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Article

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Abstract: Biodiesel, or alkyl ester, is an alternative renewable, biodegradable, and non-toxic diesel fuel produced by the catalytic transesterification of vegetable oil. Here we characterize a system for continuous transesterification of vegetable oil using five continuous stirring tank reactors (5CSTRs). We tested residence times of 16–43min, stirring speeds of 200–800rpm, a catalyst concentration (KOH) of 0.25–1 wt% of oil (in gram), different total flow rates of the oil and MeOH, and on the production performance of the 5 stage continuous reactor for transesterification of vegetable oil. Using a molar ratio of oil:methanol of 1:7 and a reaction temperature of 65 °C, we show that a high stirring speed increased the reaction rate, but an excessive stir speed decreased the reaction rate and conversion to biodiesel. Furthermore, a higher catalyst percentage significantly increased the reaction rate and production capacity. A catalyst percentage of 1 wt% of oil gave the best conversion; $99.04 \pm 0.05\%$. The resulting biodiesel esters were characterized for their physical and fuel properties including density, viscosity, iodine volume, acid volume, cloud point, pure point, gross heat of combustion, and volatility. The purity and conversion of the biodiesel was analyzed by HPLC.

Keywords: coconut oil; alkaline-catalyzed transesterification; CSTRs; high performance liquid chromatography

1. Introduction

Monoesters produced by the transesterification of vegetable oil with alcohol are known as biodiesel fuels [1]. Biodiesel is a biodegradable and renewable fuel. It contributes no net carbon dioxide or sulfur to the atmosphere and emits less gaseous pollutants than normal diesel. Carbon dioxide, aromatics, polycyclic aromatic hydrocarbons (PAHS) and partially burned or unburned hydrocarbons emissions are all reduced in vehicles operating on biodiesel [2]. Biodiesel is an environmentally friendly liquid fuel with similar combustion properties to petrol diesel. Increasing environmental concern, diminishing petroleum reserves, and the agriculture-based economy of our country are the driving forces to promote biodiesel as an alternative fuel [3]. Other advantages of biodiesel are its good lubrication properties that help to extend engine life, high cetane number, high flash point, and acceptable cold filter plugging point (CFPP), which make it very attractive as an alternative fuel [4,5]. Biodiesel derived from vegetable oil and animal fats is used in the USA and Europe to reduce air pollution and to reduce the dependence on fossil fuels. In the USA and Europe, the surplus edible oils like soybean oil, sunflower oil, and rapeseed oil are being used as feedstock for the production of biodiesel [6,7]. However, vegetable oil is about ten times more viscous than diesel. As a result, vegetable oils can cause poor fuel atomization, incomplete combustion, and carbon deposition on the injector and valve seats; all resulting in serious engine fouling. Common methods employed to reduce the viscosity of vegetable oils include blending with diesel, emulsification, pyrolysis, cracking, and transesterification. Among these, the transesterification of vegetable oils to alkyl esters appears to be the best method [8]. Use of a solid base catalyst offers several process advantages, including the elimination of a quenching step to isolate the products - and the associated contaminated water waste - as well as the opportunity to operate a continuous process [9]. Such solid bases, including zeolites [10], alkali earth oxides [11], and hydrotalcites [12] have been utilized in transesterification reactions, and recently tuneable solid bases, such as Li-doped CaO and Mg-hydrotalcites, for tributyrin transesterification [13].

Biodiesel is usually prepared in the presence of a homogenous base or acid catalyst. The acid catalyzed process often uses sulfonic acid and hydrochloric acid as the catalyst; however, the reaction time is very long (48–96 hr) even at the temperature of methanol reflux, and a high molar ratio of methanol to oil is needed (30–150:1, by mol) [14]. Potassium hydroxide, sodium hydroxide, and their carbonates, as well as potassium and sodium alkoxides such as NaOCH_3 , are the base catalysts usually used for the reaction [15,16]. Most biodiesel fuels are currently synthesized using alkaline catalysts because the transesterification reaction is much faster than with acid catalysts [17,18]. Moreover, acid catalysts are more corrosive, and thus base catalysis is preferred commercially.

