



Green heterogeneous base catalyst from ripe and unripe plantain peels mixture for the transesterification of waste cooking oil

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ABSTRACT

Economical feedstocks such as agricultural wastes, food wastes, and waste cooking oil were used for biodiesel production to expand their application. Thus, a solid base catalyst was synthesized from a mixture of ripe and unripe plantain peels at a calcination temperature of 500 °C for 4 h. The catalyst was characterized using Scanning Electron Microscope (SEM), X-ray Diffraction (XRD) analysis, Fourier Transform Infrared (FT-IR) spectroscopy, Energy dispersive X-ray (EDX) analysis, and Brunauer-Emmett-Teller (BET) method. The waste cooking oil (WCO) used in this study was first pretreated with 3% (v/v) of H₂SO₄ via esterification reaction due to its high acid value. The esterified WCO was converted to biodiesel via transesterification reaction, and the process was then modeled and optimized using Taguchi L9 orthogonal array design method considering reaction temperature, reaction time, catalyst amount, and methanol/WCO molar ratio as the input variables. Based on the results, the synthesized catalyst predominantly contained potassium phases with 45.16 wt.%. The morphology of the catalyst revealed a crystalline mesoporous nanocomposite. At the end of WCO esterification, the acidity of the oil decreased from 5 to 1 mg KOH/g. The optimal conditions established for the transesterification process were catalyst amount of 0.5 wt.%, methanol/WCO molar ratio of 6:1, reaction temperature of 45 °C, and reaction time of 45 min with a corresponding biodiesel yield of 97.96 wt.%. The quality of the biodiesel produced satisfied the specifications (ASTM D6751 and EN 14,241) recommended for biodiesel fuels. Hence, a blend of ripe and unripe plantain peels could serve as an efficient heterogeneous base catalyst in producing biodiesel from WCO.

1. Introduction

The petroleum oil price and the environmental pollution generated from fossil fuels have made biofuel production a developing area. Other alternative non-polluting technologies such as hydrogen-fueled vehicles, hybrids, plug-in electrics, compressed natural gas, and solar enable devices are possible solutions to energy and environmental problems [1]. However, biofuels are renewable energy sources with vast potential to replace fossil fuels in the present and future energy mix [2]. Biomass sources from which biofuels are naturally abundant can ensure the stability of the ecosystem. Biodiesel, a form of biofuel, is a renewable source of energy with several environmental benefits.

The methods involved in producing biodiesel depend on the catalyst used in the process. Different catalysts that have been deployed in

producing biodiesel are base catalysts (e.g., KOH or NaOH), acid catalysts (e.g., H₂SO₄), and enzymes (e.g., lipases) [3]. Currently, investment in biodiesel production is adversely affected by several factors including, production technology type, nature of raw materials, and market feedstocks price [4]. Biodiesel price doubles that of mineral diesel in the market, due to its production cost. Biodiesel production cost involves two parts, viz., raw materials (feedstocks) cost and processing cost [5].

It is argued that the biodiesel production cost depends on the technological routes; however, the cost of feedstock, regardless of the technology employed, accounts for the major production costs [6, 7]. To produce economical biodiesel would require a cheap source of feedstock such as non-edible oil, used cooking oil, and animal fats [4]. Generally, non-edible oils are not consumed by humans due to the presence of toxic

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compounds. Also, their availability and the low cost of plantation of non-edible oil crops position them as economically viable feedstocks. However, many non-edible oils are not suitable for biodiesel synthesis and require pretreatment [2, 4]. Several developed countries, due to climate change threats, are embracing alternative energy sources. The increasing interest in biofuel is due to its intrinsic advantages over fossil fuels. To this effect, the growth projection of the biodiesel industry may place huge pressure on nature and biodiversity in emergent nations [11]. One way to address this pending crisis is to adopt total biomass-dependent fuel. Waste cooking oil is a good raw material for biodiesel development since it is cheap and abundant in several countries [8]. Its usage in biodiesel synthesis will prevent competition from using edible oils [9, 10]. Apart from converting waste cooking oil (WCO) to fuel, the problems associated with its disposal are equally solved [11]. It is estimated that a minimum of 16.54 million tons of WCO is produced every year, covering major countries in Europe, Asia, and North America [12].

Conventionally, biodiesel is industrially produced using homogeneous catalysts. The use of homogeneous catalysts for the transesterification of triglycerides is not friendly to the environment and equipment, while the enzymatic transesterification process is expensive [13, 14]. The use of solid catalysts for biodiesel production has several advantages: minimum energy consumption, lower material costs, and minimum water usage [15]. This class of catalysts can easily be separated from the product and recycled many times [16, 17]. Green catalysts are mostly derived from biomass resources such as plants or animals [18, 19]. Catalysts from natural biological sources contain calcium and potassium compounds dominantly. These elements are potential solid base catalysts for the transesterification of triglycerides [17, 20]. The production of green catalysts involves a sequence of easy methods with minimal energy requirements [21]. Green feedstock and its products contribute zero carbon emission since these materials come from photosynthesis, a biological process that includes sunlight, water and CO₂ [22]. Biomass sources are preferred in developing catalysts since they are abundant, cheaper, renewable, and sustainable [15, 17]. Previous studies have reported different plant biomass in producing solid catalysts employed in the transesterification reaction [16, 23]. The rationale is to discover abundant plant biomass of high catalytic properties which can be efficiently used sustainably. Previous reports have demonstrated the use of banana and plantain wastes. These include *Musa Balbisiana* colla trunk [24], plantain peels [25], banana peels [17], *Musa Balbisiana* Colla peels [26], *Musa acuminata* [13] peduncle, and banana peduncle [27]. Also, a mixture of plant biomass has been used to develop solid catalysts; cocoa pod husk-plantain peel [28] and cocoa pod husks-plantain peel-kola nut husk [29]. The activity of a catalyst developed from ripe plantain peels is attributed to its microstructural nature and high K content [30]. Likewise, a catalyst prepared from unripe plantain peels contained a high K proportion [25]. In both studies, the presence of high K in the catalysts was responsible for the catalytic activity in the transesterification process. However, there is no information in the literature on the application of ripe and unripe plantain peels mixture as heterogeneous catalyst for biodiesel production.

This study reports the synthesis of a green solid catalyst from agricultural wastes and its subsequent application in the transesterification process. The focus was to produce biodiesel from WCO via transesterification using the base catalyst developed from ripe and unripe plantain peels mixture. The heterogeneous base catalyst was synthesized from a mixture containing an equal proportion of ripe and unripe plantain peels ashes. The biodiesel yield and the process parameters (reaction time, molar ratio of oil to methanol, reaction temperature, and catalyst loading) influencing the process were investigated using the Taguchi approach. The optimization of the transesterification of WCO was carried out to maximize the utilization of raw materials, and hence, lessen the production cost.

2. Materials and methods

2.1. Materials

The WCO, together with the ripe and unripe plantain peels used in this study, were obtained from the Cafeteria Kitchen of Landmark University, Omu-Aran, Kwara State, Nigeria. Analytical grade reagents and chemicals were used for the study.

