Influence of Torrefaction and Pelletizing of Sawdust on the Design Parameters of a Fixed Bed Gasifier

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Received: 1 May 2020; Accepted: 8 June 2020; Published: 11 June 2020

Abstract: Gasification of biomass in fixed bed gasifiers is a well-known technology, with its origins dating back to the beginning of 20th century. It is a technology with good prospects, in terms of small scale, decentralized power co-generation. However, the understanding of the process is still not fully developed. Therefore, assessment of the changes in the design of a gasifier is typically performed with extensive prototyping stage, thus introducing significant cost. This study presents experimental results of gasification of a single pellet and bed of particles of raw and torrefied wood. The procedure can be used for obtaining design parameters of a fixed bed gasifier. Results of two suits of experiments, namely pyrolysis and CO\textsubscript{2} gasification are presented. Moreover, results of pyrolysis of pellets are compared against a numerical model, developed for thermally thick particles. Pyrolysis time, predicted by model, was in good agreement with experimental results, despite some differences in the time when half of the initial mass was converted. Conversion times for CO\textsubscript{2} gasification were much longer, despite higher temperature of the process, indicating importance of the reduction reactions. Overall, the obtained results could be helpful in developing a complete model of gasification of thermally thick particles in a fixed bed.

Keywords: torrefaction; pyrolysis; gasification; fixed bed; thermally thick particle; modelling

1. Introduction

Nowadays the importance of renewable energy sources, such as biomass, is increasing due to the increased international pressure to gradually phase out the fossil fuels. Biomass is the one of non-intermittent, controllable and potentially flexible sources [1] and it can be used with sensibly low thermo-ecologic cost [2,3].

Torrefaction is a process of thermal valorization of solid fuel in order to enhance its fuel properties subsequently allowing low-grade fuels, such as biomass, to become a tradable commodity [4]. It is a process that is considered to improve the logistics and handling of the upgraded fuel [5–10] and make it more suitable for final use, such as co-firing [11–13] or gasification [14–16]. Torrefaction is
sometimes called slow pyrolysis and it typically takes place at temperatures between 250 °C and 300 °C with residence times ranging between 10 min and 60 min [4,17–26]. During torrefaction, part of the mass of the feedstock is removed. Some literature sources treat torrefaction as mild pyrolysis, which is a thermal decomposition of the organic material under the absence of oxygen [27]. Depending on the parameters of the process, pyrolysis may result in different yields of char, liquids (oils) and gases [28,29]. For lignocellulosic material reactions proceed, starting with a set of primary reactions, among which one can name char formation, depolymerization and fragmentation [30]. Released volatile organic compounds are typically unstable and are a subject of secondary reactions such as cracking, recombination or re-polymerization [30], with the latter taking place even in the pyrolysis oil [28].

Gasification is a process that converts solid fuel to gas [31,32]. Air and steam are mostly used as a gasifying agents [33]. The gas, produced by a gasifier (sometimes called producer gas or more often syngas), consists mainly of CO, H₂, CO₂, H₂O (vapor), with smaller shares of other light hydrocarbons such as methane [33,34] as well as more heavy hydrocarbons such as phenols, toluene, naphthalene, benzene along with other aromatic compounds [35], which are often referred to as tars [31,36]. After heating and drying of the solid fuel, pyrolysis takes place. In case of the fixed bed gasifiers, it is often referred to as flaming pyrolysis [31,37] as the heat for the pyrolysis is obtained from the combustion of the pyrolytic gases. Extensive amount of work has been done so far on numerical modelling of the gasification in the entrained flow reactors, working with coal [38]. Much less work was dedicated to gasification of biomass [39], although the existence of an extensive amount of pyrolysis kinetics data, using a multitude of different models shall not be overlooked [40–43]. Nonetheless, only scarce amount of work has been dedicated to numerical models of gasification of torrefied biomass in a fixed bed. However, more recently some works were dedicated to modelling of the pyrolysis of thermally thick biomass particles, which could be considered as laying the foundation for comprehensive numerical modeling of the gasification of biomass in fixed bed reactors [44–48].

There is little information, mentioned in the literature, about the influence of torrefaction of biomass on its subsequent gasification. Prins et al. suggested improvement in the efficiency of the gasification when torrefaction is applied as the valorization technique [14]. Xue et al. suggested that torrefaction might have a positive influence on gasification, due to the improved reactivity of the biomass (Miscanthus), after torrefaction [49]. Couhert et al. reported improved quality of gas, from entrained flow gasification of torrefied wood, in comparison to gasification of raw material [16]. The research, performed by Weiland et al., showed increased gasification efficiency, when comparing torrefied and raw biomass, gasified in a pilot scale entrained flow gasifier [50]. Sarkar et al. studied gasification of raw and torrefied switchgrass in an allothermal fixed bed gasifier and observed increased efficiency of the gasification of torrefied material [51]. Pawlak-Kruczek et al. showed, that valorization of sewage sludge, by means of torrefaction, resulted in decreased content of tars with high melting points, during gasification in fixed bed allothermal gasifier [24].

Gasification in a fixed bed has been a subject of extensive research [31,36,37,52–58]. However, there is still a knowledge gap in terms of influence of torrefaction on the gasification of thermally thick particles. Moreover, little is known in terms of the combined effect of torrefaction and pelleting on gasification of such particles. The aim of this paper is to fill this gap and expand knowledge on the influence of torrefaction and density of pellets on gasification of a thermally thick particle—i.e., torrefied wood pellets. The study aims to compare experimental results with model developed by Luo et al. [59], by comparing the devolatilization times of a thermally thick pellet. Finally, this study is also focused on reporting of some practical values useful as design parameters of a fixed bed gasifier.
2. Materials and Methods

2.1. Sample Preparation and Analysis of Raw and Torrefied Materials

Sawdust from mixed wood species was used for this research. Sawdust was separated into different particle size fractions by using a set of calibrated sieves and a sieve shaker. Sieving was performed for 30 min. Sieved sample was separated into two fractions according to the size, i.e. a coarse fraction between 3150 µm and 1000 µm, and fine fraction with particles smaller than 200 µm.

In order to perform pre-selection of torrefaction conditions, a sample of a raw sawdust was a subject of thermogravimetric analysis and differential thermogravimetry (TGA/DTG) that was performed using a TGA/DTG Pyris Diamond instrument from Perkin Elmer (Waltham, MA, United States). A two-step program was set. During the first step, the sample was heated in up to 105 °C with a heating rate of 10 °C/min, with a hold period of 20 min afterwards. During the second step, the sample was heated up to 850 °C with a heating rate of 10 °C/min. The nitrogen of 99.999% purity was used as inert gas for pyrolysis tests. Fine particles, of particle size smaller than 200 µm, were used.

