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Optimization Pretreatment Condition of Sweet Sorghum Bagasse for Production of Second Generation Bioethanol

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Abstract. The bagasse residue of Sweet sorghum (Sorghum bicolor (L.) Moench) cons 19 of cellulose 39.48%; hemicellulose 16.56% and lignin 24.77% that can be converted to ethanol. Pretreatment is of great importance to ethanol yield. In this study, pretreatment process was conducted in a 5-liter reactor using NaOH 10% at various temperature 110, 130, 150°C and reaction time 10, 20, 30 minutes and optimizing severity parameter (log R₀ between 1.3 - 2.9). The statistical analysis using two way anova showed that third variations of temperature give different effects significant on lignin, hemicellulose and cellulose content at 95% the confidence level. The optimum pretreatment of bagasse sorghum were obtained with Log R₀ value between 2.4-2.9. High severity value in pretreatment condition reduce lignin almost 84-86%, maximum reducing lignin content was 86% obtained at temperature 150°C for 20 minutes reaction time and cellulose increased almost two times the initial content.

INTRODUCTION

Today bioethanol has become one of prospective biofuels in the transportation sector. Most of bioethanol produced in Indonesia currently is from molasses, or starchy raw materials, such as cassava and corn, known first generation bios hanol (G1). These materials are important raw materials for producing food and supporting food industries, so will not be sufficient to meet the increasing demand for the fuel ethanol [1]. On the other side, there are large amount of abundantly lignocellulosic biomass wastes of agricultural, estate crops and forestry industries that has not been used, can be converted to ethanol, called as second generation bioethanol (G2), is a possible can be can be converted to energy.

The bagasse residue after extracting the juice from the 5 weet sorghum stalk is lignocellulosic material that consist of cellulose hemicellulose and lignin, these material can be hydrolyzed into sugar and further fermented to ethanol [2]. On the conversion of lignocellulosic to ethanol, need three key steps i.e. pretreatme 2 to modify the lignocellulose structure, enzymatic hydrolysis to obtain sugars and fermentation process. Pretreatment of lignocellulose has received considerable research globally du 2 to its influence on the technical, economic and environmental sustainability of cellulosic ethanol production. For developing countries, alkali-based methods are relatively easy to deploy in decentralized, low-tech systems owing to advantages such as the requirement of simple reactors and the easy of operation.

Alkaline pretreatment is the target to reduce lignin content to increase enzyme access to holocellulose [3,4], reduction of cellulose crystallinity [5,6], and incre 17 in the surface area [7,8] and porosity [9,10] of pretreated substrates, resulting in increased hydrolysis rate. The mechanism of alkaline pretreatment is saponification of intermolecular ester bonds crosslinking 12 lan, hemicelluloses and other component [11,12].

To fully utilize bagasse residue of sweet sorghum as a feedstock for ethanol product 21 optimal pretreatment is required to render the cellulose fiber more amenable to the action of hydrolize enzyme. The main goal of this 22 rk is to study the effect pretreatment of alkaline NaOH 10% concentration, reaction time and temperature on the sweet sorghum bagasse as raw material for ethanol production, function of pretreatment time and temperature was investigated to optimize the operating parameter of pretreatment, while glucose production is the parameter for evaluating allow enzymatic performance.

MATERIALS AND METHODS

Materials

Bagasse of Sweet Sorghum (Sorghum bicolor (L.) Moench) called (BS) was obtained from PT. Panen Energy, Malang East Java, Indonesia. After air-dried, physical pretreatment i.e. chipping and milling until 3 mm was conducted to maximize contact area of the substrate.

Alkaline Pretreatment

In this work optimization process of pretreatment was conducted using bench scale reactor a 5-liter at the Research Centre for Chemistry, Indonesian Institute of Sciences (LII 1). 250 g of BS (10% moisture content) 7 as heated using NaOH 10% (kg/L) at 110,130 and 150°C for 10, 20 and 30 minutes. A solid liquid ratio was 1:5. The pressure was controlled four bars at early heating. After pretreatment, 11 material was pressed in order to separate solidity from liquid fraction. Solidity of BS treated was washed until 4 sh water turned to pH 7 and dried in the oven at 50-60°C overnight. The composition of materials component before and after pretreatment was analyzed acc 8 ling to National Renewable Energy Laboratory (NREL) standard procedures [13].