2.2. Synthesis of catalyst from plantain peels

The two types of plantain peels used in this study were treated separately before calcination. The ripe and unripe plantain peels collected were cut into small pieces, then washed in distilled water to remove the dirt and stones. The washed peels were oven-dried at 100 °C until a constant weight was observed. The plantain peels were burnt, and the resultant ashes were mixed in equal proportion (ripe:unripe of 50:50). The mixed powdered ash was loaded in the porcelain crucible and calcinated in the muffle furnace at 500 °C for 4 h, according to Betiku and Ajala [25]. After calcination, the calcined ripe-unripe plantain ash (CRUPA) was sieved using 80 mesh size, and the obtained ash was stored for further work.

2.3. Characterization of CRUPA

The synthesized catalyst was characterized to investigate its features. Energy-dispersive X-ray spectroscopy (EDS) of CRUPA was performed to determine the elemental contents using a Thermo Fisher Nova Nano-SEM, and the diffractogram generated was recorded with an Oxford X-max 20mm² detector (Oxfordshire, U.K.). The functional groups on the active site of the CRUPA surface were determined by Fourier Transform Infrared (FTIR) analysis using a Perkin Elmer UATR Two spectrophotometer (Llantrisant, U.K.). The spectra generated were recorded over the range of 4000 to 400 cm⁻¹. The crystalline phases in CRUPA were determined using X-ray diffractometer D8-Advance from Bruker AXS (Germany). The measurement involves a continuous scan in locked coupled mode with Cu-K_α radiation with LynxEye detectors. The sample measurements run within a range of 2θ with a typical step size of 0.034° in 2θ. The surface morphological features of CRUPA were taken using Tescan MIRA3 RISE scanning electron microscopy (SEM). The Brunauer-Emmett-Teller (BET) method of adsorption/desorption on nitrogen gas was used to determine the surface area and porosity of CRUPA using a Micromeritics instrument (Tristar II 3020). The catalyst pore size distribution, width, and adsorption pore volume were determined by Barret-Joyner-Halenda (BJH) method.

2.4. Pretreatment of WCO

WCO was centrifuged to remove the solid residues and particles left after being used. Thereafter, the oil was heated at about 100 °C to evaporate any water moisture that might be trapped. After cooling, the physicochemical properties of WCO (acid value, density, kinematic viscosity, saponification value, and iodine value) were measured according to the standard methods [31]. The physicochemical characterization of the WCO had an acid value of 5 mg KOH/g oil. Hence, WCO was subjected to the esterification process to reduce its acid value [19]. The esterification of WCO was carried out in a two-neck glass reactor placed on a hotplate with a magnetic stirrer. The esterification conditions were methanol/WCO molar of 6:1, reaction temperature of 65 °C, reaction time of 45 min, and catalyst amount of 3% (v/v). For this work, 50 mL of WCO was measured, and methanol of known volume was added. After the process, the esterified oil was separated from the unused methanol and the catalyst using a centrifuge (TMT-XC-450-PRP) machine. The esterified oil was dried over a hotplate at 100 °C and stored for further use.

Table 1

Elemental constituents of CRUPA.

Composition (%)	C	O	Mg	Si	K
Mean value	12.02	35.34	3.61	3.87	45.16
Standard deviation	4.5	3.68	1.61	1.76	2.79

2.5. Experimental design and procedure for WCO transesterification

The transesterification of WCO and methanol using a synthesized catalyst (CRUPA) was carried out according to the conditions stipulated by Taguchi orthogonal array (L9) design method. The L9 design was chosen for its simplicity and ability to provide exhaustive information about the process parameters and the yield using a few experiments [32]. The chosen parameters in this present study were based on our previous studies [33, 34]. The different levels of each parameter used in this work are methanol-to-WCO molar ratio (6:1, 9:1, and 12:1), reaction temperature (45, 55, and 65 °C), reaction time (50, 70, and 90 min) and catalyst loading (0.5, 1.5, and 2.5 wt.%). The Design-Expert version 10.0 (Stat-Ease Inc., Minneapolis, MN, USA) statistical software was used for the Taguchi design, response prediction, and regression analysis in this study.

Biodiesel production from WCO was conducted in a three-necked glass reactor placed on a hotplate with a magnetic stirrer. A glass reflux condenser was mounted on the reactor to prevent methanol evaporation from the reaction mixture. For the transesterification, based on the methanol-to-oil ratio in the design, a known volume of WCO was charged into the glass reactor and a known amount of methanol was added to it. The mixture was allowed to heat to the set temperature under continuous stirring. Then, a known amount of CRUPA was added, and the reaction continued for a time suggested by the model. After the transesterification process, the mixture was centrifuged to separate methanol, biodiesel, and the residual catalyst. The biodiesel was washed three times with warm distilled water (65 °C) to remove impurities and residual methanol. The biodiesel obtained (waste cooking oil methyl esters - WOME) was dried, and the yield was gravimetrically determined using Eq. (1).

$$\text{Biodiesel yield} = \frac{\text{weight of WOME produced}}{\text{weight of WCO used}} \times 100 \quad (1)$$

2.6. Physicochemical properties of WOME

The quality of WOME produced was ascertained by determining the physicochemical properties of the fuel. The specific gravity, saponification value, acid value, density, iodine value, and kinematic viscosity were determined using the standard method [31]. Other fuel parameters such as cetane number [35] and calorific value [36] were also determined. The FTIR analysis of WOME was carried out using an FTIR-UATR spectrophotometer.

3. Results and discussion

3.1. Characterization results of CRUPA

3.1.1. Elemental composition of CRUPA

The mean (wt.%) of the elemental composition of CRUPA as revealed by the EDS analysis is presented in Table 1, and the elements present are C, O, Mg, Si, and K.

The results show a considerable presence of potassium (K) in the form of K_2O and K_2CO_3 as well as other traces of metallic oxides. This indicates that the developed CRUPA has high basicity due to the presence of metallic oxide. As a result, the synthesized catalyst could be an efficient heterogeneous catalyst in expediting the transesterification of biomass oil. The K content (45.16 wt.%) in this study was more than the K value (41.37 wt.%) found in the calcined banana peel ash [26]. However, a higher K content was earlier reported in previous studies that used either ripe plantain peels or unripe plantain peels. The K content (54.73 wt.%) was reported for a calcined sample (temperature: 500 °C) developed from unripe plantain peels [25]. Balajii and Niju [27] reported K (68.37 wt.%) as the major element in the banana peduncle calcined at 700 °C, other elements such as Mg, Ca, O were present. Similarly, Etim et al. Etim, Betiku [30] reported K (51.02 wt.%) as the dominant element, and the presence of elements such as O, P, Ca, Mg, S, Cl, Al, and Si in the calcined ripe plantain peels at 700 °C. The observed

