 was prepared at a lower pressure of about 2 bar, as opposed to up to 20 bar for other points. This might suggest that pressure is an important factor in pore size.



**Figure 19.** Frequency distribution of pore size with temperature for HTC samples.

The elemental composition of the poultry litter biochars is another key parameter in their application as soil amendments. As mentioned above, PL biochars are heterogeneous materials, with complex structure and morphology. Therefore, a heterogeneous composition is expected as well. First, SEM/EDX measurements were used to assess the elemental composition at the surface and subsurface level (about 1 micron depth) of the different biochar's samples prepared in this study. Figure S12 shows the SEM images and EDX results of the TF 200 sample, as an example of the heterogeneous nature of the PL biochar's samples.

The elemental composition of the torrefied samples was measured twice by EDX in a six-month time interval. Low-temperature chars showed little difference; however, at higher temperatures, there are big differences. In TOR 600, carbon has dropped from 70% to 47%, whilst oxygen has gone up from 22% to 28%. The most likely explanation is that the stored chars have taken up water, so oxygen increases. This alters all the percentages, but is most obvious for the largest element, carbon.

This explanation is supported by the hydrophobicity experiments, which showed that the low-temperature biochars take up little water compared to the high-temperature ones.

The mean EDX composition of the TF samples is given in Table 4.

**Table 4.** EDX composition of the TF biochar samples.

Element	Mean (Weight %)			
	Production Temperature (°C)			
	135	200	275	350
C	48.3	44.2	48.4	48.9
O	42.4	44.3	35.5	31.4
Na	0.4	0.5	0.7	0.8
Mg	0.8	1.1	1.4	1.8
P	1.2	1.6	2.0	2.6
S	0.8	0.6	0.9	0.9
Cl	0.7	0.8	1.4	1.4
K	3.8	4.5	6.8	9.0
Ca	1.3	2.0	2.5	2.9
Si	0.2	0.3	0.3	0.3
Mn	ND	ND	ND	0.1

As for the HTC biochar samples, the EDX measurements, shown in Table 5, reveal that carbon decreases slightly and oxygen increases slightly above 80 °C, but there is little change after this temperature. All other elements are present at lower levels and seem to increase slightly with temperature from 80 °C but after this stay relatively constant. This is because the chars lose water and, hence, weight, so the solid material is present at a lower level; thus, elements appear to increase. All minor elements are below 1%, apart from Ca, and so cannot be considered reliable. The tolerance of the system according to Oxford instruments is 0.1% but below 1% the peaks should be checked manually, which was not done. The higher levels of Ca may mean that Ca present on the surface is not water soluble. K shows an increase, but it is not significant and is very small compared to torrefied chars. This is presumably because the K is present as a soluble salt and dissolves into the surrounding HTC liquor.

The bulk composition of the biochar samples prepared in this study was measured by atomic absorption spectroscopy (AAS) and inductively coupled plasma mass spectrometry (ICP-MS), with the purpose to assess the presence of potentially toxic elements (heavy metals) such as Cu, Pb, and As. Both methods were considered. However, as the concentration of the heavy metals in the biochar samples determined by AAS was very low, ICP-MS was then used, as its sensitivity is better. The results of the ICP-MS measurements are shown in Table 6.

**Table 5.** EDX composition of the HTC biochar samples.

Element	Mean (Weight %)		
	Production Temperature (°C)		
	80	120	221
C	58.2	49.8	51.9
O	39.7	46.0	44.3
Na	0.0	0.0	0.0
Mg	0.1	0.2	0.2
P	0.3	0.4	0.4
S	0.4	0.6	0.5
Cl	0.1	0.2	ND
K	0.4	1.0	0.9
Ca	0.4	1.8	1.4
Si	0.1	0.3	0.2
Fe	ND	ND	ND
Mn	ND	ND	0.1
Y	ND	ND	ND
Cu	ND	ND	ND
Al	ND	ND	ND
Br	0.1	ND	ND

**Table 6.** ICP-MS results for TOR and TF biochar samples compared to PL.

Chars	Concentration Minus Blank Sample's Concentration (ppb)									
	Na	Mg	Al	K	Ca	Mn	Cu *	Zn *	Sr	Ba
TOR 350	61.3	95.0	3.1	20.5	106.6	7.0	1.6	7.9	0.4	0.2
TOR 450	24.3	131.8	4.9	0.0	157.5	9.6	2.7	6.5	0.7	0.5
TOR 550	69.5	146.0	4.4	0.0	155.5	12.1	1.3	5.2	0.7	0.5
TOR 600	77.9	138.5	3.8	0.0	177.9	10.6	0.9	3.1	0.7	0.5
TF 200	0.0	45.5	0.0	57.4	0.0	3.9	0.0	0.0	0.0	0.0
TF 275	33.1	98.1	0.0	71.5	121.0	8.0	0.6	3.0	0.3	0.2
TF 350	44.3	136.2	2.9	0.0	129.0	10.3	1.6	4.7	0.6	0.3
PL	-	83.0	4.3	-	-	4.4	1.8	9.0	0.9	0.6

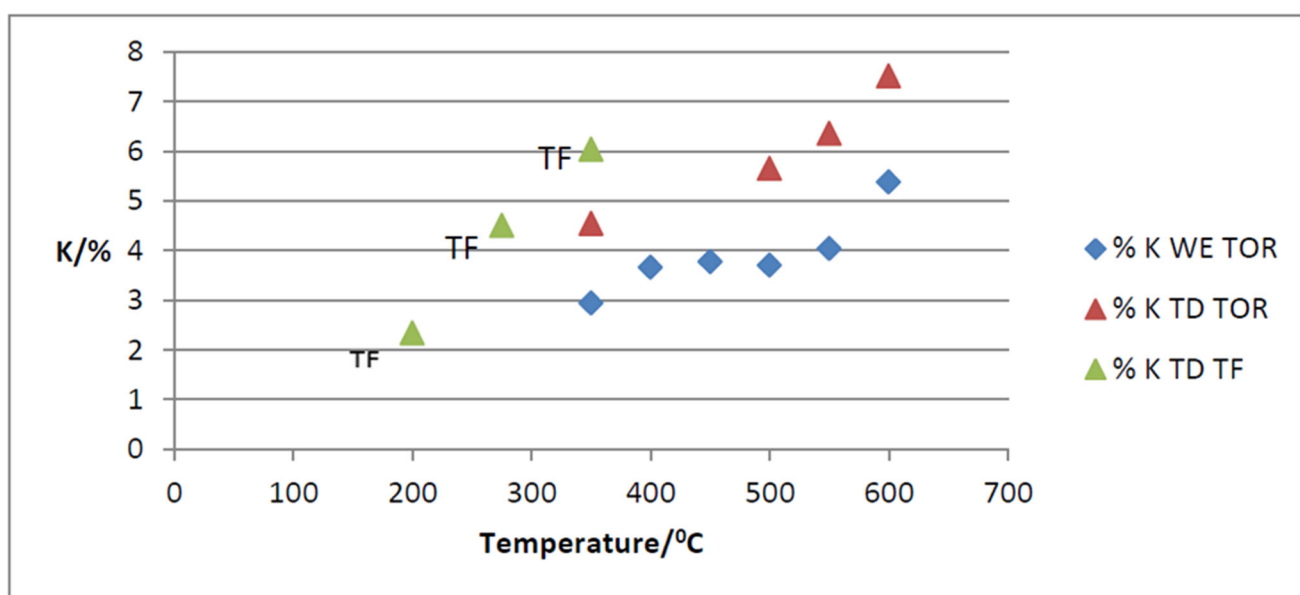
\* The Cu and Zn values reported in the table are the difference between the measured values and the respective total measurement errors.