The treatment severity was quantified by a semi-empirical parameter called severity parameter, $\log R_0$, combining pretreatment time and temperature according to the equation Overend, 1987, [14].

$$R_0 = t.e^{\left[\begin{array}{c} 18 & 0 \\ 14.75 \end{array}\right]} \tag{1}$$

where t is the time in minute and T the temperature in degree Celcius.

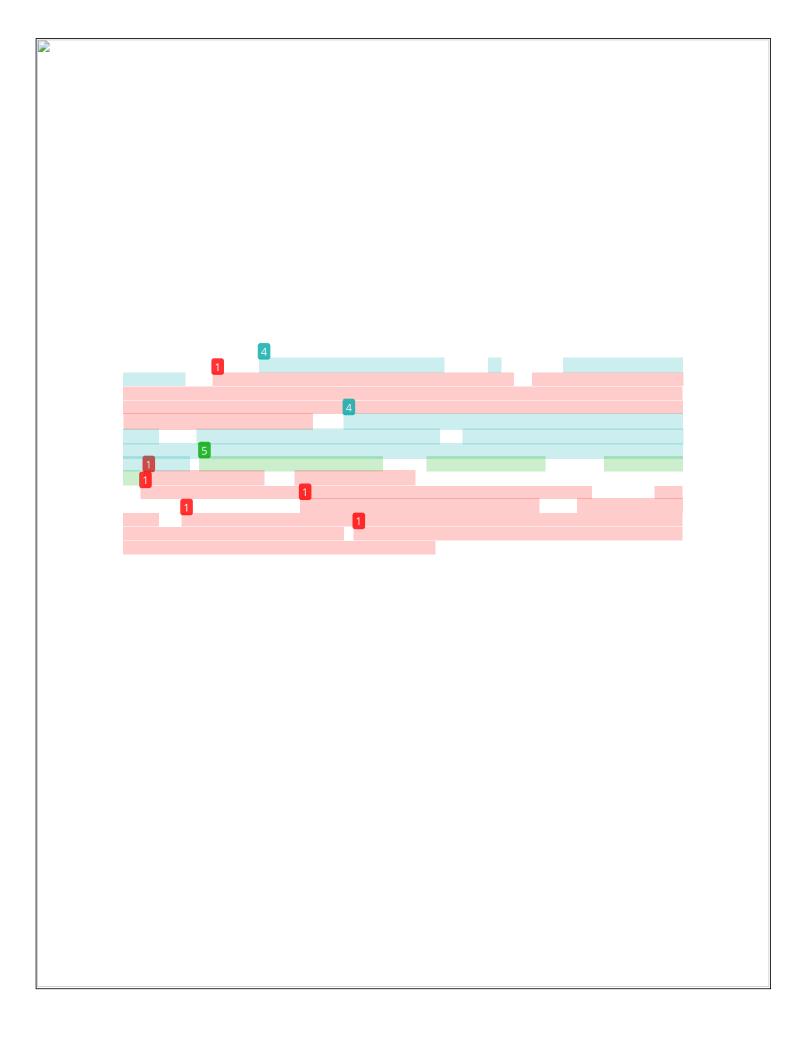


TABLE 2. Pretreated Samples Results

Pretreatm	nent	After Treatment	Weight Loss		Recovered		
Temperature (°C)	Time (min)	Charged Dry Matter (g)	(g)	Yield %	Biomass (%)		
110	10	150.27	99.73	39.89	60.11		
110	20	149.28	100.72	40.29	59.71		
110	30	103.51	146.49	58.60	41.40		
130	10	134.28	115.72	46.29	53.71		
130	20	149.34	100.66	40.26	59.74		
130	30	139.17	110.83	44.33	55.67		
150	10	154.43	95.57	38.23	63.44		
150	20	158.59	128.60	51.44	65.56		
150	30	163.89	86.11	34.44	63.44		

After pretreatment, showed that some pretreatment results in terms of pretreatment covered biomass in percentage compared to initial biomass charged into the reactor. Dilute NaOH application losens the biomass structure, separate the bonds between the lignin and the carbohydrates, increases the internal surface, decreases the degree of polymerization and crystallinity, and disrupts the lignin structure, that caused weight loss [15,16]. Highest recovered biomass was observed at temperature 150°C.

