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Modification of Shea Butter Wood Fiber for Fiber-Polymer Composites in Automotive Interior Cabinets

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Abstract

The limitations of renewable material obtain from agro-based in the manufacturing sectors indicates much prospect to substitute synthetic fibers and non-metal compounds which are the traditional materials employ at esteem years old. The request of novel organic fiber, that are price friendly, minimal-mass-volume ratio and commonly available, that is reason shea butter wood flour (SBWF) employs in this work and competes with existing fibers for alternative innovative material to solve mankind issues. This research involved determination for the composition of crude and pre-treated SBWF. The SBWF was crushed and powered. The SBWF was sieved to a specific size. The SBWF was processed under controlled parameters: concentration of NaOH from 5, 10, 15 %w/v and immersed time 2, 4, 6 hr. Before modification, the crude compositional constituent of the SBWF was obtained based on cellulose (CLC), lignin (LGC) and hemicelluloses (HMC). Then, the SBWF was pretreated at the variant concentration ant soaking period. The chemically compositions of the chemically modified SBWF was obtained and tabulated. The Fourier transform infra red (FTIR) and scanning electron microscopy (SEM) to examine the active radicals and morphology were imposed on the crude and pre-treated SBWF at ultimate condition. From the displayed outcome, the raw compositional constituents were 48.5 %, 13.27 % and 24.76 % for CLC, LGC and HMC, respectively. On the other note, the pre-treated SBWF showed a significant improvement in CLC, LGC and HMC at the variation of concentration and time. The FTIR and SEM examination showed displacement in the position of apex for the spectrum and more roughness of the micrograph due to the aid of chemical influence of the SBWF, respectively. Finally, at optimum state of 10 % w/v NaOH and 4hr immersion period, the CLC, LGC and HMC were 75.15 %, 10.28 and 4.97 %, respectively. The result exhibited by CLC from the novel SBWF is a promising engineering component for automobile parts.

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1.0 Introduction

Non-metallic substances as such as Na₂CO₃, CaCO₃, MgO, etc., were formerly used material for composite production, however, the use of these inorganic materials are very cost effective, and this led to a need to source out a cheap material from agro-plant to minimize the cost of inorganic material in composite production (Weyenberg et al., 2006; Government et al., 2017; Government et al., 2018). Generally, composite is the combination of two/ more with separate individual components characteristics yield a to single compound/mixture with qualities varies from each of component substances (Wong et al., 2010; Atuanya et al., 2014; Government & Onukwuli, 2025). It has large parts of it taking as matrix which can be polymer/ceramics/metals while the other constituents found in minute form which might be fiber which is either natural fiber/non-metals/ and coupling agents (Wong et al., 2010 Government & Onukwuli 2025). It has the distinguished following properties (Government et al., 2019; Government et al., 2020a & b); Okeke et al.,2023): it is lighter in weight, as result of this merit, it is applicable for automotive parts in fuel energy conservation, and saves the cost of materials. These also improve the specific strength, elastics stiffness, flexural strength, modulus, hardness and impact energy, etc. It can be utilized in many areas of work such as aerospace, building, electronic components, boats, railway coaches, etc. This material also acts as a perfect replacement for metals or alloys in manufacturing sector and generates urge amount money in terms material rice (Wong et al., 2010; Obasi, 2015; Government & Onukwuli, 2025).

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In this current era, plant cellulosic substances have been the main raw materials in the emerging industry for novel fiber-plastic (Obasi, 2015). composites The major productions which these can serve as purely casing, substitute are phone automobile components, door panel, containers packaging material, etc. These agro-based fibers have totally replaced non-metals from the soil and rocks for finished products. Agro-based plant yields major constituent which defines as CLC that provide strength support to the plantpolymer composite (Bogoeva-Gaceva et al., 2007; Netral et al., 2012; Wang et al., 2016; Srinivas et al., 2017). Numerous research have shown that plant-dependent-fiber with large quantify CLC is essential for fiber-polymeric composite (Bogoeva-Gaceva et al., 2007; Netral et al., 2012; Dungani et al., 2016; Government et al., 2017; Government et al., 2018), most importantly, lignin and hemicellulose face reduction to keep away from the issues of incompatibility when fully utilized commercialization (Wu et al., 2000; Rowell et al., 2000; Nachtigall et al., 2007; Setswale et al., 2017). The challenge of compatibility is quite pertinent and needs to be resolved for avoidance of the final end product to experience water absorption with time (Haristov & Vasileva, 2003; Sari et al., 2017).

