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Research Paper



Development of Avehicle Brake Pad Using Composites of PalmKernel Fiber and Groundnut Shells As Filler Material

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Abstract

This study investigated the development and assessment brake pad using palm kernel fiber (PKF)
groundnutgroundnutshellas

and

filter.Palmkernelfiber(PKF)ofdifferentparticlesizeswerecombinedwithgroundnut shellused asfiller to produce brake pads following

standard procedure. The performance of the produced brake pads was evaluated, and compared with calculated and compared with the standard procedure. The performance of the produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and compared with the standard produced brake pads was evaluated and the standard pads was evalua

ommercial(asbestosbased)brakepad.Natural waste has been used to produce fillers and fibers, including palm kernelfiber and fiber, groundnut shell, maize husk and rice straw. This study seeks to explore research using combinations of fillers and fibers at different ratios with a view to studying their effects onbrake padproperties using various mixtures for the production. Composite materials from fiber and fillers have been seen to improve compositemechanical properties, reduce costs and increase impact strength. The choice of fiber, filler, binder, particle size and composition play important responsibility in the composite of the brake pad performance. *Itwasobservedthatpalm* kernel fiberparticlesize and groundnut shell as fillerhaveasignificantinfluenceontheperformanceofpalm kernel fiberbasedbrake pads.

In order to obtain better physical properties, palm kernel fiber and groundnut fiber brake padswere studied and the composition percentage was optimized.

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I. Introduction

Thebrakingsystemisanindispensablecomponent of an automobile, and is composed of manypartsincludingbrakepads,mastercylinder,wheelcylinders, and a hydraulic control system(Maleque*et al.*,2012). The brake pad is an important part of the brakesystem and consists ofsteel braking plates with frictionmaterials bound to the surface facing the brake disc.

Thebrakepadgenerallyconsistsofasbestosfibresembeddedin polymeric matrix, along withseveral other ingredients(Olele,Nkwocha, Ekeke,Ileagu&Okeke, 2016).The use of asbestos inbrake padshas become a source of concern due to its carcinogenicnature and problem ofdisposal, consequently, it is beingphased out. The current trend in the automotive industryisthedevelopmentanduseofasbestosfreebrakepads. Brake pads are components of disc brakesused in automobiles. They are

steelbackingplateswithfrictionmaterialsboundtothesurfacefacingthebrakedisc.Brakepadsareusedin the braking systems to control the speed of the automobile (Nagesh, 2014) by converting the

kineticenergyoftheautomobiletothermalenergybyfrictionanddissipatingtheheatproducedtothes urroundings. In the recent time the production of composite materials has grown significantly worldwide, which means many industries and technology sectorsare using polymercomposite materialsto successfully replace traditional composite materials (Nagesh, 2014). The investigation of new materials, especially agricultural waste as a filler material, is providing new and low-cost materials fordevelopmentofbrakepadswhicharecommerciallyviableandenvironmentallyacceptableand which have all the required properties. Brake pads are used to control the speed of avehicleviathebraking system (Idris,

2015).Frictionmaterialsinbrakingapplicationsystems are considered as the most important sections in a vehicle performance. The brake padmaterialmust beable to sustain a higher and uniform coefficient offrictionalong side the brake disc.

Industrial and agricultural wastes are currently

receivingattentionasalterativerawmaterialstoasbestosinthemanufacture of brake pads (Leman *et al.*,2008). The use ofsuitablewastematerialscanprovideaddedvaluesandreduce environmental problems and costs associated withdisposal.

Brake pads use as automobile brakes are of two types: drum brakes and disc

brakes.Thedrumbrakeislocatedinsideadrumsothatonapplicationofthebrakes,thebrakeliningis forced outward and pressed against the drum, while disc brakes operate in similarwayexcept that theyareexposed to

theenvironment (Deepika, 2013). Asbestos had a fewengineering properties that madeit very suitable for inclusion in brake linings, and was themostpreferred filler material up till 1990. The use of asbestos isbeen avoided due to itscarcinogenic nature. Therefore, anew asbestos free friction material and brake pads hasbeendeveloped.