This study also showed a concentration effect in the char, with double the values in the char compared to the PL [49].

Concentration levels in the low-temperature chars are not much different to those in PL. However, at higher temperatures, levels of magnesium, manganese, and aluminium increase. This is presumably a simple concentration effect. Standards were taken through the whole method to give an indication of both method and machine error. Errors are due to dilutions, variable loss of water during reflux, and metals taken up from anti-bumping granules, nitric acid, and glassware. This gave a total error of about 1% for Cu, 7% for Zn and Ni, 10% for Cd, and 35% for Fe. Levels of most metals are low and are very close to detection limits, so values may not be accurate. As expected, there is good agreement between the 350 °C results for the TF and those for TOR. The highest values were for Ca, Mg, Na, and K, but these are all very susceptible to environmental contamination, and the blank is high for all four. However, K shows a very different trend. The other three elements increase and then level off; presumably this is an artefact caused by the fact that the char weighs less, so the percentage for a given weight is higher. K drops sharply, which may be important and, therefore, will be checked using a flame photometer; Zn, Al, Mn, and Cu, though present at lower concentrations, show the same trend as K.

As KCl crystals have been found on the surface of the char, it is important to know K levels; therefore, the flame photometer was used on both water-extracted and total-digestion samples.

The results are presented in Figure 20. As can be seen, over 60% of the total K in the TOR samples is water extractable (WE). This is important in terms of plant nutrients as water-extractable K is available to plants, but it may also be leached before it can be used. Soluble K may also affect germination as it is mainly on the biochar surface. This was tested using a KCl solution in the germination test. Other studies have found lower levels of water-extractable K of about 46% at 300 °C going up to 49% at 600 °C [10]. A similar trend was observed, with WE % staying fairly constant until 550 °C when it increases, whilst in this study the increase was observed above 550 °C. TF and TOR biochars show an increase in both water-extractable and total K with increased production temperature. The TF 350 value is slightly higher than the TOR 350 value.



**Figure 20.** Water-extractable (WE) and total-digestion (TD) potassium concentrations for TOR and TF samples.

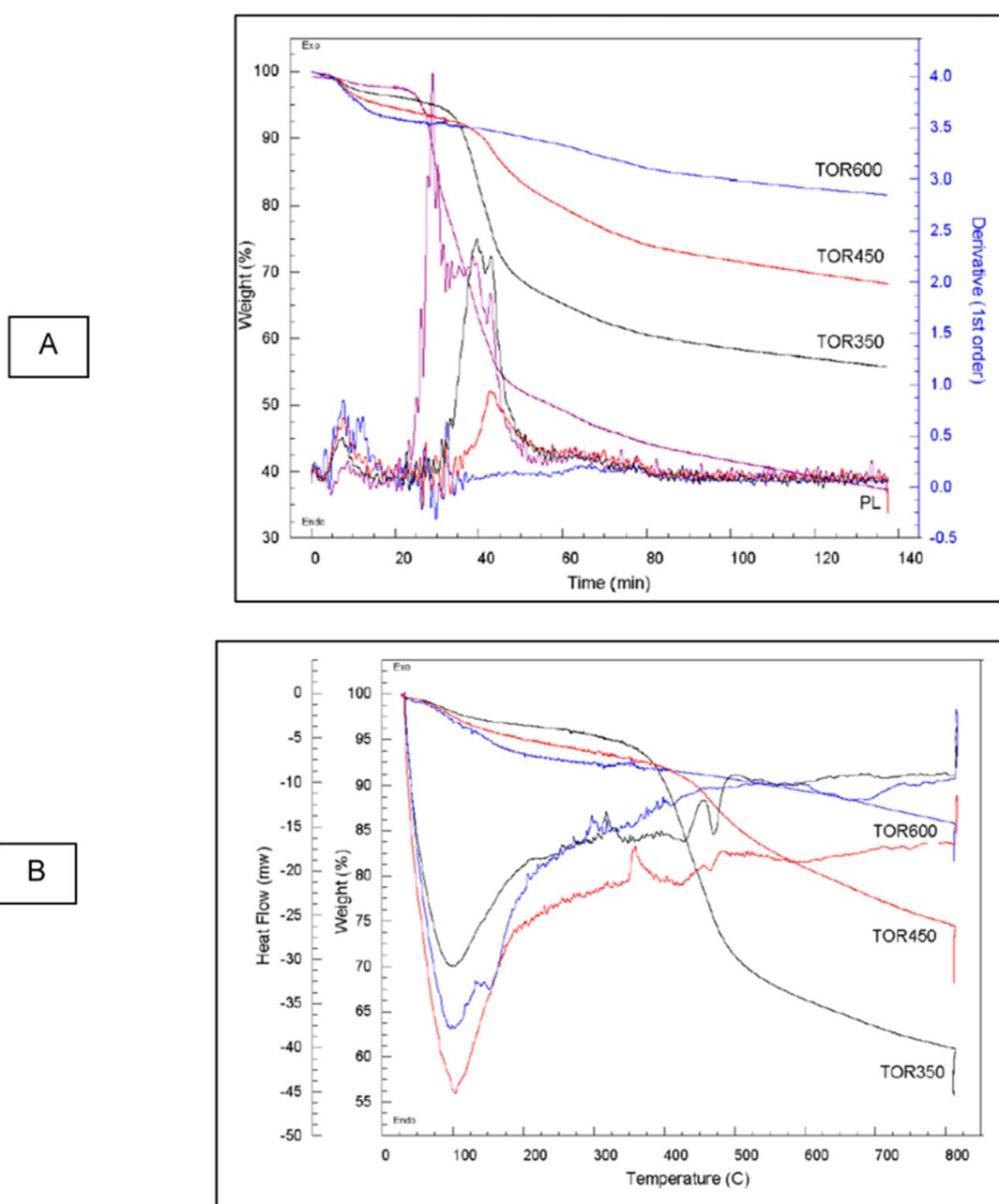
The HTC material has little K, presumably because soluble K has been extracted into the HTC liquor. Flame photometry measurements showed a maximum of 1% K (TD) for the HTC 120, while for the PL, the value was 2.3% or 22.9 mg g<sup>-1</sup>. This value is lower than the published data of 41.8 mg g<sup>-1</sup> for raw PL [10]. Another study measured between 5.65% and 7.59% total K in PL; again, the value measured in this study is lower [50], as the source of the PL is different.