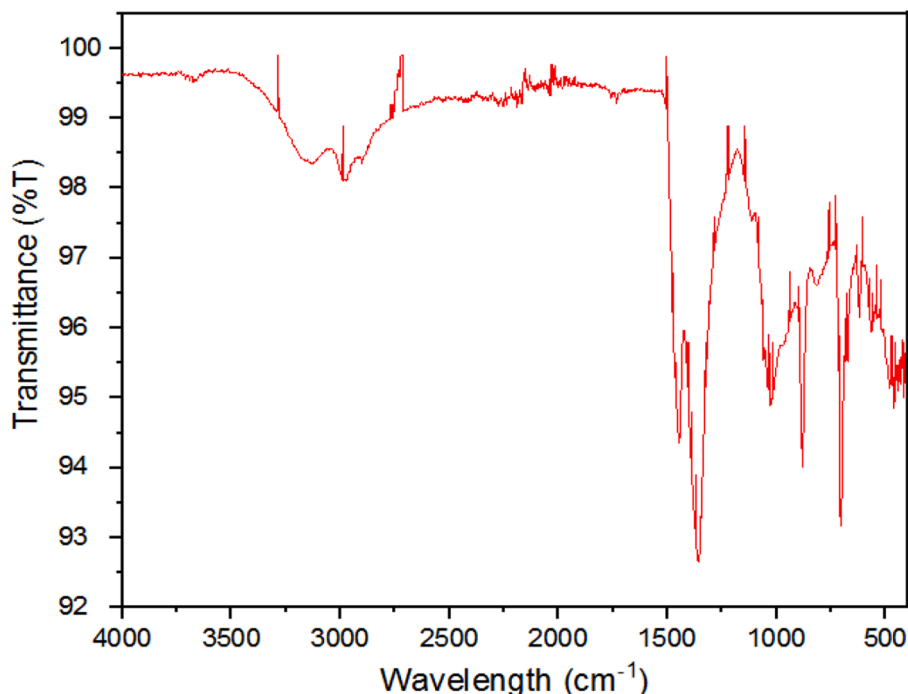


Fig. 1. FT-IR spectrum of CRUPA.

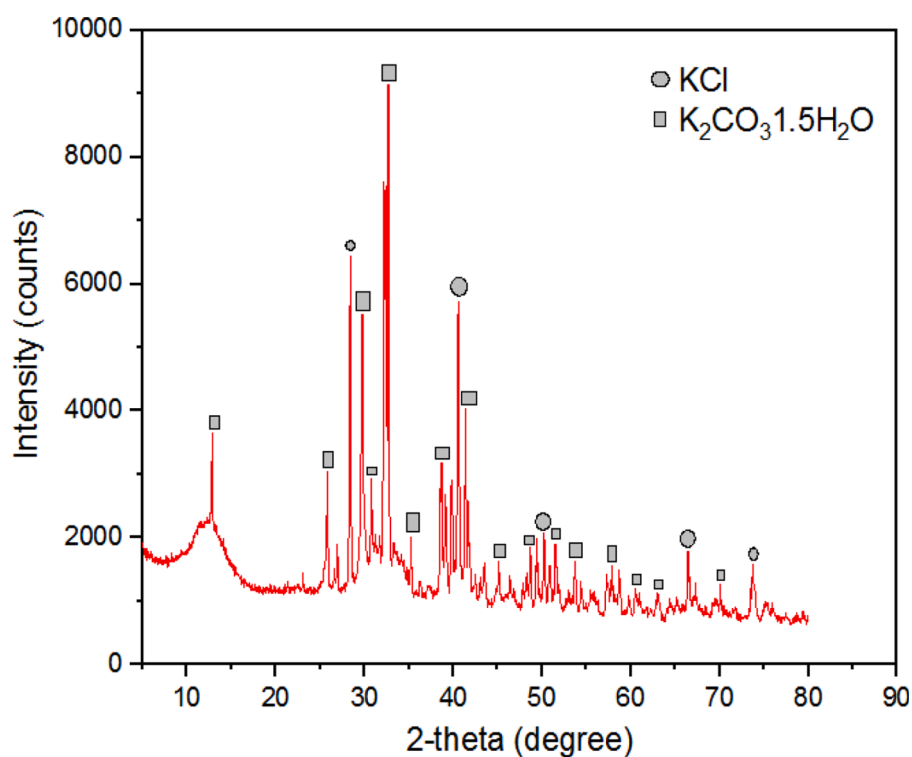


Fig. 2. XRD diffractogram of CRUPA.

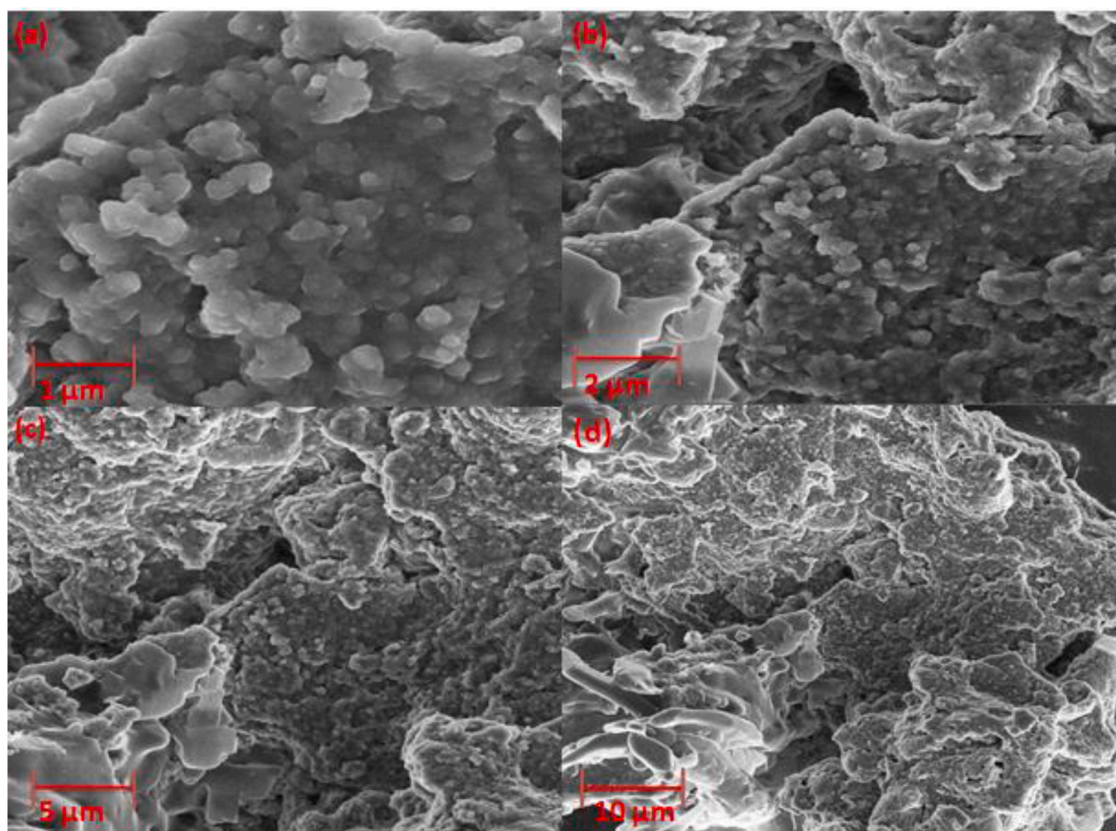


Fig. 3. (a–d) SEM images of CRUPA.

difference in the elemental constituent could be attributed to the calcination temperature used, and the blending of ripe and unripe plantain peel ashes.