The use of thermosetting resins to produce moulded brake lining instead of knittedlinings were made by combining fiber with resin and polymerizingresin under elevatedpressure and temperature. The fabrication and performance evaluation of a composite material for wear resistance application (Bashar, 2012) made use of an agro-waste (palm kernel fiber s -PKF) asfiller material with sulphur, cashew nut shell liquid, calcium carbonate, brass chips,

quartz,iron ore, ceramics, and carbon black. Similarly, coconut shells based brake pad was producedwith a formulation of grinded coconut shells (filler), epoxy resin (binder –matrix), iron chips(reinforcement), methylethylketone peroxide (catalyst), cobalt nephthanate (accelerator),ironand silica(abrasives), and brass(friction modifier) (Yawas, 2016).

The major component the brake in pad is the liningmaterials, which are categorized as metallic, semi-metallic, organic and carbon-based, depending on the composition ofthe constituent elements. Typical formulations consist of morethan10ingredients, and more than300 materials are indifferent brands (Edokpia, 2014). These ingredients classified are into

fourbroadgroups:binders,reinforcingfibresorstructuralmaterials, fillers, and frictional additives/modifiers, based onthe major function they perform apart from controlling frictionandwearperformance. The binderholds theing redient stogether, to maintain structural integrity of the brake and liningunder varving mechanical thermal stresses. The structuralmaterialsprovidethestructuralreinforcementtothecomposite matrix; fillers make up the free volume of the brakelining and friction modifiers stabilize the coefficient of frictionand wear rates. These components perform synergistically incontrollingfrictionandwear performanceofthebrakepad.

Palm Kernel fiber (PKF) is recovered as by-product in palm oilproduction. Large quantities of PKF are generated annually and only some fractions are used for fuel and other applications such as palliative for untarredroad and in producing activated carbon.

TheunusedPKFaredumpedaroundtheprocessingmill,constituting environmental andeconomic liability for the mill.Although,PKFmustbegroundintofineparticlestobesuitable for

information inclusion in brake lining, available intheliteratureareontheungroundedshellparticles.Coefficients of friction of PKF on metal surfaces were in therangeof0.37-0.52 (Lagel, 2016). In frictioncoefficient intherangeof0.30contrast, 0.70isnormallydesirablewhenusingbrakelining material (Bala, 2016). Ithas found been that incorporation

ofPKFintheproductionofstructurallightweightconcretesincreasedthemechanicalstrength.Thu s,PKFappearedsuitableforuseasbasematerialinfrictioncomposites,becausethey are subjectedtohardandvariablebrakingforces.

Mostcommercialautomotivebrakepadfrictionmaterialscontain multiple components (Kumar, 2011) and divided into four groups: fibers, fillers, binders, andfriction modifiers. Fibers provide mechanical strength in the composition. Friction modifiers the brake pads frictional properties and contain a mixture of abrasive as well aslubricants.

Fillermaterialsaremainlyusedforbrakepadproductiontoimprovebrakemanufacturability and reduce production

costs and as functional modifiers. A small amount offiller is usually added to improve or optimize performance of bake pad material. Harderparticles, for instance Al₂O₃ is added to increase the COF (μ) which is the force of frictioncausedbythescrapingthesurfaceof the materialand the disc(Bijwe, 1997).

Bindersholdallthecomponentstogetherinthebrakepadapplication,therebyreducingthe component shear rate (Blau, 2001). Binder contribute to the brake pad friction and wear rate(Rohatgi, 2012)The binder offers mechanical unity to the friction materials by firmlycombining the otherthree components in order to improve thecomposites properties.Inthepast fifty years,phenolic resins (unmodified or modified) have been employed as binders in the preparation of the friction materials due to their good thermal and mechanical properties

$in \ addition \ to \ lower costs (Chan., 2004) \\ \textbf{Methodology The Raw Materials Preparation}$

Existing agricultural waste cannot be used directly in the formulation of the final brake pad. Therefore, some treatments such as mechanical and chemical treatments are required earlier in the brake pads composition. The following are some of the natural fiber treatment methods proposed according to the literature. **PalmKernelFiber**

5kgofpalmkernelfiberwasobtainedfrom apalmoilprocessingmill. The fibers were collected using rubber bucket and thoroughly washed with water and soap to remove residual oil and extraneous materials. Thereafter, the fibers were sun dried for three (3) days. The driedfiberswerepounded using mortar and pestle

untilthedesiredparticlesizewasobtained.Thefine particles of thefiberswereclassifiedintodifferentparticle sizes.