Today, industrialists have emphasized on the substitution of synthetic fiber to organic fiber as a result of their uncommon characteristics such as insulating capacity, light weight, heat resistance, modulus, biodegradable and cost. Due to these qualities, manufacturers have also informed researchers over the world to beef up



studies for this area of research for generation of billions of dollars to them (Jawaid and Khalil 2011; Marwan & Nasim, 2016; Muryanto *et al.*, 2015).

Cellulose generates the apex components of agro-based-plant substance with hemicelluloses and lignin in correspondence, respectively (Netral et al., 2012; Dungani et al., 2016; Shuhaida and Soh, 2016). The power of cellulosic material is proportional to the characteristics of the composite material when used its constituent (Bogoeva-Gaceva et al., 2007; Netral et al., 2012; Suryanto et al., 2013). has confirmed that Research cellulosic component enhances high strength of polymer composite (Bogoeva-Gaceva et al., 2007; Netral et al., 2012; Dungani et al., 2015; Government et al., 2017; Government et al., 2018). Modification of plant - based fiber is predominantly to cut out weaken gummy constituents from the fiber which include lignin, hemicellulose and other compound that is connected the fiber (Wu et al., 2000; Rowell 1998; Nachtigall et al., 2007; Johar et al., 2012). This invariably resists the influence of incompatibility and water absorption utilization of domestic and industrial needs of the composite time (Haristov and Vasileva, 2003; Government et al., 2017; Government et al., 2018).

In this day and age, scholarly researches (Valadez-Gonzalez et al., 1999; Ray et al., 2002; Herrera-Franco and Valadez-Gonzalez 2005; Wong et al., 2010) have proven that lignocellulose compounds acknowledge chemical variation of lignin and a cellulosic fiber boosts their collaboration with polymeric arrangements. Multiple fibers previously employed after pre-treatment for mass production to solve engineering problems, such as pinewood (Chen et al., 2014), rice straw (Shuhaida & Soh 2016), waste betel nut (Lazim et al., 2014), century fiber (Reddy et al., 2013) sugarcane base (Ahmadi et al., 2016). However, SBWF has not been systematically studied for automotive composites, due to its scarcity in scientific literal work.

In this study, a novel SBWF, which is low-cost, biodegradable, and widely available, was chemically modified using NaOH, with consideration given to treatment time and concentration. The FTIR and SEM analysis were investigated on the pure and purified SBWF at ultimate condition. This is to ascertain its cellulosic constituent for its accessibility for composite production in automobile interior.

2.0 Materials and Methods

2.1 Materials

Material collection

The SBWF was derived from Wapan Nghaku in Wukari Local Government Area of Taraba State. The SBWF was washed with de-ionized H_2O , sun dried for 11 days at 8 hr before crushing and sieved using 12 mesh (1700 μ m) sieve size. The SBWF fine grain was stored in a sealed container at room temperature.

2.2 Methods

Pre-treatment process of SBWF

15 g of the crushed and sieved SBWF was soaked in NaOH solution with concentration of the alkali varied from 5-15 %w/v. The solid part of the SBWF was washed at pH 7. The time of soaking was also varied from 2-6 hr after which CLC, HMC, and LGC were examined, respectively.

Determination of CLC and LGC

The crude and chemically prepared SBWF were analyzed for its component's constituent at varying concentrations and time utilizing the Chesson's method (Mahyati *et al.*, 2013; Agu *et al.*, 2014; Government *et al.*, 2019; Government *et al.*, 2020 a, b; Okeke *et al.*, 2023) with the procedures:

(K) 1g of SBWF and 150 ml distilled H₂O was warm to attain at 100°C for 1hour and the weight recorded.