There have been several comprehensive studies on base catalyzed transesterification to produce fatty acid methyl esters (FAMEs) [19]. A higher conversion of vegetable oil has been reported when strong base catalysts such as sodium hydroxide and potassium hydroxide are used. Furthermore, basic catalysts (for example KOH) can convert 97–99 wt% in 0.5–1 hr [19]. However, the base catalyzed process still has two limitations: the base catalyzed transesterification of vegetable oil is sensitive to both moisture and to free fatty acids (FFA). This means that the raw materials must be refined; the moisture level must be no more than 0.06 wt% and FFA levels no more than 0.5 m wt% [20]. A small amount of moisture can initiate oil hydrolyzation to form soap and glycerol. Vegetable oils,

particularly cooking oil, also contain significant amounts of FFA, which incur significant processing problems in standard biodiesel manufacturing. FFA are saponified by homogenous alkaline catalysts. These soaps can bring about the emulsification between glycerol and FAME, which will make the separation even more difficult when the water washing method is used. Furthermore, the saponification leads to a loss of catalyst and an increased purification cost [21]. If the FFA content in vegetable oil is more than 0.5 wt%, the simplest approach to improve the processing is to first esterify these FFA in the vegetable oil to their alkyl esters using an acid catalyst. Such a pretreatment process adds a neutralization step as well as a separation step to the overall process. Thus, development of solid acid catalysts for esterification pretreatments is crucial to improve the efficiency of biodiesel production [22].

The present work studies the experimental conditions such as catalyst concentration, molar ratio (oil:methanol), reaction temperature, total flow rate, mixing intensity, and the residence time on biodiesel production from coconut oil, and the produced biodiesel, is analyzed by different tools.

2. Results and Discussion

2.1. Catalyst type and concentration

Catalysts used for the transesterification of vegetable oils are classified as alkali, acid, enzyme or heterogeneous catalysts; of which alkali catalysts like sodium hydroxide, potassium hydroxide, and potassium methoxide are more effective. If the oil has a high FFA content and a high concentration of water, acid catalyzed transesterification is preferred. For an alkaline catalysis, sodium hydroxide or potassium hydroxide is mostly used.

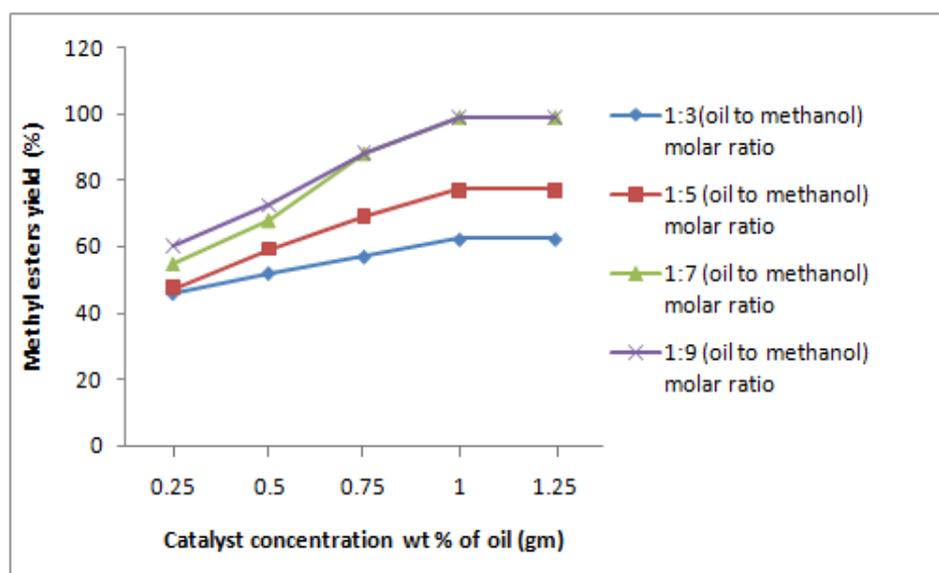
In the present work, the continuous transesterification of coconut oil is studied. For this, alkaline catalyst is more suitable due to the presence of very low FFA. For the methanolysis of coconut oil, potassium hydroxide gives the best results and viscosity of the esters. When optimizing catalyst concentration, a stirring speed of 800rpm with a mechanical stirrer, a flow rate of 142 mL/min, a temperature of 65 °C, and a molar ratio of 1:7 oil:methanol was held constant. Figure 1 summarizes the experimental effect of varying the catalyst (KOH) concentration from 0.25 to 1.25 wt% of oil. When the catalyst was 0.25 wt%, the conversion was 75%. Increasing the catalyst concentration increased the conversion rate: a 1.00 wt% of catalyst gave the best conversion rate of $99.04 \pm 0.05\%$. There are however drawbacks to the excessive use of alkaline catalysts in the transesterification reaction. A higher catalyst concentration increases the solubility of methyl esters in the glycerol phase of the final product. As a result, a significant amount of methyl esters remain in the glycerol phase after the phase separation.

