

The Effects of Sintering Temperature and Agro Wastes on the Properties of Insulation Bricks

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ABSTRACT: In this research, kaolin, ball clay, sawdust and rice husk were used to produce insulation bricks through the solid state synthesis method. Two temperatures, 1100°C and 1200°C were used to sinter the green samples. X-ray fluorescence, scanning electron microscopy, compressive strength tests etc. were used to analyse the properties of the produced bricks. Chemical composition analysis on the starting raw materials showed that SiO₂ and Al₂O₃ were the major constituents while Fe₂O₃, Na₂O, K₂O and TiO₂ were the minor constituents. As the amounts of kaolin used in preparing the samples decrease, the bulk density, modulus of rupture and cold crushing strength of the bricks decreases while the water absorption capacity, linear shrinkage increases. The thermal analysis showed that on heating the samples, the reactions were mainly exothermic with between 8 to 10 mW/mg of heat released. The morphology of the samples showed that the pores began to collapse when the amount of kaolin present is below 70 wt. %. Sintering the samples at 1100°C and 1200°C led to slightly different values in the results and is therefore very significant.

KEYWORDS: Insulation bricks, Kaolin, sawdust, rice husk, temperature effects

[Received Jan. 30, 2019, Revised Jan. 9, 2020, Accepted Feb. 22, 2020]

Print ISSN: 0189-9546 | Online ISSN: 2437-2110

I. INTRODUCTION

Refractories are materials that can withstand the effect of abrasive or corrosive materials whether solids, liquids or gases at high temperatures. They can be inorganic, non-metallic in nature and depending on how porous they are, can be classified into two categories namely dense and porous refractories. The former is used where there is need for extreme mechanical, chemical or thermal stresses while the latter is usually for insulation applications because of their low thermal conductivity and heat capacity (Aramide, 2012). In the manufacture of porous refractories, raw materials like diatomite, perlite, expanded vermiculite, calcium silicate, fireclay, kaolin, quartz, alumina etc. are usually employed. Pore formers like sawdust, foam polystyrene, fine coke, binders etc. are usually introduced to produce porous bodies which lower the density and in turn the thermal conductivity of the brick.

There are a few reports of previous studies in the literature regarding the production of insulation bricks (Bogdanov et al., 2012, Bwayo and Obwoya, 2014, Pimraksa and Chindaprasirt, 2009, Ardeshir and Ahmadi, 2016). The effect of different amounts of sawdust on the properties of Ifon clay showed that the porosity of the samples increases with increasing sawdust content with a recommendation that sawdust amounts of

between 10 and 15 wt.% are required (Aramide, 2012). Sintering studies on insulation bricks made from a blend of purified clay, alumina and sawdust indicated that above 20% alumina content, the samples crumbled at elevated temperatures. It was therefore recommended that alumina content of 10% gives the properties required for insulation bricks (Folorunso et al., 2015a). The effect of particle size of the raw materials is to increase the thermal conductivity values provided the particle size is decreasing. The thermal conductivity of bricks formed from a mixture of kaolin, ball clay and sawdust indicates a reduction from 0.1428 w/mK to 0.0417 w/mK while the samples remained stable up to 1100°C (Viruthagiri et al., 2013).

Charcoal has also been used as an admixture to produce insulation bricks where the obtained density decreased by about 10 % depending on the charcoal content in the bricks. A charcoal content of 10 wt.% in the bricks led to an increase in the water absorption capacity and porosity of the bricks (Phonphuak, 2013). Particulates from coconut shell used to produce insulation bricks showed that with 25-30 wt. % of the additives, better mechanical properties were obtained (Esezobor et al., 2015). Clay and perlite used to produce high quality bricks showed that 24-30 % of perlite gives the best thermal and physical properties (Topcu and Isikdag, 2007).

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Expanded perlite as matrix, carboxy-methylcellulose (CMC) as a chemical binder, potassium borate ($K_2B_5O_7$)/sodium borate ($Na_2B_5O_7$) as natural binders and coal powder as additive have been used to produce insulation bricks. The compressive and flexural strength of the bricks were found to increase when aged from 2 to 90 days (Celik et al., 2014).

