

Characterization of Coconut Shell Ash for Potential Utilization in Metal Matrix Composites for Automotive Applications

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ABSTRACT

Coconut shell ash is agricultural waste. The waste is produced in abundance globally and poses risk to health as well as environment. Thus their effective, conducive and eco-friendly utilization has always been a challenge for scientific applications. This paper mainly deals with identification of characteristics of coconut shell ash using spectroscopic and microscopic analysis. Density, Particle size, Refractoriness, SEM, XRD, XRF and FTIR spectroscopic methods were used for the characterization of the coconut shell ash. The results were compared and it was observed that the ash possesses nearly same chemical phases and other functional groups as reinforcement like fly ash, rice husk ash, bagasse ash that have been in Metal Matrix Composites (MMCs) specifically for automobile applications. Hence, coconut shell ash can be used as a low cost reinforcement in Metal Matrix Composites (MMCs).

Key Words: Density; microstructure; particle size; refractoriness

I. INTRODUCTION

Researches all over the world today are focusing on ways of utilizing, either industrial or agricultural wastes as a source of raw materials for the industry. These wastes utilization would not only be economical, but may also result to foreign exchange earning and environmental pollution control [1-2]

Metal matrix composites (MMCs) possess significantly improved properties including high specific strength, specific modulus, damping capacity and good wear resistance compared to unreinforced alloys [2-4]. Similarly, there has been an increasing interest in composites containing low density and low cost reinforcements. Among various discontinuous dispersoids used are fly ash, red mud, Rice husk ash [1-5] are some of the most inexpensive and low density reinforcement available in large quantities as solid waste by-product.

Coconut shell is an agricultural waste and is available in very large quantities throughout the tropical countries of the world. Moreover, coconut is becoming an important agricultural product for tropical countries around the world as a new source of energy-biofuel [6]. Previously, coconut shell was burnt as a means of solid waste disposal which contributed significantly to CO₂ and methane emissions [6]. However as the cost of fuel oil, natural gas and electricity supply has increased and become erratic, coconut shell has come to be

regarded as a source of fuel rather than refuse. Presently, the Nigeria coconut shell is used as a source of fuel for the boilers, and residual coconut shell is disposed off as gravel for plantation roads maintenance. Black smiths also buy the coconut shell as fuel material in their casting and forging operations [6].

Bamgboye and Jekayinfa [6] regretted that 90% of coconut (empty fruit bunches, fibers, fronds, trunks, shell) was discarded as waste and either burned in the open air or left to settle in waste ponds. This way the coconut processing industries waste according to him contributed significantly to CO₂ and methane emissions. Based on economic as well as environmental related issues, efforts should be directed world wide towards coconut management issues i.e. of utilization, storage and disposal. Different avenues of coconut shell utilization are more or less known but none of them have so far proved to be economically viable or commercially feasible. Hence, the objective of this present work is to characterize coconut shell in order to explore its use in metal matrix composites.

II. MATERIALS AND METHOD

Material

The coconut shell used in this work was obtained from a coconut seller in Kaduna, Kaduna state of Nigeria. The photograph of the coconut shell is shown in Plate 1.



(a) Coconutshell (b)Crushedcoconutshell

Plate1:PhotographoftheCoconutshell.

Equipment

Equipmentusedinthisresearchare-electricalresistancefurnace,X-raydiffractometer(XRD),Scanningelectronmicroscopewithenergydispersivespectrometer(SEM/EDS)Machine,X-rayfluorescentXRF

Methods

Theprocessingofthecoconutshell(Carbonization)

The coconut shell was grinded to form coconut shell powder, the powder was packed in a graphitecrucible and fired in electric resistance furnace at temperature of 1300°C to form coconut shell ash (CSAp)(seePlate2).