2.2. Molar ratio

One of the most important variables affecting the yield of ester is the molar ratio of alcohol to oil. The stoichiometric ratio for transesterification requires three moles of alcohol and one mole of triglyceride to yield the three moles of fatty acid methyl esters and one mole of glycerol. However, transesterification is an equilibrium reaction in which a high molar ratio is used to drive the reaction to

the right, enhancing the solubility and increasing the contact between the triglycerides and alcohol molecules. Experiments were conducted with molar ratios of 1:3, 1:5, 1:7 and 1:9 using a constant mixing speed of 800rpm and catalyst concentration of 1.00 wt% of oil. The results are shown in figure 1. At the molar ratio 1:7 (oil:methanol), the conversion of triglycerides was maximal $\geq 99\%$. In this experiment, methanol is fed in two parts: 5 moles in the first reactor and 2 moles in the third. Molar ratio has no effect on acid, peroxide, saponification, or iodine values of methyl esters [23]. However, the high molar ratio of alcohol to vegetable oil interferes with the separation of glycerol because there is an increase in solubility. When glycerol remains in solution, it helps derive the equilibrium back to the left, lowering the yield of esters. The production capacity of the unit was 6.4 L/hr.

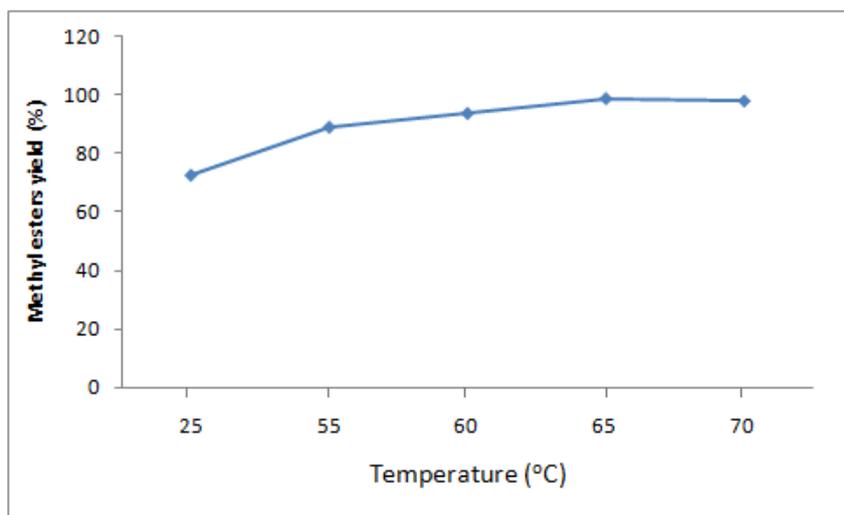
Figure 1. Effect of catalyst concentration (wt% of oil) and molar ratio (oil:methanol) on the yield of methyl esters from coconut oil (at constant temperature of 65 °C, flow rate of 142 mL/min, stirring speed of 800rpm, residence time 21.04 min).