Aggregates of pumice, expanded perlite, borax, blend of molasses and hardener ($H_3PO_4-H_4NO_3$) have been used to produce lightweight bricks. The H_3PO_4 -hardened molasses-bonded brick is preferred when 2.5% of borax was not used (Benk and Coban, 2012). 39.6% pumice was found to be effective and increased the porosity of fired bricks while the density lowered to 1.58 g/cm^3 at 900°C and increased to 1.61 g/cm^3 at 1000°C respectively (Gencel, 2015). Lime, gypsum and diatomite used to produce bricks showed a positive correlation between lime content and strength/density of the bricks. A compressive strength of 17.5 MPa, density of 0.73 g/cm^3 and water absorption capacity of 46% were obtained (Pimraksa and Chindaprasirt, 2009). The use of paper pulp as an additive (15% maximum) in the manufacture of insulation bricks led to a decrease in the thermal conductivity to 0.45 W/mk which is about 60% enhancement in properties (Munoz et al., 2013).

Most of the insulation refractory bricks currently used in the country are imported meanwhile the raw materials necessary for making insulation bricks can be found in many parts of Nigeria. In this work, kaolin and ball clay have been used as the matrix materials while sawdust and rice husk are used as pore formers. The objectives of this research are to produce insulation bricks from locally available raw materials, to check the effect of sintering at different temperatures and to investigate the variation of rice husk and kaolin contents on the properties of the produced bricks.

II. MATERIALS AND METHODS

A. Sample Preparation

The starting raw materials used in the production of the insulation bricks are kaolin, ball clay, rice husk and sawdust. The kaolin was obtained from Darazo in Bauchi State, ball clay from Ikorodu in Lagos State, rice husk from Wushishi, Niger State and saw dust from Ebute Metta, Lagos State. In order to produce 5 different compositions of the insulating bricks, the amounts of the kaolin and rice husk were varied as shown in Table 1. The raw materials were first crushed in order to reduce them to smaller sizes. They were then measured and wet-mixed in the following proportions to produce 5 different compositions.

Table 1: Table showing the amounts of the different raw materials used in the production of the insulation bricks.

Label	Raw Materials			
	Sawdust (wt. %)	Ball clay (wt. %)	Kaolin (wt. %)	Rice Husk (wt. %)
S1	5	5	90	0
S2	5	5	80	10
S3	5	5	70	20
S4	5	5	60	30
S5	5	5	50	40

The raw materials were ball milled until their average particle sizes were less than $150 \mu\text{m}$. For ease of representation, the samples have been labelled. The mixture was poured into already prepared moulds of dimensions $220*60*60 \text{ mm}$ for the bricks and $60*60*20 \text{ mm}$ for test samples and pressed to produce the green samples. 3 samples for each composition were produced in this study. The bricks were placed in an oven operating at 110°C and left to dry for 7 days and two temperatures, 1100°C and 1200°C were used to sinter the samples. The heating rate is $2^\circ\text{C}/\text{min}$ and the samples were held at these temperatures for 1 h and allowed to cool to room temperature in the furnace. A picture of the insulation bricks while still inside the oven is shown in Figure 1.



Figure 1: A picture showing the different sizes of the insulating refractory bricks (a) while still inside the oven at 110°C and (b) after sintering in the furnace.

B. Sample Characterisation

The chemical composition analysis of the raw materials was determined with an X-ray Fluorescence Analyser at Venarum Scientific Materials Laboratory Limited Apapa, Lagos. The dimensions of the samples before and after sintering were measured and used to determine the physical properties of the bricks. To characterize the samples, a number of tests were carried out in accordance to the ASTM C134–95(2016) and ASTM C20-00(2015) which are standard methods used to measure size, dimension, bulk density and apparent porosity, water absorption and shapes by boiling water respectively. The linear shrinkage (LS) of the samples were measured by measuring the length of the bricks before and after sintering and using the formula in eqn (1).

$$\text{Linear Shrinkage (\%)} = \frac{DL-FL}{FL} * 100 \tag{1}$$

where D_L stands for dry length and F_L is fired length.