### 3.4. Effect of Production Method and Temperature on Samples' Thermal Stability

As is well known, the conversion of poultry litter, a nutrient-rich organic residue, into biochars is an alternative to stabilize its carbon through conversion to aromatic and recalcitrant forms, because the ability of biochar to retain carbon in relatively stable forms is considered one of its most important properties, either for carbon sequestration or soil conditioner use. However, despite the widespread expectation of high biochar stability in the environment, a considerable portion of its carbon content is labile, and it can be decomposed under its usage conditions [51]. The evaluation of the biochars' carbon stability was done by TGA analysis under an inert (Ar) environment at 800 °C.

Figure 21 shows the TGA curves for PL and three of the TOR samples, i.e., 350, 450, and 600, respectively. There is a slight loss in weight at a consistent rate for all chars up to 150 °C due to water loss. Although the chars and PL had previously been heated, so

should have been dry, they were stored for 6 months, so would have taken up water. It is noticeable that TOR 600 lost the most weight of the chars in this initial phase, presumably because higher-temperature chars have a larger surface area and are less hydrophobic, so take up more water whilst stored. The second weight loss of volatile substances (organic carbon) occurred at different temperatures for the different samples. It started at about 280 °C for PL, 370 °C for TOR 350, and 430 °C for TOR 450—all fitting well with the fact that these chars had already lost volatiles up to their torrefaction temperature. There is no steep drop for TOR 600, just a smooth curve, showing it had already lost most volatiles. TOR 350 lost the most weight of the TORS, as would be expected, as it had not lost as many volatiles due to torrefaction. The last section indicates the further weight loss during the 60 min residence time. The final weight value would be expected to be residual ash.



**Figure 21.** Thermogravimetric analysis (TGA) under N<sub>2</sub> atmosphere of PL and TOR 350, TOR 450, and TOR 600 sample, respectively: (A) thermograms and first derivatives of the thermograms; (B) heat flow.

As for the first derivatives of thermograms, the first small peak at 100 °C is due to water loss [52]. Then, the derivative shows a major exothermic reaction starting at 300 °C for TOR 350 and a lower peak at 430 °C for TOR 450. TOR 600 does not show this peak, indicating that this component had already been lost during torrefaction. The PL peak starts at about 210 °C, and it is higher than the TOR 350 peak. The peak for TOR 350 may be a double peak, with the second peak coinciding with the exothermic heat flow peak at 450 °C. The peaks are caused by the thermal degradation of hemicellulose, cellulose, and lignin. The 300 °C peak is hemicellulose, the 350 °C peak is cellulose, and lignin peaks at 450 °C [52]. Chen and Kuo [53] studied the thermal degradation of pure hemicellulose, cellulose, lignin, xylose, and glucose. They found that the torrefaction of hemicellulose at 230 °C gave an exothermic peak. Hemicellulose contains oxygen, and has a general formula of  $(C_5H_8O_4)_n$ —this oxygen is used in thermal degradation. At temperatures below about 230 °C, some hemicellulose in the chars will have been broken down during the torrefaction process; at higher temperatures, above 310 °C, the hemicellulose is totally broken down. For cellulose, total breakdown occurs at 361 °C. Lignin pyrolysis is an exothermic reaction and occurs at about 400 °C. The peak for PL starting at 210 °C is hemicellulose. For TOR 350, hemicellulose will have broken down, so this peak is not seen, but cellulose and lignin have not broken down. Consequently, when the char was reheated in TGA, the weight loss was due to the decomposition of cellulose and lignin. For TOR 450, all components should have been lost and further weight loss should be due to inorganic components, such as carbonates. However, there was a larger drop for TOR 450 at a higher rate than for TOR 600, suggesting that not all organic components had been lost. This suggests that the components left in the different temperature chars are different. Their properties as fuels will, therefore, be different, as shown by the decrease in calorific value (CV) with increased production temperature. Xylan is a component of hemicellulose, and it shows a decrease in weight prior to 100 °C, so it may be that the drop in weight previously attributed to water might be due to xylan. However, this would not fit with TOR 600 losing the most weight, as it will have the least xylan, since the pyrolysis temperature for xylan is 300 °C, so it should not be present in the TOR 600.

The same decomposition behaviour was observed for the TF samples.

Figure 22 presents the TGA curves for TOR 350 in comparison with TF 350. As can be seen, the TF 350 results agree well with those obtained for TOR 350; this would be expected as they are produced at the same temperature. Heat flows and derivatives are very similar; however, TOR 350 shows a higher peak, which starts slightly earlier. Also, TF loses more weight at above 100 °C and then drops more gradually, so, by the end of the process, the weight loss is very similar.

HTC biochar samples showed a different behaviour, although the feedstock decomposition is dominated by reaction mechanisms similar to those in dry pyrolysis, which include hydrolysis, dehydration, decarboxylation, aromatization, and recondensation [33]. However, the hydrothermal degradation of biomass is initiated by hydrolysis, which exhibits a lower activation energy than most of the pyrolytic decomposition reactions.

As seen in Figure 23, the weight loss seems to be more gradual than for the torrefied samples, with less of a distinct step change. The very broad derivative peak explains why there is not such an obvious step change.

Therefore, the principal biomass components are less stable under hydrothermal conditions, which leads to lower decomposition temperatures. Hemicellulose decomposes between 180 and 200 °C, most of the lignins between 180 and 220 °C, and cellulose above approximately 220 °C. So, the bulk of the polymer left will be cellulose [33]. The 400 °C peak is presumably lignin that has not yet decomposed, but the broad peak up to 300 °C is cellulose. There is less hemicellulose present.