PalmKernelfiber asBrakeLiningIngredient

Palm Kernel fiber (PKF) is recovered as by-product in palm oilproduction. Large quantities of PKS are generated annually and only some fractions are used for fuel and other applications such as palliative for untarredroad and in producing activated carbon.

TheunusedPKFaredumpedaroundtheprocessingmill,constituting environmental and economic liability for the mill.Although,PKFmustbegroundintofineparticlestobesuitable forinclusion in brake lining, available informationintheliteratureareontheungroundedshellparticles.

Coefficients of friction of PKF on metal surfaces were in therangeof0.37-0.52 (Deepika,2013)In contrast, frictioncoefficient intherangeof0.30-

0.70isnormallydesirablewhenusingbrakelining material. It has been found (Nagesh, 201) that incorporation of PKS in the production of structural lightweight concretes increased the mechanical strength. Thus,

PKFappearedsuitableforuseasbasematerialinfrictioncomposites, because they

subjected to hard and variable braking forces. Reported that PKF did not change significantly in

physicalstructureandweight,forappreciabletimeduration,whenexposed to organic solvent. It is also important that the frictionmaterials experience very little or no changes on

contacting varying environmental conditions: we tordry weather, or hydraulic fluid spilling over.

These observations therefore, stimulated the interest in considering PKF for use as friction material in brake lining. Hence, the aim of this research is to develop a new asbest osfree brake pad from Palmkernel fiber, which is readily available and nontoxic.

Brakepadformulation

Production of the brake pad consists of a series of unit operations including mixing, cold and hot pressing, cooling, post-curing and finishing (Fono-Tamo., 2013). The samples were produced using compression

moulding machine. EachofthethreePKSgritsizeswasusedtoformulate brake pad by mixingwith brass chips, steelfibre,graphite,latexrubber,calciumcarbonate,resinbinder, carbonblackpowder and groundnut shell was used as the filler.Aftermixing, the mixture was compacted in a mould to assume the required shape. A compressive force of 50 KN wasappliedthroughapunch,byahydraulicspress,foraperiod of 20 min.The brake pad produced was curedby heating in a wooden oven at a temperature of 80^oC for45 minutes, and allowed

to cool. Afterward the brake pad samplesweresubjected toperformancetests. The produced brake pad samples and a commercial brake pad (produced from as bestos) were tested for wear/application, disc temperature rise, and disc stopping time (braking efficiency) at different speeds. **GroundnutShell**

Groundnut shell was collected and washed with water to remove sand and put in a

are

sodiumhydroxide(NaOH)solutionhavingacompositionratioof1:20withwatertoremoveimpurit iessuchas lignin and pectin. After this, the groundnut shell was washed in distilled water to reduce the sodiumhydroxideinthe shelland sundried tillallthe moisture content in the shelldried up. Thedried shell was ground in a grinding machine to reduce its size, after which the shell wassieved into a 200 and 400 um sieve size. The shell particles were then mixed in a two-rollmixingmill togetherwithotherconstituents of filler the brake pad with palm kernel shell.



Fig. 1: Diagrams of materials used for the formulation of PKF + GS brake pad

Fig. 2: Diagram of materials combination for the production of brake pad

Microstructure analysis

The microstructure analysis of the samples was carried out by grinding the samples

using 200, 300 and 500 grit papers respectively. Dry polishing was then carried out on

 $these samples and the internal structures we reviewed under the computerized Metallurgical micros\ cope.$

Brinellhardnesstest

The resistance of the composites to indentation was carried outusing the Brinellhardnesstesting equipment of BS240, a Tensometer (M500-25kN, hardened steel ball ofdiameter D to indent the test specimen. Based on ASTM specification, a steel ball of D = 10mm diametersteel ball was used, and the load applied P was kept stable at 3000 kgf. Thediameter of the indentation, d, was measured along two

perpendicular directions, using anopticalmicrometerscrewgauge.ThemeanvaluewasusedtoobtaintheBrinellHardness Number(BHN)usingequationbelow. BHN = 2^{P}

 $\pi D \left(\overline{D - \sqrt{D^2 - d^2}} \right)$

WhereP=appliedload,D=diameterof hardenedsteel ball, d =diameter of indentation.