A Perkin Elmer (Waltham, MA, United States) 2400 analyzer was used for the ultimate analysis, which was performed in compliance with the procedure set in the standard EN ISO 16948 [60]. Each of the tested materials was milled and sieved through a sieve with aperture size of 200 µm, using mortar and pestle, prior to ultimate analysis. Friedl Equation [61–63] was used to estimate the Higher Heating Value (HHV) of both raw and torrefied wood:

\[
HHV = 3.55 \cdot C^2 - 232 \cdot C - 2230 \cdot H + 51.2 \cdot C \cdot H + 131 \cdot N + 20,600
\]  

(1)

where, C, H, N represent carbon, hydrogen, and nitrogen, respectively, in dry biomass (i.e. values of C, H and N are substituted, without % sign, using values from the ultimate analysis). The result of the calculation gives HHV with kJ/kg unit. This equation was chosen as it had been validated for many different types of biomass, including woody biomass [61]. Friedl et al. reported a standard error of calibration of 337 kJ/kg and an \( R^2 \) coefficient of 0.943 achieved during validation of Equation (1) against the experimental results [61]. Moreover, Gucho et al. [64] compared results obtained by calorimetric bomb with the results obtained by this formula for torrefied beech wood and Miscanthus, concluding that results obtained by Equation (1) varied less than 5% from calorimetry results [64].

2.2. Torrefaction and Pelletizing

Torrefaction of fine particles (particle sizes smaller than 200 µm) and coarse particles (particle sizes between 1000 µm and 3150 µm) was performed in closed ceramic dishes of ellipsoidal footprint, with wall thickness of approx. 4 mm and volume of approx. 100 ml, length of 10 cm, width of 6 cm, and height of 4 cm. Prior to the experiment, each dish was filled with material, which was poured from the height of approx. 5 cm, above the upper rim of the container. Pouring was performed, until a conical shaped heap formed, with the bottom of the heap aligned with the rim of the dish. Then the excess material was removed by shuffling a rigid scantling over the edge of the container in a manner similar to bulk density measurement, according to EN ISO 17828. This allowed the material to fill the container completely, without significant compaction. Closed containers were subsequently inserted into hot laboratory furnace, pre-heated to 300 °C. Torrefaction was performed, in duplicate, with two distinct residence times of 15 min and 30 min, in order to obtain distinctly different samples, with respect to the severity of the torrefaction process. Temperature and residence time were selected based on a typical range of torrefaction conditions, reported in the literature [4,8,23–26,65–68]. Selection of the temperatures from the higher end and residence time from lower end of reported conditions was dictated by the way these parameters influence productivity of torrefaction reactors. Containers were closed in order to prevent access of the air, as torrefaction in slightly oxidizing conditions can have significant influence on the product [18,69–77]. Closed containers were subsequently taken out of the furnace and placed into a desiccator, filled with silica gel, in order to cool down.
Pelletizing was performed, using a laboratory pelletizer, producing one pellet at a time. Pellets were pressed, using fine particles, in a dye of 12 mm diameter. Pressure of 60 bars was applied constantly for 1 min, using manual hydraulic press. Pressure of the hydraulic fluid was measured by an analogue gauge. Pelletizing was repeated at least 10 times for each material.

2.3. Devolatisation Model for Thermally Thick Particles

The model was originally developed by Luo et al. [59] to investigate spherical wood devolatilization at high temperature conditions in a single particle combustor. It is further modified to simulate biomass devolatilization of cylinder wood particles in this work.

The model is a one-dimensional model for cylinder wood particles and the anisotropy of the wood is neglected. Both internal and external heat transfer are included, and the mass transfer of the released volatiles is assumed to be unlimited. Radiation between wood particles and reactor walls is considered. The particle shrinkage changes linearly with the devolatilization degree. A fully devolatilized particle has a shrinkage factor of 0.2 (volume based).

Based on the assumptions, mentioned above, the heat balance of thermally-thick biomass particle can be expressed by the following Equation:

\[
C_p \rho_p \frac{dT(t, r)}{dt} = \frac{1}{r} \left( \lambda_{eff} \frac{dT(t, r)}{dr} \right) - \frac{dX_{w}}{dt} \rho_{DB} Y_w Q_w - \frac{dX_{vol}}{dt} \rho_{DB} Y_{vol} Q_{pyro}
\]

where, \( C_p \), \( \rho_p \), \( \lambda_{eff} \) are the specific heat capacity (J/(kgK)), particle density (kg/m\(^3\)), and effective thermal conductivity (W/(mK)) of the biomass particle at time \( t \), respectively; \( \rho_{DB} \) is the initial particle density (kg/m\(^3\)) at \( t = 0 \) s; \( r \) indicates the radial position (m), \( t \) indicates time (s), whereas \( T \) is the particle temperature (K) and \( Y_{w0} \) is the initial water mass fraction (dry basis), and \( Y_{vol0} \) is the initial mass fraction of volatiles (dry basis). \( Q_w \) and \( Q_{pyro} \) are the evaporation heat and devolatilization heat (J/kg), respectively. \( dX_{w}/dt \) and \( dX_{vol}/dt \) are conversion rates described by a drying model and a devolatilization model. More details on drying and devolatilization models, and physical properties (such as conductivity of biomass, heat capacity, enthalpy of pyrolysis etc.) can be found in Luo et al. [59]. The boundary conditions at particle center and the external particle surface are described by Equation (3) and Equation (4), respectively:

\[
\left. \frac{dT}{dr} \right|_{r=0} = 0, \quad \text{all} \ t
\]

\[
\left. \lambda_{eff} \frac{dT}{dr} \right|_{r=R} = h_c (T_g - T_s) + \xi \sigma (T_s^4 - T_{w}^4) \quad t > 0
\]

where, \( h_c \) is the convective heat transfer coefficient, W/(m\(^2\)K) determined by the correlation of Churchill and Bernstein [78], \( T_g, T_w \) and \( T_s \) are gas, wall and particle external surface temperatures (K) respectively; \( \xi \) is the particle emissivity, which depends on many factors (e.g. temperature, wood type, surface structure). Following Luo et al. [59] it was assumed to be 0.85. \( R \) is the particle radius (m).

2.4. Pyrolysis and Gasification Experiments Using Isothermal Furnace

Pyrolysis is one of the stages of a gasification process [31,33,38]. Therefore, a pyrolysis zone can be distinguished in all types of fixed bed gasifiers [31,33]. Consequently, it influences design parameters, such as the residence time of the particles in the bed of a gasifier and its size [37]. However, Boudouard reaction along with other reduction reactions also play role in gasification [33,37]. Therefore, sizing of the reduction zone should not be overlooked in a design of a fixed bed gasifier. Due to these reasons two suits of experiments were performed, within the course of this study. Firstly a suite of pyrolysis experiments was performed at temperature of 600 °C, in N\(_2\) atmosphere (impurities <1%). Subsequently gasification tests were performed at temperature of 950 °C, in CO\(_2\) atmosphere.
(impurities <1%). Both temperatures were selected, as representative for the two distinct zones, based on temperature distribution in each of the zones, during fixed bed gasification of wood [31]. Each time a sample of approx. 1 g was used, either in a form of a pellet or as a bed of particles, which were introduced in a basket made of heat resistant steel mesh. Experiments were performed in duplicates.