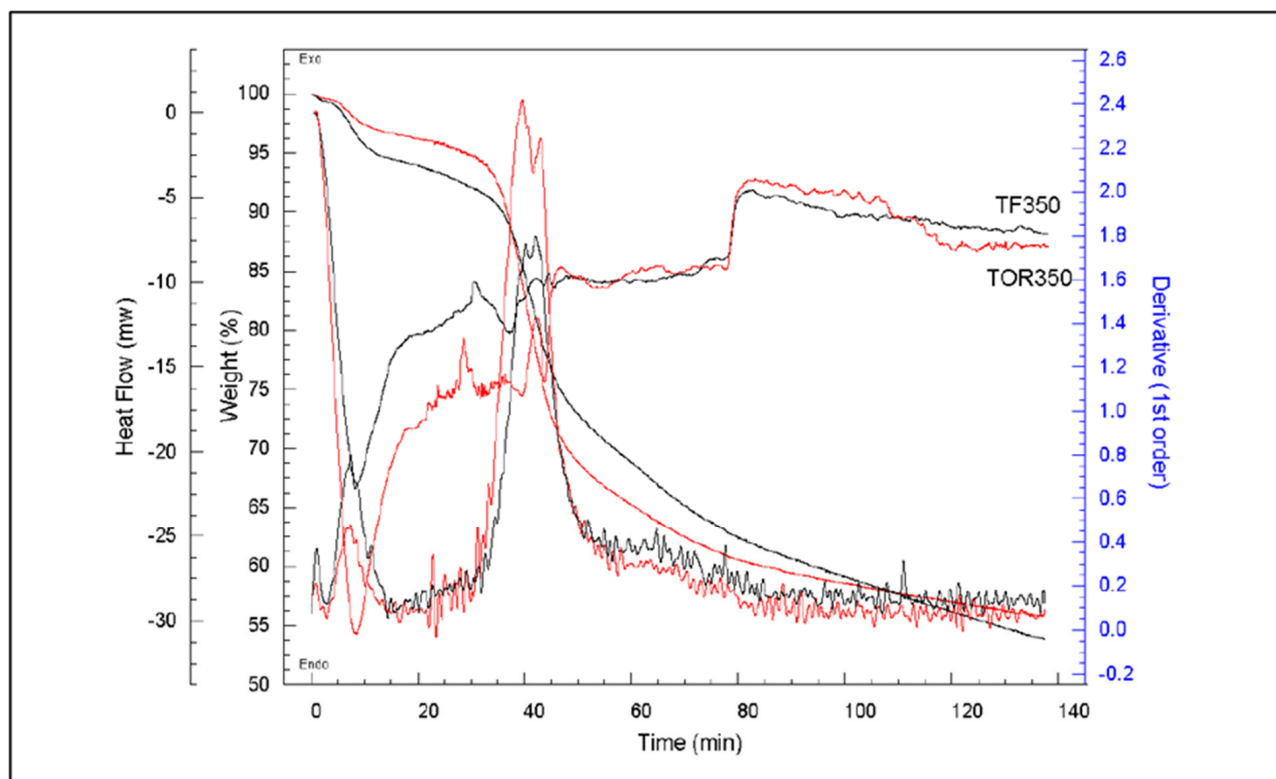


Figure 22. TGA curves (TG, DTG, heat flow) for TF 350 and TOR 350 samples.

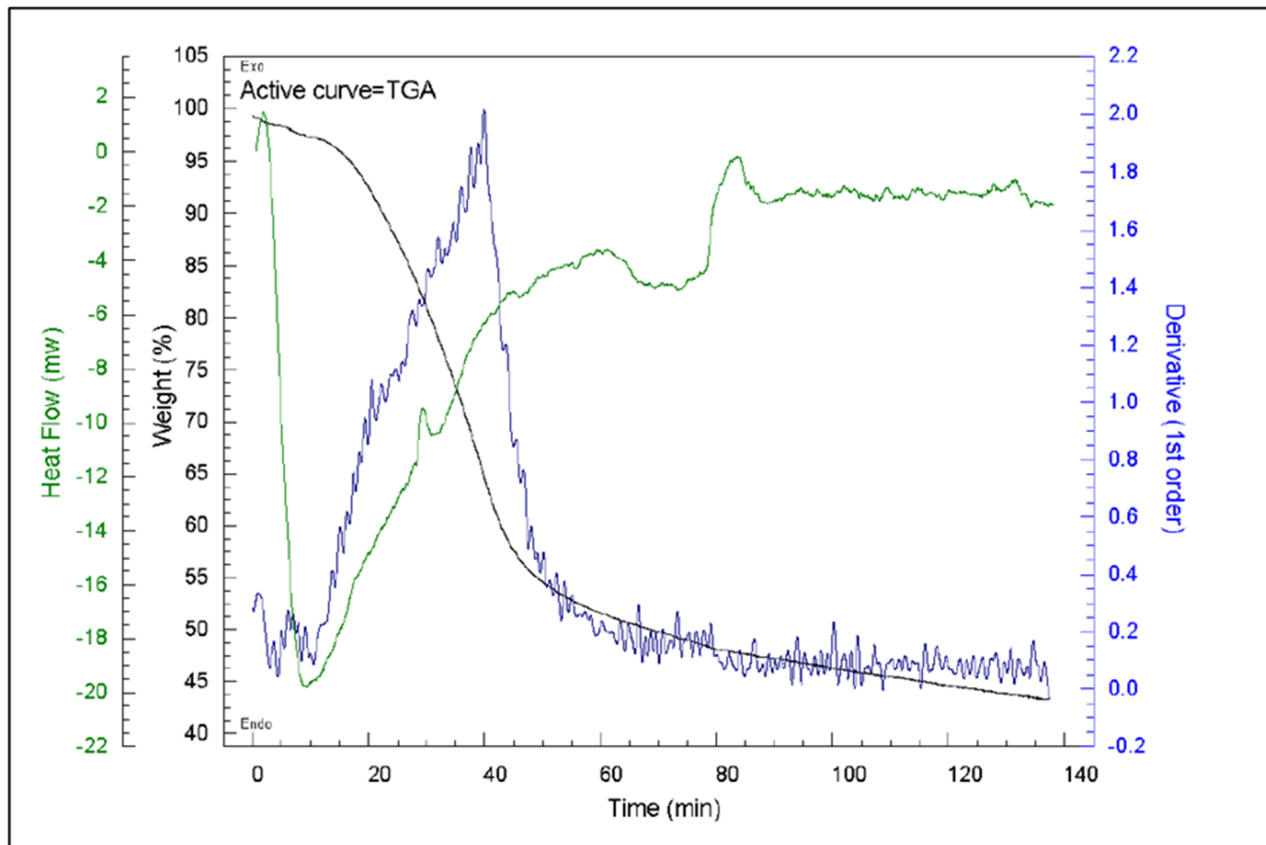


Figure 23. TGA curves (TG, DTG, heat flow) for HTC 216 biochar sample.

It is possible to attach an MS to a TGA device to see what gases are evolved at the different temperatures [52]. The results for PL suggest that the main gas lost at 375 °C is water, so dehydration reactions are occurring, whilst the second peak at 450 °C coincides with CO<sub>2</sub>, so suggests higher losses of hydrocarbons (C<sub>2</sub>H<sub>2</sub>+, C<sub>2</sub>H<sub>3</sub>+) and aldehydes. This gives more information about the reactions occurring and the gases being lost during torrefaction.

