

---

# EFFECT OF CONCENTRATION AND TIME OF ALKALI TREATMENT ON THE MOISTURE CONTENT AND REGAIN OF NIGERIAN SISAL FIBRE

C.S. EZEANYANASO AND E.S NWADIOKWU  
POLYMER AND TEXTILE DIVISION

CHEMICAL, FIBRE AND ENVIRONMENTAL TECHNOLOGY DEPARTMENT  
FEDERAL INSTITUTE OF INDUSTRIAL RESEARCH, OSHODI, P.M.B. 21023, IKEJA,  
LAGOS, NIGERIA.

[esnwadiokwu@gmail.com](mailto:esnwadiokwu@gmail.com)

## ABSTRACT

*The effect of concentration and time of alkali treatment on the moisture content and regain of Nigerian sisal fibres has been studied. The sisal fibres were chemically modified using different concentrations of caustic soda (4%, 8%, 12% and 16%) while varying the treatment time (4mins, 8mins and 12mins). Treated and untreated samples were investigated for their moisture properties. The moisture regain and moisture content of the fibres were substantially influenced by the alkali treatments which increased with increasing concentration at constant time. The increase in time of treatment at constant concentration had less significant effect on the moisture properties investigated.*

**Keywords:** Concentration, alkali treatment, moisture content, moisture regain, sisal fibres

## 1.0 INTRODUCTION

Sisal fibre is a hard fibre extracted from the leaves of sisal plant (*Agave sisalana*). Though native to tropical and subtropical North and South America, sisal plants are widely grown in tropical countries of Africa, the West Indies and the Far East (Bisanda et al., 1994).

Tanzania and Brazil are the two main producing countries (Chand et al, 1998). Using sisal fibre as reinforcement in composites has raised great interest and expectations among material scientists and engineers (Bledzki et al, 1996). However, sisal fibre reinforced composites generally have poor interface (Bledzki et al, 1999).

Moisture content and regain of the sisal fibres determine to a large extent the nature of the fibre surface and the surrounding matrix. Several fibre surface treatment methods have been studied to improve the adhesion properties between sisal fibres and a surrounding matrix. An effective method includes alkali treatment to improve the moisture properties of fibre.

In this research, alkali treatment was employed to chemically modify sisal fibres using different concentrations of NaOH (4%, 8%, 12% and 16%) and at different times (4mins, 8mins and 12mins). The effects of this modification on the moisture properties were assessed and analyzed.

## 2.0 MATERIALS AND METHOD

### 2.1 Materials

The main materials for this study are sisal fibres obtained from the Botanical garden of the Ahmadu Bello University, Zaria.

### 2.2 Chemicals

Aqueous sodium hydroxide solution (4, 8, 12 and 16% by weight) and acetic acid (2% by weight).

### 2.3 METHOD

#### 2.3.1 Extraction of the Sisal Fibres

The leaves were crushed and beaten manually by a smooth edged stick so that only fibres remain. After extraction, the fibres were washed thoroughly in plenty of water to remove surplus wastes such as chlorophyll, leaf juices and adhesive solids (hemicelluloses). The fibres were then dried in open air. Dried sisal fibre strands are usually creamy white in colour.

#### 2.3.2 Chemical Modification of the Fibres

5 grams of the sisal fibres were soaked in 4, 8, 12 and 16% NaOH solution for 4, 8 and 12 minute. These fibres were further rinsed with water followed by neutralization in 2% acetic acid solution. A final rinse in water and then dried at room temperature.  $\text{Sisal-OH} + \text{NaOH} \rightarrow \text{Sisal-O}^- \text{Na}^+ + \text{H}_2\text{O}$

#### 2.3.3 Determination of Moisture Content and Moisture Regain

The sisal fibres were weighed and dried in an oven at a temperature of 103°C for 30 minutes, followed by cooling for 30 minutes and then weighed again. This step was repeated until the

weight was constant. The moisture regain of the test specimen was expressed as a percentage loss in weight of the final oven-dry weight using the following equation;

$$\text{Moisture Content} = \frac{W_0 - W_1}{W_1} \times 100 (\%)$$

Where  $W_0$  is the weight of fibre before dried in oven and  $W_1$  is weight of fibre after dried in oven. Also moisture regain was calculated using the following formula (Booth J.E, 1968);

$$R = \frac{M}{1 + \left[ \frac{M}{100} \right]}$$

Where  $R$  =Moisture regain,  $M$ =Moisture content. The effects of concentration and time on the moisture regain and moisture content were determined for both the modified and unmodified sisal fibres. Ten tests were carried out and the mean values are reported in table 1.