Experiments were performed in a custom build isothermal furnace (Figure 1). The furnace has three distinct heating zones. Each of the zones consists of a band heater with ceramic insulation. Centre of the furnace is made of heat resistant steel tube, 2 m long, with an inner diameter of 60 mm. The furnace allows feeding various gases in and taking samples of the gases out at three different heights. Overall length of the tube is significant enough, with respect to its diameter, to preheat the gases relatively quickly. Thus, the temperature gradient in the furnace is considered negligible. Two fittings are used to close the top and the bottom of the pipe. The top fitting has a 10 mm hole in the center axis of the furnace. The drive shaft mounted at the top of the furnace can move vertically. A heat resistant wire is attached to the top of the drive shaft on one side, and to the sample basket on the other. Sample basket is made of heat resistant steel plate and mesh. Samples can be loaded by moving the drive shaft all the way to the bottom, with the bottom fitting taken off, thus moving the sample basket into the loading zone (Figure 1).

Figure 1. Isothermal furnace—diagram of the experimental rig.
The composition of permanent gases in both cold, dry producer gas and pyrolytic gas was determined on-line using the Gas 3100P analyzer (manufactured by G.E.I.T Europe bvba, Bunsbeek, Belgium and supplied by Atut Sp. z O.O. Lublin, Poland). This analyzer uses non-dispersive infra-red (NDIR) sensors for measurements of $\text{CO}_2$, $\text{CO}$, $\text{CH}_4$ and $\text{C}_x\text{H}_y$ (light hydrocarbons, given as an methane equivalents). A thermal conductivity detector (TCD) sensor is used to measure the $\text{H}_2$ content, whereas an electrochemical sensor is used for the determination of the $\text{O}_2$ content. The $T_{90}$ response time of the analyzer is 15 s. Due to this reason time step for recording of the measurement of composition was set to be 15 s.

Conversion during the pyrolysis experiment was determined indirectly, using measurement of the composition of gaseous products, according to the following Equation:

$$C_{\text{ti}} = \frac{\int_{t_i}^{t_\Omega} (u_{\text{CO}_2} + u_{\text{CO}} + u_{\text{H}_2} + u_{\text{CH}_4} + u_{\text{C}_x\text{H}_y})}{\int_{t_\Omega}^{\infty} (u_{\text{CO}_2} + u_{\text{CO}} + u_{\text{H}_2} + u_{\text{CH}_4} + u_{\text{C}_x\text{H}_y})}$$

(5)

where $u_{\text{CO}_2}$, $u_{\text{CO}}$, $u_{\text{H}_2}$, $u_{\text{CH}_4}$, $u_{\text{C}_x\text{H}_y}$ is a respective share of $\text{CO}_2$, $\text{CO}$, $\text{H}_2$, $\text{CH}_4$ and $\text{C}_x\text{H}_y$ in the measured gas in the time step $t_i$ and $t_\Omega$ is the time, when concentrations of the respective compounds were below detectability of the gas analyzer. Method could not be used for determination of the degree of conversion, during gasification, as it was impossible to distinguish between $\text{CO}_2$ delivered to the process and produced during pyrolysis stage.

2.5. Uncertainty of Measurements

For the determination of the density of pellets maximum permissible error of the scale was 0.01 g, whereas, maximum permissible error of the caliper was 0.02 mm. Each measurement was repeated at least 10 times. For A type uncertainty confidence level of 95% was assumed, along with t-student distribution of the results. B type uncertainty was calculated according to Equation (6):

$$u_B = \frac{\Delta_g}{\sqrt{3}}$$

(6)

where, $\Delta_g$ was respective standard permissible error.

Combined standard uncertainty for respective measurements of mass, length and diameter of produced pellets was calculated according to the following Equation:

$$u_C = \sqrt{u_A^2 + u_B^2}$$

(7)

where, $u_A$ and $u_B$ were A and B type uncertainties for measurement of each value (mass, length, diameter) respectively.

Combined standard uncertainties for measurements of length, diameter and mass of pellets were subsequently used to calculate systematic uncertainty for density of pellets, with the following Equation:

$$u_\rho = \sqrt{\left(\frac{d_p}{d_l}u_{Cl}^2 + \frac{d_p}{d_\varphi}u_{C\varphi}^2 + \frac{d_p}{d_m}u_{Cm}^2\right)}$$

(8)

where indexes $l$, $\varphi$ and $m$ indicate length, diameter and mass of pellets, respectively.

The maximum linearity error of Gas 3100P is 2% of the full measuring range. Measuring ranges were as follows: $\text{CO}_2$ 20%, $\text{CO}$ 40%, $\text{CH}_4$ 10%, $\text{C}_x\text{H}_y$ 5%, $\text{H}_2$ 55% and 25% in the case of $\text{O}_2$. The Gas 3100P device has a linearity drift of 1% of measuring range per week, both for zero and for span. Uncertainty for the time measurement was assumed to be 15 seconds as this was the time step set for the gas analyzer.
3. Results and Discussion

Results of the ultimate analysis (Table 1), show increased carbon content of the torrefied biomass, when comparing to untreated wood. This trend has been confirmed by many different studies, for many different types of feedstock [62,79–84]. Similarly, an increase in HHV of torrefied wood was observed, which is similar to results reported different types of biomass in different studies [85–88]. The increase of carbon content is not significant, in comparison to results reported for torrefaction of wood at 300 °C [4,21–23]. This is especially the case for the torrefaction performed within the course of this study, with residence time of 15 min. This could be attributed to the way of performing torrefaction—i.e., bed of torrefied material was not packed and there was no flow of gas through the bed of material. Thus conduction was the only mean of heat transport throughout the bed of material.

| Sample Description             | C % dry | H % dry | N % dry | S % dry | Ash % dry | HHV kJ/kg |
|--------------------------------|---------|---------|---------|---------|-----------|-----------|
| Raw Sawdust                    | 44.75   | 6.24    | 0.59    | 0.19    | 0.61      | 17,786    |
| Torrefied; 300 °C; 15 min; φ < 200 µm | 46.21   | 4.81    | 0.28    | 0.13    | 0.78      | 18,150    |
| Torrefied; 300 °C; 15 min; φ > 1000 µm | 46.59   | 4.56    | 0.14    | 0.09    | 0.84      | 18,224    |
| Torrefied; 300 °C; 30 min; φ < 200 µm | 48.08   | 4.72    | 0.42    | 0.14    | 1.09      | 18,801    |
| Torrefied; 300 °C; 30 min; φ > 1000 µm | 52.84   | 4.92    | 0.28    | 0.21    | 1.24      | 20,629    |

1 φ—particle size.

This heat conduction limitation had more profound effect on smaller particles, which could be attributed to more void spaces between the particles. Typically gases are bad conductors of heat, therefore void spaces between particles have detrimental effect on heat conduction within a bed of particles.

