Development of Solid Acid Catalyst from Biomass Obtained from Cake of *Vitellaria paradoxa* and Its Application in Biodiesel Production Via a Two-Step Reaction System

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Abstract

An integrated approach is needed to make biodiesel production cost-effective and a suitable choice of fuel. This study was therefore designed to utilize the seed cake of *Vitellaria paradoxa* obtained during oil extraction stage for the synthesis of a solid acid catalyst capable of hydrolysis and/or esterification reaction. The seed cake was pyrolyzed in an inert tube furnace at 420°C for 5 h, and then sulfonated in fuming sulphuric acid (15 wt.%) at 150°C for 10 h to obtain a biomass-derived catalyst. The catalyst was characterized by elemental analysis, FT-IR spectroscopy, X-ray diffraction (XRD) spectroscopy and total acid density. The hydrolytic activity of the catalyst was performed using vegetable oil obtained from *V. paradoxa* biomass and the reaction was optimized using a randomized central composite rotatable design (CCRD). The optimization results showed that catalyst derived from seed cake of *V. paradoxa* was active in hydrolysis reaction and a conversion of 34.3% was obtained and it was also observed that reaction time and weight of catalyst have significant effect on the reaction. When the catalyst was further employed for esterification of free fatty acids in *V. paradoxa* oil, higher conversion (70%) was achieved at a prescribed reaction condition. The approach of utilizing the waste generated during oil extraction from plants could make the process much greener when viewed from the context of resource efficiency and has the potential to kick start the commercialization of biodiesel production especially in developing countries.

Keywords: *Vitellaria paradoxa*, Solid acid catalyst; Hydrolysis; Esterification; Biodiesel; Optimization

Introduction

Biodiesel, mainly derived from oil-bearing plants, has the potential to replace petro-diesel in parts or whole in the energy sectors of both developed and developing countries. In the automobile industries, biodiesel can be used with little or no modification of engines with improved combustion because of its oxygen content and can improve engine performance due to its higher lubricity [1]. In addition, biodiesel can be used in engines in its pure form (B100) or may be blended with petroleum diesel in different ratios [2].

There are many potential feedstocks available for use in biodiesel, which include vegetable oils as well as animal fats. Common raw materials for biodiesel production include soybean oil, sunflower, corn and olive oils, rapeseed oil, castor and lesqureira oils, milkweed seed oil, *Jatropha curcas* oil, *Calophyllum inophyllum* oil [3-9]. Algae and waste vegetable oils are also being considered as suitable feedstock for biodiesel production [10,11].

Transesterification reactions are commonly employed to convert vegetable oils to biodiesel in the presence of either alkali or acid catalysts. Qiu et al. [12] utilized mixed soybean and rapeseed oil with sodium hydroxide as catalyst and 94% yield of biodiesel was reported at the optimized conditions. Dawodu et al. [13] also used *Sesamum indicum* L. as feedstock to produce biodiesel in the presence of sodium methoxide (NaOCH₃) catalyst and a yield of 87.8% was reported at the optimum reaction condition. Alkali-catalyzed reactions are only possible with feedstock having low free fatty acids, thus edible vegetable oils are preferred for biodiesel production since alkali-catalyzed reaction is much faster than acid-catalyzed reaction [14]. However, acid-catalyzed reaction requires high reaction temperature, retention time, molar ratio of vegetable oil to methanol/ethanol and catalyst weight to achieve high conversion of vegetable oils to biodiesel [9,15,16].

Therefore, new methods are required to convert feedstock with high free fatty acids to biodiesel and the methods have to be energy efficient in terms of recovery, recycling and re-use. A two-stage solid acid/alkali-catalyzed reaction is now preferred due to easy catalyst separation and purity of the product [15]. Cavalcanti-Oliveira et al. [17] proposed a hydro esterification route to convert triglycerides in soybean oil to free fatty acids (FFAs) via hydrolysis reaction, which was followed by esterification of FFAs with alcohol to biodiesel in the presence of niobic acid in pellets.