3.1.2. FT-IR results of CRUPA

FT-IR spectroscopy was used to identify the vibration of chemical bonds in the analyzed sample. Each vibration uniquely describes a

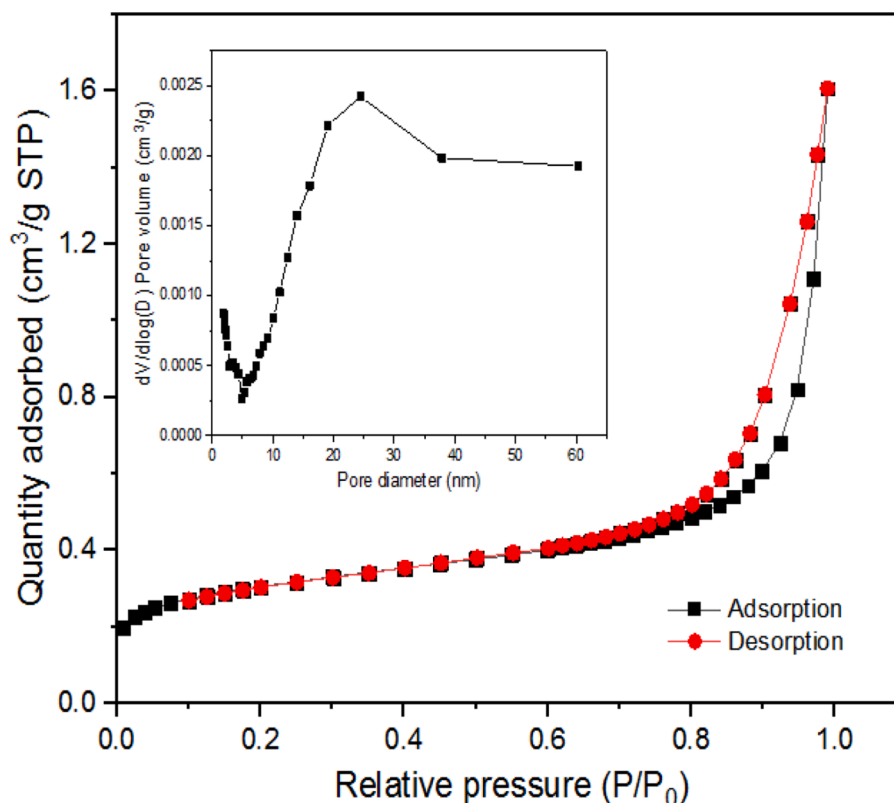


Fig. 4. BET adsorption isotherm and BJH pore-size distribution (*inset*) of CRUPA.

certain functional group; hence, the feature of the spectrum gives structural information on the catalyst molecules. The FT-IR spectrum of CRUPA is shown in Fig. 1.

The absorption vibration at 1380 cm^{-1} was assigned to asymmetric stretching of carbonate functional group while the band at 900 cm^{-1} was due to out-of-plane bending vibration of the C–O group [19, 37], hence, indicating the presence of carbonate compound (K_2CO_3) which has been identified around this band [23]. The band around 1430 cm^{-1} can be ascribed to atmospheric CO_2 adsorption onto metal oxide [27]. The absorption at 1000 cm^{-1} can be attributed to Si–O–Si vibration, which is prominent in the calcined material [26]. The weak adsorption around $3100\text{--}2900\text{ cm}^{-1}$ may be assigned to the M–K–O group, where ($M = \text{Mg}$ or Si). The results in this section corroborate the EDS results.

3.1.3. XRD results of CRUPA

The bulk phases of the CRUPA were identified using XRD analysis. The XRD peak of the synthesized catalyst is depicted in Fig. 2. According to the results, the two phases identified in this catalyst are potassium chloride (KCl) and potassium carbonate hydrate ($\text{K}_2\text{CO}_3 \cdot 1.5\text{H}_2\text{O}$).

The lattice structure of KCl was face-centered cubic while that of potassium carbonate hydrate was monoclinic. The XRD pattern of KCl phase showed clear diffraction peaks at $2\theta = \sim 15, 28, \sim 42, 50$, and 67° while $\text{K}_2\text{CO}_3 \cdot 1.5\text{H}_2\text{O}$ exhibits strong intensity at $2\theta = 26, 30, 33, 40$, and 43° . In the XRD diffractogram of calcined banana peels, KCl and $\text{K}_2\text{CO}_3 \cdot 1.5\text{H}_2\text{O}$ are the dominant two crystalline phases [17]. Meanwhile, different crystalline compounds were identified in calcined ripe plantain peels [30]. Only KCl was among the crystalline phases.

3.1.4. SEM results of CRUPA

The surface morphology of CRUPA was examined by SEM. The SEM images of the catalyst taken at different micrometer (μm) scales are shown in Fig. 3(a–d), which revealed the microstructure similar to the morphology earlier reported [17, 28].

Fig. 3a shows a large aggregate of spherical particles clumped

together. The SEM image at (3b) shows increasing agglomerated particles with regular size distribution. The SEM image becomes clearer in (3c) with a highly ordered surface with large cracks across the image [30]. At $10\text{ }\mu\text{m}$ (3d), the porosity of CRUPA was highly noticeable, having a glossy-like surface [26]. The mesoporous and microporous structures of the developed catalyst become clearer as the microscale increases.

3.1.5. Surface area, pore-volume, and pore width of CRUPA

The synthesized catalyst (CRUPA) had a BET surface area of $1.10 \pm 0.0050\text{ m}^2/\text{g}$, pore volume of $0.002366\text{ cm}^3/\text{g}$, and pore size width of 13.15 nm . The surface area compared favorably to the earlier reported results; $1.487\text{ m}^2/\text{g}$ from *Musa Balbisiana* trunk [24]. The BET isotherm of CRUPA is displayed in Fig. 4. The adsorption isotherm exhibited by CRUPA is a typical Type IV isotherm. This means that adsorbate would condense in the tiny capillary of the catalyst at a pressure below saturation pressure of the gas [38]. At the lower pressure, the inflexion region represents monolayer formation while capillary condensation or multilayer formation occurs after monolayered surface coverage of the pore walls [38, 39].