2.3. Reaction temperature

To test the effect of reaction temperature, experimental trials were carried out at room temperature (25 °C), and at 55 °C, 60 °C, 65 °C and 70 °C. In all experiments, a constant methanol to oil ratio of 7:1 and a 1 wt% KOH catalyst concentration was used, based on the optimal conditions in the previous section). As shown in figure 2, the conversion of coconut oil into fatty acid methyl esters increased with reaction temperature. The yield of methyl esters at 25 °C, 55 °C, and 60 °C was 60, 85, and 94%, respectively. The maximum yield of $\geq 99\%$ was obtained at the temperature 65 °C. Methanolysis of vegetable oil is normally performed near the boiling point of the alcohol. A reaction temperature above the boiling point of alcohol, 64 °C, was ignored, because high temperatures lead to the acceleration of the saponification of glycerides in the presence of a base catalyst before completion of the methanolysis.

Figure 2. Effect of reaction temperature on yield of methyl esters from coconut oil. (constant catalyst concentration of 1 wt% of oil, molar ratio methanol:oil of 7:1, flow rate 142 mL/min, stirring speed 800rpm, residence time 21.04 min).



2.4. Flow rate

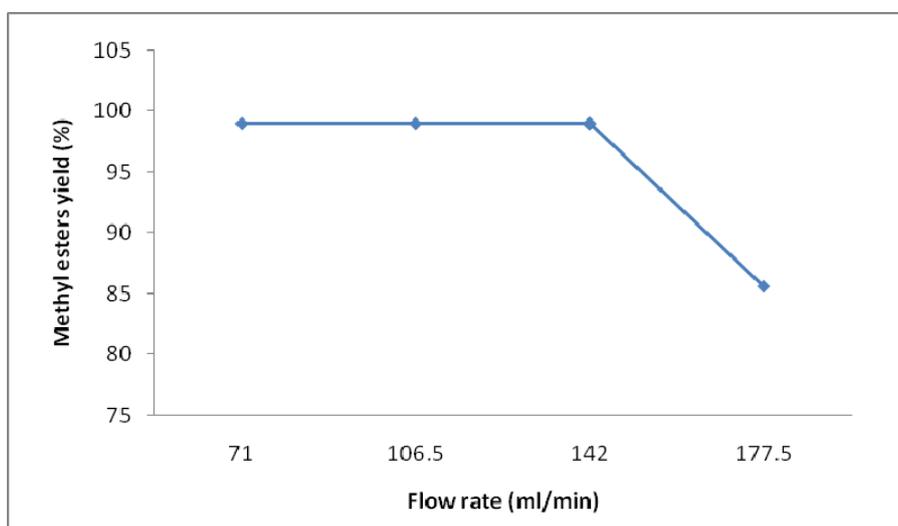
To optimize the flow rate for the process of continuous transesterification of coconut oil, total flow rates ranging from 70 mL/min to 180 mL/min were tested. All experiments were conducted with a molar ratio of methanol:oil 7:1, a 1 wt% KOH catalyst concentration, and a reaction temperature of 65 °C. The effect of flow rate on the biodiesel yield is presented in Figure 3 and the total flow rate of oil and methanol is given in Table 1. A combined total flow rate of oil and methanol was 142 mL/min (100 mL oil and 42 mL methanol) gave the best conversion of ≥ 99 .

Table 1. Effect of flow rate and residence time on yield of methyl esters from coconut oil. (Molar ratio oil:methanol 1:7, catalyst concentration 1 wt% of oil, mixing intensity 800 rpm, reaction temperature 65 °C).

Flow rate (ml/min)			Residence time (min)					Conversion %			
Oil (ml)	MeOH (ml)	Total (ml)	Reactor R1	Reactor R2	Reactor R3	Reactor R4	Reactor R5	Total	S1	S2	S3
50	21.0	71.0	8.07	8.07	8.4	8.4	9.85	42.79	82.5	97.0	99
75	31.57	106.5	5.1	5.1	5.6	5.6	6.57	27.97	82.5	96.8	99
100	42.0	142.0	3.84	3.84	4.22	4.22	4.92	21.04	82.6	96.8	99
125	52.5	177.5	3.07	3.07	3.38	3.38	3.94	16.84	75.2	80.0	85.6

S1 = Separating funnel first, S2 = Separating funnel second, S3 = Separating funnel third.