Composition components of Pretreated substrates

Before pretreatment the carbohydrate fraction of BS (cellulo 3 and hemicellulose) were 56.04% of total biomass, and the major component were cellulose (39.48%). Meanw 9e, lignin content of BS was 24.77%, comparable to the lignin content of empty of fruit bunch (25.83%) [17]. Lignocellulosic biomass requires pretreatment, mainly because the lignin in plant cell walls forms a barrier against enzyme attack. 11 deal pretreatment reduces the lignin content and crystallinity of the cellulose and increase surface area [18]. The loss of lignin in the pretreatment is one of the most important indicators of pretreatment effectiveness because the presence of lignin impedes enzymatic hydrolysis of the carbohydrates [15]. In order to de ade and reduce the lignin and non-cellulose component, BS was treated using alkaline solution NaOH 10%, nine samples were selected for the following stage, in the range Log R₀ 1.3-2.9. The effect of NaOH 10% at different temperature and different times pretreatment. After treatment at all condition, the BS was quantified for lignocellulose components, the results as shown in Tabel 3.

Pretreatment changed the BS composition. From Table 3 it can be 24 en during pretreatment delignification was occured and the process was effective for reduction of hemicellulose, so the proportion of lignin decressed, and the cellulose content after pretreatment was increased in all pretreatment condition. The results shows the optimum pretreatment condition was NaOH 10% at 150°C for 10-30 minutes, with the optimum loss of lignin and hemicellulose was 86%, and 553 respectively. Further increasing reaction time also affected in increase of cellulose content from 10 to 20 minutes and there was no significant effect reaction time after 20 minutes. Optimum cellulose content was 86.85% was achived at 150°C for 20 minutes. Comparing the two parameter, we observe an approximately constant trend which does influence of recovered biomass in line with cellulose content after treatment. The increase of cellulose content and the decrease of hemicellulose and lignin content can facilitate the process of enzymatic hydrolysis.

TABLE 3. Chemical Composition of Bagasse Sweet Sorghum

Pretreatment	Callulaca 0/	Hamicallulana 0/	Lionin 0/	A -1- 0/
Temperature (°C), Time (min)	Cellulose %	Hemicellulose %	Lignin %	Ash %
Untreated	39.48	16.56	24.77	4.20
110, 10	76.52	10.03	8.61	0.16
110, 20	78.06	9.94	8.31	0.32
110, 30	84.03	9.93	8.08	0.28
130, 10	81.74	8.87	6.16	0.25
130, 20	83.32	9.37	6.11	0.33
130, 30	88.25	8.57	5.45	0.55
150, 10	86.72	7.37	3.76	0.25
150, 20	86.85	7.45	3.51	0.52
150, 30	86.44	7.50	3.49	0.53

Further investigation for pretreatment parameters were carried out, using the analysis of two way ANOVA, to identify which parameter significantly affect the component cellulose, hemicellulose and lignin.

TABLE 4. ANOVA

Source	DF -	Lignin 25			Hemicellulose				Cellulose				
		SS	MS	F	Sig	SS	MS	F	Sig	SS	MS	F	Sig
Corrected Model	8	68.16	8.52	2479	0.00	21.43	2.68	419.25	0.00	268.83	33.60	100.89	0.00
Intercept	1	636.40	636.40	185200	0.00	1380.17	1380.17	216000	0.00	125646.25	125646.25	377200	0.00
Temperature	2	67.19	33.59	9774	0.00	20.74	10.37	1623	0.00	159.75	79.87	239.80	0.00
Time	2	0.73	0.36	105.51	0.00	0.20	0.10	15.83	0.00	68.67	34.33	103.08	0.00
Tempera- ture*Time	4	0.25	0.06	18.33	0.00	0.49	0.12	18.97	0.00	40.42	10.11	30.34	0.00
Error	9	0.03	0.00			0.06	0.01			3.00	0.33		
Total	18	704.60				1401.66				125918.08			
Corrected Total	17	68.19				21.49							
R Squared (Adjst. R Squared)		1.00 (0.999)			0.997 (0.995)			0.989 (0.979)					