The adsorption of the adsorbate on the pore surface steadily increases until it jumps at the relative pressure (P/P_0) equals 0.9. The BJH pore-size distribution of CRUPA is depicted in Fig. 4(*inset*). The plot exhibits a strong interaction between adsorbate and calcined sample surface between the pore diameters of 5–25 nm. The synthesized catalyst in this study can be classified as a mesoporous material, that is, material or particles having a pore diameter within 2–50 nm [29, 39].

3.2. Results of wco transesterification

The pretreatment of high acid value oil is essential in producing biodiesel [40]. The esterification reaction of WCO using H_2SO_4 as a catalyst proved effective in reducing the acid value of the oil from 5 mg KOH/g oil to 1 mg KOH/g oil, which meant that 80% acid value

Table 2
Taguchi model of actual and predicted biodiesel yield.

Run	A: MeOH/ WCO ratio	B: Temp. °C	C: Cat. loading wt.%	D: Time min	Actual WOME wt.%	Predicted WOME wt.%
1	9:1	65	0.5	70	62.37	62.49
2	9:1	45	1.5	90	66.02	65.98
3	9:1	55	2.5	50	60.35	60.27
4	6:1	65	2.5	90	90.56	90.48
5	6:1	45	0.5	50	96.39	96.51
6	6:1	55	1.5	70	87.34	87.31
7	12:1	65	1.5	50	81.94	81.90
8	12:1	55	0.5	90	76.07	76.19
9	12:1	45	2.5	70	84.51	84.43

reduction was achieved within the experimental conditions. The esterified WCO was directly transesterified to biodiesel since its acid value was within an allowable range. The actual yields and predicted biodiesel yields obtained from the transesterification of WCO under 9 experimental conditions are presented in Table 2. It was observed that the actual and predicted biodiesel yields are comparable, demonstrating the accuracy of the model in predicting the responses.

Run 5 at the lowest level of the operating parameters gave the highest biodiesel yield of 96.39 wt.%. The lowest biodiesel yield (60.35 wt.%) was recorded at Run 3, with the lowest reaction time, the upper level of catalyst loading, and the middle level of temperature and methanol/oil molar ratio. The results obtained from the transesterification of WCO were fed into the software as the responses for the experimental conditions generated by the Taguchi L9 design to obtain the regression model equation for the process. The linear regression model in terms of coded factors that described the transesterification of WCO is presented in Eq. (2). This model equation is useful in predicting biodiesel yield at a given level of each parameter.

$$WOME = +78.39 + 13.04A[1] - 15.48A[2] + 3.91B[1] - 3.81B[2] - 1.17D[1] - 0.32D[2] \quad (2)$$

where A[1] and A[2] represent methanol/WCO molar ratio at 1st and 2nd level, B[1] and B[2] represent the temperature at 1st and 2nd level, while D[1] and D[2] are the reaction time at 1st and 2nd level, respectively. The relative impact of an individual parameter on the yield can be identified by comparing the coefficient of the regression equation.

An analysis of variance (ANOVA) was used to statistically examine the model equation to determine the relative influence of process parameters considered. The p-value of the model and each parameter at a 95% confidence level (p-value ≤ 0.05) was used to determine the significance of these terms. From the results (Table 3), the model is significant, including the methanol/WCO molar ratio, temperature, and reaction time. The impact of catalyst loading on the transesterification process cannot be captured statistically; hence, it is insignificant within the range investigated. The relative contribution of the individual parameter was quantified by the ratio of the pure sum of squares to the total sum of squares. The methanol/WCO molar ratio is the most influential parameter, followed by temperature. Also, reaction time had minimal effect on the biodiesel yield since it has the lowest F-value among the parameters investigated.

Table 3
ANOVA results of model and process parameters.

Source	Sum of Squares	Degree of freedom	Mean Square	F-value	p-value
Model	1342.92	6	223.82	6880.67	0.0001
A-MeOH/WCO	1247.01	2	623.51	19,167.85	< 0.0001
B-Temperature	89.37	2	44.68	1373.69	0.0007
D-Time	6.54	2	3.27	100.47	0.0099
Residual	0.0651	2	0.0325		
Corrected Total	1342.98	8			

The statistical fitness of this model was evaluated (Table 4). The coefficient of determination (R^2) in this study was 0.9998, which indicates that the experimental data obtained from the chosen model fitted perfectly. The R^2 value is close to 1, an indication that 99.98% of the variation observed in biodiesel yields with respect to the process parameters can be explained by the model. The adjusted R^2 and predicted R^2 values are 0.9993 and 0.9990, respectively. These values are in reasonable agreement with one another and less than the maximum allowable difference of 0.2. The standard deviation of this analysis was 0.1804, which shows that the results obtained from this study are reliable. The coefficient of variation (C.V.%) was 0.2301; the lower the value, the better the capability of the model in optimizing the process [41]. The adequate precision of the model was 227.82; a value greater than 4 is considered adequate [40]. Meanwhile, the value obtained is very high, which indicates that the model has sufficient signal to optimize the transesterification of WCO.

3.3. Parametric influence of reaction parameters

3.3.1. Effect of methanol/WCO molar ratio on WOME yield

The influence of methanol/WCO molar ratio on WOME yield at a temperature of 45 °C, a reaction time of 50 min, and 0.5 wt.% catalyst loading is displayed in Fig. 5.

Generally, transesterification reaction according to the stoichiometric ratio requires a methanol/oil molar ratio of 3:1. However, excess methanol is needed to shift the equilibrium position to favor forward reaction. In this study, maximum biodiesel yield (96.39 wt.%) was reached at a methanol/oil molar ratio of 6:1. Oladipo and Betiku [42] obtained 96.97 wt.% of *Hevea brasiliensis* oil methyl ester using a methanol/oil molar ratio of 6:1. Increasing the molar ratio to 9:1 led to a sharp reduction in biodiesel yield. The reason could be due to excessive dilution of oil by the methanol, thereby decreasing the rate of transesterification reaction [43]. A marginal increase in biodiesel yield was observed when the molar ratio was later increased to 12:1. A similar observation was reported in Oraegbunam, Oladipo [32], the maximum biodiesel yield was reached at a methanol/oil molar ratio of 6:1, and excess methanol to the reaction mixture led to a reduction in biodiesel yield. Excess methanol was reported to have increased glycerol solubility, which interferes with the ester phase and subsequently lowers biodiesel yield [11].

3.3.2. Effect of reaction temperature on WOME yield

The influence of temperature on WOME yield at methanol/WCO molar ratio of 6:1, catalyst loading of 0.5 wt.%, and reaction time of 50

Table 4
Quality of statistical parameters.