(a) Coconutshellpowder (b)Coconutshellash

Plate2:Photographofcoconutshellash

Particle size analysis

The particle size analysis of the coconut shell ash particles was carried out in accordance with BS 1377: 1990 [7]. About 100g of the coconut shell ash particles was placed into a set of sieves arranged in descending order of fineness and shaken for 15 minutes which is the recommended time to achieve complete classification. The weight retained on each sieve was taken and expressed as percentages of the total sample weight. From the weight retained, the grain fineness number (AFS) was computed [8].

Density Measurement

Density measurements were carried out on the coconut ash sample using Archimedes's principle. The buoyant force on a submerged object is equal to the weight of the fluid displaced. This principle is useful for determining the volume and therefore the density of an irregularly shaped object by measuring its mass in air and its effective mass when submerged in water (density = 1 gram/cc). This effective mass under water was its actual mass minus the mass of the fluid displaced. The difference between the real and effective mass therefore gives the mass of water displaced and allows the calculation of the volume of the irregularly shaped object. The mass divided by the volume thus determined gives a measure of the average density of the sample [8].

Refractoriness

The Pyrometric Cone Equivalent (PCE) as recommended by ASTM Test C-24 was used in the determination of the refractoriness of the sample [2].

Mineralogical Characterization of the Coconut Shell ash

Mini Pal compact energy dispersive X-rays spectrometer (XRF) was used for the elemental analysis of the coconut shell ash. The system is controlled by a PC running the dedicated Mini Pal analytical software [8].

The XRD analysis of the coconut shell ash was carried out using Philips X-ray diffractometer. The X-ray diffractogram was taken using Cu K α radiation at a scan speed of 3 $^{\circ}$ /min. The samples were rotated at precisely one-half of the angular speed of the receiving slit, so that a constant angle between the incident and reflected beams is maintained. The receiving slit is mounted in front of the counter on the counter tube arm, and behind it is usually fixed a scatter slit to ensure that the counter receives radiation only from the portion of the specimen illuminated by the primary beam. The intensity diffracted at the various angles was recorded automatically on a chart and the appropriate (θ)

and (d) values were then obtained [7-9].

Microstructural Analysis

The microstructure and the chemical composition of the phases present in the coconut shell ash was studied using a JOEL JSM 5900LV Scanning Electron Microscope equipped with an Oxford INCATM Energy Dispersive Spectroscopy system. The sample was placed on sample holder and the images were captured under various magnifications. Prior to it, sample was applied with the gold coating to avoid charge effect, so to obtain clear images. The SEM was operated at an accelerating voltage of 5 to 20 kV [2].

Fourier Transform Infrared Spectroscopy (FTIR)

FTIR-8400S Fourier transform infrared spectrophotometer (SHIMADZU) was used for the functional groups present in the coconut shell ash. Spectrometer and detector, capable of measuring functional group to the predetermined minimum detectable level. The system includes a personal computer with compatible software that provides real-time updates of the spectral profile during sample collection and spectral collection using FTIR system using 1 cm $^{-1}$ resolution, 22 meter path length, and a broad band MCT detector. The Data analysis was performed using appropriate reference spectra whose concentrations can be verified using CTS spectra [7].

III. RESULT AND DISCUSSION

From the particle size analysis results, it is shown that, the coconut shell ash has a Grain Fineness Number (GFN) of 75.08. The sample can be considered to be fine as GFN value of 100 is ranked the finest. Also, the sample can be considered to have met the AFS specification since four sieve-size, has the bulk of the retained sample on four consecutive sieves corresponding to 355 μ m, 180 μ m, 125 μ m, and 63 μ m size fractions respectively [2].

The density of the coconut shell ash is 2.05 g/cm 3 which means that coconut shell ash is very light material. The value obtained falls within the range of density of fly ash, bagasse and silica which is 1.8 and 2.2 g/cm 3 respectively [1, 2] currently used in metal matrix composites. The coconut shell ash was observed to have Seger Cone No. 22, with equivalent temperature of 1500 $^{\circ}$ C. This means coconut shell ash can withstand operating temperature of 1500 $^{\circ}$ C without load [1-4].