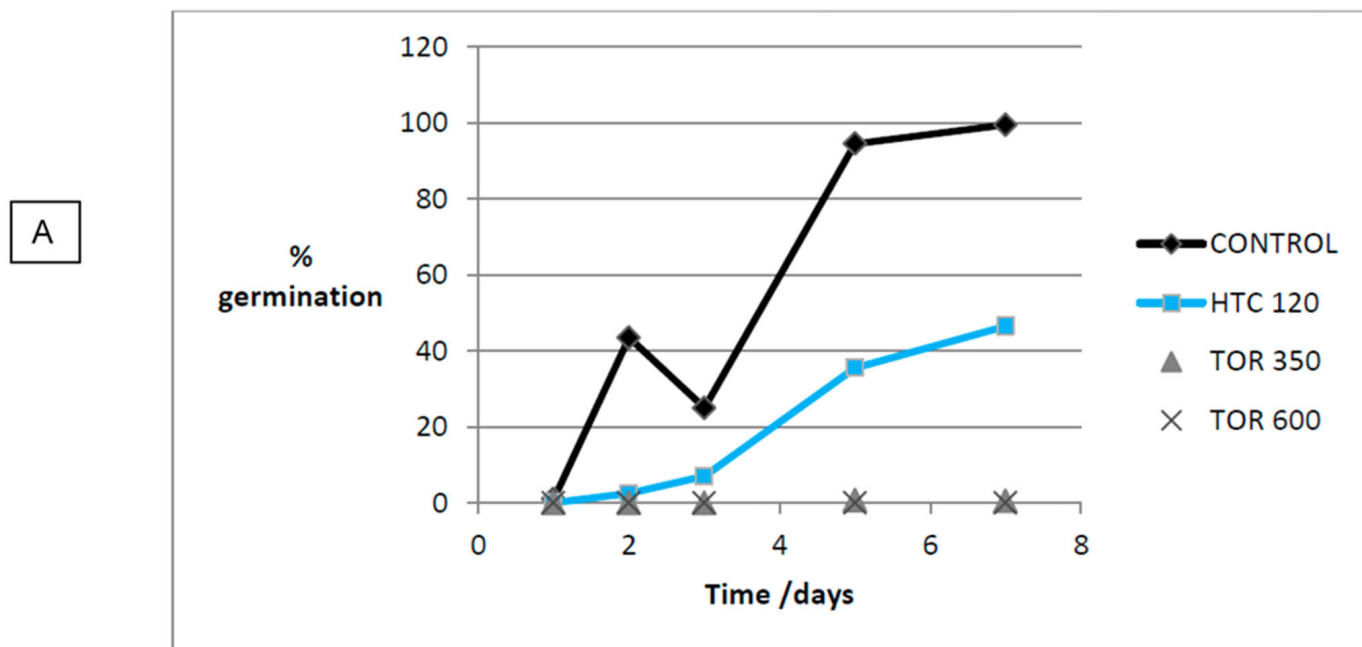
The weight losses for the TOR, TF, and HTC samples measured in this study are presented in Table 7.

**Table 7.** TGA results for PL, TOR 350, TOR 450, TOR 600, TF 275, TF 350, and HTC 216 samples, respectively.

Sample	Weight Remaining at 800 °C (Weight %)	Weight Remaining after 60 Min Run at 800 °C (Weight %)	Loss Over 60 Min Residence Time at 800 °C (Weight %)	Residual Ash (Weight %) after 60 Min Run at 800 °C	Loss of Volatiles at 400 °C (Weight %)
PL	26.0	22.0	4.0	22.0	55.0
TOR 350	62.0	56.0	6.0	56.0	10.0
TOR 450	76.0	69.0	7.0	69.0	8.0
TOR 600	87.5	82.5	5.0	82.5	8.0
TF 275	45.0	37.5	7.5	37.5	30.0
TF 350	64.0	55.0	9.0	55.0	10.0
HTC 216	50.0	37.5	7.5	37.5	28.0

### 3.5. Effect of Production Method and Temperature on Germination

Germination was tested in two separate experiments. The data cannot be amalgamated as these experiments were not carried out in controlled environmental conditions. Consequently, germination must be compared to the relevant control (Figures 24 and 25).



**Figure 24.** Cont.

B

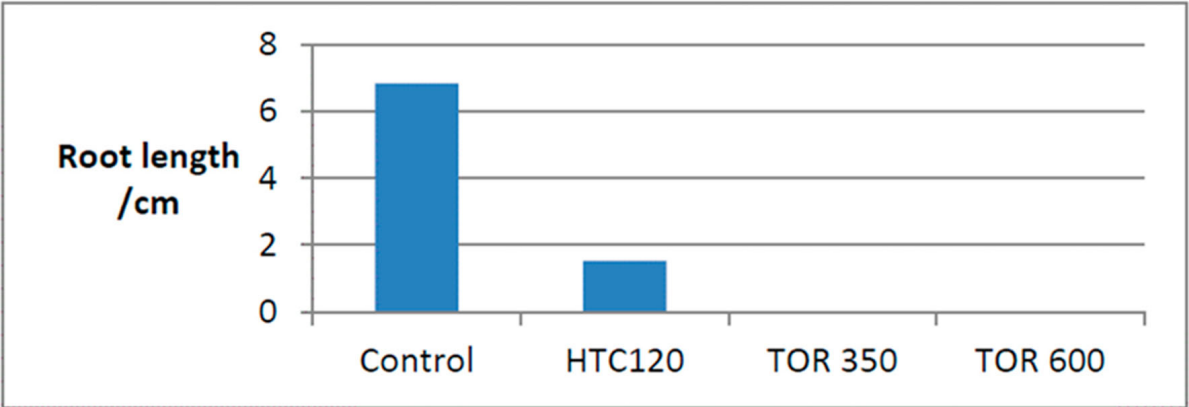
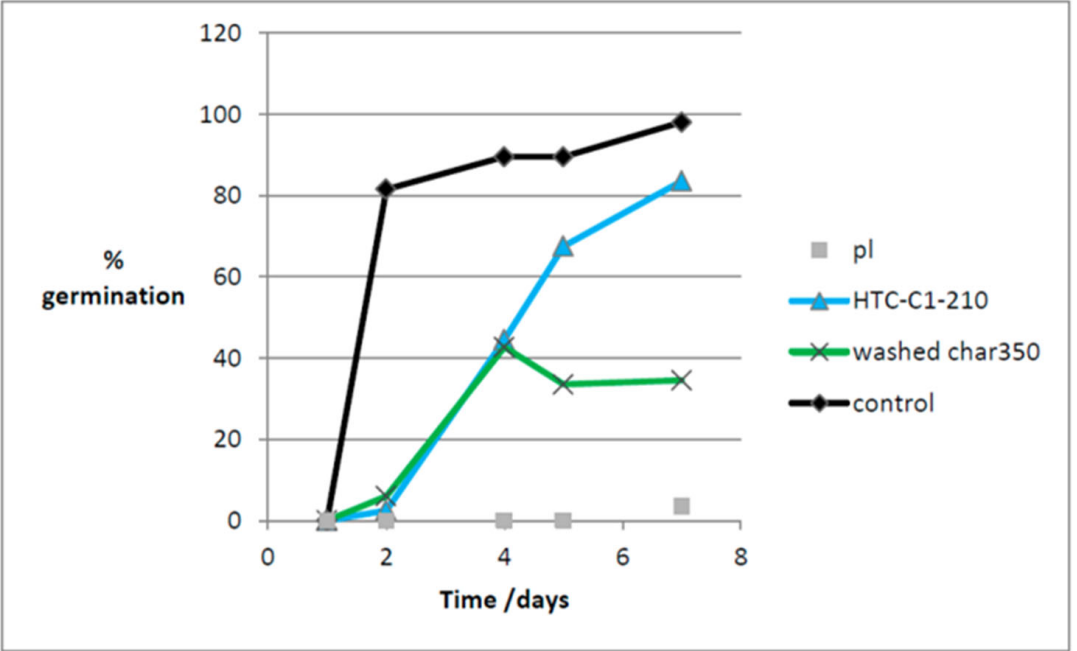


Figure 24. Mean germination percentage over time (A) and root length at 7 days (B) for experiment 1.