Parameters	Value
Standard deviation	0.1804
Mean	78.39
Coefficient of Variation (%)	0.2301
R^2	0.9998
Adjusted R^2	0.9993
Predicted R^2	0.9990
Adequate precision	227.8172

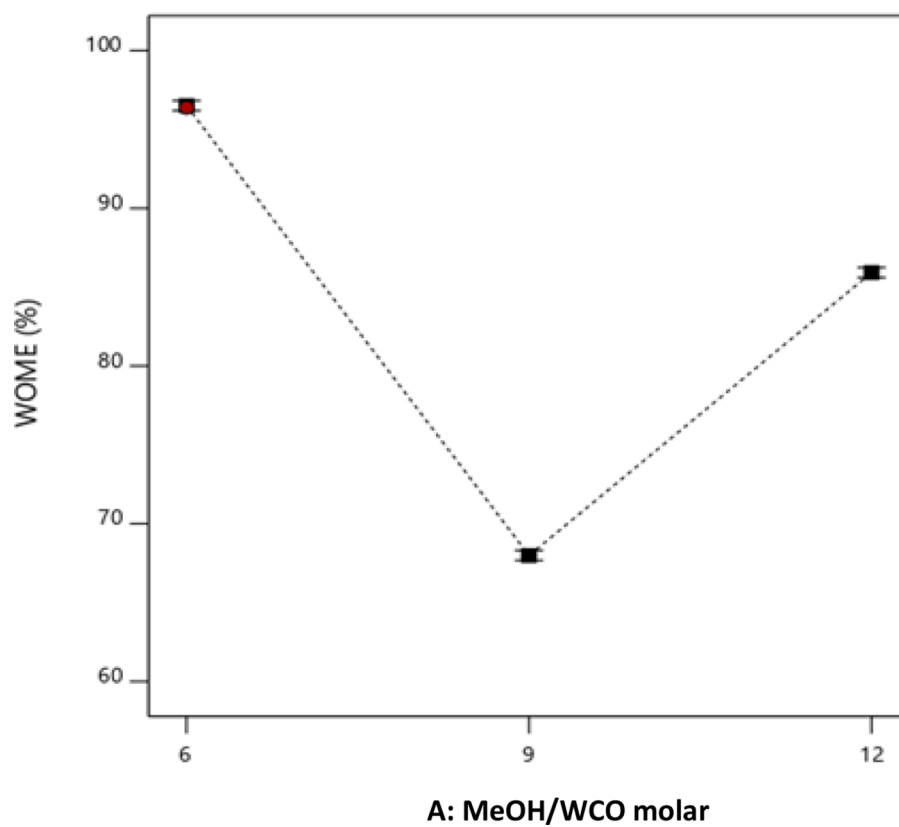


Fig. 5. Plot of methanol/WCO molar ratio effect on WOME yield (wt.%).

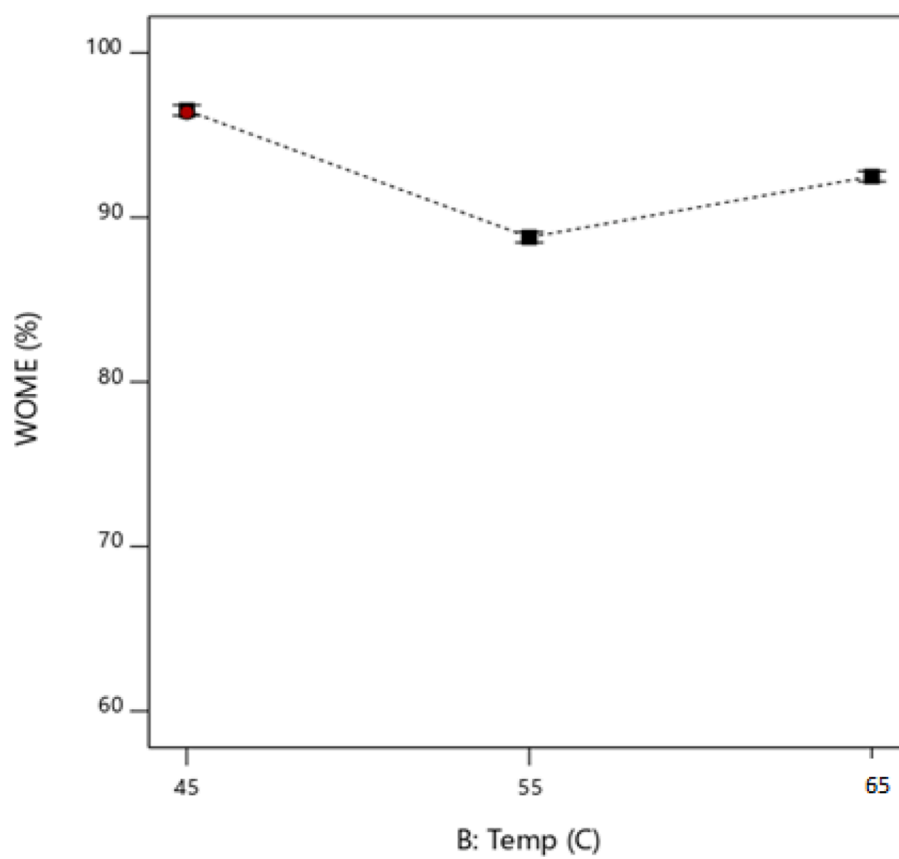


Fig. 6. Influence of temperature on WOME yield (wt.%).

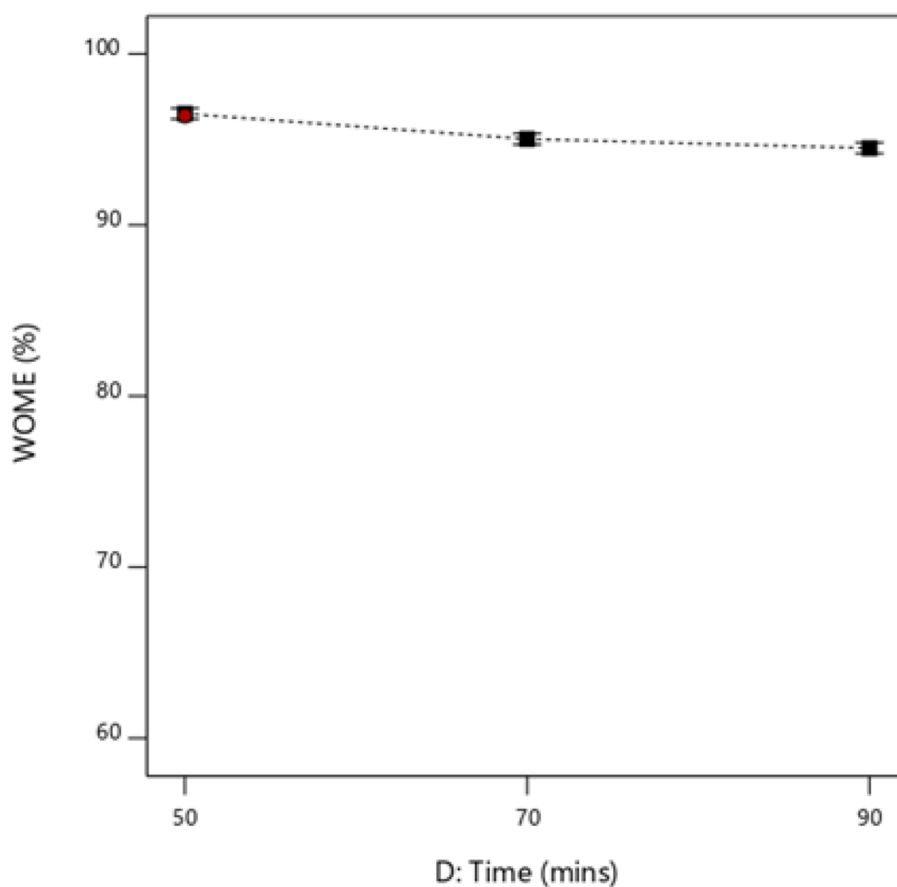


Fig. 7. Influence of reaction time on WOME yield (wt.%).

