Journal of Materials Science Research and Reviews

Effect of Surface Treatment on the Mechanical Properties of Epoxy Filled with Dates Palm *(Phoenix dactylifera)* **Particulates Composite**

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Authors' contributions

This work was carried out in collaboration among all authors. The work is part of a Ph.D research work conducted by author SAK under the supervision of author AD with the assistance of authors USI and BMD. All authors read and approved the final manuscript.

Article Information

Editor(s): (1) Dr. Yang Qu, Associate Professor, Key Laboratory of Functional Inorganic Material Chemistry, Ministry of Education of the People's Republic of China, Heilongjiang University, China. *Reviewers:* (1) Antonio L. Beraldo, Campinas University, Brazil. (2) J. Dario Aristizabal-Ochoa, Universidad Nacional de Colombia, Colombia. Complete Peer review History: http://www.sdiarticle3.com/review-history/50781

Original Research Article

Received 02 June 2019 Accepted 11 August 2019 Published 17 August 2019

ABSTRACT

Treatments of lignocellulosics material have been found to be useful in preparation of polymer composites since such treatment help to improve their properties. Epoxy filled with untreated and 5% NaOH treated date particulates of particle size 150 µm composites were prepared with filler loadings ranging from 10 wt% to 50 wt% using hand layup techniques. Mechanical properties of the composite samples were determined in accordance with ASTM standards. Tensile and flexural tests were measured with the aid of universal testing machine in accordance with ASTM D3039 and ASTM D790 standard tests for tensile and flexural respectively, for polymer composite. SEM morphology and water absorption were equally studied. Results showed that the introduction of 10 wt% filler loading of the untreated and treated date filler reduced the tensile strength of the unfilled epoxy resin by 20.2% and 11.1% respectively. The date pits (DTP) composites gave maximum and minimum tensile strength values of 29.35 MPa at 10 wt% of date pits/epoxy (DTP/EP). Composites produced from the treated filler showed appreciable properties that is better than the untreated filler and can be used to produce particleboard, interior parts of automobile, ceiling and tiles in building application.

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Keywords: Lignocellulosics; composites; particulates; tensile strength; flexural strength.

1. INTRODUCTION

Quest for a better life makes man to always research and think of the best way life of comfort could be achieved. As a result, most research and development are focusing on the development of new composite materials. The use of polymer matrix composite has found wide application in our modern day world. This is because of the combination of properties which these materials possess. Some of the properties of polymer matrix composites include specific strength, high modulus, good fracture and fatigue properties as well as corrosion resistance [1]. The need for polymer composite with better properties increases the quests of researcher to employ several ways in which properties of such composite could be enhanced. One of such way is chemical treatment such as sodium hydroxide treatment. The interfacial adhesion between the filler and polymer has a determining influence in the mechanical properties of composites; however, to improve mechanical properties through the use of fillers is by treating them with coupling agents besides by changing its particular size. i.e improvement of interfacial bonding can be achieved by addition of coupling agent, that is compactibilizer and by changing the particles size. The ability to convert date palm pits filler into useful engineering materials with a better quality sharpened the focus of this present research work.

Date palm is a monocotyledonous plant belonging to the Arecaceae family, cultivated in dry tropical regions worldwide for its edible sweet fruit. It contains a single seed (kernel) about 2– 2.5 cm long and 6–8 mm thick. The kernel is a major by-product of the date palm-processing industry. They contained 7.1–10.3% moisture,

5.0–6.3% protein; 9.9–13.5% fat; 46–51% acid detergent fibre; 65–69% neutral detergent fibre; and 1.0–1.8% ash. Date pit is mainly used as animal feed [2].

Plate 1. Date palm pits

Epoxies are thermosetting resin materials characterised by two or more oxirane rings or epoxy groups within their molecular structure. The commonest epoxy resin is the diglycidyl ether of bisphenol A (DGEBA), which is prepared by the reaction of epichlorohydrin (ECD) and bisphenol A (BPA). ECD is prepared from polypropylene (PP) by reacting chlorine with sodium hydroxide [3]. Thus, epoxy resins are available in various consistencies from low viscous liquid to a tack-free solid [4].

Epoxies are among the most important classes of thermosetting polymer which are widely used as matrices for fibre-reinforced composite materials and as structural adhesive. Epoxies are amorphous, highly cross-linked polymer and this structure result in the materials possessing various desirable properties such as high tensile strength and modulus, uncomplicated processing, good thermal, chemical and corrosion resistance, and dimensional stability [5]. The resins equally possess outstanding adhesion properties, low shrinkage upon cure and good electrical properties [6].

