

# Original Research Article

## Effect of Dilute Acid and Alkaline Pretreatment of *Typha australis* (Typha grass) for Bioethanol Production

### ABSTRACT

*Typha australis* (Typha grass) obtained from Kware Lake was used in this research to produce bioethanol. Different pretreatment methods including dilute acid (0.2M H<sub>2</sub>SO<sub>4</sub>), dilute alkaline (0.2M NaOH) and liquid hot water pretreatments were used to pretreat the Typha grass sample before enzymatic saccharification for 7 days using *Aspergillus niger* and the hydrolysate was seeded with *Sacchromyces cereviceae* to produce bioethanol. HPLC was used to ~~determine~~ **analyse** bioethanol product. The result showed that pretreatment with 0.2M H<sub>2</sub>SO<sub>4</sub> removed more hemicelluloses (7.0%) when ~~compare~~ **compared** with other pretreatment methods used, but pretreatment with 0.2M NaOH and liquid hot water ~~remove~~ **removed** more lignin (14.29%) than dilute acid pretreatment. The highest percentage reducing sugar concentration of 0.58% was obtained from lower part of the sample pretreated with liquid hot water while Typha grass pretreated with 0.2M H<sub>2</sub>SO<sub>4</sub> and 0.2M NaOH produced the highest percentage reducing sugar concentration of 0.32% each from the upper part of the sample. Also, the highest Bioethanol concentration of 2.07% was obtained at day 6 of fermentation from the Typha grass pretreated with liquid hot water while Typha grass pretreated with 0.2M H<sub>2</sub>SO<sub>4</sub> and 0.2M NaOH produced highest Bioethanol concentration of 0.43% and 0.54% respectively. The results indicate that Typha grass can be harnessed for bioethanol production thereby reducing their negative impact on Lakes.

**Keywords:** Typha grass, Pretreatment, Bioethanol, HPLC,

### Introduction

The search for renewable biomass sources that do not serve as human food and animal fodder have focused primarily on plant biomass that possesses mainly cellulose and lignocellulosic materials [1]. Many agricultural waste in the form of plant materials such as grass, wood, crop

29 residues and some plants such as water hyacinth [2], ~~offers~~ **provide** the possibility of ~~–a~~  
30 ~~renewable energy and relatively greenhouse gas favoring source of sugars that can be converted~~  
31 ~~to~~ ethanol **sources**. The agricultural wastes are abundant and have disposal problems. An  
32 alternative is to utilize these agricultural wastes so that the competition between food and fuel  
33 can be minimized [3]. *Typha* grass has been identified as a particularly suitable biomass crop for  
34 wetland, because of their superiority, productivity, pest resistance, adaptability and chemical  
35 composition [4].

36 Kware lake serves a lot of functions to the nearby communities including a source of water for  
37 domestic activities such as washing and bathing. *Typha* grass have prolific vigorously, nearly  
38 impenetrable stand, getting to close the view of the water and take over the lake. This has raised  
39 concern that it may take over the lake and prevent a lot of activities from the lake. The present  
40 research is aimed at pretreating and hydrolyzing *Typha grass* for bioethanol production thereby  
41 adding more value to the grass, increase conservation and sustainability of the plant and lake,  
42 contributing towards alternative energy supply and also create profit and jobs opportunity.

## 43 **Methodology**

### 44 **Collection and Preparation of Samples**

45 The *Typha* grass sample was collected from Kware Lake in Kware local government area,  
46 Sokoto State, Nigeria. The sample was washed under tap water and then cut into two separating  
47 the upper part that grow above the water level and the lower part that grow inside water. The two  
48 portions were shade dried separately for 14 days. The dried samples were then grounded into  
49 powder and stored at room temperature for further analysis.

### 50 **Isolation and Identification of *Aspergillus niger* and *Saccharomyces cereviceae***

51 *Aspergillus niger* was isolated from soil sediment collected from kware lake and identified base  
52 on microscopic (morphological) and macroscopic characteristics (colour, texture appearance and  
53 diameter of colonies) according to Sourza-motta *et al.* [5] and *Saccharomyces cereviceae* was  
54 isolated from palm wine sample collected from Giginya Barrack market, Sokoto and identified  
55 by the standard morphological and physiological test and identification keys described by  
56 Barnett *et al.* [6].

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## 60 **Determination of Structural Composition of Typha Grass**

61 The percentage of acid – insoluble lignin, which is defined as the residue, was determined  
62 according to TAPPI procedure (T224 om-88). The holocellulose content, which is the  
63 combination of hemicellulose and cellulose, was determined in order to find the total amount of  
64 cellulose and hemicellulose in Typha grass. The holocellulose content was determined according  
65 to DIN 2403.  $\alpha$ -Cellulose is the pure cellulose content of the materials which was extracted from  
66 holocellulose using alkali solution. The  $\alpha$ -Cellulose content of Typha grass was determined as  
67 the residue insoluble in the 17.5 % NaOH solutions according to TAPPI 203 om-93 method.

