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Kinetic Modeling of Rice Husk Components Pyrolysis Based on Independent Parallel Reactions

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Abstract

This study presents the pyrolysis decomposition mechanism of rice husk in nitrogen atmosphere by thermogravimetric analysis (TGA). The thermal decomposition was carried out in three stages: moisture removal (180-200 °C), main devolatilization (200-400 °C) and continuous slight devolatilization (400 °C). The weight loss of rice husk was modeled based on the assumption that lignocellulose components (cellulose, hemicellulose and lignin) undergo pyrolysis independently in parallel first-order reactions. The kinetic parameters of the three lignocellulose components were determined by means of Microsoft Excel Solver tool using least square algorithm. The result of thermal degradation of rice husk samples shows that the model predictions of the cellulose, hemicellulose and lignin components agreed with the experiments. The activation energy and pre-exponential factors of cellulose, hemicellulose and lignin are187kJ/mol and 1.2×10¹⁵min⁻¹, 29kJ/mol and 1.6×10⁷min⁻¹ and 90 kJ/mol and 1.8×10¹min⁻¹ respectively. The results suggest that thermal decomposition rate of cellulose was found to be higher, whereas that of lignin decomposition rate was lower. The thermal decomposition of hemicellulose decomposition was intermediate.

Keywords: Rice husk, pyrolysis, thermogravimetric analysis, lignocellulose, kinetic model.

Introduction

The depletion of fossil fuel reserves and its associate environmental problems are the major reasons for the shift and focus on sustainable energy resources. Biomass is alternative an as extensively used renewable energy source which is in abundance and in most cases considered to be a waste. It is the world fourth energy resource after oil, natural gas and coal (Vamvuka et al., 2003). Lignocellulose materials are unlike carbonhydrate which belong to food that cannot be digested. The use of lignocellulose biomass as a source of chemical, transportation and energy fuels has been growing at a very rapid rate because lignocellulose biomass does not compete with the food supply chain (Basu, 2006).

Nigeria like many other African countries is blessed with abundant biomass because of its huge agricultural resources and this sector has been reported to provide employment to over 60% population. Agricultural activities generate huge waste providing a large potential source of lignocellulosic materials. Among the various lignocellulose species, rice husk has potential for chemical energy. Rice husk is used as a source of energy in rice mill, cooking, furnace and boiler (Garba et al., 2006). In Nigeria, about 664,000tonnes of rice husk is derived from about 3.32 million tons of paddy rice per year (Rainer).

Conversion of lignocellulosic material to efficient fuel remains a burden due to their structural complexity which is responsible for the challenges in separating lignocellulose components.

Thermochemical conversion processes such pyrolysis, combustion, gasification and liquefaction are considered as the converting of alternative means lignocellulose biomass to efficient fuel. However, pyrolysis remains the focal point conversion thermochemical processes as in the case of combustion process in which the first step is pyrolysis followed by the reaction of pyrolysis residue with oxygen. The pyrolysis process is the thermal decomposition of presence of inert biomass in the atmosphere to form liquid (bio oil), char (charcoal) and gas. Lignocellulose biomass is composed of hemicellulose, cellulose and lignin (Garba etal.. 2012). Experimental results have shown that pyrolysis of organic matter is a complex process involving parallel and series reactions (Garba et al., 2013). The decomposition of biomass has been described by three-component independent each corresponding to the reaction, decomposition of the hemicellulose, cellulose and lignin. The weight loss kinetic model developed for biomass is of two types. The single component overall model (SOM), which considered the biomass as being composed of a single component and uses its char-volatile reaction to describe the weight loss kinetics(Rogers et al., 1980; Cordero et al., 1989; Cordero et al., 1990). The second one is the multi-component overall model (MOM) considering biomass as being composed of hemicellulose, cellulose and lignin (Orfao et al., 1999; Font et al., 19991). In this case, the components are modeled separately and the biomass decomposition is the summation of all the three components. Generally, the weight loss kinetics change as the temperature increases and the weight loss is controlled

mainly by cellulose, hemicellulose and lignin. Since the weight loss depends on hemicellulose and cellulose. component, SOM cannot account for the possible change in weight loss kinetics. MOM can reveal possible Contrary, weight loss kinetics. in change Koufopanos et al. (1989) developed a modeling approach which described the overall rate of decomposition as a sum of corresponding rates the hemicellulose, cellulose and lignin using nonlinear regression method. In a related work, Rao et al. (1998) developed a kinetic model in which the pyrolysis of lignocellulose components is described by single reaction first order kinetics. In recent years MOM has been used successfully to describe the weight loss processes of bagasse (Liangfeng et al., 2001;), jatropha residue (Rajeev et al., 2016), switch grass (Vamvuka et al., 2011) and wood pine and cotton stalk (Wang et al., 2012). The result of almost all the previous studies have shown good agreement between experimental and calculated weight loss pattern.