The apparent porosity (AP) of the samples was determined using the boiling water method. The sintered samples were put inside a water filled container and boiled for 2 hours, allowed to cool to room temperature after which they were removed and the porosity measured. The apparent porosity of each test specimen was calculated using eqn (2).

$$\text{Apparent Porosity(\%)} = \frac{W-D}{W-S} * 100 \tag{2}$$

where: D = Weight of fired specimen, S =Weight of fired specimen suspended in water, and W = Weight of soaked specimen suspended in air.

For water absorption (WA) capacity tests, the weight of each sintered sample was recorded and then they were immersed in water at 30 and 45°C for 24 h. The surfaces of the samples were dried and re-weighed after the immersion. The water absorption capacity of the samples was calculated using the formula in eqn (3).

$$\text{WaterAbsorption (\%)} = \frac{W_2-W_1}{W_1} * 100 \tag{3}$$

where W_1 = weight of dry specimen, W_2 = weight of wet specimen after 24 h submersion in water.

The bulk density (BD) of the bricks was determined using the boiling water method. The Archimedes method was used to calculate the density by placing the samples in water, boiling for 2 h and subsequently allowing then to cool to room temperature. They were removed from the water, cleaned and

re-weighed in air. The formula used to calculate the density is shown in eqn (4).

$$\text{Bulk Density} = \frac{D}{W-S} \tag{4}$$

where: D = Weight of fired specimen, S =Weight of fired specimen suspended in water, and W = Weight of soaked specimen suspended in air.

Modulus of Rupture (MOR) was carried out using a mechanical device where the specimens were placed on the bearing edges of the compression tester positioned 70 mm apart. Loads were then applied uniformly at the rate of 12.26 N/min to the specimen until failure occurs; the formula used to calculate MOR is shown in eqn (5).

$$\text{MOR} = \frac{3WL}{2bt^2} \tag{5}$$

where L = the distance between bearing edges (mm), b = width of the specimen (mm), t = thickness of the specimen, and W = load at which the specimen failed.

The thermal analysis on the samples was carried out using a differential thermal analyser (DTA). The measurement was made on heating the samples from room temperature to 1100°C. The cold crushing strength (CCS) of the samples was obtained by placing it between two plates on the compression tester and steadily increasing the compressive load applied to the specimen until it fails. The load value at which cracks begin to appear the bricks fails while operating at room temperature is the CCS. The CCS was calculated using the relation in eqn (6).

$$\text{CCS} = \frac{\text{Maximum Load (KN)}}{\text{Cross Sectional Area (m}^2\text{)}} \tag{6}$$

The compressive strength of the samples was determined using the cold compression strength test where the sintered samples were placed between two plates between the equipment followed by the application of a uniform load. The load where a visible crack appears on the sample is noted and divided with the cross sectional area of the sample. The morphology of the samples was measured with a scanning electron microscope (VEGA3 TESCAN, Czech Republic) at the Tshwane University of Technology, South Africa. The samples to be measured were prepared by sputtering their surface with carbon to ensure electrical contact. The energy dispersive X-ray spectroscopy (EDS) was obtained by selecting an area 50*50 mm. The amounts of the quantitative EDS analysis (Oxford Instrument, UK) on the major elements present in the samples are shown.

III. RESULTS AND DISCUSSION

A. Presentation of Results

The results of the chemical composition analysis on the raw materials are shown in Table 2. SiO_2 is the main compound that is present in all the raw materials. Kaolin and ball clay contains significant amounts of Al_2O_3 while its quantity in the inorganic constituent of rice husk and sawdust are 4.49 wt. % and 11.8 wt. % respectively. Fe_2O_3 is also present in all the raw materials as a minor constituent. The SiO_2 and Al_2O_3 contents of kaolin are 51.38 and 42.05 wt. % respectively and in ball clay, 63.59 and 23.37 wt. % respectively. The impurities present in significant amounts in all the raw materials include Fe_2O_3 , Na_2O , K_2O and TiO_2 . They are present in amounts ranging from as low as 0.43 wt. % to as high as 10.44 wt. %.