The XRD pattern (see Figure 1) obtained reveals that, the major diffraction peaks are 20.75 $^{\circ}$, 10.22 $^{\circ}$ and 35.40 $^{\circ}$ and their inter-planar distance, 3.87 Å , 8.66 Å and 2.54 Å , and their relative intensity of X-

rayscatteringare100.00,61.90and8.44andphasesat these peaks as Quartz(SiO_2),Cordierite, syn ($\text{Mg}_2\text{Al}_4\text{Si}_5\text{O}_{18}$)andMoissanite(SiC),whileeachof

these phaseshaveascoreof44,32and16respectively(seeT able1).

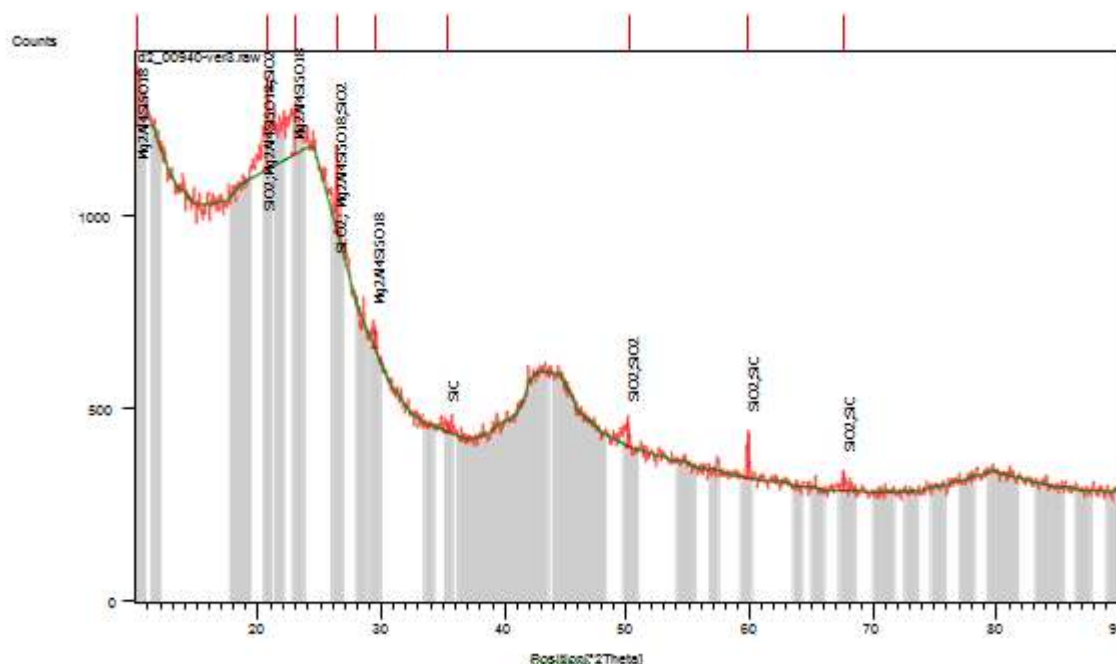


Figure1:XRDpatternofCoconutshellash

Table1:Identifiedpatternslistofcoconutshellash

Visible	Ref.Code	Score	CompoundName	Displac	ScaleFactor	ChemicalFormula
*	00-043-0596	44	SiliconOxide	0.000	0.912	SiO_2
*	00-013-0294	32	Cordierite,syn	0.000	0.590	$\text{Mg}_2\text{Al}_4\text{Si}_5\text{O}_{18}$
*	01-089-1961	26	Quartzlow, dauphinee-twinned	0.000	0.642	SiO_2
*	01-075-1541	16	Moissanite	0.000	0.207	SiC