The understanding of biomass pyrolysis is essential for accurate prediction of pyrolysis rates require for optimum design of pyrolyzer. In order to optimize process variable and obtain quality pyrolysis products, there is the need for more profound knowledge of pyrolysis kinetics. In the present work, the pyrolysis of rice husk components is analysed and pyrolysis process is described by independent parallel reactions of first order using nonlinear square methods. The design of equipment for the conversion of biomass through thermochemical processes requires knowledge of pyrolysis kinetic and modeling. Many literatures reported the kinetic investigation however; little

knowledge is available on the modeling aspect. Thermogravimetric curves was used to analysis biomass decomposition, the pyrolysis kinetic model of rice husk based on three-component independent parallel first-order reactions was established by non-linear least squares algorithm. The kinetic parameters obtained for rice components were compared with each other and meaningful results achieved were discussed.

Methodology

Proximate and ultimate analysis

The rice husk used in this experiment was collected from Minna, Niger State. The experiments were divided into two steps; the first step was to study the pyrolysis characteristics of rice husk by using proximate, ultimate analysis and thermal decomposition. The second step was to study thepyrolysis kinetics of rice husk. The proximate analysis of the rice husk sample was carried out according to ASTMD 3172 method and ultimate analysis was done according to ASTMD 3176. The thermal degradation was carried out by with the aid of thermogravimetric analysis (TGA). The sample of a ground rice husk was obtained by grinding the rice husk to an average particle size of less than 150 μm . 20 mg of the rice husk sample was used in the experiment. The weight loss of the sample as a function of temperature was measured.

Kinetic modeling of rice husk pyrolysis

The approach by most of these kinetic models simulate the chemical reactions that take place during pyrolysis and the chemical reaction scheme is presented in equation (1).

Biomass
$$\rightarrow$$
 volatile + char+gas (1)

The kinetic modeling was performed on the pyrolysis profile provided by TGA. The total mass measured by TGA were assumed to be summation of thepyrolyzable materials such as cellulose, hemicelluloses, lignin, charcoal, and ash (Liangfeng et al., 2011).

$$m = m_p + m_c + m_a$$

where m is the total mass of biomass by TGA, m_p is the mass of pyrolyzable (cellulose, hemicelluloses, lignin components) organic materials, m_c is the mass of produced char, and m_a is the ash mass.

Assuming the ash is ignored, the mass of biomass is a sum of the pyrolyzable organic chemical and the produced charcoal (Liangfeng et al., 2011):

$$m_i = m_{ip} + m_{ic}$$
 for i=1,2,3 (2)

the pyrolyzable organic chemicals are mainly cellulose, hemicelluloses, lignin components.

During the pyrolysis process, the degree of conversion and conversion rate for each component in biomass sample are defined as follows:

$$\alpha_i = \frac{m_{io} - m_{ip}}{m_{ip}}$$
 for i=1,2,3 (3)

Where m_{io} is the intial weight of biomass and α_i is the conversion of each lignocellulose components.

For the three lignocellulose components, the overall conversion rate for each reactioncan be expressed as

$$\frac{dm}{dt} = \sum_{i=1}^{3} c_i \frac{d\alpha}{dt} \tag{4}$$

Where c_1 , c_2 and c_3 are the partial contributions of the overall weight loss:

$$c_i = m_{io} - m_{ichar} \tag{5}$$

components lignicellulose presumed to decompose by obeying firstorder reaction according to

$$\frac{d\alpha_i}{dt} = k_{i0}.e^{-E_i/RT}(1 - \alpha_i)$$
(6)

where E_i , k_{io} , T and R are the activation energy (kJ/mol), frequency factor (min⁻¹), temperature (K) and respectively.