### 68 **Dilute Acid Pretreatment**

69 Approximately 2 g of *Typha* grass sample was mixed with 0.2M H<sub>2</sub>SO<sub>4</sub> solution in a 250 ml  
70 flask with a stopple and then allowed to suck for 24hrs. The mixture was subsequently autoclave  
71 at 121 °C for 15mins and then cooled and filtered through a Whatman filter paper to separate the  
72 solid residue. The residue was washed with distilled water until neutral pH. The sample was air  
73 dried and stored in tightly sealed plastic bag in a refrigerator for further use [7].

### 74 **Dilute Alkaline Pretreatment**

75 Approximately 2 g of *Typha* grass sample was soaked in 0.2M NaOH solution in a 250 mL flask  
76 for 24hrs and then autoclave at 121 °C, for 15min. The solid residue was separated from the  
77 mixture by filtration with Whatman filter paper and thoroughly washed with distilled water to  
78 neutralize its pH. Finally, the filtrate was dried and stored as above [7].

### 79 **Liquid Hot Water Pretreatment (LHW)**

80 In liquid hot water pretreatment of the *Typha* grass sample, the grounded powdered of the plant  
81 was slurried with distilled water using a solid to liquid ratio of 10% (w/w) and autoclaved at 121  
82 °C for 15 min. After autoclaving, the sample was filtered and the solid residue was air dried and  
83 stored for further use [8].

### 84 **Enzymatic Saccharification of Pretreated Typha Grass**

85 The enzymatic saccharification of *Typha* grass was carried out using *Aspergillus niger* isolated  
86 from soil sediment as described by Gupta [9]. In this method, the pretreated *Typha* grass samples  
87 were inoculated with 0.5 ml suspension of 96 hours culture of *Aspergillus niger*. Hydrolysis was  
88 carried out at room temperature for 7 days. Samples were taken daily for reducing sugar

89 determination using 1,4-dinitro salicylic acid (DNS) method to find out the net yield of  
90 fermentable sugars. The samples were then filtered using Whatman filter paper No. 1 and the  
91 filtrates were used for fermentation.

## 92 **Fermentation of the Hydrolysate and Bioethanol Production**

93 The fermentation studies were carried out using *Saccharomyces cerevisiae* isolated from palm  
94 wine. The hydrolysates were autoclaved at 121 °C for 15 min and the flasks were then cooled to  
95 room temperature. The pH of the fermentation medium was adjusted to 6.5 and then 1ml of  
96 prepared suspension of yeast isolated was added in the hydrolysate [10]. The fermentation was  
97 allowed for 7 days and samples from the medium were withdrawn periodically at 24 hrs interval  
98 from the flasks to determine ethanol quantity using UV-visible spectrophotometer.

## 99 **Distillation**

100 The fermented broth was filtered using whatman filter paper. Each sample was weighed into  
101 Microkjeldahl flasks and then heated at 78 °C on the Microkjeldahl apparatus until the solution  
102 turned colourless. The presence of bioethanol was determined using high performance liquid  
103 chromatography (HPLC).

## 104 **Results and Discussion**

### 105 **Structural Composition of Typha Grass Obtained from Kware Lake Before Pretreatment**

106 The structural composition of both the upper and lower part of Typha grass before pretreatment  
107 show that, hemicelluloses has a highest composition of 28.70% and 30.00% while Lignin has a  
108 composition of 28.15% and 24.93% respectively, and  $\alpha$  cellulose has least composition of  
109 12.30% and 10.00% respectively in the Typha grass collected from Kware Lake. The result of  
110 structural composition of hemicelluloses and lignin is in conformity with the report of Bajpai  
111 [11] who reported that hemicelluloses compose between 35-50% and lignin compose 10-30% of  
112 grasses. However, this current study disagrees with his report that  $\alpha$ -celluloses compose of 25-  
113 40%. This research shows that the upper part of Typha grass has higher structural composition  
114 than the lower part of the plant (Table 1).