It is well known that thermal processing influences any subsequent pelletizing process [89–92]. First of all, pellets of raw biomass are limited, in terms of their density, by the upper limit of the plant cell wall density, ranging between 1450 kg/m³ and 1500 kg/m³ [93]. This is determined by the density of the polymers the cell wall consists of, i.e., cellulose and hemicellulose [93–95]. During torrefaction this structure is effectively broken down, but also significant part of hemicellulose is lost due to thermal decomposition [19,22,66,96]. Nonetheless, true density reported for particles after torrefaction, can be higher, ranging between 1525 kg/m³ and 1640 kg/m³ [77]. Therefore, effectively the density of pellets obtained from torrefied biomass is a result of an intertwined relationship between the strengths and structure of material as well as the availability of surfaces that can be potentially bonded to one another, as well as the presence of the binder itself. Pellets made of torrefied wood within the course of the study are not significantly different to pellets of torrefied wood, presented in other studies. Comparisons with some existing literature results are made in Figure 2. Stelte et al. obtained pellets with densities, ranging between 700 kg/m³ and 830 kg/m³, for wood torrefied at temperatures lower than in this study (from 250 °C to 270 °C) [97]. However, severity of torrefaction might have not been as significant as suggested by the process temperature, as torrefaction was performed with relatively long residence time of 120 min [97]. Higher densities (between 950 kg/m³ and 1000 kg/m³) were achieved by Wang et al. [77], for similar torrefaction times and temperature ranging between 250 °C and 290 °C [77]. Torrefaction in this study was performed in the presence of oxygen [77]. However, this should not be considered as a cause of higher density of pellets, as similar density was also achieved by Wang et al. for material torrefied under an atmosphere with 0% oxygen content [77] (see Figure 2). Use of a hot die pre-heated to 170 °C during the experiments [77], seems to be the most plausible explanation. Another study, by Gaitan-Alvarez et al. [98] seems to confirm this hypothesis. The study reported densities of pellets ranging between 900 kg/m³ and 1300 kg/m³. Different species of tropical wood from Costa Rica were torrefied at relatively low temperatures (from 200 °C to 250 °C) and residence times (from 8 min to 12 min) [98]. However, temperature of 180 °C was applied during pelleting
process [98]. Overall, in comparison to untreated biomass, torrefaction introduces additional difficulty in terms of its pelletizing. However, in practice this could be off-set by decreased energy consumption for comminution, which is a pre-requisite for pelletizing or any other form of densification [99].

At first glance, the density of pellets does not seem to be an important design parameter of the fixed bed gasifier. However, density, shape, and the way pellets are stacked determine bulk density of the bed. Any particular size of the gasifier’s bed will only be able to hold certain volume of the material and higher density will undoubtedly imply higher mass of the bed. The mass of the bed, in conjunction with the time necessary to devolatilize and gasify a portion of that mass will determine the required size of the bed.

Additional practical implications will concern the mass of the bed that can be held, without collapsing. Typically, char after complete devolatilization becomes brittle. Thus, too high mass of the bed, pressing from the top, might introduce unnecessary compaction of the bed, leading to increase of the pressure drop. Pressure drop over the bed is a very important parameter of fixed bed gasification, as it significantly influences the flow rate of the air [100,101]. Moreover, density is an important parameter for thermally-thick particle devolatilization models [45,59].

The ability to correctly predict thermal decomposition of thermally-thick particle is crucial for estimation of the basic design parameter for a fixed bed gasifier—i.e., the residence time. This in turn allows estimation of the total size of the reactor, as well as the pyrolysis zone. Mass loss during pyrolysis, measured within the course of this study, is compared against the results obtained by the model (Figure 3).
The degree of conversion, predicted by the model, was slightly underestimated for raw wood pellets, in comparison to experimental results (Figure 3a). On the other hand, the model overestimated the degree of conversion ($\alpha$) for torrefied wood, with the difference increasing with the severity of torrefaction (Figure 3b and Figure 3c). Regarding the modeling side, it seems plausible that the model gives satisfactory predictions, which could be used for estimation of the design parameters of fixed bed gasifiers. It seems prudent to mention that estimation of the mass loss history indirectly (Figure 5) is prone to some errors, both on the modeling and experimental side.
torrefaction (Figure 3b,c). Regarding the modeling side, it seems plausible to state that the observed differences were probably caused by two reasons:

1. The devolatilization kinetic used here was derived from raw wood [102] which may be not a good choice for torrefied wood particles.
2. The model is 1D model, which assumes that the cylinder particle should be infinitely long (length/diameter >4). However, the experimental cylinder wood particle’s length/diameter is in range of 1–2, which may lead to an overestimation of devolatilization rates [103].

Regarding the mass loss history, obtained from the experiments, the underlying assumption behind the indirect estimation, according to Equation (5), is the plug flow along the sampling train. Velocity of the gases in the tubes connecting the outlet of the gas (Figure 1) from the series of impinger bottles with the gas analyzer was approx. 1 m/s, which is not sufficient to obtain a fully developed turbulent flow. However, the gas sample was the subject of mixing in impinger bottles. Moreover, the hose was not perfectly straight, thus introducing additional sources of turbulence. Furthermore, the results were adjusted, by moving the starting point, by the time necessary for the gas to travel between the sampling point (Figure 1—outlet located at heating zone 3) and the inlet to the analyzer, also adjusting for the time necessary for the analyzer to react ($t_{90}$). Finally, an indirect determination of the degree of conversion by measuring exclusively main permanent gases does not take into the account condensable compounds. However, the extent of the influence of the condensable compounds, produced during pyrolysis, on the overall result is not likely to be profound for high values of $\alpha$, which is exactly the time that is the aim of the estimation. Overall, only results of experiments performed with pellets could be compared with the model (Figures 3 and 4), as the model cannot be used to make prediction for beds of material [59].

Figure 6 shows the amount of residue (fixed carbon) that remained after each pyrolysis experiment. Two important observations could be made, based on the obtained results. Firstly, the amount of fixed carbon is higher in torrefied samples, in comparison to raw wood. Moreover, increased amount of fixed carbon can be observed with increasing severity of torrefaction. This is sensible, as part of the volatile matter is removed during torrefaction. However, difference can be observed between the results of pyrolysis for pellets and non-packed bed of raw sawdust (Figure 6). It could be stated without any doubt, that the pyrolysis of raw wood pellet was complete, as could be observed in Figure 3a. Entrainment of the bed of particles also does not seem to give plausible explanation, as it would’ve been a factor also for the case of torrefied material. Moreover, the bed of particles was held together by the basket made of a heat resistant steel mesh. It seems plausible to hypothesize that densely packet structure of the pellet restricted the mass flow of the pyrolysis products within the inside of the pellet. This, in conjunction with relatively high amount of volatile matter, could’ve caused secondary reactions to happen inside of the pellet, which would include re-polymerization of some of the products of primary reactions. More research is needed to confirm this hypothesis and unequivocally state that re-polymerization is responsible for results observed in Figure 6, for raw wood pellets.
Experimental study on gasification of raw and torrefied wood yielded interesting results, as shown in Figures 7–9. Peak times for CO were relatively shorter for pelletized and torrefied material, in comparison to their respective non-pelletized equivalent, whereas release times were generally longer, for the dominant species (Figure 7). This could be caused by an impeded flow of hot gas through the bed of non-pelletized material. In conjunction with poor heat transfer within the bed of sawdust it most likely resulted in higher thermal gradient within that bed, thus causing a longer release time of the gases. From the practical point of view of some designs of fixed bed gasifiers, incorporating constrictions and nozzles, such as Imbert gasifier, it implies that the use of pelletized biomass might decrease the problem with limited penetration of the bed by the gasifying agent. Peaks for both H2 (Figure 8) and CH4 (Figure 9) appeared sooner than respective CO peaks.