### 3.0 RESULTS AND DISCUSSION

#### 3.1 Moisture Content and Regain

Usually, alkali modification causes appreciable changes in the moisture absorption for all sisal fibres. Modified sisal has higher regain compared to the unmodified ones. It can be seen from table 1 that increases in the concentration of NaOH at constant time, increases the moisture content and regain as compared to the unmodified fibre. This increase may be due to the fact that increase in concentration of NaOH, increases the accessible surface of the sisal fibre due to increase in the amorphous region owing to the swelling of the fibres (Fengel and Wengener, 1983).

Moreover, alkali treatment increases the number of possible reactive sites and allows better fibre wetting. The effect of alkali on sisal fibres is a swelling reaction, during which the natural crystalline structure of the cellulose relaxes. Native cellulose shows a monocyclic crystalline lattice of cellulose I which can be changed into different polymeric forms through chemical treatments (Wenyanberg *et al.*, 2006).

Increase in the concentration of alkali will influence the degree of swelling and hence the degree of lattice formation into cellulose II (Fengel and Wengener, 1983). Studies have shown that  $\text{Na}^+$  has got a favourable diameter, able to widen the smallest pores in between the lattice planes and penetrates into them. Consequently, NaOH treatment results to higher degree of swelling (Wenyanberg *et al.*, 2006). However, increase in time of treatment at constant concentration does not have much effect on the moisture properties. The increase in alkali concentration at constant time and increase in time of modification at constant concentrations are shown in the tables 1 and 2.

**Table 1: Increase in Alkali Concentration at Constant Time**

	<b>Moisture Content (%)</b>	<b>Moisture Regain (%)</b>
<b>Untreated</b>	6.90	7.80
<b>4% for 4mins</b>	9.15	10.30
<b>8% for 4mins</b>	9.95	11.10
<b>12% for 4mins</b>	10.89	12.09
<b>16% for 4mins</b>	11.90	13.10
<b>4% for 8mins</b>	9.60	10.86
<b>8% for 8mins</b>	10.76	11.95
<b>12% for 8mins</b>	11.78	13.03
<b>16% for 8mins</b>	12.68	13.90
<b>4% for 12mins</b>	10.58	11.55
<b>8% for 12mins</b>	11.64	12.06
<b>12% for 12mins</b>	12.73	13.08
<b>16% for 12mins</b>	13.53	14.81

**Table 2: Increase in Time of Alkali Treatment at Constant Concentration**

	Moisture Content (%)	Moisture Regain (%)
<b>Untreated</b>	6.90	7.80
<b>4mins at 4%</b>	9.15	10.3
<b>8mins at 4%</b>	9.60	10.86
<b>12mins at 4%</b>	10.58	11.55
<b>4mins at 8%</b>	9.95	11.10
<b>8mins at 8%</b>	10.76	11.95
<b>12mins at 8%</b>	11.64	12.06
<b>4mins at 12%</b>	10.89	12.09
<b>8mins at 12%</b>	11.78	13.03
<b>12mins at 12%</b>	12.73	13.08
<b>4mins at 16%</b>	11.90	13.10
<b>8mins at 16%</b>	12.68	13.90
<b>12mins at 16%</b>	13.53	14.81

## CONCLUSION

Treated and untreated samples were investigated for their moisture properties. The moisture regain and moisture content of the fibres were substantially influenced by the caustic soda treatments which increased with increasing concentration at constant time. The increase in time of treatment at constant concentration had less significant effect on the moisture properties investigated.

## REFERENCES

- Bisanda E. T. N. and Ansell, M. P. (1994). *Journal of Material Science*, pp. 27.
- Bledzki, A.K., Reihmane, S. and Gassan, J. (1996). *Applied Polymer Science*, 5, pp.1329
- Bledzki, A. K., and Gassan, J. (1999). Natural Fiber Reinforced Plastics, in *Handbook of Engineering Polymeric Materials*, ed. N. P. Cheremisinoff, Marcel Dekker, New York, pp. 787–810.
- Chand N., Tiwary, R. K. and Rohatgi, P. K. (1998). *Journal of Material Sciences*, pp.23
- Fengel, D. and Wegener G. (1983). *Wood: Chemistry, Ultrastructure, Reactions*, de Gruyter, Berlin, 482.
- Wenyanberg, I.V., Truong T.C., Vangrimde, B. and Verpoest. (2006). *Compos. A*, 37, 1368