*Vitellaria paradoxa*, commonly known as shea tree is a tree of the Sapotaceae family. It is the only species in genus *Vitellaria* and is indigenous to Africa. *V. paradoxa* fruit consists of a thin, tart, nutritious pulp that surrounds a relatively large, oil-rich seed from which shea butter is extracted and the oil is of interest in biodiesel production. Two-stage process involving hydrolysis and esterification reactions is rarely used to produce biodiesel and even when used, edible vegetable oils were the preferred feedstock. Therefore, this study...
adopts two-step process to catalyze *V. paradoxa* oil to biodiesel. The catalyst was developed through pyrolysis and sulfonation of *V. paradoxa* seed cake to obtain a solid acid catalyst. The catalyst was utilized in the hydrolysis of *V. paradoxa* oil to FFAs and the reaction was optimized using a randomized central composite rotatable design (CCRD). The product from hydrolysis experiment was further converted to FAME (biodiesel) using sulfonated *V. paradoxa* biochar catalyst.

**Experimental**

**Materials**

Dried Shea butter (*Vitellaria paradoxa*) kernels were collected from Fufu village in kwara state, the Southern part of Nigeria. The seeds were manually removed from the kernels. The seeds and the shaft were dried to acceptable moisture content, pulverized to increase the surface area and then stored in airtight bags prior to further use.

**Oil extraction and characterization**

The dried seeds of *V. paradoxa* were extracted by means of soxhlet apparatus using n-hexane (99.0%). After the extraction, the set up was dismantled and the mixture of oil and n-hexane was distilled to obtain the *V. paradoxa* oil and waste product (seed cake) while the recovered solvent was used for subsequent extraction. Fatty acids profile of *V. paradoxa* oil was determined using GC-FID equipped with capillary fused silica column SPTM 2380 (30 m × 0.25 mm × 0.2 mm thickness). Fatty acids identification was done using standards of palmitic (99.0%), stearic (99.0%), oleic (99.0%), linoleic (99.0%) and arachidic (99.0%) fatty acids. Saponification value, iodine value, acid value, free fatty acid, viscosity, ash, moisture content was determined by standard methods described by the Association of Official Analytical Chemist [18].

**Catalyst synthesis**

The pulverized *V. paradoxa* seed cake was sieved to 0.6 m mesh size and oven dried at 105°C for 10 h. Thereafter, the sample was carbonized at 420°C in a tube furnace under nitrogen for 5 h. The process is a form of destructive distillation in which a carbonaceous residue of highly porous structure is obtained.

The obtained carbon residue (10 g) was immersed in 100 ml of fuming sulphuric acid for 10 h at 150°C. The mass ratio of acid to shaft was 18.4:1; the resultant sample was washed with distilled water until a neutral pH was achieved (litmus paper test). The sulfonated *V. paradoxa* seed cake was then dried at 110°C for 24 h. The scheme 1 is shown below.

![Scheme 1: Sulfonated V. paradoxa seed cake.](Image 45x151 to 283x199)

**Scheme 1: Sulfonated V. paradoxa seed cake.**

**Catalyst characterization**

The carbon, hydrogen, nitrogen, sulfur and oxygen contents of the sulfonated *V. paradoxa* catalyst were determined by Elemental Analyzer (Vario cube). The Fourier transform infrared (FT-IR) spectroscopy was obtained using Perkin Elmer FTIR instrument. Spectra were taken in the range of 350 to 4000 cm using KBr pellet method (solid sample) or Nujol method (liquid sample). X-ray diffractometer (XRD) pattern was obtained using Rigaku D/Max-III C X-ray diffractometer operated at 40 kV and 20 mA using Cu Ka radiation source. The sample was scanned in the range of 20 of 5-80°C at a scanning speed of 2°/min. The surface morphology of the catalyst was done using JOEL JSM-6400 microscope. The total acid density of the catalyst was determined using the standard acid-base back titration method. The catalyst samples were pre-dried in the oven at 110°C for 2 h, and then ~ 0.10 g of catalyst was added into 60 ml of 0.0080 mol/L NaOH and stirred for 30 minutes before back-titrination with 0.02 mol/L HCl. For the sulfonic group density determination, it is assumed that all the sulfur content of the catalyst is in the form of -SO$_3$H and thus sulfonic group density was calculated based on the weight percentage of S content obtained from elemental analysis.