![Fixed carbon left after pyrolysis at 600 °C](image)

**Figure 6.** Fixed carbon left after pyrolysis at 600 °C (% mass left after pyrolysis, in comparison to the initial sample mass).

![CO concentration for performed gasification experiments](image)

**Figure 7.** CO concentration for performed gasification experiments: (a) average concentrations of carbon monoxide; (b) release time and the time of peak concentration (Torr. & Pell.—torrefied and pelletized).
were left at the end of gasification, for all of the cases. This is probably due to the Boudouard reaction, according to the following Equation [33]:

\[ C + CO_2 \leftrightarrow 2CO \]  

(9)

It seems perfectly reasonable, taking into account the temperature selected for gasification experiments and the abundance of CO₂. It seems important to note that the release time for hydrogen and methane were much longer, than conversion times recorded for pyrolysis in relatively lower temperatures (600 °C for pyrolysis and 950 °C for gasification). It can be partially contributed to impurities in the gasification agent (impurities <1%). However, the influence of the impurities is limited and cannot be used as the only explanation of the presence of H₂ and CH₂. The reason for these impurities is imperfect separation of CO₂ from other compounds, such as N₂ or water vapors. Drying of the gas is relatively easier than separation of CO₂ and N₂. Moreover, N₂ can be considered inert, so only water vapors, which would be a minor part of the remaining 1% of the gas, could take part in generation of additional CH₄ and H₂ by steam gasification of carbon (10) and methanation reactions (11, 12 and 13) [33]:

\[ C + H_2O \leftrightarrow CO + \text{H}_2 \]  

(10)

\[ 2CO + 2H_2 \leftrightarrow \text{CH}_4 + \text{CO}_2 \]  

(11)

\[ CO + 3H_2 \leftrightarrow \text{CH}_4 + \text{H}_2O \]  

(12)

Overall, it could be stated that both pelletizing and torrefaction had influence on the composition of the gas. The release time of CO was somewhat longer than release time of CH₄ and significantly longer than release time of H₂ for corresponding samples. Moreover, negligible amounts of sample were left at the end of gasification, for all of the cases. This is probably due to the Boudouard reaction, according to the following Equation [33]:

\[ C + CO_2 \leftrightarrow 2CO \]  

(9)

It seems perfectly reasonable, taking into account the temperature selected for gasification experiments and the abundance of CO₂. It seems important to note that the release time for hydrogen and methane were much longer, than conversion times recorded for pyrolysis in relatively lower temperatures (600 °C for pyrolysis and 950 °C for gasification). It can be partially contributed to impurities in the gasification agent (impurities <1%). However, the influence of the impurities is limited and cannot be used as the only explanation of the presence of H₂ and CH₄. The reason for these impurities is imperfect separation of CO₂ from other compounds, such as N₂ or water vapors. Drying of the gas is relatively easier than separation of CO₂ and N₂. Moreover, N₂ can be considered inert, so only water vapors, which would be a minor part of the remaining 1% of the gas, could take part in generation of additional CH₄ and H₂ by steam gasification of carbon (10) and methanation reactions (11, 12 and 13) [33]:

\[ C + H_2O \leftrightarrow CO + \text{H}_2 \]  

(10)

\[ 2CO + 2H_2 \leftrightarrow \text{CH}_4 + \text{CO}_2 \]  

(11)

\[ CO + 3H_2 \leftrightarrow \text{CH}_4 + \text{H}_2O \]  

(12)
\[ \text{CO}_2 + 4\text{H}_2 \leftrightarrow \text{CH}_4 + 2\text{H}_2\text{O} \]  

(13)

Water gas shift reaction (14) is less likely to occur, as \( \text{CO}_2 \) content of the gas was initially high, due to the use of \( \text{CO}_2 \) for gasification:

\[ \text{CO} + \text{H}_2\text{O} \leftrightarrow \text{CO}_2 + \text{H}_2 \]  

(14)

Therefore it seems plausible to hypothesize that the composition of the gasification agent and the composition of the surrounding atmosphere, which is a subject of change during the pyrolysis stage, can have influence on the composition of the gas. It should be noted that respective peaks of \( \text{CH}_4 \) and \( \text{H}_2 \) occurred relatively early, in comparison to the respective peaks of \( \text{CO} \) (Figures 7–9).

Interesting observation could be made, based on the offset between this negative peak and peak of \( \text{CO} \) (Figure 10) increased with growing severity of torrefaction. However, that was the case for non-pelletized torrefied material only. For torrefied and pelletized material the \( \text{CO} \) peak appeared much closer to the \( \text{CO}_2 \) negative peak. This could be attributed to the already mentioned heat transfer issues of non-pelletized material. The much shorter time of the \( \text{CO}_2 \) negative peak, when comparing raw and torrefied sawdust, was probably caused by much quicker release of the gases during pyrolysis phase, which could be attributed to a higher reactivity of the torrefied material.

![Figure 10. Off-set between negative peak of CO\(_2\) and peak of CO for performed gasification experiments (Torr. & Pell.—torrefied and pelletized).](image)

Overall, it could be stated that both physical (pelletizing) and thermal (torrefaction) valorization has an influence on the performance of the feedstock during gasification in a fixed bed gasifier, which in turn will have influence on both composition of producer gas and the amount of unreacted carbon, remaining after the process. In order to obtain satisfactory performance, unreacted carbon loss should be minimized. Reduction zone of sufficient size is needed to achieve that.

The work of Reed et al. [37] reported propagation front values in the order of magnitude of 1 cm/min, for raw wood gasification. Looking at the results obtained within the course of this study, much more time is needed for complete conversion of char in the reduction reactions, for torrefied material, which could be attributed to the higher content of fixed carbon in pellets made of torrefied wood, when comparing to raw feedstock. It does not seem plausible at this point to state, that propagation velocity for the gasification of torrefied wood would be similar to raw wood. Nevertheless, for a single particle at least 25% up higher residence time is needed, in order to avoid losses caused by chemical energy left in unreacted carbon. This should be also taken into account, in the design
of gasifiers aiming at using torrefied wood pellets. Moreover, proper consideration is advised for the projects where torrefied wood pellets are to replace raw wood pellets.