The resultsshowedthat SiO_2 has the highest percentage of all the compound and element present as revealed by the XRD analysis. Complete Mineralogical analysis carried out by X-ray diffraction also revealed that the ash contains each of these elements C, O, Mg, Al, Si, Fe, Na, K, Zn and none of these elements H, He, Li, Be, B, N, F, Ne, P, S, Cl, Ga, Ge, As, Se, Br, Kr, Rb, Sr, Y, Nb, Mo, Tc, Ru, Rh, Pd, Ag, Cd, In, Sn, Sb, Te, I, Xe, Cs, Ba, La, Ce, Pr, Nd, Pm, Sm, Eu, Gd, Tb, Dy, Ho, Er, Tm, Yb, Lu, Hf, Ta, which means that with the absence of all these other elements the coconut shell ash may not contain radioactive materials. This is in par with the earlier of other biomass by [1-2]. The XRD result showed that both

SiO_2 and SiC have a fine structure, the former having a finer one. This could be associated with pore size [2].

The XRF chemical composition of the coconut shell ash is represented in Table 2. XRF analysis confirmed that SiO_2 , Al_2O_3 , MgO and Fe_2O_3 were found to be major constituents of the ash. Silicon dioxide, iron oxide and alumina are known to be among the hardest substances. Some other oxides viz. CaO , K_2O , Na_2O and MnO were also found to be present in traces. The presence of hard elements like SiO_2 , Al_2O_3 and Fe_2O_3 suggested that, the coconut shell ash can be used as particulate reinforcement in various metal matrixes. This result of XRF is in agreement with the result of XRD obtained. Therefore, the

presentwork suggests the possibility of using coconut shell ash as particulate in metal matrix composites since thechemical composition has

close similarity with the XRF analysis of rick husk ash, bagasse ash and fly ashcurrentlyusedinmetal matrix composites[1-4].

Table2:XRFanalysisofCoconutshellash

Element	Al ₂ O ₃	CaO	Fe ₂ O ₃	K ₂ O	MgO	Na ₂ O	SiO ₂	MnO	ZnO
%	15.6	0.57	12.4	0.52	16.2	0.45	45.05	0.22	0.3

Particle Morphology of coconut shell ash can be seen in back scattered electron (BSE) as shown inFigure 2.Coconut shell ash particles were observed to be solid in nature, but irregular in size. Somespherical shape particles can also be seen in the Figure 2.The chemical analysis

of the coconut shellash morphology consists mainly of Si, C, O, Mg, Al with small amounts of Fe as shown in the EDSscan(seeFigure2).TheresultsareconsistentwithXRDandXRFanalysisofotherbiomassby[1-5].

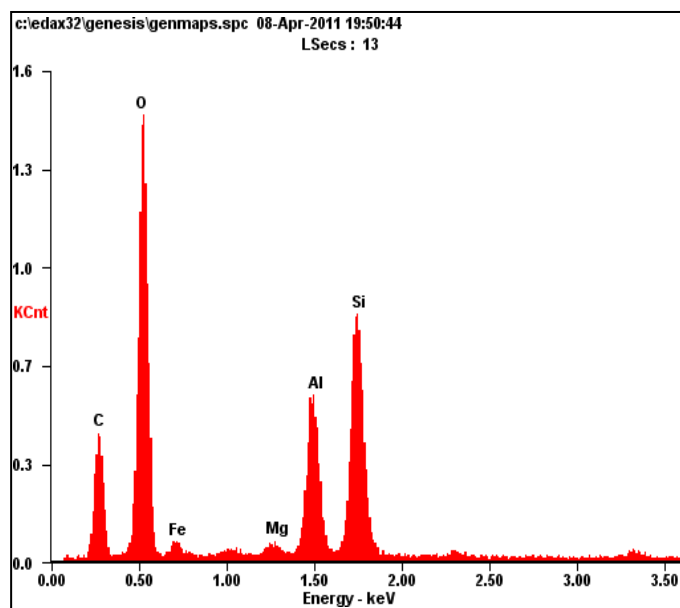
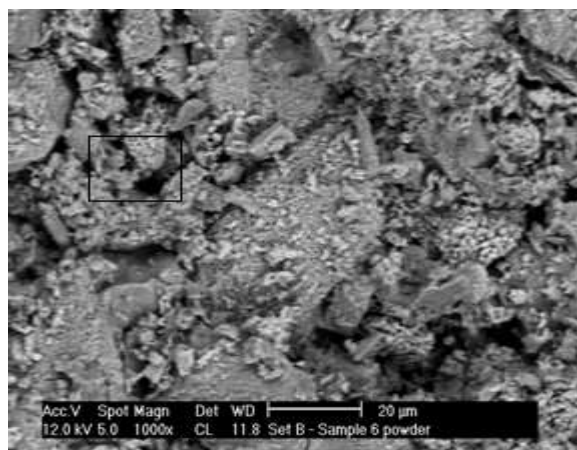