**Catalyst activity: Hydrolysis and esterification of *Vitellaria paradoxa* oil**

Hydrolysis reaction was carried out in a quick fit round bottom flask of 250 ml equipped with a hot plate and magnetic stirrer. 5 g of *V. paradoxa* oil, 50 g of distilled water, and catalyst of 0.1 g were charged into the reactor for 5 h at 140°C. At the end of the reaction, the reactor was cooled, and catalyst was separated by filtration. The reaction product consisted of two layers, an upper oily layer of fatty acids, catalyst and unreacted triglycerides and a lower, aqueous layer of water and glycerol which was separated by funnel.

Esterification of fatty acids with alcohols in the presence of sulfonated *V. paradoxa* catalyst was carried out. 4 g of hydrolyzed oil, 40 g of methanol and 0.2 g of catalyst were charged inside a 250 ml reactor which was then placed in an oil-bath at 65°C, magnetic stirring (600 rpm) for 5 h under a reflux condition. The reaction was terminated by placing the reactor in a cool water bath, excess methanol recovered, and the catalyst was phase separated by decantation. The reaction schemes 2 and 3 are shown below.

![Scheme 3: Esterification of Fatty acids.](Image 312x219 to 551x275)

**Scheme 2: Hydrolysis of Triglycerides.**

![Scheme 3: Esterification of Fatty acids.](Image 312x219 to 551x275)

**Scheme 3: Esterification of Fatty acids.**

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The percentage conversions for both hydrolysis and esterification reactions were calculated by titrimetry method. Titration was carried out using 0.25 N NaOH solution and phenolphthalein as indicator. The percentage conversion of triglyceride was calculated using Equation 1.

\[
\text{FFA conversion}=a_i-a_i\times 100 \quad \text{eqn}(1)
\]

Where \(a_i\) is the initial acid value of the \(V.\) \textit{paradoxa} oil and at is the final acid value at the end of the reaction.

### Optimization study

Central Composite Rotatable Design technique (CCRD) was used for optimization of the hydrolysis reaction. The experimental design employed was a full 32 factorial design (2 factors each at 3 levels) with 4 centre points making a total of 12 runs (Table 1). Experiments were randomly carried out in order to minimize errors from systematic trends in the variables. Optimization conditions chosen were: molar ratio of oil/water: 1:10; catalyst concentration 1-3 wt.% based on the weight of oil; reaction temperature of 150°C. The reaction was timed between 4-6 h soon as the catalyst was added. After a specific time, interval, the reaction was quenched in a cold-water bath. The percentage conversion of triglyceride was determined by titration and percentage converted was calculated using Equation 1.

The Design Expert 6.0 software was therefore used for regression, analysis of variance (ANOVA) and graphical analysis of the data obtained. A quadratic polynomial was developed to predict the response as a function of independent variables and their interactions and a second-order polynomial equation was used [19,20].

| Variables      | Levels               |
|----------------|----------------------|
| Wt% catalyst (A) | 0.59 1 2 3 3.41      |
| Time (B)       | 3.59 4 5 6 6.41      |

Table 1: Coded levels and real values of tested variables in the CCRD.