The bulk density values as well as the linear shrinkage in the samples are shown in Figure 2. Samples sintered at 1100°C had slightly higher bulk density and lower linear shrinkage compared to those sintered at a temperature of 1200°C . As the amount of kaolin present in the bricks decreased (increasing rice husk), the density of the samples decreased while the linear shrinkage values increased. When the kaolin content is 90 wt. %, the density value is 1.49 g/cm^3 and increases to 1.61 g/cm^3 when it is 80 wt. % before gradually decreasing to 1.24 g/cm^3 when the kaolin content is 50 wt. %. The reason for this behaviour in density could be due to the volatile pore formers which escape leaving pores at their initial location. The density values in this work is similar to the literature report by (Folorunso et al., 2015a), (Chemani and Chemani, 2012) as well as (Tonayopas et al., 2008) who obtained values between 1.4 and 1.6 g/cm^3 for their insulation bricks.

Sintering at both 1100°C and 1200°C respectively did not alter the trend in the linear shrinkage values. Sintering at 1200°C gave slightly higher linear shrinkage values compared to that at 1100°C . Sintering at 1100°C resulted in shrinkage values ranging from 8.21% for S1 to 11.48% for S5 while the shrinkage at 1200°C is 8.23% for S1 to 11.91% for S5. It seems that kaolin inhibits shrinkage because the higher the kaolin content, the lower the shrinkage. This is slightly lower than

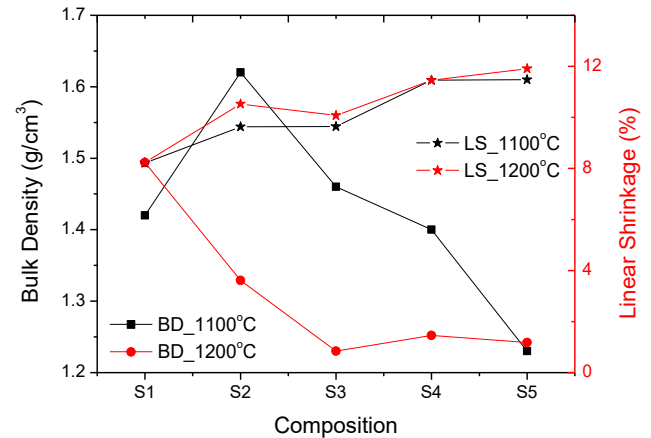


Figure 2: Graphs of bulk density and linear shrinkage (%) as a function of composition of the insulation bricks.

the 14 -17% reported by (Chemani and Chemani, 2012) for Clay-sawdust bricks but similar to that reported by (Ugheoke and Mamat, 2012).

The water absorption and apparent porosity values of the samples are shown in figure 2. The samples sintered at 1100°C have a lower water absorption capacity compared to the samples sintered at 1200°C . The observed trend is the increase in water absorption as the amount of rice husk increases. At 1100°C , the water absorption value is 28.07% for sample S1 and gradually increases to 37.46% for sample S5. At 1200°C , the value is 29.81% for sample S1 and gradually increases to 38.63% for sample S5. As the samples become more porous, the rate of water absorption increases.

The apparent porosity (%) of the samples shows a gradual increase from sample S1 to sample S3 and a decrease in the values afterwards. For samples sintered at 1100°C , the initial porosity was 24.25% and gradually increased to 29.87% before dropping to 27.55%. For samples at 1200°C , a similar trend was observed. For sample S1, the porosity value is 27% and gradually increased to 33% for sample S3 and decreases to 31.58%. The reduction in porosity values with increasing amount of rice husk above 30 wt. % may be due to collapse in the pores structure within the samples.

Table 2: Chemical composition analysis of the raw powders using X-ray Fluorescence.

Composition (wt. %)	SiO_2	Al_2O_3	Fe_2O_3	MgO	CaO	Na_2O	K_2O	TiO_2	BaO	ZnO	P_2O_5	MnO
Kaolin	51.38	42.05	1.39	0.02	0.16	1.73	0.43	2.37	0.001	0.005	-	0.007
Ball Clay	63.59	23.37	3.02	-	0.57	0.87	1.18	0.93	0.039	0.001	-	-
Rice Husk	68.85	4.49	3.56	1.36	5.33	0.68	10.44	1.01	-	-	1.04	-
Sawdust	51.24	11.8	7.45	0.95	15.1	-	5.58	1.73	-	0.77	1.09	-