Scheme 1. Reaction for the synthesis of DGEBA-type epoxy resin

Chemical treatments have been done to improve adhesion or interfacial bonding between natural fibres and synthetic polymers [7]. These treatments will inevitably enhance the basic properties of natural fibres reinforced polymer composites [8,9]. Alkaline treatment, bleaching, acetylation and steaming are such various processes applied to improve fibre matrix interaction [10,11,12]. The primary drawback of the use of natural fibre is the lower processing temperature due to the possibility of lignocellulosic degradation, lack of good interfacial adhesion and the poor resistance to moisture absorption. Studies have shown that the effect of moisture absorption on mechanical properties of composites can be improved modifying reinforcing fibres by chemical treatment. The coupling agent contains chemical groups, which can react with the fibre and the polymer. The bonds formed are covalent and hydrogen bonds that improve the interfacial adhesion. The use of compatibilizers, the surface modification techniques such as treatments, acetylation, graft co-polymerisation or the use of maleic-anhydride–polypropylene copolymer has been reported to overcome the incompatible surface polarities between the natural fibre and polymer matrix. The influence of surface treatments of natural fibres on the interfacial characteristics was also studied [13]. Investigations that have been carried out on the use of natural fibres/fillers with polymeric materials are endless, and several others researches have shown the various properties such as physical, mechanical, thermal, water absorption etc, obtained from the natural fiber/
filler composites. Thus, the use and filler composites. Thus, the use and enhancement of these date pit fillers will add more information to those provided in the literature.

2. EXPERIMENTAL

2.1 Materials

Epoxy Resin (commercially available epoxy resin $(3554A)$ of density 1.17 $g/cm³$ and Polyamine amine (Hardener3554B) of density 1.03 g/cm³ were procured from a local supplier in Ojota-Lagos, Nigeria. The date palm fruits were obtained from Gwagwalada market in Gwagwalada, Area Council, F.C.T; Nigeria.

2.2 Methods

The date pits (DTP) were separated from their fruits manually, thereafter; they were washed,

cleaned, dried and grounded with hammer mill to obtain filler powder. The fillers were made to pass through wire mesh screen to obtain particle size of 150 μm. The particulate fillers were thereafter modified by alkali treatment. The treated fillers were obtained by soaking in 5% NaOH for 4 hrs and thereafter washed with water, followed by neutralising the basic solution with few drops of acetic acid and carefully monitored using litmus paper.

The fillers were then oven dried for 24 hrs at temperature of about 70° C before use to reduce the moisture content. Samples were thereafter stored in a sealed container prior to compounding.

2.3 Compounding

The composites with varying degrees of filler percentage (i.e. 10 wt%, 20 wt%, 30 wt%, 40 wt%, & 50 wt%) were prepared. This was achieved by mixing the various filler ratios with the epoxy to form homogenous blends. The mixing was achieved via manual stirring method for 10 min. The volume ratio of resin to hardener was 2:1, and after thorough mixing with the filler, the resin was poured onto the cavity of glass mould of dimensions 160 mm x 70 mm x 4.5 mm overlaid with aluminium foil to serve as releasing agent. The mixture was allowed to cure at room temperature for 24 hrs before removal from the mould. Neat resins without filler were equally prepared to serve as control. The block composites were thereafter machined into shapes for analyses.

2.4 Characterization

Determination and evaluation of mechanical properties through tensiles test (ASTM D3039), flexural test (ASTM D790), water absorption capacity and surface morphorlogy of the prepared polymer composites.

2.5 Tensile Test

The test was carried out on a Universal testing machine (TIRA test 2810) with maximum load of 10 KN in accordance with ASTM D3039. Samples with dimension 140 mm, 20 mm, and 4.5 mm of length, breadth and height respectively were used for the test. A cross – head speed of 2 mm/min was used. Tensile strength and tensile modulus were expressed as:

Tensile strength (MPa) =
$$
\frac{P}{bh}
$$

Where,

 $P =$ Pulling force (N) , b = Specimen Width (m) $h =$ Specimen thickness (m)

Tensile modulus (MPa) = $\frac{\sigma}{s}$

Where, σ = Stress (N/m²) ε = Strain

Five specimens for each composite were tested and statistical average for each set of results was recorded.

2.6 Flexural or 3-point Bending test (ASTM D790)

The 3-point bending test was carried out on a universal testing machine (TIRA test 2810) as shown below with maximum load of 1 KN in accordance with ASTM D790. A cross speed of 5 mm/min was used with sample dimension 140 mm, 20 mm by 4.5 mm i.e length, breadth by height respectively. The flexural strength of composites was found using the following equation.

$$
F
$$
lexural strength =
$$
\frac{3FL}{2bd^2}
$$

Where,

 $F =$ Maximum load applied on test specimen (N) .