Figure2:SEM/EDSspectrumofCoconutshellash

Mainly twenty peaks were detected in the FTIR Analysis of the coconut shell ash as visible in

Figure 3.These peaks are shown in Table 3. This result has shown that the presence of quartzinthe

original ash gives rise in the IR spectrum to a series of bands located at 1132 and 443.64 cm^{-1} . The presence of mullite, in turn, is responsible for a series of bands at around 3797 cm^{-1} . The presence of carbon group is present in series of bands at around 4091.15-4617.74 cm^{-1} . Quartz, mullite and the vitreous phase of the ash

overlap in the area between 1220 cm^{-1} and 1434.12 cm^{-1} . Hence Quartz, Mullite, carbon and vitreous phases are confirmed to be present. More over the peaks in treated and untreated cases does not show any variations. This is in agreement with the earlier work of [4,9].

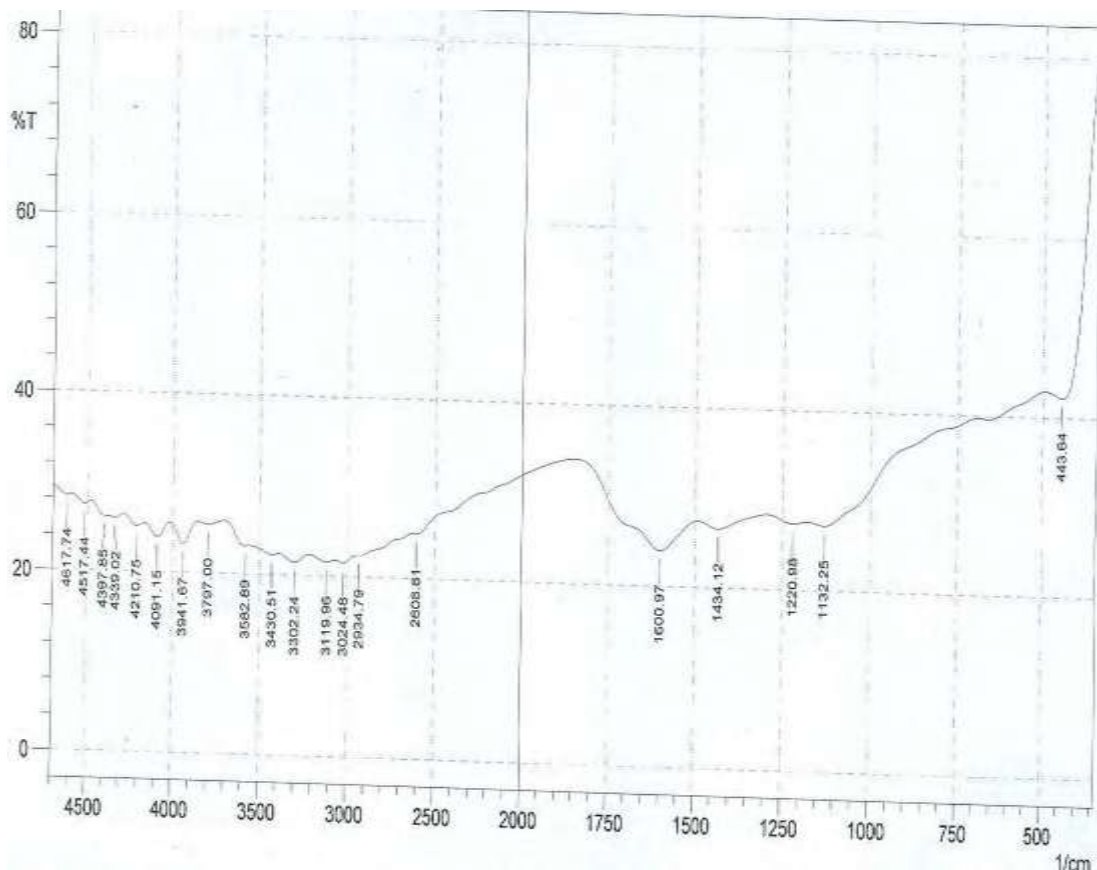


Figure 3: FTIR spectrum of Coconut shell ash

Table 3: Identification Peaks of the FTIR Analysis

	Peak	Intensity	Corr. Intensity	Base (H)	Base (L)	Area	Corr. Area
1	443.64	42.2149	13.1434	404.76	339.48	45.0051	10.1546
2	1132.25	27.1942	1.5636	1179.51	692.47	231.175	2.8009
3	1220.98	27.3523	0.4785	1295.21	1180.47	64.5302	0.4347
4	1434.12	26.5234	1.1098	1492.95	1297.17	110.3001	1.6248
5	1600.97	23.953	5.3532	1848.83	1493.92	198.1139	14.825
6	2608.81	24.9516	0.268	2630.03	1849.8	414.8633	0.2663
7	2934.79	22.3717	0.092	2944.44	2630.99	196.7375	0.5234
8	3024.48	21.4832	0.5281	3070.76	2945.4	82.8073	0.5844
9	3119.96	21.5287	0.4499	3218.34	3071.74	96.8457	0.6531
10	3302.24	21.4044	0.9435	3390.01	3219.3	112.6911	1.9018