Figure 3. Effect of flow rate on yield of methyl esters from coconut oil (Catalyst concentration 1 wt% of oil, molar ratio methanol:oil 7:1, temperature 65 °C, stirring speed 800 rpm, residence time 21.04 min).



2.5. Mixing intensity

Mixing is a very important factor in the transesterification reaction. Without mixing, the reaction occurs only at the interface between the two layers and is considered too slow to be feasible. In the present work, the effect of mixing intensity at various catalyst concentrations was examined. Methanolysis was conducted with stirring rates of 200, 400, 600, 800 and 1,000 revolutions per minute (rpm) and catalyst concentrations of 0.25%, 0.50%, 0.75%, 1% and 1.25%. Based on the observations above that 1wt% of catalyst give the best conversion, the yield of methyl ester at different mixing intensities (rpm) using the catalyst concentration of 1wt% is shown in Figure 4. The methanolysis reaction was partially incomplete at 200 rpm, only exhibiting a conversion of 85.5%. The conversion increased as mixing intensity was accelerated. The best yield of methyl esters was achieved at 800 rpm and 1% catalyst, up to 99%.

2.6. Residence time

The residence time of the reaction is the effective volume of the reactors divided by the total flow rate. The effective volume of the five CSTRs is 500 mL, 500 mL, 600 mL, 600 mL, and 700 mL, respectively. The desired total flow rate of oil and alkaline methanol were 71 mL/min, 106.5 mL/min, 142 mL/min and 177.5 mL/min for residence times of 42.79min, 27.97 min, 21.04 min and 16.84 min, respectively. The calculated molar ratio of methanol to oil in all experiments was 7:1, the flow rate was calculated based on the available density of coconut oil at 20 °C (0.92 gm/mL) and the density of methanol at 20 °C (0.792 gm/mL). The yield of biodiesel at different residence times is shown in Figure 5. A flow rate of 142 ± 0.5 mL and a residence time of 21.04 gave the best conversion of $\geq 99\%$. When the total flow rate of oil and methanol was 142 mL/min, the residence time of each reactor was 3.84 min, 3.84 min, 4.22 min, 4.22 min and 4.92 min, respectively.

Figure 4. Effect of stirring speed on yield of methyl esters from coconut oil (catalyst concentration 1wt% of oil, molar ratio methanol:oil 7:1, temperature 65°C, flow rate 142 mL/min, residence time 21.04 min).