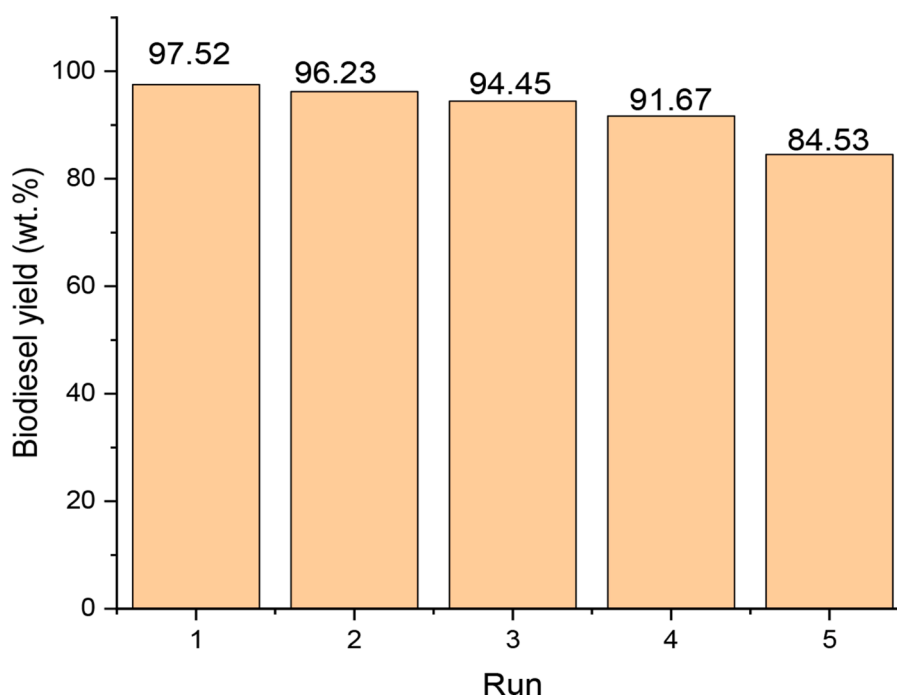


Fig. 8. Reusability results of CRUPA.

min is depicted in Fig. 6.

The biodiesel yield was at the peak at the lowest level of the temperature (45 °C). It can be seen that the biodiesel yield decreased when

the temperature was increased, and the yield again marginally increased with a further increase in temperature to 65 °C. The response of biodiesel yield to changes in temperature demonstrated the effect of

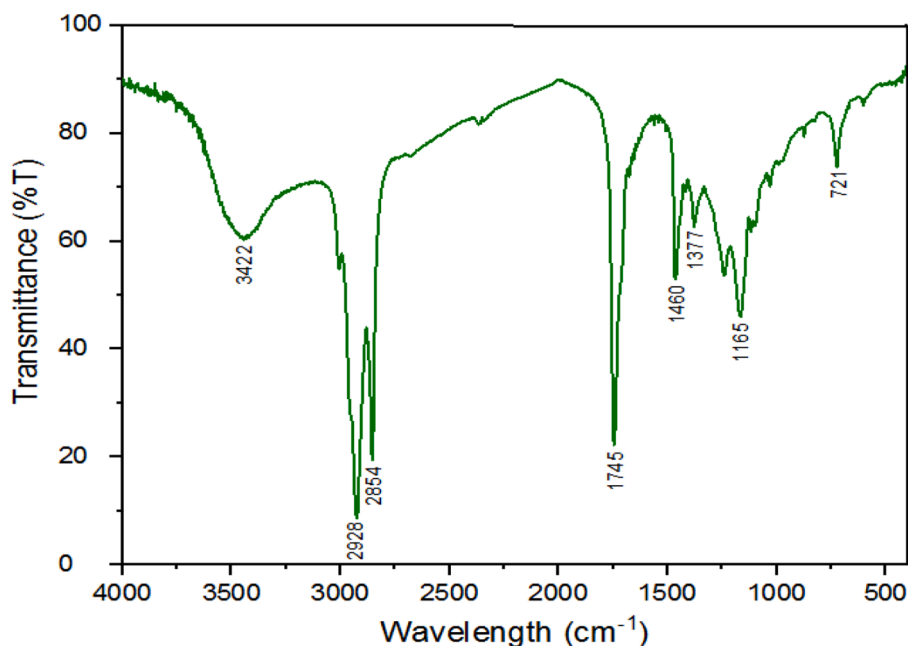


Fig. 9. FT-IR spectrum of WOME.

Table 5

Wavenumber and vibration types present in WOME.

Wavenumber (cm ⁻¹)	Functional group	Vibration type	References
3422	-O-H	Stretching	[18, 28]
2928	-CH ₂	Asymmetric stretching vibration	[33, 49]
2854	-CH ₂	Symmetric Stretching vibration	[28, 47]
1745	-C = O	Carbonyl stretching vibration	[33, 46]
1460	-CH ₂	Shear-type vibration	[23, 45]
1165	-C-O-C	Anti-symmetric stretching vibration	[46, 49]
721	-CH ₂	Plane rocking vibration	[34, 45]

temperature on the equilibrium constant of the transesterification process. The influence of temperature observed in this study aligns with earlier studies. Soji-Adekunle, Asere [10] reported that biodiesel yield decrease with an increase in reaction temperature. The authors produced a maximum biodiesel yield of 94 wt.% at a temperature of 55 °C. Similarly, a biodiesel yield of 96.43 wt.% was obtained at the reaction temperature of 35 °C. A marginal increase in biodiesel yield was observed at the reaction temperature (65 °C) close to the boiling point of

methanol. This could be attributed to the turbulence in the system, which subsequently strengthens the mass transfer rate of reacting molecules [43].

3.3.3. Effect of reaction time on WOME yield

The impact of reaction time on the WOME yield at methanol/WCO molar ratio of 6:1, a temperature of 45 °C, and 0.5 wt.% CRUPA is shown in Fig. 7.

At the lowest level of reaction time, 96.39 wt.% of biodiesel yield was obtained. Further increment in reaction time led to a marginal reduction in biodiesel yield. According to the results, increasing the reaction time from 70 to 90 min at a constant temperature, CRUPA loading, and MeOH/WCO molar ratio had minimal effect on the biodiesel yield. Oladipo, Ojumu [33] reported that 40 min was sufficient to produce 96.53 wt.% of biodiesel from moringa oil. Since transesterification is a reversible reaction [44], produced biodiesel can be converted to triglycerides at a longer reaction time. This could be the reason the biodiesel yield decreases with increasing reaction time.