 $L =$ Length of support span (mm),

 $b =$ Width of specimen tested (mm),

d = Thickness of specimen tested (mm)

The flexural modulus can also be found using the following equation:

Flexural modulus =
$$
\frac{ML^3}{4bd^3w}
$$

Where,

- M =Maximum load applied on test specimen (N)
- $L =$ Length of support span (mm)

b = Width of Specimen,

d = Thickness of specimen

w = Deflection at maximum force.

2.7 Water Absorption Test (ASTM D570)

The specimens for the test were machined into 40 mm, 20 mm and 4.5 mm of length, breadth and thickness respectively. Three specimens for

each composite were tested and the average values were taken for each composite. The specimens were dried in an oven for 4 hrs at 70° C and then placed in desiccators to cool for 2 hours. Immediately, upon cooling the specimens were weighed. The materials were then immersed in water at room temperature of 25 °C for 24 hours. Specimens are removed, patted dry with a lint free cloth, and weighed immediately in an analytical weighing balance to the nearest milligram. The hydration capacity of the composites was then monitored for period of 32 days.

2.8 Surface Morphology

The fractured surface after the flexural test of composites was examined using an SEM microscope (PHENOM PRO X). Prior to the analysis, the specimens were placed on a stub and were coated with a thin layer of gold using a sputter coater to avoid charging under the electron beam. The inner conditions of the scanning barrel were vacuumed to prevent interferences of scanning picture due to the presence of air. Magnification, focus, contrast and brightness of the result were adjusted to produce the best micrographs.

3. RESULTS

3.1 Tensile Test

The results of the effect of filler loading on the tensile strength and modulus of epoxy filled with untreated and treated date pits particulate composites are depicted in Fig. 1.

3.2 Elongation at Break

The results of the effect of filler loading on the elongation at break of epoxy filled with untreated and treated date pits particulate composites are shown in Fig. 2.

3.3 Flexural Test

The results of the effect of filler loading on the flexural strength and modulus of epoxy filled with untreated and treated date pits particulate composites are depicted in Fig. 3.

3.4 Water Absorption

The effect of time on hydration capacity of epoxy/untreated and treated date pits particulate composite are illustrated in Figs. 8 and 9, respectively.

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Fig. 1. Effect of filler loading on the tensile strength and modulus of NaOH treated and untreated date pits/ epoxy composites

Fig. 2. Effect of filler loading on the percentage elongation at break of NaOH untreated and treated date pit/ epoxy (DTP/ EP) composites

Fig. 3. Effect of filler loading and particle size on the flexural strength and modulus of treated and untreated date pits/ epoxy composites

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Fig. 4. Water absorption curves of untreated date pits/ epoxy composite

Fig. 5. Water absorption curves of treated date pits/ epoxy composite

Plate 2. SEM micrograph of the fracture surface of 10% date pits filler /epoxy composite at 1500x magnification

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Plate 3. SEM micrograph of the fracture surface of 10% treated date pits filler /epoxy composite at 1500x magnification

Plate 4. SEM micrograph of the fracture surface of 10% date pits filler/epoxy composite at 5000x magnification

Plate 5. SEM micrograph of the fracture surface of 10% treated date pits filler /epoxy composite at 5000x magnification

Study of the morphology of the composites was carried out and the results are as shown in plates 2 to 5. The scanning was done on the fracture surface of the composites.

4. DISCUSSION

Fig. 1 shows the effect of filler loading on the tensile strength and modulus of untreated and

treated date pits/ epoxy composites. It can be seen generally that the tensile strength decreases with increase in filler loading. The minimum and maximum value was observed for treated sample at 50 wt% and 10 wt% filler ratio with 15.44 MPa and 29.35 MPa respectively. The NaOH treatment improved the tensile strength of date pits/ epoxy (DTP/EP) composite by 11% and 15% at 10 wt% and 50 wt% filler loading, respectively. The higher mechanical properties of the samples due to chemical modification were an indication of improved interaction and stress transfer between the particles and the binder.

The modulus of the treated sample also showed a corresponding increase as the filler loading increases. The modulus of treated date pits/epoxy was at maximum and minimum value at 50 wt% and 10 wt% with 0.85 GPa and 0.72 GPa respectively, compare to the neat resin value of 0.67 GPa.

The figure compares the effects of filler loading on the tensile strength and modulus of the treated and untreated filler of date pits/ epoxy (DTP/EP) composites. The maximum tensile strength of untreated date pits/ epoxy (DTP/EP) composite increased from 26.34 MPa to highest value of 29.35 MPa; On the other hand, the minimum value increased from 13.44 MPa to 15.44 MPa when the filler was treated with 5 % NaOH at the same filler loading. The addition of 10% wt and 50% wt ratio of untreated date pits filler into epoxy resin reduces the tensile strength by 20% and 59% respectively, while the flexural strength was reduced by 5.2% and 32.7%, respectively, as shown in Fig. 3. Incorporation of the filler into the resin reduces the ductility of the resin. However, treatment of the filler with 5 % NaOH helps to improve such mechanical properties. The treatment improved the tensile strength of untreated DTP/EP at 10 % filler loading by 11.4%. In addition, the treatment equally improved the ductility of the composites. For example, elongation at break of DTP/EP at 10%, 20% and 50% improved by 10.5%, 10.2% and 9.9%, respectively as shown in Fig. 2.