Therefore, it seems to be sensible to recommend further work on development of a model for fixed bed gasification. In order to give satisfactory results for carbon conversion, such model would need to take into account reduction reactions (especially Boudouard reaction). Such a model would also need to incorporate features of a thick-particle model, as the obtained results clearly show the influence of the particle, especially its density, on the gasification process. The model used in this study seems to be a good foundation for such development. Moreover, it could be used as a first approximation of the size of the pyrolysis zone.

4. Conclusions

Of three major compounds released during gasification, release of CO was the slowest one, therefore this release time should be used as a required residence time, if maximum carbon conversion is to be achieved. Both release time and the time of achieving a peak concentration of CO increased with increased severity of the torrefaction. Both release time and the time of achieving a peak concentration of CO were significantly lower for pelletized torrefied biomass in comparison to their respective non-pelletized equivalents. Concentrations of all the measured compounds were higher for pelletized torrefied sawdust in comparison to the respective non-pelletized equivalents.

Peak times were relatively shorter for pelletized and torrefied material, in comparison to their respective non-pelletized equivalent, whereas release times were generally longer, for the dominant species (CO). This suggests that pelletized biomass is potentially less susceptible for potential problems with penetration of the gasifying agent in gasifiers with constrictions, due to better heat transfer within individual particles. With respect to the design parameters of the stratified gasifiers, for torrefied pellets, at least 25% higher residence time is advised, in comparison to the use of raw wood pellets. This would imply increase in the assumed height of the bed of material above the grate. However, more research is needed to confirm this, as the gasification propagation velocities within the bed also need to be determined for the case, where torrefied pellets are used.

The offset between the negative peak of CO$_2$ and the peak of CO concentration was rising with increased severity of torrefaction, however, for non-pelletized torrefied material only. For torrefied and pelletized material the CO peak appeared much closer to the CO$_2$ negative peak. This could be attributed to the already mentioned heat transfer issues for non-pelletized material.

Further work is recommended on development of the model that could work for a bed of material, taking into account all stages of gasification. Such a model would be a pre-requisite for tailoring designs of fixed bed gasifier allowing the most optimal use of valorized fuels. This would eliminate the trial and error phase of research and development process and speed up the effort aiming for optimization of the design of gasifiers.

Author Contributions: Conceptualization: H.L., H.P.-K. and L.N.; methodology: H.L., H.P.-K. and L.N.; validation: K.K. and M.B.; formal analysis: H.L. and L.N.; investigation: L.N., H.L., K.K., M.T., T.S. and A.S.; resources: H.P.-K. and A.A.; data curation: M.T., T.S., A.S., K.K. and M.B.; writing—original draft preparation: L.N., H.L. and A.A.; writing—review and editing: L.N., Z.L., and H.L.; supervision: H.P.-K., A.A. and K.M.; project administration: L.N., K.M.; funding acquisition: L.N. and H.P.-K. All authors have read and agreed to the published version of the manuscript.

Funding: Support of this research from the National Science Centre, Poland, project no. 2019/33/N/ST8/02641, is gratefully acknowledged.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Pawlak-Kruczek, H.; Niedźwiecki, L.; Ostrycharczyk, M.; Czep, M.; Plutecki, Z. Potential and methods for increasing the flexibility and efficiency of the lignite fired power unit, using integrated lignite drying. Energy 2019, 181, 1142–1151. [CrossRef]
25. Pawlak-Kruczek, H.; Arora, A.; Gupta, A.; Azam, M.; Niedz wiecki, L.; Andrews, G.; Phylaktou, H.; Gibbs, B.; Newlaczyl, A.; Livesey, P.M. Biocoal - Quality control and assurance. Biomass Bioenergy 2020, 135, 105509. [CrossRef]

26. Pawlak-Kruczek, H.; Arora, A.; Mo´scicki, K.; Krochmalny, K.; Sharma, S.; Niedzwiecki, L. A transition of a domestic boiler from coal to biomass – Emissions from combustion of raw and torrefied Palm Kernel shells (PKS). Fuel 2020, 263. [CrossRef]

27. Jahirul, M.I.; Rasul, M.G.; Chowdhury, A.A.; Ashwath, N. Biofuels production through biomass pyrolysis—A technological review. Energies 2012, 5, 4952–5001. [CrossRef]

28. Bridgwater, A.V. Review of fast pyrolysis of biomass and product upgrading. Biomass Bioenergy 2012, 38, 68–94. [CrossRef]

29. Butler, E.; Devlin, G.; Meier, D.; McDonnell, K. A review of recent laboratory research and commercial developments in fast pyrolysis and upgrading. Renew. Sustain. Energy Rev. 2011, 15, 4171–4186. [CrossRef]

30. Collard, F.X.; Blin, J. A review on pyrolysis of biomass constituents: Mechanisms and composition of the products obtained from the conversion of cellulose, hemicelluloses and lignin. Renew. Sustain. Energy Rev. 2014, 38, 594–608. [CrossRef]

31. Reed, T.B.; Das, A. Handbook of Biomass Downdraft Gasifier Engine Systems; SERI a Division of Midwest Research Institute: Golden, CO, USA, 1988.

32. Reed, T.B.; Markson, M. Biomass Gasification Reaction Velocities. In Fundamentals of Thermochemical Biomass Conversion; Overend, R.P., Milne, T.A., Mudge, L.K., Eds.; ElsevierScience Publishing Co. Inc.: Barking, UK, 1985; pp. 951–965.

33. Mularski, J.; Pawlak-Kruczek, H.; Modlinski, N. A review of recent studies of the CFD modelling of coal gasification in entrained flow gasifiers, covering devolatilization, gas-phase reactions, surface reactions, models and kinetics. Fuel 2020, 271, 117620. [CrossRef]

34. Ramos, A.; Monteiro, E.; Rouboa, A. Numerical approaches and comprehensive models for gasification process: A review. Renew. Sustain. Energy Rev. 2019, 110, 188–206. [CrossRef]

35. White, J.E.; Catallo, W.J.; Legendre, B.L. Biomass pyrolysis kinetics: A comparative critical review with relevant agricultural residue case studies. J. Anal. Appl. Pyrolysis 2011, 91, 1–33. [CrossRef]