Compressivestrengthtest

The compressive strength test was done using the Tensometric Machine. The samplesof diameter 29.40mm were subjected to compressive force, loaded continuously until failureoccurred. The load at which failure occurred was then recorded.

Ashcontent test

About 1.30 g \pm 0.2 g of the samples were weighed in a cooled crucible which was ovendriedbyheatinginafurnaceto600°Cfor30 minutes. Then the samples were charred byheating in a hot plateafter which the charred samples were placed in a furnace and heatedat600°Cfor minutes. Then cooled in a desiccator and weighed. This process of heating, cooling and reweighing were repeated until a constant

weight was obtained the ash content was calculatedusingequation below.

Ashcontent = $\frac{W^2 - W^0}{W^2 - W^0} x \ 100(2)$ W1-W0

 $Where W_0 = weight of empty crucible, W_1 = weight of crucible and sample, W_2 = weight of crucible and dresidue after cooling. \\ \textbf{Densitytest}$

The density of the samples was determined by weighing the samples on a digitalweighing machine and their volumes determined by liquid displacement method. The density was determined using equation below.

Density, $\rho = \frac{M}{V}$

 $where M = mass of test piece(g), V = volume of the test piece(cm^3) by liquid displacement method. \\ \textbf{We arratetest}$

Thewearrateforthesamplesweremeasuredusingpinondiscmachinebyslidingit over a cast ironsurface

at a load of 10N, sliding speed of 125rev/min and sliding distance

of2000m.Theinitialweightofthesampleswasmeasuredusingasinglepanelectronicweighing machine with an accuracy of 0.02g.During the test, the pin was pressed againstthecounterpart rotating cast iron disc of Rockwell hardness 65HRC of counter surfaceroughnessof 0.3µm by applying the load. A friction detecting arm connected to a straingauge held andloaded the pin samples vertically into the rotating hardened cast iron disc.After runningthrough a fixed sliding distance, the samples were removed, cleaned withacetone, dried, andweighed to determine the weight loss due to wear. The difference inweight measured

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beforeandafterthetestsgive thewear of the samples and thewear rate is calculated by equation

Wear rate = ΔW S

 $Where \Delta W = weight difference of the sample before and after the test (mg), S = is the total sliding distance (m). \\ \textbf{Waterabsorptiontest}$

The samples were weighed on a digital weighing machine and soaked in water at roomtemperature for 24 hours. The samples were then removed, cleaned and weighed. The waterabsorptionrate was calculated thus: Waterabsorption $=\frac{M2M1}{2}$

 M_1

x 100 %

where M_1 =mass of the sample(g), M_2 =mass of the sampleafter absorbing water(g).

FlameResistanceTest

Weigh about $1.30g \pm 0.2g$ of the samples in a cooled crucible previously oven dried byheating in a furnace at 600 0 C for 30 minutes. Then the samples were charred by heating in a hot platethereafter, the charred samples were taken into the furnace and heated at 600 0 C for 30 minutes. Then the samples were charred by heating in a hot platethereafter, the charred samples were taken into the furnace and heated at 600 0 C for 30 minutes. Then cooled inadessicator and weigh. This cycleofheating, cooling and weighing were repeated until a constant weight was obtained. **Calculation:**

 $\text{%ash} = \frac{W2 - W0}{W1 - W0}$

x 100

 $Where W_0 = weight \ of \ empty \ crucible W_1 = weight \ of \ crucible sample W_2 = weight \ of \ crucible and \ residue \ i.e. after cooling.$

Density(ρ)= ^M

	4.0	
х	10	

Where M is the of piece and V is the mass test (g) measuringvolumeoftestpiece(cm³)byliquiddisplacementmethod ${\bf Specific Gravity Test}\ {\bf Subsequently their specific gravities were determined by dividing the unit weight of the sample in a indicating the sample i$ rbytheunitweight of the sample in air and water. The formula is show below. Specificgravity(sg)=<u>Wa</u> $W_{a-}W_{b}$

Where Wais the weight of sample in air (g); and W bis the weight of sample in water (g).