### Results and Discussion

#### Properties of \(V.\) \textit{paradoxa} oil

Vegetable oils generally used for biodiesel should have low moisture content since alkali catalyzed transesterification reactions are susceptible to water which usually lead to soap formation [21]. The properties of \(V.\) \textit{paradoxa} oil are shown in Table 2. The oil yield is considerably high thus making the oil feedstock economically viable for biodiesel production. Also, low contents of ash and moisture made the oil a suitable feedstock for biodiesel production. In addition, the fatty acids profile of \(V.\) \textit{paradoxa} oil are palmitic (6.5%), stearic (28.7%), oleic (55.5%), linoleic (6.2%) and arachidic (0.7%) making the oil a suitable feedstock for soap formation. An alternative conversion route is then required to convert the oil to biodiesel. The oil has a relative density of 0.9 g/cm³ which showed that it is less dense than water and a kinematic viscosity of 9.2 centipoises which showed that is not as thick as most of drying oils.

| Properties         | Values     |
|--------------------|------------|
| Moisture (%)       | 1.2 ± 0.4  |
| Ash (%)            | 0.2        |
| Colour             | Light yellow |
| State of the oil at room temperature | Solid |
| Saponification value (mg KOH/g) | 160.2 ± 0.9 |
| Relative density (g/cm³) | 0.9 |
| Iodine value (I₂/100 g) | 42 |
| Dynamic viscosity (cP) | 2.4 |
| Free fatty acid (%) | 4.7 ± 0.1 |
| Acid value (mg KOH/g) | 9.4 |

Table 2: Proximate and physico-chemical properties of \(V.\) \textit{paradoxa} oil.

#### Catalyst characterization

X-ray diffraction (XRD) pattern of sulfonated \(V.\) \textit{paradoxa} catalyst (Figure S1) exhibited diffraction peaks around the 2θ=15-30° region which is due to the presence of an amorphous carbon structure with randomly oriented aromatic carbon sheets [23-25]. The XRD pattern obtained was similar to that obtained from seed cake of \(Calophyllum\) \textit{inophyllum} [26].

The functional groups present on \(V.\) \textit{paradoxa} biochar and sulfonated \(V.\) \textit{paradoxa} catalyst contained aromatic and aliphatic hydrocarbons, as well as some functional oxygen-containing groups, including -OH and C=O [27]. The peaks at around 3440 cm are more intense in \(V.\) \textit{paradoxa} biochar than in sulfonated \(V.\) \textit{paradoxa} catalyst (Figure S2). Therefore, \(V.\) \textit{paradoxa} biochar contains larger phenolic OH groups. Both \(V.\) \textit{paradoxa} biochar and sulfonated \(V.\) \textit{paradoxa} catalyst spectra contain peaks attributable to aromatic ring modes, at 1463 and 1467 cm respectively [28]. After sulfonation of the carbonized materials prepared at 400°C, the vibration bands observed at 1079 cm (SO₃ stretch) and 1216 cm (S=O asymmetric vibration in SO₃H) confirmed that the catalyst possess SO₃H groups [29]. The sulfonated \(V.\) \textit{paradoxa} catalyst also exhibited peaks at 1702 cm and 1608 cm which can be attributed to C=O of a carboxylic acid and the stretching vibration of a polycarboxylic aromatic compound containing carboxylic groups [15]. However, the peaks attributed to carboxylic groups at (1700 cm) and polycarboxylic aromatic compounds (1608 cm) were absent in the spectra of \(V.\) \textit{paradoxa} biochar. These peaks could have been overlapped by the peak at 1631 cm. The band at 2300-2700 cm was as a result of an overtone due to –OH =O(bend) and this is linked by a strong hydrogen bond, which suggests that some SO₃H groups are within a short distance to each other [9,29].

The morphological analysis of the sulfonated \(V.\) \textit{paradoxa} biochar catalyst by SEM revealed that the material has a loose irregular network structure (Figure S3). The main determinant of the structural...
properties of a carbon-based material is the nature of the starting material. A char obtained by slow pyrolysis retains the fibrous structure while char produced by rapid pyrolysis has a porous structure. Removal of cell contents and consequently, the opening-up of cellular structures give rise to a porous network of distinctive features, which depends on the heating rate and the raw material [30].