The higher tensile strength of the samples due to chemical modification was an indication of improved interaction and stress transfer between the particles. The treatment on the filler helps to reduce the effect of impurities such as oil and fat on the interaction between matrix (resin) component and filler thereby improving the stress

transfer process that will ultimately culminate into higher strength as seen in Figs. 1 and 3. The improved strength of the composite as a result of sodium hydroxide treatment, can also be linked with reduction in lignin content of the filler, as lignocellulosic material are originally composed of cellulose in lignin matrix, reduction of the natural matrix (lignin) in the filler will help the resin to bind better with the cellulose in the filler, thereby giving an improved strength.

On the other hand, the modulus increased as the filler loading increases. However, it can be seen from the Fig. 1 that the modulus of the untreated sample composite improved greatly at 40 wt% filler loading more than that of the treated composite sample. Modulus of untreated date pits/ epoxy composites at 40 wt% was 1.28 GPa while that of treated date pits/ epoxy composite at the same filler weight percent was 1.09 GPa.

4.1 Water Absorption Capacity

The percentage hydration of untreated date particulate / epoxy at room temperature of about 25° C has the following values after 24 hrs of absorption as shown in Fig. 8: 1.13%, 1.16%, 1.44%, 1.58%, and 1.91% at 10 wt%, 20 wt%, 30 wt%, 40 wt%, 50 wt% filler loading, respectively, while the unfilled epoxy resin gives 0.789%. The test shows that the absorption continues to increase daily and after 768 hours (32 days), the following values were obtained: 3.82%, 4.71%, 6.48%, 7.56%, and 9.82% for the corresponding 10 wt%, 20 wt%, 30 wt%, 40 wt%, and 50 wt% filler weight content, respectively. The daily absorption is primarily due to the hydrophilic nature of the lignocellulosic filler. The unfilled epoxy reached maximum absorption value of 1.61% after 648 hours. However, the rate of absorption for all the composites was at maximum after the first 24 hrs of absorption. In comparison to the result shown in Fig. 8, it was possible to conclude that treated DTP/EP composite is stronger as shown in Fig. 9, that gives 1.13%, 0.97%, 1.21%, 1.28%, and 1.51% at 10 wt%, 20 wt%, 30 wt%, 40 wt%, 50 wt% filler loading, respectively.

4.2 Morphology

Plates 2 and 3 reveals the state of dispersion of 10% of untreated and treated date pits particulate in epoxy composites (DTP/EP) at 1500x and magnification, respectively. It can be

observed in the SEM micrograph in plate 3 that the filler dispersed uniformly in the matrix and a strong interfacial bonding exits between the filler and the resin except the line cracks seen. Thus, the line cracks can be as result of manual mixing employed during fabrication. Also, plates 4 and 5 further shows the interaction of 10% of the untreated and treated filler with the epoxy resin at higher magnification. From the results, it can be seen that the interfacial bonding between the filler and matrix was higher in plate 5 that might be due interaction between filler and the resin as a result of the filler treatment [14]. Plate 4 equally showed the presence of pulled out traces, voids which are indicative of weak interfacial adhesion at the interface which further confirmed the reduced tensile and flexural strength observed in the untreated filler composite.

5. CONCLUSION

Date pits particulates have been used successfully as fillers in the preparation of epoxy composites. The addition of the filler increased the bulk of the composite. Properties such as tensile and flexural modulus, hardness were improved while properties such as tensile strength and flexural strength were
affected negatively. Incorporation of 10% Incorporation of 10% untreated filler into epoxy resin improved the tensile and flexural modulus of DTP/EP composites by 8% and 1.6%, respectively.

The addition of 10% wt and 50% wt ratio of untreated date pits filler into epoxy resin reduces the tensile strength by 20% and 59%, respectively, while the flexural strength was reduced by 5.2% and 32.7%, respectively. Incorporation of the filler into the resin reduces the ductility of the resin. However, treatment of the filler with 5% NaOH helps to improve such mechanical properties. The treatment improved the tensile strength of untreated DTP/EP at 10 wt% filler loading by 11.4%, Also, the treatment equally improved the ductility of the composites thereby increasing the impact strength. For example, elongation at break of DTP/EP at 10 wt%, 20 wt% and 50 wt% improved by 10.5%, 10.2% and 9.9%, respectively.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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