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118 **Table 1:** Structural Composition of Typha Grass Obtained from Kware Lake Before  
 119 Pretreatment

<b>Contents</b>	<b>Sample A (%)</b>	<b>Sample B (%)</b>
Holocellulose	41.00	40.00
Hemicelluloses	28.70	30.00
$\alpha$ cellulose	12.30	10.00
Lignin	28.15	24.93

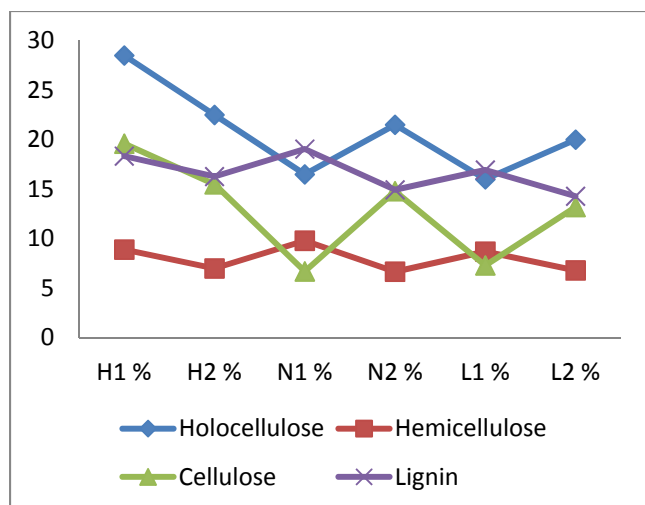
Key: Sample A: Upper part of Typha grass that Grows above water level

Sample B: Lower part of Typha grass that grow inside water

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121 **Structural Composition of Typha grass Obtained from Kware Lake after Pretreatment**

122 Pretreatment of Typha grass with 0.2M H<sub>2</sub>SO<sub>4</sub> produced lowest lignin content of 16.28%  
 123 cellulose content of 15.5% and hemicelluloses content of 7.0%. Pretreatment with 0.2M NaOH  
 124 produced lowest lignin content of 14.94% cellulose content of 13.8% and hemicelluloses content  
 125 of 7.67%. Typha grass sample pretreated with liquid hot water has lowest lignin content of  
 126 14.29% cellulose content of 12.2% and hemicelluloses content of 7.8%. The result indicates that  
 127 0.2M H<sub>2</sub>SO<sub>4</sub> remove more hemicelluloses content than the other pretreatment process (figure 1).  
 128 This is in agreement with the result of many other researches such as [4] and [13]. Mosier *et al.*,  
 129 [12] also reported that hemicelluloses are removed when dilute H<sub>2</sub>SO<sub>4</sub> is added and this enhances  
 130 digestibility of cellulose in the residual solid. Also, dilute acid pretreatment was not good for  
 131 lignin removal when compare with dilute alkaline and liquid hot water pretreatment. Chang *et*  
 132 *al.*, [14] reported that Alkaline pretreatment removes amorphous substances lignin, which  
 133 increases the crystallinity index of lignocellulosic materials.



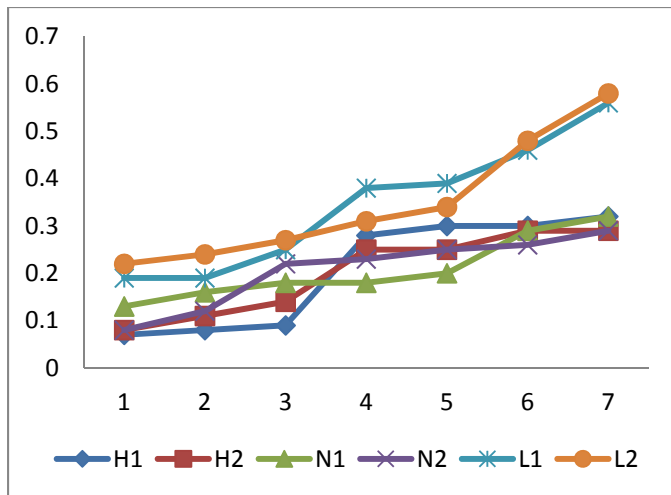
Key: H1: Upper part of Typha grass pretreated with 0.2M H<sub>2</sub>SO<sub>4</sub>  
 H2: Lower part of Typha grass pretreated with 0.2M H<sub>2</sub>SO<sub>4</sub>  
 N1: Upper part of Typha grass pretreated with 0.2M NaOH  
 N2: Lower part of Typha grass pretreated with 0.2M NaOH  
 L1: Upper part of Typha grass pretreated with liquid hot water  
 L2: Lower part of Typha grass pretreated with liquid hot water

134

135 Figure 1: Structural Composition of Typha Grass after Pretreatment with 0.2M H<sub>2</sub>SO<sub>4</sub>, 0.2M  
 136 NaOH, and Liquid Hot Water

137 **Reducing Sugar Concentration of Typha Grass**

138 Pretreatment with 0.2M H<sub>2</sub>SO<sub>4</sub>, 0.2M NaOH and liquid hot water produces highest reducing  
 139 sugar concentration of 0.58% at day 7 when liquid hot water was used to pretreat 2g of the  
 140 sample (figure 2). Chemicals used in this research produce small quantity of reducing sugar. This  
 141 agrees with the work of Arumugan and Manikandan, [8] who said significant sugar production  
 142 was not recorded from pretreatment with dilute chemicals, but the current studies disagree with  
 143 their findings that said more reducing sugar from dilute acid pretreatment was produce than the  
 144 liquid hot water pretreatment. Also, the production of low reducing sugar in the present study  
 145 from dilute chemicals can be attributed to the washing of the chemical after pretreatment in order  
 146 to neutralize their pH. It can also be as result of the solubilization of the carbohydrate by the  
 147 chemical during pretreatment [13]. According to Ogier *et al.*, [15] and Laser *et al.*, [16], liquid  
 148 hot water pretreatment can be a promising pretreatment method that present elevated recovery  
 149 rates of sugars which does not generates inhibitors.