Elemental analysis on *V. paradoxa* seed cake, *V. paradoxa* biochar and sulfonated *V. paradoxa* catalyst are presented in Table 3. It can be seen from Table 3 that carbonization of biomass results in an increase in the H/C ratio of the material. This is due to the removal of oxygen from the biomass in the form of carbon dioxide (and carbon monoxide) during the carbonization process and successful incorporation of SO$_3$H group was also confirmed. In addition, sulfonated *V. paradoxa* catalyst SO$_3$H density as determined from the EA was 3.35 mmol/g and this was absent from *V. paradoxa* seed cake and *V. paradoxa* biochar (Table 4). However, the total acid density for the catalyst was higher than biomass-derived solid acid catalysts obtained in our previous studies [15, 26].

| Material                  | Elemental Analysis Results (%) | Composition                   | H/C   |
|---------------------------|--------------------------------|-------------------------------|-------|
|                           | C     | H    | O     | N     | S     | Total b | Acid Density (mmol/g) | Conversion (wt.%) c | Acid value (mg KOH/g) |
| *V. paradoxa* seed cake   | 52.1  | 0.72 | 41.68 | 5.5   | -     | 0.28     | 0.84              | -                  | 12.84               | 10.67               |
| *V. paradoxa* Biochar     | 61.8  | 2.56 | 30.14 | 5.5   | -     | 0.50     | 0.37              | 0.50               | 21.26               | 11.46               |
| Sulfonated *V. paradoxa*  | 62.08 | 4.12 | 17.58 | 5.5   | 10.72 | 0.80     | 0.21              | 0.21               | 34.32               | 12.7                |

*Calculated based on the sulphur content from elemental analysis. * Obtained by titration. *Reaction Conditions: Oil/water: 1:10, Catalyst: 2 wt.% of oil, temperature: 150°C, Stirring speed: 600 rpm, Time: 5 hrs.*

Table 3: Elemental composition of catalyst materials.

Catalytic activity of sulfonated *Vitellaria paradoxa* catalyst

**Hydrolysis of Vitellaria paradoxa oil:** The activity of the catalyst was tested under pre-determined reaction conditions. The yield obtained was compared to those obtained without the catalyst and the *V. paradoxa* biochar. The results of hydrolysis of *V. paradoxa* oil with or without catalyst are presented in Table 4. Blank experiment in the absence of catalyst, at the reaction condition (Temperature: 150°C; Oil/ molar: 1:10; Catalyst: 2 wt.% of oil; Stirring speed: 600 rpm; Time: 5 h), showed that the hydrolysis reaction takes place, to a certain extent. This may be due to the presence of free fatty acids in the oil which acts as weak acid catalyst for the reaction [31]. The result also showed that a higher conversion was obtained with *V. paradoxa* biochar. The catalytic activity of the non-sulfonated carbon material is due to presence of -COOH and phenolic -OH groups which confer acidic properties on the material. Table 4 also shows the total acid density on the carbon material to be 0.28 mmol/g, which is a combination of the -COOH and phenolic -OH group acidic group. The presence of FFAs in the oil acting as weak acid catalyst also complement the activity of the acidic group in the non-sulphonated carbon material in driving the hydrolytic reaction forward. Under the same reaction conditions, sulfonated *V. paradoxa* catalyst gave 34.32% conversion of triglycerides to FFAs. Despite high acid density of the catalyst, which should have translated to a higher catalytic activity of the catalyst, the obtained conversion was relatively low. Comparing the conversion rate with similar work, Satyararthi et al. [32] were able to achieve a triglyceride conversion of up to 93.9% employing a solid Fe-Zn double-metal cyanide (DMC) complex catalyst (acid density of 0.84 mmol/g) in a Teflon-lined, stainless steel autoclave at a temperature of 190°C and autogeneous pressure. Since the rate of the forward reaction of an endothermic reaction is favoured by an increase in temperature, the reaction yield is significantly high. The pressure generated within the system is also of greater magnitude compared to that obtained under the reflux condition under which this study was done.