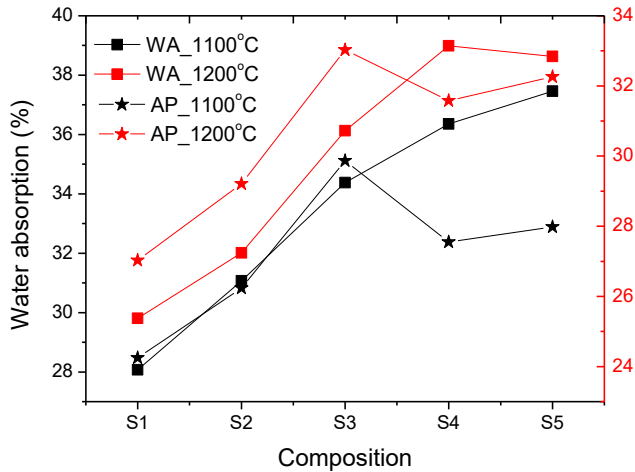


Figure 3: A plot of water absorption (%) and apparent porosity (%) as a function of sample composition.

Figure 4 shows the values of the modulus of rupture (MOR) and cold crushing strength (CCS) as a function of sample composition. The values of MOR generally decreased as a function of composition. Samples sintered at 1100°C generally have slightly higher values of MOR compared to samples sintered at 1200°C. As the amount of rice husk added to the insulation brick increased, the MOR values decreased from 3.52 MPa for sample S1 to 1.76 MPa for sample S5. The values of MOR for samples sintered at 1200°C followed the same trend from 3.18 MPa for sample S1 to 1.51 MPa for sample S5.

The values obtained in this research are similar to that reported by Folorunso et al., 2015 for bricks containing Al₂O₃ and sawdust (Folorunso et al., 2015b). The CCS values decreased with increasing rice husk content from 2.5 MPa to about 1 MPa. The value obtained here is about an order of magnitude lower than that reported for bricks produced using Clay and Rice husk ash (Tonnyapas et al., 2008). The difference in values could be due to differences in compositions used for the respective researches.

The plot of the thermal analysis as a function of composition using a differential thermal analyser is shown in Figure 5. The graph shows that the measurements for all the samples are in the exothermic range. At a temperature of about 50°C, exothermic reactions could be observed and these continued until about 1000°C. The explanation for this behaviour can be that as the samples are being heated; volatile or low melting temperature materials in the brick are gradually being burnt up giving up heat in the process. The trend in the thermal behaviour of the samples is similar for samples sintered at 1100°C and 1200°C respectively. As the quantity

of pore formers in the bricks increase, the exothermic reactions increased.

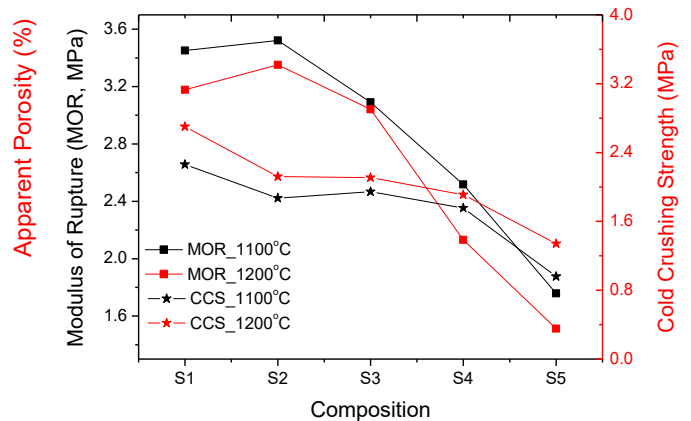


Figure 4: A graph of modulus of rupture (MOR) and cold crushing strength as a function of sample composition and sintering temperature.