The nonlinear least square algorithm has been used to identify the kinetic parameters that show the lowest values of the objective function (O.F):

$$O.F = \sum \left[\left(\frac{dm}{dt} \right)^{\exp} - \left(\frac{dm}{dt} \right)^{sim.} \right]^{2}$$
 (7)

the experiment DTG (dm/dt)^{exp} is curverate of rice husk and (dm/dt)sim. is the simulated DTG curve rate of rice husk.

Kinetic constants from modeling

thermal the shows Equation4 decomposition rates of the rice husk components during the heating process was a function of temperature and degree of conversion. Equation (5) is a nonlinear ordinary first-order differential equation. The degree of conversion is the dependent variable and absolute temperature is the kinetic Two variable. independent constants for each component, exponential factor kio and activation energy Ei, are assumed to be independent of the degree of conversion. The numerical solution could be obtained by solving Equation (6) for the known values of kio and Ei respectively. These constants are determined from the least-square method implemented by the Microsoft Excel Solver tool where the difference between experimental and calculated weight loss were minimized.

Results and Discussion

Characterization of fuel

The proximate and ultimate analysis results are presented in Table 1. The result show that the rice husk was characterized by high volatiles content of about 66% which makes them desirable for a good regulation of gasification processes. The high percentage of ash in rice husk indicates potential slag or foul formation during combustion. However, the content of sulfur is very low, which means that there will be fewer emissions or corrosion whenutilization for power generation (Koufopanos et al., 1989; Rao et al., 1998).

Table 1. Ultimate and proximate analysis of rice husk

	analysis	Ultimat analysis				
(wt.%)	3.81	С	Н	0	N	S
Moisture	-	•	6.76	46.07	0.93	0.94
Volatile matter (VM)	66	43.83	0.70	40.07		
Fixed carbon (FC ^a)	22.01					
Ash	8.18					

Thermal decomposition characteristic

derivative and thermogram The thermogram (TG/DTG) curves in Fig. Iwere obtained for the pyrolysis of the rice husk at a heating rate of 10 °C/min. The TG/DTG curves represent the weight loss of the tested biomass with respect to temperature. The curves are generally stages divided into different three regardless of what biomass sample is tested. The first stage is drying and release of some light volatiles resulting into slight weight loss at temperature below 200 °C as shown in the TG curve. The second stage decomposition the thermal hemicellulose, cellulose and lignin where a significant weight loss was observed at a temperature between 200 and 400 °C.In the third stage, the weight loss occurs

slowly over wide range of temperature higher than 500 °C mainly due to thermal decomposition of heavy components. In the previous studies (Liangfeng et al., 2011; Vamvuka et al., 20011; Wang et al., 2012), DTG has been reported to qualitatively identifies the components of lignocellulose structures. Generally, hemicellulose decomposition occurs at temperatures between 150 and 350 °C; cellulose decomposition occurs at the temperatures range of 275-350 °C, and lignin is decomposed slowly over a wider temperature range of 250-500 (Vamvuka et al., 20011; Wang et al., 2012). The distribution of lignocellulose components in the DTG curve of rice husk is presented in Fig. 1.The DTG curve of rice husk exhibited only one peak. The main peak at temperature of 316 °C stems from cellulose decomposition and its shoulder peak in the DTG curve (which is almost merged to cellulose)is due to hemicellulose decomposition. The curve after the main peak (cellulose) is the decomposition of lignin whose intensity is smaller than that of the hemicellulose and cellulose peaks. It is important to note that one small peaks induced between the temperatures of 800°C and 850°C are due to the fact that rice husk includes another important component which is reactive at a higher temperature .In addition, from DTG curve in Fig. 1 it can be observed that two bumps developed at the temperature between 70 °C and 150°Cas a results of moisture removed from the rice husk.

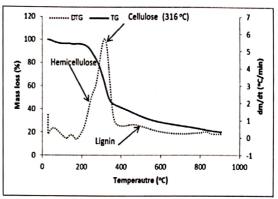


Fig. 1: TG/DTG curves of rice husk at heating rate of 10 °C /min showing the three lignocellulose components.