**Optimization study**

An optimization study of the hydrolysis reaction was carried out in order to study the effect of temperature and amount of catalyst on the reaction yield, while keeping other factors affecting the reaction constant. This optimization was carried out using a Randomized Central Composite Rotatable Design (CCRD). The design matrix and the results obtained are shown in Table 5. The best conversion of FFAs was obtained in Run 1, with a catalyst ratio of 3% and a reaction time of 6 h. From the model, variables with P-values lower than 0.05 were considered statistically significant (Table 6). Based on the statistical analysis of the results, an empirical model was constructed to describe the variation in the free fatty acid concentration as a function of catalyst ratio (A) and reaction time (B) as shown in equation (2). Only terms A and B were significant. A Fisher F-test with a P-value lower the 0.05 indicates a high significance for the model.

| Material                  | Elemental Analysis Results (%) | Composition                   | H/C   |
|---------------------------|--------------------------------|-------------------------------|-------|
|                           | C     | H    | O     | N     | S     | Total b | Acid Density (mmol/g) | Conversion (wt.%) c | Acid value (mg KOH/g) |
| *V. paradoxa* seed cake   | 52.1  | 0.72 | 41.68 | 5.5   | -     | 0.28     | 0.84              | -                  | 12.84               | 10.67               |
| *V. paradoxa* Biochar     | 61.8  | 2.56 | 30.14 | 5.5   | -     | 0.50     | 0.37              | 0.50               | 21.26               | 11.46               |
| Sulfonated *V. paradoxa*  | 62.08 | 4.12 | 17.58 | 5.5   | 10.72 | 0.80     | 0.21              | 0.21               | 34.32               | 12.7                |

*Calculated based on the sulphur content from elemental analysis. * Obtained by titration. *Reaction Conditions: Oil/water: 1:10, Catalyst: 2 wt.% of oil, temperature: 150°C, Stirring speed: 600 rpm, Time: 5 hrs.*

Table 4: Properties and activities of derived catalyst.
%Conversion = 33.80 + 5.65A + 4.34B + 1.00AB + 1.79A^2 - 0.83B^2 \ldots \text{eqn}(2)

| Standard order | Run Order | % wt of catalyst | Time (Hrs) | % Conversion |
|----------------|-----------|------------------|------------|--------------|
| 4              | 1         | 3                | 6          | 45.4         |
| 3              | 2         | 1                | 6          | 30.9         |
| 5              | 3         | 2                | 5          | 34.5         |
| 1              | 4         | 1                | 4          | 28           |
| 2              | 5         | 3                | 4          | 38.5         |
| 9              | 6         | 2                | 6.41       | 40           |
| 10             | 7         | 2                | 5          | 36.4         |
| 7              | 8         | 3.41             | 5          | 43.6         |
| 8              | 9         | 2                | 3.59       | 22.4         |
| 6              | 10        | 0.59             | 5          | 29.3         |
| 11             | 11        | 2                | 5          | 33.8         |
| 12             | 12        | 2                | 5          | 30.5         |

Table 5: Design matrix and table of result for optimization experiments.

| Source | Sum of Squares | Df | Mean Square | F value | P-value |
|--------|----------------|----|-------------|---------|---------|
| Model  | 440.1          | 5  | 88.02       | 9.35    | 0.0085  |
| A      | 255.64         | 1  | 255.64      | 27.15   | 0.002   |
| B      | 150.43         | 1  | 150.43      | 15.98   | 0.0071  |
| AB     | 4              | 1  | 4           | 0.42    | 0.5387  |
| A2     | 20.59          | 1  | 20.59       | 2.19    | 0.1896  |
| B2     | 4.42           | 1  | 4.42        | 0.47    | 0.5187  |
| Residual | 56.49        | 6  | 9.41        |         |         |
| Lack of Fit | 38.35    | 6  | 12.78     | 2.11    | 0.2772  |
| Pure Error | 18.14      | 6  | 3.05       |         |         |
| Cor Total | 496.59        | 11 |             |         |         |

Table 6: ANOVA for response surface quadratic model.