Considering parameter condition, optimal pretreatment seem to move towards higher Log R_0 2.4. In these study best results in terms of raw material (cellulose content 39,48%) was obtained at temperatur 150°C. These results in line with the statistical analysis using two way anova showed that third variations of temperature give different effects significant on lignin, hemicellulose and cellulose content at 95% the confidence level.

Considering the overall process, optimum pretreatment of bagasse sorghum were obtained with Log R_0 value between 2.4-2.9. High severity value in pretreatment condition resulted high in reduce lignin almost 84-86%, maximum reducing lignin content was 86% obtained at temperatur 150°C for 20 minutes reaction time and cellulose increased almost two times the initial content.

Enzymatic hydrolysis of treated substrate

Enzymatic l 3 rolysis is a step after pretreatment for bioethanol production. The BS substrate treated by NaOH 10% at 150°C was subjected to enzymatic saccharification. This condition was selected based on the highest component of celullose of BS treated for 0, 20 and 30 minutes was 86.44-86.85% and the lowest component of lignin at 3.49-3.76%. The BS substrate was treated with commercial cellulase preparation at enzyme loading 30 FPU/g substrate. The result can be seen in Figure 1.

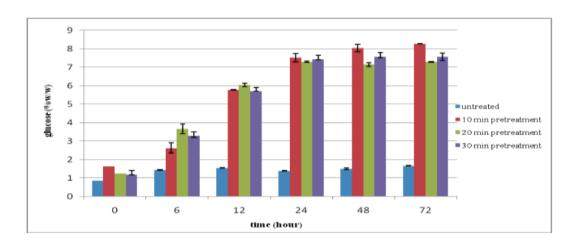


FIGURE 1. Glucose concentration from enzymatic hydrolysis of pretreated subtrates with alkaline pretreatment at 150°C and reaction time 10, 20 and 30 minutes

Figure 1 showed that glucose concentration from each substrates increased for 72 h of enzymatic hydrolysis. The treated substrates produced higher glucose concentration than untreated substrate. It is because of reduction of lignin content, reduction of cellulose crystallinity and increasing porosity of treated substrates. The highest concentration of glucose concentration can be obtained by pretreatment substrate at 150oC, 10 minutes. The optimum time for enzymatic hydrolysis was 48 h that produce glucose almost similar with 72 h, i.e. above 8% w/w. From the data can be conclude that alkaline pretreatment at 150oC, 10 minutes has been enough to produce high glucose concentration.

CONCLUSIONS

Considering the overall process, optimum pretreatment of bagasse sorgles in were obtained at Log R0 value between 2.4-2.9. Based on the results, pretreatment at 150oC gives the highest of cellulose content and the lowest of lignin content. The pretreatment condition of bagasse sorghum at 150oC for 10 minutes produces the optimal glucose concentration for bioethanol production.

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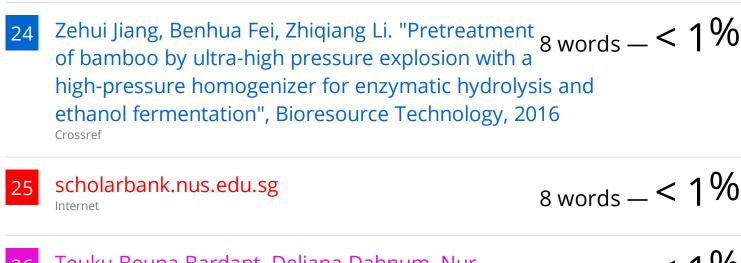
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