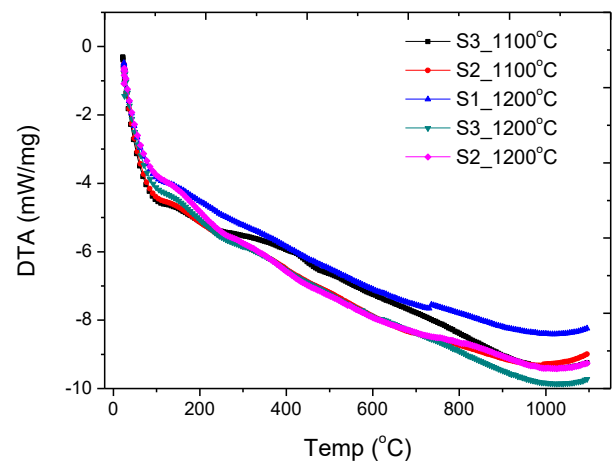


Figure 5: A plot of differential thermal Analysis (DTA) as a function of temperature for sintered at 1100 and 1200°C and measured on heating from room temperature.

The electron microscope images of the samples which reveal their morphology is shown in Figure 6. Two distinct phases with dark and bright features respectively can be observed in the micrographs. The microstructure of sample S1 reveals that in addition to the phases in the samples, pores could clearly be observed. As the amount of the pore formers increased in sample S2, the average size of the grains increased as well as the size of the pores. The size of the grains as well as the pores increased with higher amount of the pore formers in sample S3. As the amount of the pore formers increased further in samples S4 and S5, the size of the grains as well as the pores decreased. This observation is in agreement with the apparent porosity graph in Figure 3 where the porosity began

to decrease after 20 wt. % of rice husk. An explanation for this behaviour could be that with more pore formers, instead of the grains to further increase, the pores begin to collapse thereby decreasing the porosity in the samples.

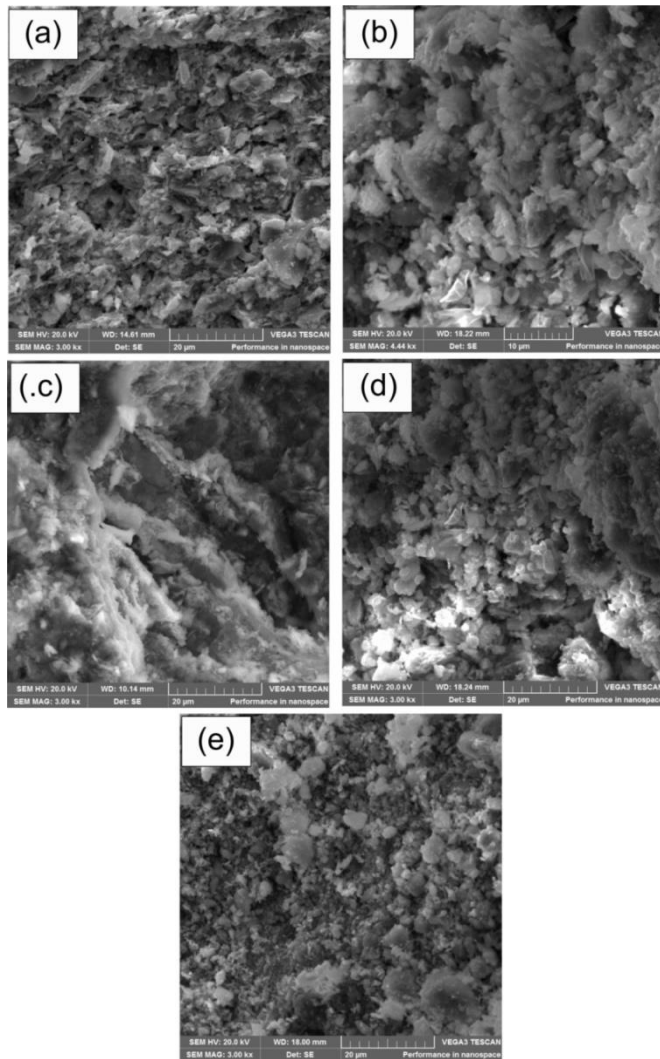


Figure 6: Scanning electron micrographs of samples (a) S1 to (e) S5 respectively sintered at 1100°C.

Energy dispersive X-ray spectroscopy was carried out on the samples and a representative of the measurements is shown in Figure 7. An area scan of 50 μm by 50 μm was carried out to quantitatively determine the amounts of the major elements in the samples which are basically O, C, Si and Al. The carbon is mainly from the coating to ensure electrical contact during the measurements. A summary of the quantitative analysis is shown in Table 3.