Kinetics analysis

Figure 2 and 3 shows TG and DTG curve of the modeling for the rice husk. Figure 2 shows that the model simulation gives prediction good compared experimental data for the TG curve of rice husk sample. The model simulation of the DTG curve (Fig. 3) also shows good prediction compared to the experimental Fig. 4 shows the individual contribution of each lignocellulose to the total decomposition rate. With regard to each lignocellulose biomass component, hemicellulose proved to be least stable component lignocellulose and it decomposes first and fast at low temperature. Lignin is more stable lignocellulose component decomposition takes place over wider temperature range. Finally, the cellulose decomposition started last and that is the reason why kinetic value has high values (Table 2). Among the lignocellulose component, cellulose shows the maximum decomposition. total Generally, the differences between the thermal behaviours of lignocellulose components are attributed to their different in the intrinsic chemical structures. Hemicellulose is least thermally stable

component due to its amorphous nature and cellulose is the next thermally stable than hemicellulose due its strong intramolecular bonds. Finally, Lignin is more thermally stable than the other two lignocellulose components and the reason for this is that it has strong cross-linked polymer (Pasangulapati et al., 2012; Xiong et al., 2015). The thermal behavior of each lignocellulose components agreed with other works in the literature (Jeguirim et al., 2001; Vamvuka et al., 2003).

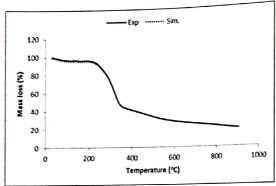


Fig. 2. Comparison of experimental and simulated TG pyrolysis profile of rice husk.

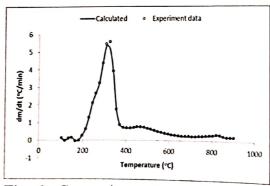


Fig. 3. Comparison of experimental and simulated DTG pyrolysis profile of rice husk.

Kinetic parameters are illustrated in Table 2. In the cellulose decomposition, kinetic constants appeared to have higher values than hemicellulose and lignin. Activation energy is the minimum energy required for each decomposition starts. Subsequently,

the higher its value is, the higher is the the figure for the initiation of each decomposition. Cellulose requires more compared to the other energy lignocellulose components because of its intra-molecular bonds, which strong decomposition in lower its prevent temperatures (Xiong et al., 2015). Preis another important exponential factor indicates constant indicate kinetic molecular collisions that lead to chemical reaction. A higher value of the preexponential factor, the more the molecular collisions that can lead to decomposition.

Lignin shows lowest values of activation energy and pre-exponential factor in Table 2. This fact could be observed from the (Fig. 4), where the curves DTG decomposition is seen to have started at lower temperatures and its decomposition rate was lower, compared to that of the other decomposition reactions. In term of thermal stability, lignin is the most stable component, therefore it is not easily decomposed. The observation made herein have been confirmed by previous studies (Xiong et al., 2015; Jeguirim et al., 2009).

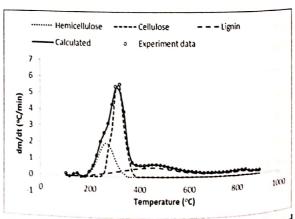


Fig. 4. Comparison of experimental and simulated pyrolysis profile of rice husk and rice husk components.

The calculated pyrolysis kinetic parameters of rice huskare presented in Tables 2. The activation energy of cellulose decomposition higher (187kJ/mol), whereas that of lignin decomposition lower (29kJ/mol). The activation energy of hemicellulose decomposition is intermediate (90 kJ/mol). The pre-exponential factors of cellulose, hemicellulose and lignin are in the ranges of 1.2×10^{15} , 1.6×10^{7} , $1.8 \times 10^{1} \text{min}^{-1}$, respectively.

Table 2. Content of lignocellulose and the kinetic parameters of the pyrolysis process.

		100	1
Component	Kinetic pa	arameters	
	C(%)	A(min)	E (kJ/mol)
Hemicellulose	26.9	1.6×10 ⁷	90
Cellulose	48.9	1.2×10 ¹⁵	187
Lignin	24.2	1.8×10 ¹	29

Conclusions

In this study, the thermal decomposition and pyrolysis kinetics were studied. The thermal decomposition shows that rice husk was divided into moisture removal, main devolatilization and continuous slight devolatilization stages. The mass loss process of rice husk was modeled by assuming cellulose, hemicellulose and lignin undergo pyrolysis independently and in parallel first-order reactions. The kinetic parameters of the three components were determined by means of Microsoft Excel Solver tool using least square algorithm. The simulation rate of thermal decomposition is very close to the experimental data.

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