- (L) The mixture (SBWF at 150 ml of distillation H₂O) was dried, filtered, and weighed until the weight was constant.
- (M) The ruminant in B was mixed with 150 ml of 0.5 M H₂SO₄ was passed through heating for 100 °C at 1hr. The fusion was filtered, dried and weighed till reading is fixed
- (N) The residue was mixed with 10 ml of 72 %w/v H₂SO₄ to maintain normal temperature for 4 hr and then subject to heating for 1 hr and weighed.
- (O) The final remains were further subjected to heat until it turned to ash and weighed. The CLC and LGC were estimated as viewed in Eq. 1 and Eq. 2, respectively.

$$CLC(\%) = \frac{(M-N)}{K} \times 100 \tag{1}$$

LGC (%) =
$$\frac{(N-0)}{K} \times 100$$
 (2)

Determination of HMC

2 g of the SBWF deposit was injected in the beaker and 10 ml aqueous NaOH was discharged in the containing material. The mixing was maintained at a vibration of 300rpm with the aid of orbital shaker with the alkali solution. The alkaline solution was field frequently to maintain at 20°C. Distilled H₂O of 33 ml was introduced in the container and placed at 1 hr. The weight of the ruminant after the process was evaluated as (L), The remains in A was move into a crucible where 100ml NaOH was introduced again followed by 200ml distilled H₂O, and 15ml CH₃COOH. The total mixture was finally dehydrated to the weight (L) to determine the SBWF HMC employing Eq. 3. The cubicle with the residue was dried and weighed (L), the HMC of SBWF was computed using Eq. (3).

$$\frac{\text{HMC(\%)} = }{\text{residue weight (K)- residual weight in cubicle (L)}}$$

$$\frac{\text{residue weight (K)}}{\text{residue weight (K)}}$$
(3)

FTIR Investigation

The virgin and the alkalized SBWF with the best components composition were examined by Shimadu 8400s FTIR facility. The specimen to be injected into this facility involved combination of 50mg of KBr + 1.5mg NLWBF. The combined compound was pulverized, passed into a die for formulation of pellet which was moved into the machine for scrutinization to obtain the FTIR spectrum. The equipment has TGS detector, placed in a frequency of 0.1 cm⁻¹ and the spectra were measured at 4 cm⁻¹ precision varied from 800 to 3500 cm⁻¹.

SEM examination

1 g of crude and alkali-purified in each specimen was ground and transferred in the SEM equipment. After a period of 5 s, the micrograph was generated. The model employed was Phenon ProX with energy of 15 KeV.

3.0 Result and Discussion

The characteristic components of SBWF as captured in Table 1. It was noticed that the CLC content for crude SBWF was 48.50 %. The CLC of SBWF was proportional to that of kenaf (45-57 %) (Taj et al., 2007; Agu et al., 2014), oilpalm (44.20-49.60 %) (Abdul-khalil et al., 2006; Dungani et al., 2016) as stated by previous study. Also, the SBWF HMC content was observed to be 13.27 %. SBWF HMC showed close connection with that of jute (12.55-13.65) (Mwaikambo et al., 2002; Dungani et al., 2016), and ramie fiber (13-16 %) (Hattallia et al., 2002; Wambua et al., 2003; Hoareau, 2004). Furthermore, the SBWF LGC content was observed to be 24.76 %. This is also similar to other obtainable fiber such as bamboo fiber (21-31%) (Hattallia et al., 2002; Hoareau, 2004), sugarcane (24.35-26.30%) (Wambua et al., 2003; William & Patrick 2017, Hemmasi et al., 2001; Dungani et al., 2016), oil-palm (Abdulkhalil et al., 2006; Dungani et al., 2016). In the view of the fact that the composition of SBWF demonstrated parallel properties with that of other accessible natural fiber; SBWF is proposed for automotive parts etc. (Dungani et al., 2016; Agu et al., 2014).