The model predicts that percent yield of 33.8% could be obtained under the optimum operating conditions. To test the fit of the model, residual distribution graph was plotted, and it is observed that the residual distribution does not follow a particular trend with respect to the predicted values (Figure 1A). This test the assumption of constant variance and should be random scatter (constant range of residuals across the graph). This shows the relationships between the experimental response values and predicted response values. This graph helps to detect a value, or group of values, that are not easily predicted by the model. To test the fit of the model, the data points should be split evenly by the 45-degree line. The graph agrees with this assumption and the fit of the model was further confirmed by normal plot of residuals (Figure 1B).

Figure 1: Residual distribution plot of the optimized hydrolysis reactions, i.e., (A) Residuals vs Predicted Plot (B) Predicted vs Actual Plot.

The normal probability plot indicates whether the residuals follow a normal distribution, in which case the points will follow a straight line (Figure 2). This indicates that the errors are normally distributed for all the responses. It can thus be concluded that the proposed empirical model adequately correlates the relationship between the reaction variables and the conversion of triglycerides to FFAs.

Figure 2: Normal probability plot of the optimized hydrolysis reaction.

Contour plots and three-dimensional plots for the interaction between the variables and percentage conversion of triglycerides are shown in Figures 3 and 4.

Figure 3: Contour plot of weight of catalyst vs reaction time.
Theoretical value at 30%.

Reaction time is also positive, though it is slightly not significant.

Comparison of the acid value of the oil before and after esterification of FFAs to biodiesel was carried out setting a target of 2 wt.% catalyst and a reaction time of 4 h. A conversion of 27% was obtained, with the theoretical value at 30%.

Esterification of FFAs to biodiesel

The hydrolyzed V. paradoxa oil was subjected to esterification reaction with methanol in the presence of sulfonated V. paradoxa catalyst. The conversion of FFAs to biodiesel was calculated by determining the triglyceride conversion.

The biodiesel produced in the present work. Aliphatic esters produce two most polar bonds in esters which are C=O and C-O bands at 1750-730 cm⁻¹ and 1300-100 cm⁻¹, respectively.

| Wave number (cm⁻¹) | Present work | Group Assignment | Vibration type |
|--------------------|--------------|------------------|---------------|
| 720                | 722.08       | 0                | Plane rocking |
| 1112070            | 1097.96      | -C=O             | Stretching    |
| 1300100            | 1117.76      | -C=O             | Stretching    |

Table 7: Characteristic infrared bands of biodiesel.

Conclusion

A promising heterogeneous catalyst for hydrolysis of triglycerides has been prepared through the sulfonation of Vitellaria paradoxa biochar with fuming sulphuric acid. Structural study through SEM, elemental analysis and FT-IR spectroscopy suggests that biochar-based catalyst consists of polycyclic aromatic carbon sheets bearing three different acidic groups of phenolic, carboxylic, and sulphonic. The catalytic activity of the developed biochar-based catalyst was evaluated in a two-step reaction involving hydrolysis of V. paradoxa oil and subsequent esterification of FFAs to FAME (biodiesel). The hydrolysis reaction showed that the catalyst increased the conversion of V. paradoxa oil to FFAs when compared to the reaction carried out without the catalyst. A percent conversion of 34.32% was obtained with the catalyst compared to the 12.84% obtained without the catalyst. Optimization study of the hydrolysis reaction using a randomized central composite rotatable design showed that a percent yield of 33.80% was obtainable at optimum reaction condition of the study. Evaluation of the esterification activity of the catalyst showed a FFAs conversion of 70%.

Conflict of Interest

The authors declare no conflict of interest.

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