11	3430.51	22.1843	0.3604	3553.96	3390.97	105.2941	0.4467
12	3582.89	23.0736	0.6718	3703.45	3554.93	91.4995	0.8093
13	3797	25.3657	0.3682	3854.87	3704.41	89.1332	0.5007
14	3941.67	23.1722	2.4099	4016.89	3855.83	98.5847	3.24
15	4091.15	23.8574	1.5527	4165.42	4017.66	89.7741	1.9765
16	4210.75	24.9963	0.6758	4287.9	4166.38	72.2261	0.7304
17	4339.02	25.8493	0.3262	4381.45	4288.86	54.1344	0.2576
18	4397.85	26.0397	0.2859	4476.93	4382.42	54.2572	0.3228
19	4517.44	27.3549	0.5835	4588.81	4477.89	51.741	0.4816
20	4617.74	28.3067	0.3104	4700.68	4589.77	60.1765	0.3021

IV. CONCLUSIONS

From the analysis of the results and discussion given above, the following conclusions can be made.

- 1) XRD analysis of the coconut shell ash reveals Silicon Oxide: (SiO₂), Corderite, syn: (Mg₂AlSi₅O₁₈), Quartz: (SiO₂) and Moissanite (SiC) as the primary compound with SiO₂ as the highest percentage of all the compound and element present.
- 2) XRF studies revealed the presence of hard element like SiO₂, Al₂O₃, MgO and Fe₂O₃ as major constituents which can be used as particulate reinforcements in MMCs for automotive applications.
- 3) FTIR graphs showed that Quartz, Mullite and Vitreous, carbon phases were present in coconut shell ash powder and proposed to use coconut shell ash as particulate reinforcement in MMCs.
- 4) The coconut shell ash can withstand a temperature of up to 1500°C with a density of 2.05g/cm³. That means this ash can be used in production of lightweight MMCs component with good thermal resistance.

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