A



B

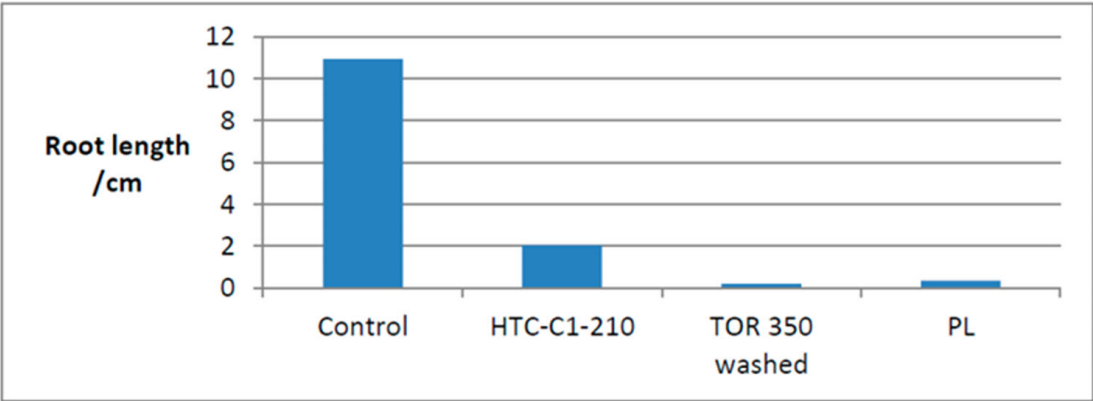


Figure 25. Cont.

C



**Figure 25.** Mean germination percentage (A), root length (B), and germination on PL on day 7, (C) for experiment 2.

In both experiments, germination was poor in TOR samples, and although HTC samples showed germination, root growth seemed abnormal. In experiment 2, washed TOR 350 char was used, and this provoked some germination (34%), suggesting that washing possibly removed some inhibition.

All the germination results can be compared using a germination index (GI) as shown in Table 8.

**Table 8.** Germination results compared by using germination index (GI) at day 7.

Substrate	Germination (%)	Germination (% of Control)	Mean Root Length (cm)	Mean Root Length (% of Control)	GI
Control 1	99.5		6.8		
Poultry litter	3.5	3.5	0.3	3.2	0.1
TOR 350	0.5	0.1	0	7.3	0.01
TOR 600	0	0	0	0	0
Control 2	98.0		11.0		
Washed TOR 350	34.5	32.2	0.2	1.8	0.6
HTC 120	46.5	46.7	1.5	22.4	10.5
HTC-C1-210	83.5	85.2	0.3	18.6	15.8
Extract					
KCl	0.0	0.0	0.0	0.0	0.0
Washed TOR 350	99.3	101.4	3.7	33.8	34.3
HTC 250	0.0	0.0	0.0	0.0	0.0
HTC 87	89.0	90.8	0.3	2.6	2.4

The literature suggests that biochars do not inhibit germination, but that hydrochars do inhibit germination [54]. These results appear to contradict this, with germination occurring in hydrochars but not in biochars. However, the high germination index in HTC 120 can be explained by the fact that it was a low-temperature hydrochar that had been washed. This washing should remove soluble substances and some toxic oils from the surface, allowing germination. Initial seed germination is mainly dependent on water

availability for imbibition, and the hydrochar showed good water retention, allowing good germination. However, after initial germination, substrate toxicity was demonstrated as reduced root growth showed that toxic substances still remained, even after washing.

Germination in the TOR samples was almost completely inhibited, probably due to a lack of water for imbibition by the seeds. As water was physically present, a lack of water for imbibition could be due to two factors. Firstly, the lower-temperature chars are very hydrophobic, and when water was added, it formed discrete bubbles on the surface, meaning the water did not disperse; consequently, only some seeds had access to water. However, hydrophobicity is more pronounced in low-temperature chars, whereas the high-temperature TOR 600 showed poorer germination; indicating that toxic compounds might be more important. Secondly, physiological drought can occur, i.e., water is present, but the concentration of soluble salts is so high that osmosis cannot occur into the seeds to allow germination. It is thought that the total lack of germination was due to toxic salts and physiological drought.

It is possible that soluble salts, such as KCl, could be inhibiting germination by either of these mechanisms. EDX showed about 10% of K present on the surface of chars, probably as KCl. So, a 10% solution of KCl was used in an extra experiment along with a TOR 350 wash extract. The KCl completely inhibited germination.

The total lack of germination in the high-temperature HTC extract and low GI for the low-temperature liquor also supported the toxicity of the HTC extracts. This liquor toxicity could be due to a number of factors. The liquor is acid (the pH for HTC 87 is 3.27, while for HTC 250 it is 4.57), which may be a problem because pH affects germination. Acidity also increases the solubility of many metals, which can also cause germination inhibition, although levels of heavy metals (as shown by AAS) were generally low. There is also a complex mix of organics including phenols and PAHs in HTC liquor. TOR 350 and HTC 87 extracts showed good germination at almost the same rate as the control, although root growth was decreased in both, with HTC 87 showing almost no root growth. Roots were so short they were difficult to measure.

In summary, biochars totally inhibit germination, whilst hydrochars slightly inhibit germination. However, after germination, hydrochars inhibit root growth. Washing of both biochars and hydrochars reduces germination inhibition, but root inhibition remains. It is suggested that the initial germination inhibition in biochars is due to the presence of soluble salts such as KCl. Root inhibition is probably due to insoluble organic compounds deposited on the char surface. Surprisingly, PL also showed almost total germination inhibition. The fact that PL is used successfully as a fertiliser suggests that lower concentrations do not adversely affect plant growth; possibly, the chars could be used as soil amendments if used in low concentrations.

#### **4. Preliminary Kinetic Analysis of Poultry Litter's Thermochemical Conversion via Torrefaction**

Kinetic studies of the thermal decomposition of poultry litter are a matter of major interest for investigating its use for energy.

Kinetic studies allow a better understanding of the mechanism and provide the basis for the most appropriate reactor design.

As the SEM micrographs show, the poultry litter used in this study is a mixture of poultry manure, feathers, spilled feed, and bedding materials, which increased the complexity of the torrefaction process. The main components of poultry litter are lignin, cellulose, and hemicellulose (as in any other biomass) plus protein.

TGA analysis evidenced that a high temperature starts to break the bonds of groups bound to carbon, as carbon polymers such as hemicellulose, cellulose, and lignin are broken down. Figure 26 below shows the varied reactions involved in the torrefaction of the lignin present in the poultry litter.

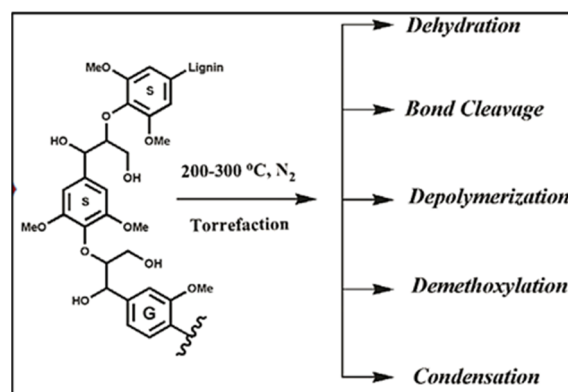


Figure 26. Lignin torrefaction reactions. Reprinted with permission from [42], Elsevier Ltd.