Table 1: Individual components of raw SBWF

Wood Fiber	CLC (%)	HMC (%)	LGC %)
SBWF	48.5	13.27	24.76

The consequence of NaOH on the CLC of SBWF is described in Figure 1. It was observed that as the NaOH concentration increased from 0 to 15% w/v, the CLC of SBWF was enhanced from 48.5 to 78.43 % at soaking time of 2 hr. The percentage increase between the unrefined and refined **SBWF** the at these variant concentrations was found to be 29.93 % at this resident time. This trend occurs during immersion of SBWF in alkali solution, the bonds that hold the constituent of the fiber were broken. This allows some of the unneeded component to remove from SBWF as an impurity in the immersion solution (Oushabi et al., 2017). Also, when the resident time for the absorption of SBWF further increases from 4hr and 6hr, the CLC of the fiber improves when compared than the purified fiber from 48.5 to 72.58 % and 48.5 to 65.62 %, respectively. Furthermore, as the absorption time of SBWF in NaOH solution increase, the CLC reduces.

Notwithstanding, at 4 hr of soaking, the CLC increases from 70.68 to 73.58 % and decreases from 75.15 to 73.58 % when the concentration increases from 5 to 10 % and reduction can be seen from 10 to 15 % increase of NaOH concentration in SBWF absorption solution. Similar observation was recorded at 6 hr soaking time with total reduction of the cellulose from 67.74 to 65.62 %. The occurrence as a result of at higher concentration of NaOH increment in SBWF immersion, part of the cellulosic composition of the fiber has been annihilated (Oushabi et al., 2017; Palamae et al., 2017). However, the optimum CLC was viewed at 75.15 % which corresponds at the concentration of NaOH 10 % wt. and treatment time of 4hr. it can be concluded that, any further increment in concentration of NaOH than this concentration, will lead to reduction of the CLC of SBWF than this concentration and time. This analysis studied toe the trend of earlier works (Government et al., 2019; Oushabi et al., 2017; Palamae et al., 2017; Shuhaida & Soh 2016; Barlianti et al., 2015).

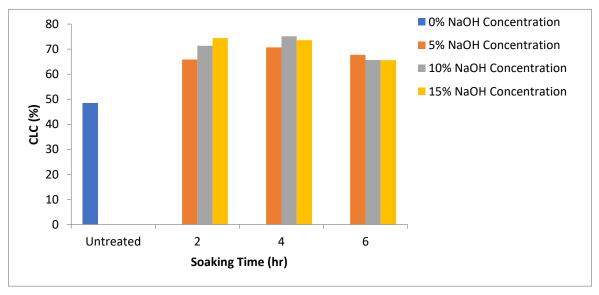


Figure 1: Influence of NaOH pre-treated variables on the CLC of SBWF

Figure 2 illustrates the effect of treatment parameter on the HMC of SBWF. From Figure 2, a noticeable removal of the HMC of SBWF was felt when the concentration of SBWF

improves from 0 to 15 %. At this state, the HMC was reduced from 13.27 % to 5.26 % at a constant time of 2 hr. The efficiency of the SBWF HMC reduction between the crude and



the pre-treated fiber was found to be 60.36%. This may be attributed after modification of SBWF in the solution of NaOH, the extractive and other additional components including the HMC has been removed to the solution of soaking. In this process, the SBWF has been fully activated by hydroxyl group that led to the introduction of alkoxide in composition of the fiber (Oushabi *et al.*, 2017; Palamae *et al.*, 2017). Similar observation was recorded at 4hr and 5hr when the immersion time increases. For

4hr and 5hr, the percentage removal between the unmodified and chemically modified SBWF was discovered to be 50.94 and 31.42 %, respectively. In conclusion, it can be stated that the percentage removal of HMC increases at the absorption time of soaking; this is as a result of too much soaking time reduces the HMC removal of SBWF. Scholarly studies have displayed similar trend (Government et al., 2019; Oushabi *et al.*, 2017; Palamae *et al.*, 2017; Shuhaida and Soh 2016; Barlianti, *et al.*, 2015).