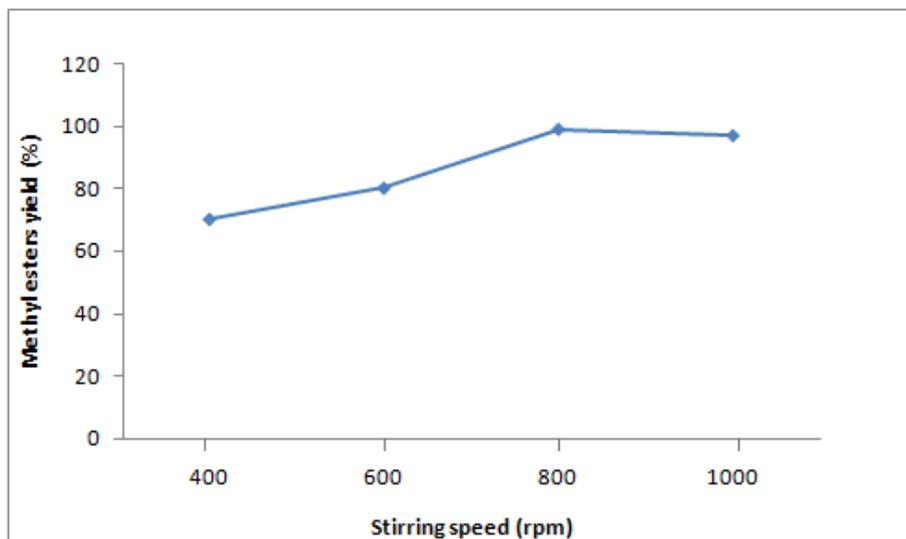
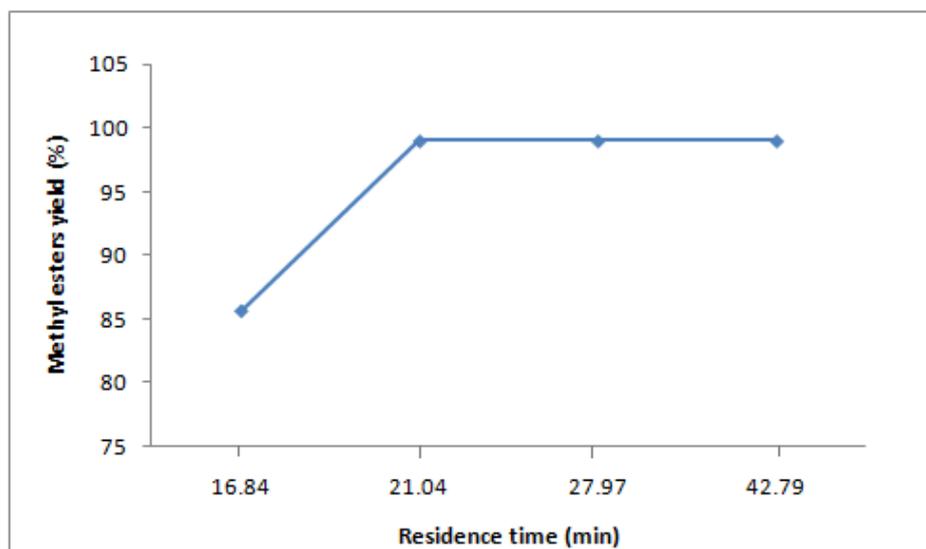


Figure 5. Effect of residence time on yield of methyl esters to coconut oil. (catalyst concentration 1 wt% of oil, molar ratio methanol:oil 7:1, temperature 65 °C, flow rate 142 mL/min, stirring speed 800 rpm).



2.7. HPLC analysis

Reverse phase high performance liquid chromatography (RP-HPLC) was used to analyze the purity, conversion and FAME composition of the biodiesel esters produced. RP-HPLC separates different components according to their polarity. The chromatographic apparatus was from Waters, Milford MA, USA. It consisted of a model waters 600 pump with waters 600 controller, waters 2,996 photodiode array detector, a nova-pack®, 3.9 × 150 mm column with guard column of dimension 3.9 × 20 mm, both packed with C18 particle with diameter 4 μm. The RP-HPLC conditions used for the separation

and determination of the compounds produced during the methanolysis of coconut oil in all the experiments were: a flow rate of 1 mL/min, an injection volume of 5 μ L, a column temperature of 45 °C, UV detection at 215 nm and a 40 min gradient mobile phase 15% H₂O + 85% CH₃OH in 10 min, 100% CH₃OH in 0 min, 60% CH₃OH + 15% hexane + 25% propane-2-ol in 30 min and for the last 10 min system back to initial state 15% H₂O + 85% CH₃OH. Figure 6 shows the HPLC chromatogram of coconut oil. The FAME present in coconut oil-based biodiesel are presented in Table 2.

Figure 6. HPLC Chromatogram of coconut oil.