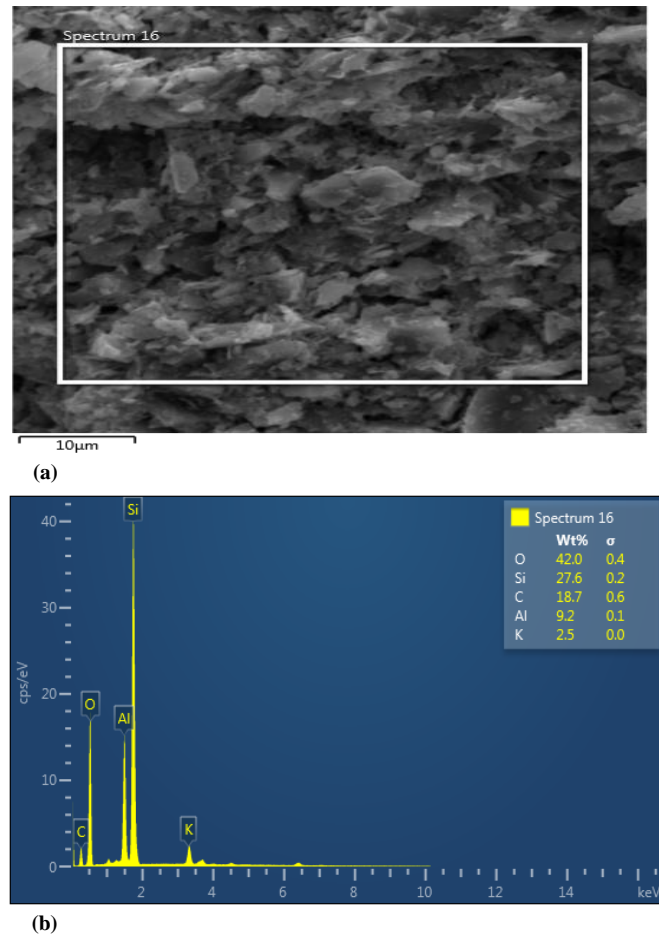


Figure 7: (a) Representative area scan of the energy dispersive X-ray spectrograph of the samples, and (b) the diffraction spectrum.

Table 3: Table showing the values of the quantitative energy dispersive X-ray spectroscopy on the major elements present in the samples

Elements (wt. %)	O	Si	C	Al	K
S1	42±0.4	27.6±0.2	18.7±0.6	9.2±0.1	2.5±0.0
S2	47.5±0.2	42.2±0.1	-	10.3±0.1	-
S3	45.9±0.3	16.7±0.1	20.5±0.4	16.9±0.1	-
S4	45.2±0.2	16.5±0.1	21.3±0.3	17.1±0.1	-
S5	46±0.2	16.3±0.1	21.1±0.3	16.5±0.1	-

The amounts of the elements in sample S1 show that silicon is 27.6% while Al is 9.2%. Carbon was not detected in sample S2 which increased the silicon present to 42%. Samples S3 to S5 are basically similar where the amount of silicon decreases to 16.5% while the Al increases to approximately 16.8%.

IV. CONCLUSION

Investigation into the effect of sintering temperatures and the amounts of pore formers has been carried out in this research. The temperature of sintering has a slight effect on the physical and mechanical properties of the insulation bricks. The samples that were sintered at 1200°C had slightly better properties compared to those sintered at 1100°C. As the amount of rice husk increases with decreasing kaolin content, the density values decreased from approximately 1.6 g/cm³ to 1.2 g/cm³.

The linear shrinkage, water absorption capacity, Apparent Porosity, Modulus of Rupture and Cold Crushing Strength of the bricks generally increased. It was observed that the Cold Crushing Strength and Water Absorption Capacity results were up to the standard required for refractory bricks. The SEM images show that the structure of the samples contained adequate pores for insulation. The pores however, begin to collapse when the amount of rice husk is more than 30 wt.%. Great potentials therefore exist in the use of these raw materials to produce insulation bricks.

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