36. Gómez, M.A.; Forteiro, J.; Fatino, D.; Miguez, J.L. Fast-solving thermally thick model of biomass particles embedded in a CFD code for the simulation of fixed-bed burners. Energy Convers. Manag. 2015, 105, 30–44. [CrossRef]
47. Sieradzka, M.; Gao, N.; Quan, C.; Młonka-Medrala, A.; Magdziarz, A. Biomass thermochemical conversion via pyrolysis with integrated CO2 capture. *Energies* 2020, 13, 1050. [CrossRef]
48. Młonka-Medrala, A.; Magdziarz, A.; Dziok, T.; Sieradzka, M.; Nowak, W. Laboratory studies on the influence of biomass particle size on pyrolysis and combustion using TG GC/MS. *Fuel* 2019, 252, 635–645. [CrossRef]
49. Xue, G.; Kwapisinska, M.; Kwapisinski, W.; Czajka, K.M.; Kennedy, J.; Leahy, J.J. Impact of torrefaction on properties of Miscanthus × giganteus relevant to gasification. *Fuel* 2014, 121, 189–197. [CrossRef]
50. Weiland, F.; Nordwaeger, M.; Olofsson, I.; Wiinikka, H.; Nordin, A. Entrained flow gasification of torrefied wood residues. *Fuel Process. Technol.* 2014, 125, 51–58. [CrossRef]
51. Sarkar, M.; Kumar, A.; Tumuluru, J.S.; Patil, K.N.; Bellmer, D.D. Gasification performance of switchgrass pretreated with torrefaction and densification. *Appl. Energy* 2014, 127, 194–201. [CrossRef]
52. Reed, T.B.; Levie, B.; Graboski, M.S. *Fundamentals, Development and Scaleup of the Air-Oxygen Stratified Downdraft Gasifier*; Solar Energy Research Institute: Golden, CO, USA, 1988.
53. Hasler, P.; Nussbaumer, T. Gas cleaning for IC engine applications from fixed bed biomass gasification. *Biomass Bioenergy* 1999, 16, 385–395. [CrossRef]
54. Mandl, C.; Obernberger, I.; Biedermann, F. Modelling of an updraft fixed-bed gasifier operated with softwood pellets. *Fuel* 2010, 89, 3795–3806. [CrossRef]
55. Zhang, Z.Z.; Zhu, M.M.; Liu, P.F.; Wan, W.C.; Zhou, W.X.; Chan, Y.L.; Zhang, D.K. Effect of Biochar on the Cracking of Tar from the Pyrolysis of a Pine Sawdust in a Fixed Bed Reactor. *Energy Procedia* 2015, 75, 196–201. [CrossRef]
56. Sharma, A.K. Experimental investigations on a 20 kWe, solid biomass gasification system. *Biomass Bioenergy* 2011, 35, 421–428. [CrossRef]
57. Werle, S. Gasification of a Dried Sewage Sludge in a Laboratory Scale Fixed Bed Reactor. *Energies* 2015, 8, 8562–8572. [CrossRef]
58. Werle, S. Impact of feedstock properties and operating conditions on sewage sludge gasification in a fixed bed gasifier. *Waste Manag. Res.* 2014, 32, 954–960. [CrossRef]
59. Luo, H.; Lu, Z.; Jensen, P.A.; Glarborg, P.; Lin, W.; Dam-Johansen, K.; Wu, H. Experimental and modelling study on the influence of wood type, density, water content, and temperature on wood devolatilization. *Fuel* 2020, 260, 116410. [CrossRef]
60. CEN (European Committe for Standardisation). EN ISO 16948:2015 Determination of Total Content of Carbon, Hydrogen and Nitrogen—Instrumental Methods; European Committe for Standardisation: Brussels, Belgium, 2015.
61. Friedl, A.; Padouvas, E.; Rotter, H.; Varmuza, K. Prediction of heating values of biomass fuel from elemental composition. *Anal. Chim. Acta* 2005, 544, 191–198. [CrossRef]
62. Sidiras, D.K.; Nazos, A.G.; Giakoumakis, G.E.; Politi, D.V. Simulating the effect of torrefaction on the heating value of barley straw. *Energies* 2020, 13, 736. [CrossRef]
63. Wilk, M.; Magdziarz, A. Hydrothermal carbonization, torrefaction and slow pyrolysis of Miscanthus giganteus. *Energy* 2017, 140, 1292–1304. [CrossRef]
64. Gucho, E.M.; Shahzad, K.; Bramer, E.A.; Akhtar, N.A.; Brem, G. Experimental study on dry torrefaction of beech wood and miscanthus. *Energies* 2015, 8, 3903–3923. [CrossRef]
65. Pawlak-Kruczek, H.; Wnukowski, M.; Niedźwiecki, L.; Kowal, M.; Krochmalny, K. Gasification of torrefied sewage sludge with the addition of calcium carbonate. *J. Energy Resour. Technol.* 2020, 142, 070904-1–070904-8. [CrossRef]
66. Khalsa, J.H.A.; Leistner, D.; Weller, N.; Darvell, L.I.; Dooley, B. Torrefied biomass pellets—Comparing grindability in different laboratory mills. *Energies* 2016, 9, 794. [CrossRef]
67. Medic, D.; Darr, M.; Shah, A.; Rahn, S. The effects of particle size, different corn stover components, and gas residence time on torrefaction of corn stover. *Energies* 2012, 5, 1199–1214. [CrossRef]
68. Pawlak-Kruczek, H.; Krochmalny, K.; Mościcki, K.; Zgórna, J.; Czerep, M.; Ostrychaczyk, M.; Niedźwiecki, L. Torrefaction of Various Types of Biomass in Laboratory Scale, Batch-Wise Isothermal Rotary Reactor and Pilot Scale, Continuous Multi-Stage Tape Reactor. *Eng. Prot. Environ.* 2017, 20, 457–472. [CrossRef]
69. Lasek, J.A.; Kopczyński, M.; Janusz, M.; Iluk, A.; Zuwala, J. Combustion properties of torrefied biomass obtained from flue gas-enhanced reactor. *Energy* 2017, 119, 362–368. [CrossRef]
70. Rousset, P.; Macedo, L.; Commandré, J.M.; Moreira, A. Biomass torrefaction under different oxygen concentrations and its effect on the composition of the solid by-product. *J. Anal. Appl. Pyrolysis* 2012, 96, 86–91. [CrossRef]

71. Uemura, Y.; Omar, W.; Othman, N.A.; Yusup, S.; Tsutsui, T. Torrefaction of oil palm EFB in the presence of oxygen. *Fuel* 2013, 103, 156–160. [CrossRef]

72. Uemura, Y.; Saadon, S.; Osman, N.; Mansor, N.; Tanoue, K.I. Torrefaction of oil palm kernel shell in the presence of oxygen and carbon dioxide. *Fuel* 2015, 144, 171–179. [CrossRef]

73. Lu, K.M.; Lee, W.J.; Chen, W.H.; Liu, S.H.; Lin, T.C. Torrefaction and low temperature carbonization of oil palm fiber and eucalyptus in nitrogen and air atmospheres. *Bioresour. Technol.* 2012, 123, 98–105. [CrossRef] [PubMed]

74. Chen, W.H.; Zhuang, Y.Q.; Liu, S.H.; Juang, T.T.; Tsai, C.M. Product characteristics from the torrefaction of oil palm fiber pellets in inert and oxidative atmospheres. *Bioresour. Technol.* 2016, 199, 367–374. [CrossRef] [PubMed]

75. Joshi, Y.; Di Marcello, M.; Krishnamurthy, E.; De Jong, W. Packed-Bed Torrefaction of Bagasse under Inert and Oxygenated Atmospheres. *Energy Fuels* 2015, 29, 5078–5087. [CrossRef]

76. Chen, W.H.; Lu, K.M.; Liu, S.H.; Tsai, C.M.; Lee, W.J.; Lin, T.C. Biomass torrefaction characteristics in inert and oxidative atmospheres at various superficial velocities. *Bioresour. Technol.* 2013, 146, 152–160. [CrossRef]