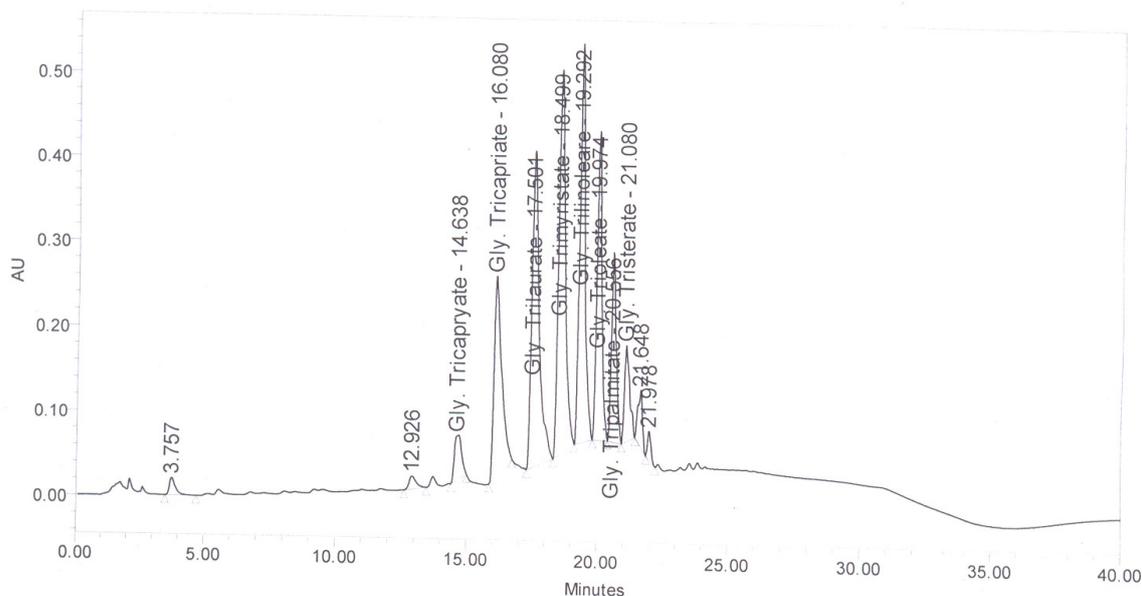


Table 2. Fatty acid methyl esters (FAME) composition of coconut oil-based biodiesel.

Fatty acid	Percentage
Caprylic (C8:0)	8.86
Capric (C10:0)	6.17
Lauric (C12:0)	48.83
Myristic (C14:0)	19.97
Palmitic (C16:0)	7.84
Stearic (C18:0)	3.06
Oleic (C18:1)	4.44
Linoleic (C18:2)	0.76

2.8. Characterization

Kinematic Viscosity, Acid number and oxidation stability

All the physicochemical properties of the coconut-based biodiesel are given in Table 3. The kinematic viscosity of biodiesel at 40 °C was determined following ASTM D 445 [24] using a Rheotek AKV 800 automated kinematic viscometer (Poulten Selfe and Lee Ltd., Essex, England). The acid number of biodiesel was determined according to ASTM D 664 [25] using a Brinkman/Metrohn 809 Titrand

(Westbury, NY). The oxidation stability of biodiesel was determined as the induction period (IP) according to EN 14112 [26] using a Metrohn 743 Rancimat instrument (Herisau, Switzerland).

Table 3. Physical properties of coconut biodiesel.

Property	ASTM method	ASTM Specification	Oil	Biodiesel
Viscosity 40 °C (mm ² /s)	D 445	1.9–6.0	27.23	2.83
Acid number (mgKOH/g)	D 664	0.5	2.1	0.18
Free glycerin (mass %)	D 6584	0.020	----	0.008
Total glycerin (mass %)	D 6584	0.24	----	0.24
Oxidation stability (IP, h)	EN 14112	3 minimum	----	9.2
Cloud point	D 2500-05	–3 to +12		–3
Pour point	D 97-96a	–15 to +10	–6	–12
Cetane number	D 613	>47	60	51
Cold filter plugging point	D 6371	–4 to –9	----	–5
Density at 15 °C (Kg/m ³)	D 976	0.575 to 0.900	0.9251	0.8733
Flash point (°C)	D 93	>100	266	110
Carbon residue (wt%)	D 4530	<0.02	0.01	0.005
Copper strip corrosion (3 hrs at 100 °C)	D6751	<No. 1.0	1.0	1.0
Water content (ppm)	D 95	<500	1,293	230

Cold flow properties:

The cloud point (CP), pour point (PP) and cold filter plugging point (CFPP) measurements were done as per ASTM standards, D 2500-05 [27], D 97-96a [28] and D 6371-05 [29], respectively. A Lawler model DR-34H automated cold properties analyzer (Lawler Manufacturing Corporation, Edison, NJ) was used to measure the cold flow properties.