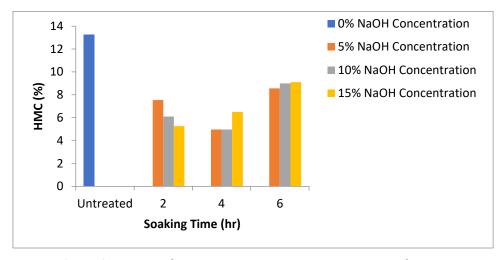


Figure 2: Impact of NaOH treatment parameter on HMC of SBWF

Figure 3 shows the effect of treatment parameter on the LGC of SBWF. From this Figure, there was drastic removal of the LGC from SBWF when the concentration of SBWF improves from 0 to 15 %. At this state, the LGC was reduced from 13.27 % to 5.26 % at a soaking time of 2 hr. The effectiveness of LGC removal between the pre-treated and the unpurified SBWF was found to be 56.74475. This occurs after purification of SBWF in the solution of NaOH, extra components including the LGC has been removed into the soaking solution.

From the process, the SBWF has been fully activated by a hydroxyl group which leads to the introduction of alkoxide in content of the fiber (Oushabi *et al.*, 2017; Palamae *et al.*, 2017). Similar observation was recorded at 4 hr and 5 hr when the soaking time increases, respectively. For 4 hr and 5 hr, the removal percentage between the unpurified and chemically treated

SBWF was observed to be 58.36026 % and 28.19063 %, respectively. The best LGC removal was observed at 15 % wt and 4 hr soaking time. The values of LGC between the modified and unmodified SBWF abridged from 24.76 to 10.31 % at that condition. This corresponds to preceding studies by other research fellows ((Brinchi *et al.*, 2013; Ahmadi *et al.*, 2016; Government *et al.*, 2019; Oushabi *et al.*, 2017; Palamae *et al.*, 2017; Shuhaida and Soh 2016; Barlianti *et al.*, 2015).

The FTIR of unpurified and purified SBWF is represented in Figure 4 (a-b). The peak at 3414.58 cm⁻¹ is a character of OH compound in the unmodified SBWF fiber as observed in Figure 4a. In Figure 4b, the same peak was situated at 3439.28 cm⁻¹ with a reduction in the transmittance from 106 to 93.5 %; this is an influence of the NaOH on the composition of SBWF. In the same vein, the peak from the



spectrum for the untreated as observed between (2882.01 to 2818.05 cm⁻¹) is a (-COOH-) bond found in the SBWF. This reappeared in the purified SBWF between the peaks 2994.19 to 2808.51 cm⁻¹ maintaining reduction in the transmittance which is a further reason for the deduction of unwanted components in the SBWF. The presence of (P-OH) compound was crested at 2352.03 cm⁻¹ in Figure 4a after NaOH purification, this peak was noticed to be invisible in Figure 4b. The pyridine compound was traced

at 1612.42 cm⁻¹ in the untreated spectrum. For the pre-treated SBWF, the compound was found with a functional group of (C=N) and (C=C) at this time maintaining a jump in the transmittance. Also, at 1379.71 cm⁻¹ is an indication of (C=benzene ring stretching), this is due to the presence of HMC and pectin in the SBWF before treatment. In the pre-treated SBWF as situated in Figure 4b, the compound was unnoticed due to the influence of NaOH on the fiber.