Several methods of studying torrefaction kinetics have been postulated by different researchers. The methods are classified as model-fitting and model-free methods, respectively. In model-fitting methods, different types of kinetic models are fitted into experimental data until the best statistical fit is obtained. Kinetic parameters are then calculated based on the best fit model.

The Coats–Redfern method is the most popular model-fitting technique [55]. In contrast, model-free methods allow the estimation of kinetic parameters without a priori knowledge of the kinetic model by utilising TGA-measured data generated at different heating rates. Common examples of model-free methods include Kissinger–Akahira–Sunose (KAS), Ozawa–Flynn–Wall (OFW), Friedman, Starink, and Distributed Activation Energy [56]. These methods are also called iso-conversional because they use the iso-conversional principle to analyse the experimental data. The iso-conversional principle states that the reaction rate at a constant extent of conversion is only a function of temperature. Although the model-free methods are relatively simple, they are sensitive to the accuracy and precision of the analytical techniques employed.

As TGA experiments were performed at only one heating rate, i.e.,  $10\text{ }^{\circ}\text{C min}^{-1}$ , the iso-conversional methods were not appropriate to be used for the kinetic parameters' determination. Instead, the Coats–Redfern method was used in this study. The method is described in detail in [42], while the application of this method to poultry litter pyrolysis can be found in [57].

The TGA data of PL are presented in Figure 21A.

From the recorded mass loss curve (mass loss versus temperature), the reaction conversion curve (the extent of the reaction versus temperature) was obtained easily.

Briefly, the method consists of the following steps [58].

The pyrolytic conversions ( $Y$ ) for each temperature were determined using Equation (1).

$$Y = \frac{w_i - w}{w_i - w_f} \quad (1)$$

where  $w$  represents the mass loss; initial ( $i$ ), at a certain temperature, and final ( $f$ ), respectively.

The pyrolytic decomposition of chicken waste in the reactor can be modelled as a function of conversion as follows.

$$\frac{dY}{dt} = kF(Y) \quad (2)$$

The conversion function can be written as follows.

$$F(Y) = (1 - Y)^n \quad (3)$$

where  $n$  is the reaction order.



By substituting Equation (3) and the general form of the Arrhenius equation into Equation (2), we obtain

$$\frac{dY}{dt} = A \exp\left(\frac{-E_a}{RT}\right) (1 - Y)^n \quad (4)$$

If the reactor heating rate  $\beta$  ( $^{\circ}\text{C}/\text{min}$ ) given by Equation (5) is constant, we can make use of Equation (4) to obtain Equation (6).

$$\beta = \frac{dT}{dt} \quad (5)$$

$$\frac{dY}{dt} = \left(\frac{A}{\beta}\right) \exp\left(\frac{-E_a}{RT}\right) (1 - Y)^n \quad (6)$$

The Coats–Redfern method is a model fitting that requires the integration of Equation (6). As a result, we obtain the following expressions in Equations (7) and (8), for  $n = 1$  and  $n = 2$ , respectively.

$$\ln\left[\frac{-\ln(1 - Y)}{T^2}\right] = \ln\frac{AR}{\beta E_a}\left[1 - \frac{2RT}{E_a}\right] - \frac{E_a}{RT} \quad (7)$$

$$\ln\left[\frac{(1 - Y)^{-1} - 1}{T^2}\right] = \ln\frac{AR}{\beta E_a}\left[1 - \frac{2RT}{E_a}\right] - \frac{E_a}{RT} \quad (8)$$

where  $T$  is the absolute temperature (K),  $A$  is the pre-exponent factor in the Arrhenius equation,  $R$  is the universal gas constant ( $0.0083 \text{ kJ}/(\text{mol} \cdot ^{\circ}\text{C})$ ), and  $E_a$  is the activation energy ( $\text{kJ}/\text{mol}$ ).

The pyrolysis kinetic parameters, that is, the activation energy and the pre-exponential factor, can then be evaluated by linearising Equations (7) and (8) and determining the slope and y-intercept of the resultant plots. Furthermore, the rate constant ( $k$ ) can be calculated by Equation (9). In this case,  $T$  is taken as the median value of the selected temperature range.

$$k = A e^{\frac{-E_a}{RT}} \quad (9)$$

Following the above steps, the apparent activation energy was  $112 \text{ kJ}/\text{mol}$ , assuming that the pyrolysis process follows first-order kinetics, and  $153 \text{ kJ}/\text{mol}$ , assuming second-order kinetics, which is in good agreement with the literature data.

## 5. Conclusions

The aim of this study was to investigate the effect of temperature and treatment on the morphology (most and foremost), structure, and composition of PL chars. It has succeeded, as an almost comprehensive overview of the most important parameters in PL chars has been produced, which does allow the linking of various parameters to char functions and will allow recommendations to be made as to the best production methods for a particular application. It has also discovered several interesting new facts in certain areas, although further research will be needed to elucidate the exact importance of these discoveries.

The main new discoveries are the presence of sylvite (KCl) crystals on the surface of TOR samples; the variation in pore structure leading to the significant variation in pore size at a given temperature; the inhibition of germination in TOR biochars; and the suppression of growth in HTC hydrochars. The hydrophobicity differences between different temperatures of TOR chars have also been demonstrated.

Basic tests were performed on the biochars. Some of these parameters measured are dependent on the feedstock, which was also analysed in this study. The moisture content of the PL used in this paper was 30%. Biochar pH, residual ash, and weight loss increased with increasing temperature. Higher-temperature chars have been shown to be less hydrophobic and have a larger surface area for adsorption of water. This is important in char storage and may also affect analysis.

In terms of morphology, the major characteristic of the biochars is their heterogeneity. Many distinct structures were observed and identified in PL using the SEM technique. These included wheat and barley leaves, with round siliceous structures on their surface, roots, feathers, and phytoliths. These phytoliths are very resistant and remain in ash. Phytoliths can be a serious problem when chars are used as fuels [37]. As well as these discrete multicellular structures, there is obvious cellular structure present in the plant material. Gross and cellular structures found in the PL morphology remain in the biochars. However, the structure of these areas changes as the cell wall materials are converted to carbonised material to produce the characteristic honeycomb structures in the chars.

These honeycomb structures are the most obvious characteristic of the biochars and are found at all temperatures. The pore sizes of these honeycombs and other structures containing pores were investigated. Pore size decreases with temperature because of the shrinkage caused by the heating. The bulk of the macropores in biochars are derived from the cellular structure of the plant material, and the honeycomb structure of pores of about 10 µm is the origin of most of the macropores in biochars [2]. Certainly, this study showed the massive, interconnected network in some biochars that could not be measured simply by pore size; in fact, it is difficult to even define pore size in this context.