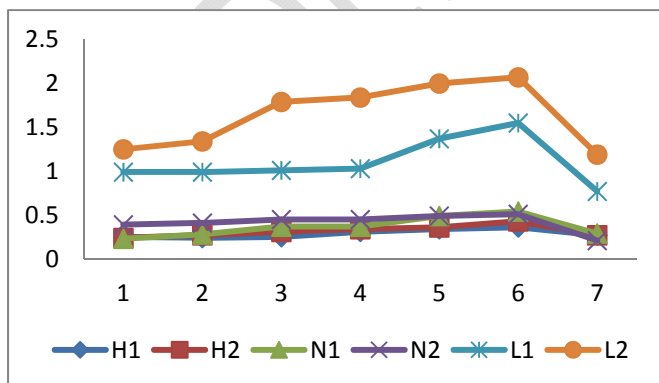
Key: H1: Upper part of Typha grass pretreated with 0.2M H<sub>2</sub>SO<sub>4</sub>  
 H2: Lower part of Typha grass pretreated with 0.2M H<sub>2</sub>SO<sub>4</sub>  
 N1: Upper part of Typha grass pretreated with 0.2M NaOH  
 N2: Lower part of Typha grass pretreated with 0.2M NaOH  
 L1: Upper part of Typha grass pretreated with liquid hot water  
 L2: Lower part of Typha grass pretreated with liquid hot water

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151 Figure 2: Results of Reducing Sugar Produce after Pretreatment of 2g of Typha Grass with  
 152 Liquid Hot Water Pretreatment, 0.2M H<sub>2</sub>SO<sub>4</sub> and 0.2M NaOH

153 **Bioethanol concentration of Typha grass**

154 A highest bioethanol concentration of 2.07% was produced from the sample that was pretreated  
 155 with liquid hot water and sample pretreated with dilute NaOH and H<sub>2</sub>SO<sub>4</sub> produces highest  
 156 bioethanol concentration of 0.54% and 0.43% respectively (figure 3). The result of ethanol yield  
 157 from dilute NaOH and H<sub>2</sub>SO<sub>4</sub> pretreated sample is almost the same with 0.5% reported by Fish  
 158 *et al.*, [17]. Also, Grous *et al.*, [18] reported that 90% efficiency of enzymatic hydrolysis was  
 159 achieved in 24 h for poplar chips pretreated by liquid hot water, compared to only 15%  
 160 hydrolysis of untreated chips.



Key: H1: Upper part of Typha grass pretreated with 0.2M H<sub>2</sub>SO<sub>4</sub>  
 H2: Lower part of Typha grass pretreated with 0.2M H<sub>2</sub>SO<sub>4</sub>  
 N1: Upper part of Typha grass pretreated with 0.2M NaOH  
 N2: Lower part of Typha grass pretreated with 0.2M NaOH  
 L1: Upper part of Typha grass pretreated with liquid hot water  
 L2: Lower part of Typha grass pretreated with liquid hot water

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162 Figure 3: Percentage Concentration of Bioethanol Produce from the Sample after Pretreatment  
 163 with Liquid Hot Water, 0.2M H<sub>2</sub>SO<sub>4</sub> and 0.2M NaOH

164 **Conclusion**

165 From the result of this research, it has been find out that pretreatment using dilute acid and  
166 alkaline were very effective in removing lignin and hemicelluloses from typha grass.  
167 Pretreatment with dilute acid removed hemicelluloses by 77% which proved to be better than  
168 dilute alkaline which removed hemicelluloses by 75%. Dilute alkaline removed more lignin in  
169 typha grass by 60% than dilute acid pretreatment which removed lignin by 35%. However,  
170 cellulose content of sample pretreated with dilute H<sub>2</sub>SO<sub>4</sub> increased by 59% which is higher than  
171 50% increase by sample pretreated with dilute NaOH. Also, sample pretreated with dilute NaOH  
172 produce highest bioethanol concentration of 0.54% than H<sub>2</sub>SO<sub>4</sub> which produce 0.43%.  
173 Determination of bioethanol by HPLC conclude that bioethanol can be produced from Typha  
174 grass and pretreatment of the sample can increase accessibility to cellulose and increase reducing  
175 sugar production thereby increasing bioethanol production.

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