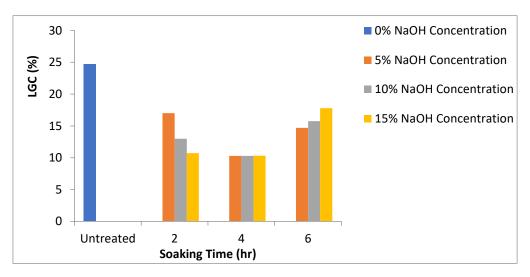


Figure 3: Influence of NaOH treatment on values of LGC of SBWF

At 1200 cm⁻¹ in the Figure 4a, the peak indicates the presence of LGC and also 1107.74 cm⁻¹ shows C-O-C indicating the presence of CLC in the pre-treated SBWF (Ikramullah et al., 2018). When the SBWF was modified by NaOH, the LGC was found to shrink thereby increasing the CLC as cited at the apex of 1108.11 cm⁻¹. The frequency of (798.06 and 987 to 745.88 cm⁻¹ with ⁻C-H branch) was captured in crude and the purified SBWF which were the evidence of CLC, respectively. The C-Cl active groups of alkyl halide were also observed in 618.35 cm⁻¹ and (618.73 to 521.73 cm⁻¹) for the pure and the mercerized SBWF, respectively. Finally, the movement of the spectrum peak, departure of the active radicals of some compounds between the unmodified and the modified SBWF exposes the alleviation of HMC and LGC under the control of NaOH which amplified the CLC. This corresponds to prior studies by scholarly journals (Ikramullah *et al.*, 2018; Oushabi *et al.*, 2017).

The SEM micrograph for un-modified and chemically modified SBWF is illustrated in Figure 5 (a-b) above. Smooth array of SBWF particulates was found in the SEM micrograph. Also, more pore space of SBWF was also traced when the fiber is not modified. This leads to lower reaction site when the SBWF is to be used reinforcement to another material. As it was observed in Figure .5b, the particles of the fiber clustered to each other. This leads to better interlocking when the SBWF is used with a substrate for production of new material. This is as a result of after pre-treatment with NaOH, the reaction site when the SBWF is mixed with a substrate in manufacturing of a new material for industrial application has increased; thereby creating



better adhesion with the substrate. When compared with untreated SBWF. Therefore, pretreated SBWF will have lower water absorption

and higher strength. Close result was analyzed by previous authors (Oushabi, *et al.*, 2017; Reddy *et al.*, 2013).

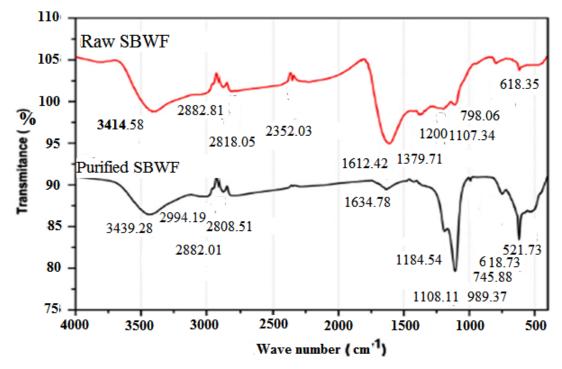


Figure 4: FTIR spectra of SBWF (a) raw (b) after treatment at 10 %w/v NaOH and 4 hr soaking time.

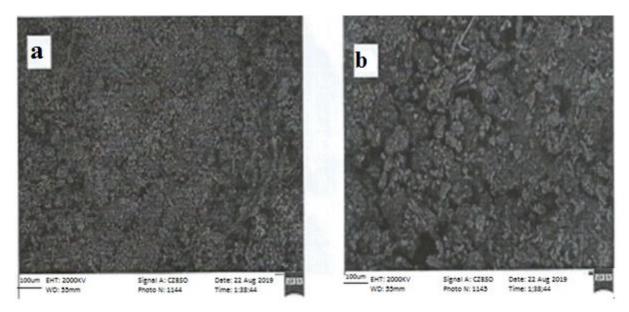


Figure 5: SEM diagram for (a) raw (b) purified SBWF at 10 %w/v and 4 hr

4.0 Conclusion

The research on the composition for the unrefined and refined novel SBWF with the action of NaOH at distinguished concentration and period has been developed. The variation of time and concentrated SBWF enhanced the CLC, diminished LGC and HMC, respectively. The increase in cellulose and reduction in lignin

and hemicelluloses were confirmed on the FTIR and SEM, respectively. The higher concentration of NaOH and moderate exposure due to soaking of SBWF yielded optimal CLC and minimal LGC and HMC, respectively. With this out, the SBWF is a novel component for polymeric-material-fiber composite for car parts production due to amplified amount of CLC.



Recommendations

The novel low priced SBWF is recommended for the constituent of fiber-polymer composite for interior door of a car cabinet.

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