One of the most interesting findings in this study was the presence of large numbers of crystals on the surface of the chars. These crystals had a regular structure, although size was very variable. The most obvious contained large amounts of K and Cl, so were probably KCl. This was confirmed by the presence of sylvite (KCl) in the XRD results. These crystals help to explain other properties of the chars, including some of the germination inhibition and the water extractability of K from the biochars. There were many other crystals present, and those confirmed by XRD include calcite and quartz.

In terms of composition, the elemental composition is mainly C and O, as seen by EDX. EDX cannot detect lighter elements, so there are no data for hydrogen, which means the H/C ratio could not be determined. Neither does EDX detect nitrogen, which is a major omission if the char is to be used as a soil amendment because nitrogen is a major plant nutrient. In the case of minor elements detected, there was mostly K, with about 10% at high temperatures, followed by Ca 5%, P 3%, Mg 2%, Cl 3%, and Na 1%. Many temperature effects are simply concentration effects caused by mass loss; thus, all elements appear to increase with increased torrefaction temperature. However, K levels increased dramatically after 400 °C. This is because the K is present as crystals on the surface and, therefore, will show high levels using EDX, which only samples the top few microns of the surface. These crystals have a major effect on biochars used as soil amendments.

ICP-MS shows a similar trend, with levels in the biochars not much different to those in PL for low-temperature biochars.

In TGA, as oven-dried material was used, weight loss above 100 °C was due initially to further water evaporation and then to the loss of volatiles. The high temperature started to break the bonds of groups bound to carbon, as carbon polymers such as hemicellulose, cellulose, and lignin were broken down and the products were lost as gases and oils. At 260 °C, 38% of hemicellulose weight was lost; by 290 °C, 58%. Most cellulose had degraded by 360 °C; lignin degrades at higher temperatures. Thus, in terms of lignocellulose content (sawdust, wood shavings, straw) most weight loss by biochars in TGA will be due to breakdown of lignin. Lower-temperature chars will also have some cellulose and so will lose more weight than higher-temperature chars. Inorganics such as carbonates start to break down above 450 °C and are lost. In high-temperature biochars such as TOR 600, only small quantities of lignin and inorganics should be left to cause weight decreases, so weight loss was small at 12.5%.

The germination results were not what would be expected from the literature, where it is stated that biochars do not significantly affect germination [9]. In this study, biochars showed almost total germination inhibition, which needs explanation. As seen in other parts of the study, volatile compounds are lost during torrefaction. When cellulose and other compounds break down, they form low-molecular-weight compounds such as organic

acids, alcohols, phenols, and polyaromatic hydrocarbons (PAHs). These can recondense on the surface of the char and can be toxic [59]. These volatile organic compounds (VOCs) are not important if the char is to be used as a fuel but become potentially very important if the char is to be used as a soil amendment.

The most likely factor causing germination inhibition is an osmotic effect as initial germination is dependent on water availability. The presence of large amounts of soluble salts including KCl crystals on the surface of the biochar, as seen in the SEM study, would cause a major osmotic effect. This could inhibit germination by reducing the imbibition of water. As seen here, washing the soluble salts off the biochar increased germination, but then another toxicity factor became important, with the germinated seeds showing severe root inhibition. So, germination inhibition is not removed by thirty minutes of washing, although further soluble toxic compounds could be removed by additional washing.

Hydrochar samples are markedly different to biochars in gross properties. They are softer, lighter in colour, and smear rather than fragment when pulverised. Morphologically, hydrochar is fairly similar to biochar, with the same gross morphology, though having more fibrous structures visible and more cell wall debris on the hydrochar surface. Pore size showed little variation with temperature; however, there were only a small number of samples, and the temperature range was not large. The mean pore size for low-temperature hydrochars is 10  $\mu\text{m}$ —smaller than that for the lowest-temperature biochar, which was 14  $\mu\text{m}$ . The pore structure is also different; true honeycombs are found in HTC120, but lower-temperature honeycombs could be considered different: there are no true pores as there is no cavity in the centre. It is possible that the pores are in the process of forming, or this too could be connected to the different decomposition sequence of cell wall components. There are some crystalline structures present, but they seem small and diffuse, unlike the defined structures in the chars, presumably because many crystals are soluble and dissolve in the liquor.

Hydrochars are acidic, in this case because citric acid was used as the liquid medium. However, pH rises with production temperature, so high-temperature hydrochars are pH 4 and above. This is still quite acidic and could have detrimental effects on plant growth. EDX shows far lower levels of K in hydrochar, presumably because the soluble K salts are in the liquor. All elements appear to increase slightly with temperature, presumably due to the fact that the hydrochars lose weight. The only element found by EDX on the surface was Ca, possibly because it is present as an insoluble salt such as calcium carbonate ( $\text{CaCO}_3$ ). Total K values in the HTC material are very low, presumably because the K is in the liquor.

The TGA for HTC char samples shows a different curve to torrefied samples. The curve is far smoother, with no distinct step changes. There is a very broad derivative peak, which starts earlier, but at a lower level, which suggests that there is less cellulose and hemicellulose and fits with the fact that HTC breaks down hemicellulose at low temperatures. The residual ash's value, at 46%, was higher than the value for TF 275 (37.5%) but lower than that for TOR 350 (56%), so it was relatively high considering it was produced at a lower temperature (216 °C).

**Supplementary Materials:** The following supporting information can be downloaded at <https://www.mdpi.com/article/10.3390/reactions5030020/s1>, Figure S1. SEM/EDX results from wheat samples; Figure S2. SEM/EDX results from barley leaves; Figure S3. SEM of barley leaves showing phytolith structures; Figure S4. SEM images: (left) feathers from TF 135 (its provenience is PL); (right) actual feather; Figure S5. SEM/EDX results from pure KCl; Figure S6. SEM images of TOR 400 sample showing complex macroporosity: (A) pores formed by the open ends of cells; (B) size range of pores compared to crystal size; Figure S7. SEM images of TOR 450 sample showing complex macroporosity: (A) pore size; (B) pore size at 2 x higher magnification; Figure S8. SEM images of TOR 500 sample showing complex macroporosity: (A) true honeycomb porous structure with mean pore size of 7.9  $\mu\text{m}$ ; (B) non-empty pores; Figure S9. SEM images of TOR 550 sample showing complex microporosity; Figure S10. SEM images of TOR 600 sample showing complex macroporosity: (A) pore size measurements; (B) true honeycomb of the cellular structure; Figure S11. SEM images

and pore size measurements of the TF 135 biochar sample; Figure S12. SEM images and pore size measurements for the TF 200 sample: (A) pores with appearance and sizes different from TF 135 sample; (B) pores with similar appearance and sizes similar to TF 135 sample; Figure S13. SEM images and pore size measurements for the TF 275 sample; Figure S14. SEM images and pore size measurements for the TF 350 sample; Figure S15. SEM images and pore size measurements for the HTC 80 sample; Figure S16. SEM images and pore size measurements for the HTC 95 sample; Figure S17. SEM images and pore size measurements for the HTC 120 sample: (A) open pores; (B) pores in the process of forming; Figure S18. SEM images and pore size measurements for the HTC 221 sample.

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