3.4. Establishment of optimal condition and model validation

The optimal condition for the transesterification of WCO to WOME by CRUPA was established by solving Eq. (2) using the numerical optimization method of the Design-Expert statistical software. For the

Table 6

Properties of biodiesel synthesized from waste cooking oil using different catalysts.

Properties	Unit	This study		Gohain, Laskar [50]	Niju, Ajieth Kanna [51]	Soji-Adekunle, Asere [10]	Standards	
		WCO	CRUPA	Calcined pawpaw stem	Calcined sugarcane bagasse	KOH	EN 14,214	ASTM D6751
Density (25 °C)	kg/m ³	981	890	870	878	852	860 - 900	880
Acid value	mg KOH/g oil	5.00	0.46	0.06	0.45	0.32	0.5 max	0.5 max
Kinematic viscosity	mm ² /s	6.22	3.90	3.10	4.75	5.87	3.5 - 5	1.9 - 6
Iodine value	g I ₂ /100 g oil	122.02	71.28	—	—	74.67	< 120	NS
Flash point	°C	—	—	150	158	176	—	—
Saponification value	mg KOH/g	186.45	154.5	—	—	58.10	NS	NS
Cetane number	—	48.19	60.12	57	—	58.64	51 min	47 min
Calorific value	MJ/kg	39.97	42.34	40.30	—	45.93	35	NS

process optimization, the catalyst loading was set to its lowest (0.5 wt. %) because it is an insignificant parameter in the process. The other parameters were chosen within the range investigated while the WOME yield was set at maximum. The optimum values predicted were catalyst loading of 0.5 wt.%, methanol/WCO molar ratio of 6:1 (v/v), reaction temperature of 45 °C, and reaction time of 45 min with a corresponding biodiesel yield of 97.96 wt.%. The performance of the model was validated by conducting the transesterification of WCO under optimum conditions predicted by the regression model. The experiment at the optimum values was repeated thrice, and the average value of biodiesel yield obtained was 97.52 wt.%. The average biodiesel yield is close to the predicted biodiesel yield, which signifies that the model used in this study is capable of predicting the optimum values and adequately describes the process.

3.5. Results of CRUPA reusability

One of the advantages of the heterogeneous catalyst over the homogeneous catalyst is recycling the catalyst and reapplying it in another production cycle. The reusability test was conducted to determine the number of production cycles this catalyst can be reused. This test was conducted at the optimum conditions of the model. At the end of each run, the solid catalyst was separated and applied to another transesterification set-up without any treatment. A plot of biodiesel yields against production run is presented in Fig. 8. It was observed that the synthesized catalyst could be reused for five production cycles without losing a substantial amount of biodiesel. At the end of the 5th cycle, the biodiesel yield was reduced by 13.32%. The loss of catalytic activity of the catalyst can be attributed to partial blocking of catalyst pores by reacting molecules which subsequently prevent other triglyceride molecules from accessing the active catalyst surface [45].

3.6. FT-IR spectrum of WOME

The FT-IR spectrum of the WOME is shown in Fig. 9. The absorption bands present and the corresponding vibrational types are depicted in Table 5. The stretching vibration of the hydroperoxide functional group was assigned to the weak band at 3422 cm^{-1} . The two strong absorption bands that characterized the presence of ester: the carbonyl group ($\text{C}=\text{O}$, 1745 cm^{-1}) and anti-symmetric axial stretching ($\text{C}-\text{O}-\text{C}$, 1165 cm^{-1}) [46, 47] are present. The signals at 2928 and 2854 cm^{-1} are assigned to the asymmetric and symmetric stretching vibration, respectively [18, 47]. The FT-IR spectra obtained has a notable fingerprint region similar to spectrum observed in biodiesel produced from *Canarium odontophyllum* [48]. Thus, the FT-IR spectrum obtained indicates that the transesterified WCO possessed characteristic signals of fatty acid methyl esters, which confirms that the transesterification process was successful.

3.7. Physicochemical properties of WOME

The physical and chemical properties of WCO and biodiesel produced using CRUPA were characterized, and the results, along with the recommended standards (ASTM D6751 and EN 14,214) for biodiesel, are presented in Table 6. Based on the results, there was an improvement in the physicochemical properties of the biodiesel over WCO. This further highlights the importance of transesterification in the conversion of triglycerides to methyl esters. The acid value of WOME synthesized in this study compared with the value reported in other studies [10, 50] indicates that the biodiesel is stable and will minimize the deposit rate in the fuel system. The kinematic viscosity, cetane number, calorific value, and density of the produced WOME show that the biodiesel has a good ignition quality and sufficient energy capacity. The values also compared favorably with the biodiesel produced from WCO using different catalysts and the values recommended for biodiesel standards.

4. Conclusion

A mixture of ripe and unripe plantain peels used in synthesizing solid base catalyst is a viable feedstock. The catalyst (CRUPA) had high catalytic properties to transesterify WCO to biodiesel. The synthesized solid base catalyst predominantly contains potassium compounds; the K content was 45.16 wt.%. Factors such as temperature, reaction time, catalyst loading, and methanol/oil molar ratio affect the response of the transesterification process. The optimal condition established for WOME production from WCO using CRUPA was catalyst loading of 0.5 wt.%, methanol/oil molar ratio of 6:1, a temperature of 45 °C, and reaction time of 45 min, with a corresponding biodiesel yield of 97.96 wt.%. The biodiesel produced (WOME) had physicochemical properties which met biodiesel standards (ASTM D6751 and EN 14,241). Thus, a blend of calcined ripe-unripe plantain peels could be utilized as an efficient heterogeneous catalyst for biodiesel production.

Ethical approval

The research results presented in this manuscript are the original works of the authors. This paper has not been published elsewhere in any form or language. The manuscript has not been submitted to another journal for consideration. The authors of this work have presented their results clearly without any falsification or misrepresentation of data. All references used in this study were properly acknowledged and all rules guiding a good scientific practice were maintained to ensure professionalism.

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Author contributions

Falowo Olayomi Abiodun and Olaiya Tomiwa Ayomiposi contributed to the study conception and design. Material preparation, data collection and analysis were performed by Falowo Olayomi Abiodun, Oladipo Babatunde, Oyekola Oluwaseun, Taiwo Abiola Ezekiel and Betiku Eriola. The first draft of the manuscript was written by Falowo Olayomi Abiodun and Oladipo Babatunde, while Oyekola Oluwaseun and Betiku Eriola commented on and proof-read the previous versions of the manuscript. All authors read and approved the final manuscript.

Availability of data and materials

The authors confirm that the data supporting the findings of this study are available within the article.

Declaration of Competing Interest

The authors have no relevant financial or non-financial interest to disclose.

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