Result

Table 1: Brake pad Formation from the combination of PKF and GS

S/N	MATERIAL	FUNCTION	QUANTITY (g)	PROPORTION
1	Palm Kernel fiber	Base material	300	30.5
2	Groundnut Shell	Filler	80	10.2
3	Steel Fibre	Reinforcement	60	7.5
4	Calcium Carbonate	Hardening Agent	330	31.5
5	Resin Binder	Binding Agent	200	25.3
6	Graphite	Lubricant	32	4.5
7	Brass Chips	Abrasive	50.5	6.2

Table 2: Brake pad application with speed

WEAR RATE (mg)	DISC TEMPEATURE (⁰ C)	STOPING TIME (sec)	SPEED (Km/h)
10	2	2	10
15	4	2.5	20
20	6	5.0	30
25	8	6.5	40
30	10	7.0	50
35	12	12	60
40	14	10	70
45	16	12	80
50	18	14	90

V



Fig. 3: Graph of Brake pad application Vs speed

Discussion

Effect of epoxy resin binder and palm kernel fiber brake on pad performanceThepresenceofPKFandlowbindercontent(epoxyresin)inthemixtureincreasedth ewearratesof the pad. Thewaterandoilabsorption rate forGS-basedbrakepadwasclosedtotheconventional brakepad witha%deviationof0.001 and0.022.GS-based brake padsareenvironmentalfriendlyandcost effective. Whenthewt%ofPKFincreased in the composition, the compressive strength of the sample also increased. Thereason for the increase may beattributed to the percentage and ratio of the composite in thepad. The pad' hardness values increased when the wt% of PKFsamples increased in the composition. Assessment of the wear behaviour of the brakepadsampleswhensubjectedtodifferentspeedsispresented in Fig. 1. All the samples including the asbestosbased exhibited marginal increase in wear rate

80km/h. The asbestos brake pad lowest with speedup to had the we arrate followed by the PKF sample. The PKF based samples showed different resistances to we ar.The of

presence

PKS particles provides a higher thermal stability, increased a brasion and sliding we arresist ance and delay sthe transition of the trafrom mild to severewear (Olele, 2016).

However, when the brakepads amples were tested at speeds above 80 km/h, they presented sharp in creases in wear rates. It is well known that wear processinvolves fracture, tribochemicaleffects and plastic flow.Transitions between regions dominated bv each of

the secommon lygiver is eto changes in wear rate. This behaviour beyond 70 km/h speed could be due to subsurface the second sedeformation of the brake pad as a result of hightemperature.

The PKF brake pads generally did not show anydifference in behavior in terms of disctemperature risewith speed. They maintained the same Temperaturechange. However, at speeds below 40 km/h the asbestosbrake pad had a lower disc temperature rise while the PKFsamples maintained lower values of temperature change. The Asbestos sample was higher beyond 30 km/h speed. Thus, the PKF brake pads are a better choice ahead of theasbestos in applications where disc temperature rise is of great concern.

Atspeedbelow60km/hallthePKFpadswiththe exception of sample A, had lower stoppage time (thatis, better braking efficiency) when compared with the asbestos pad beyond 60 km/h and up to 90 km/h.

Conclusion

Although asbestos brake pads have good tribological and mechanical properties, they arecarcinogenicinnature.Fromthestudies,betterphysicalpropertieswereobtainedwithpalm

as the percentage kernel fiber brake pads of composition was optimized. In palmkernelfiber, both filler materials and binder improved the mechanical, tribological, and physic alproperties of the brake pads ampled eveloped. These studies also indicate that chemicaland physical treatments also used to subdue poor wettability and enable highermoistureabsorption. The wear rate improved with the addition of filler material in the brake padsamplepreparation. Wear rate of the developed composite also affected by the speed significantly. The smaller quantity of the filler, binder and fiber material had effect on the composite wearrateandgavebetter. The performance of the composite material such as mechanical and physical propertieshave been reported to be affected by the filler materials in the composition. In addition, various studies so far have found that as the composite filler material content decreases the properties for example shardness, compressive strength, thermal conductivity and tensile streng the strength of the streth, of the composite produced increase, while the density, oil and water absorption of the developed brack of the second seconakepadsampleincreaseswhen the filler content of the composite increases. Reference

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