77. Wang, C.; Peng, J.; Li, H.; Bi, X.T.; Legros, R.; Lim, C.J.; Sokhansanj, S. Oxidative torrefaction of biomass residues and densification of torrefied sawdust to pellets. *Bioresour. Technol.* 2013, 127, 318–325. [CrossRef]

78. Churchill, S.W.; Bernstein, M. A correlating equation for forced convection from gases and liquids to a circular cylinder in crossflow. *J. Heat Transfer* 1977, 99, 300–306. [CrossRef]

79. Li, J.; Brzdekiewicz, A.; Yang, W.; Blasiak, W. Co-firing based on biomass torrefaction in a pulverized coal boiler with aim of 100% fuel switching. *Appl. Energy* 2012, 99, 344–354. [CrossRef]

80. Park, S.; Kim, S.J.; Oh, K.C.; Cho, L.H.; Kim, M.J.; Jeong, I.S.; Lee, C.G.; Kim, D.H. Characteristic analysis of torrefied pellets: Determining optimal torrefaction conditions for agri-byproduct. *Energies* 2020, 13, 423. [CrossRef]

81. Świechowski, K.; Liszewski, M.; Babelewski, P.; Koziel, J.A.; Białowiec, A. Oxytree pruned biomass torrefaction: Mathematical models of the influence of temperature and residence time on fuel properties improvement. *Materials* 2019, 12, 2228. [CrossRef]

82. Bajcar, M.; Zagula, G.; Saletnik, B.; Tarapatskyy, M.; Puchalski, C. Relationship between torrefaction parameters and physicochemical properties of torrefied products obtained from selected plant biomass. *Energies* 2018, 11, 2919. [CrossRef]

83. Sh, L.; Lee, B.H.; Lee, Y.J.; Jeon, C.H. Comparing the physicochemical properties of upgraded biomass fuel by torrefaction and the ashless technique. *Appl. Sci.* 2019, 9, 5519. [CrossRef]

84. Dudek, M.; Świechowski, K.; Manczarski, P.; Koziel, J.A.; Białowiec, A. The effect of biochar addition on the biogas production kinetics from the anaerobic digestion of brewers’ spent grain. *Energies* 2019, 12, 1518. [CrossRef]

85. Stepień, P.; Świechowski, K.; Hnat, M.; Kugler, S.; Stegenta-Dąbrowska, S.; Koziel, J.A.; Manczarski, P.; Białowiec, A. Waste to carbon: Biocoal from elephant dung as new cooking fuel. *Energies* 2019, 12, 4344. [CrossRef]

86. Sygula, E.; Koziel, J.A.; Białowiec, A. Proof-of-concept of spent mushrooms compost torrefaction—Studying the process kinetics and the influence of temperature and duration on the calorific value of the produced biocoal. *Energies* 2019, 12, 3060. [CrossRef]

87. Poudel, J.; Karki, S.; Oh, S.C. Valorization of waste wood as a solid fuel by torrefaction. *Energies* 2018, 11, 1641. [CrossRef]

88. Wilk, M.; Magdziarz, A.; Kalemba, I. Charactetisation of renewable fuels’ torrefaction process with different instrumental techniques. *Energy* 2015, 87, 259–269. [CrossRef]

89. Distler, T.; Sitzmann, W. An investigation on additives for pelletizing highly torrefied biomass. *Biofuels Bioprod. Biorefining* 2018, 1–8. [CrossRef]

90. Stelte, W.; Nielsen, N.P.K.; Hansen, H.O.; Dahl, J.; Shang, L.; Sanadi, A.R. Pelletizing properties of torrefied wheat straw. *Biomass Bioenergy* 2013, 49, 214–221. [CrossRef]

91. Reza, M.T.; Uddin, M.H.; Lynam, J.G.; Coronella, C.J. Engineered pellets from dry torrefied and HTC biochar blends. *Biomass Bioenergy* 2014, 63, 229–238. [CrossRef]
92. Białowiec, A.; Micuda, M.; Koziel, J.A. Waste to carbon: Densification of torrefied refuse-derived fuel. *Energies* 2018, 11, 3233. [CrossRef]

93. Stelte, W.; Holm, J.K.; Sanadi, A.R.; Barsberg, S.; Ahrenfeldt, J.; Henriksen, U.B. Fuel pellets from biomass: The importance of the pelletizing pressure and its dependency on the processing conditions. *Fuel* 2011, 90, 3285–3290. [CrossRef]

94. Dinwoodie, J.M. *Timber: Its Nature and Behaviour*, 2nd ed.; Routledge: Abingdon, UK, 2000; ISBN 978-0419235804.

95. Forest Products Laboratory—USDA. *Wood Handbook: Wood as an Engineering Material*; United States Department of Agriculture: Madison, WI, USA, 2010.

96. Acharya, B.; Dutta, A.; Minaret, J. Review on comparative study of dry and wet torrefaction. *Sustain. Energy Technol. Assess.* 2015, 12, 26–37. [CrossRef]

97. Stelte, W.; Clemons, C.; Holm, J.K.; Sanadi, A.R.; Ahrenfeldt, J.; Shang, L.; Henriksen, U.B. Pelletizing properties of torrefied spruce. *Biomass Bioenergy* 2011, 35, 4690–4698. [CrossRef]

98. Gaitán-Alvarez, J.; Moya, R.; Puente-Urbina, A.; Rodriguez-Zuñiga, A. Physical and compression properties of pellets manufactured with the biomass of five woody tropical species of Costa Rica torrefied at different temperatures and times. *Energies* 2017, 10, 1205. [CrossRef]

99. Shang, L.; Nielsen, N.P.K.; Dahl, J.; Stelte, W.; Ahrenfeldt, J.; Holm, J.K.; Thomsen, T.; Henriksen, U.B. Quality effects caused by torrefaction of pellets made from Scots pine. *Fuel Process. Technol.* 2012, 101, 23–28. [CrossRef]

100. Donaj, P.; Izadpanah, M.R.; Yang, W.; Blasiak, W. Effect of pressure drop due to grate-bed resistance on the performance of a downdraft gasifier. *Energy Fuels* 2011, 25, 5366–5377. [CrossRef]

101. Gunarathne, D.S.; Chmielewski, J.K.; Yang, W. Pressure drop prediction of a gasifier bed with cylindrical biomass pellets. *Appl. Energy* 2014, 113, 258–266. [CrossRef]

102. Wagenaar, B.M.; Prins, W.; van Swaaij, W.P.M. Flash pyrolysis kinetics of pine wood. *Fuel Process. Technol.* 1993, 36, 291–298. [CrossRef]

103. Momeni, M.; Yin, C.; Kaer, S.K.; Hansen, T.B.; Jensen, P.A.; Glarborg, P. Experimental Study on Effects of Particle Shape and Operating Conditions on Combustion Characteristics of Single Biomass Particles. *Energy Fuels* 2013, 27, 507–514. [CrossRef]

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