3. Experimental

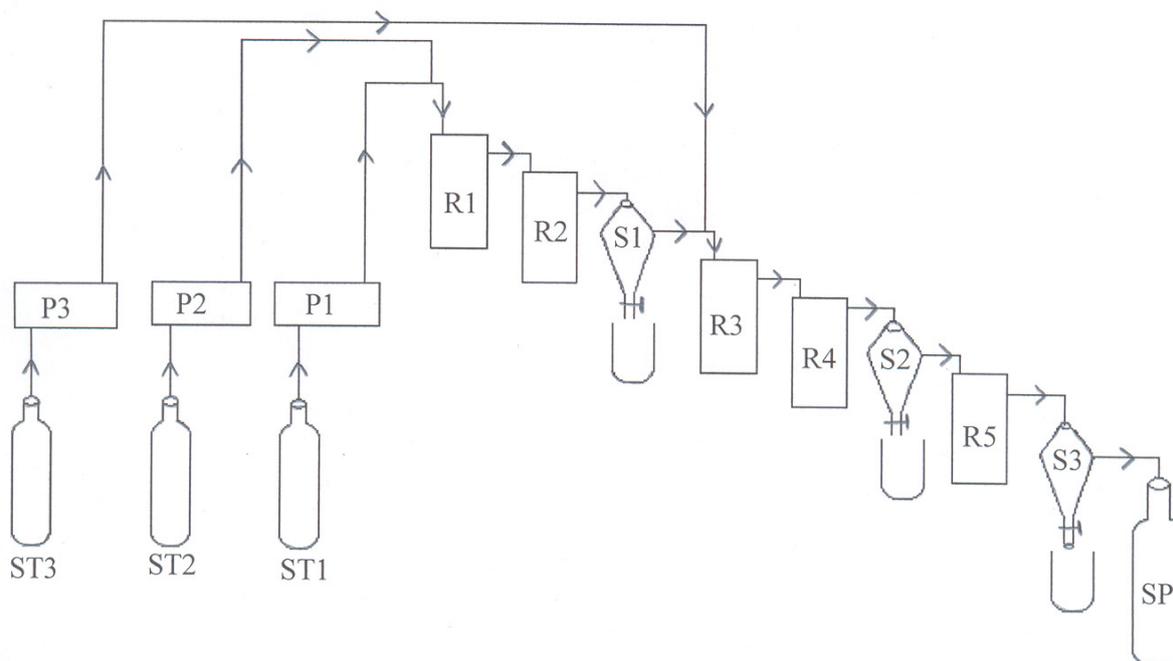
3.1. Material

Commercial grade coconut oil was procured from local markets, and methanol (purity 98%) and KOH (purity 85%) were used from M/S Ranbaxy. The free fatty acid content of oil was determined according to ASTM. According to ASTM, the acid value of the coconut oil was 1.3–1.8 mg KOH/gm sample. Initial composition of oil analyzed by HPLC was triglycerides (TG) 96.5 wt%, diglycerides (DG) 3 wt% and monoglycerides (MG) 0.5 wt%.

3.2. Experimental set-up

The experimental set-up for continuous transesterification of coconut oil is shown in Figure 7. Three storage tanks were used; one to store oil, and two to store two solutions of KOH at different concentrations (0.75% and 0.25wt%) in methanol. Three peristaltic pumps that could run at various constant speeds were used to pump reactants, oil, and methanol to the insulated reactors. In this experiment 5CSTRs (continuous stirring tank reactors) are joined in series at different height and three separating funnel were attached after the second, fourth and fifth reactors for the separation of glycerol. Each reactor was equipped with a mechanical stirrer, thermometer and reflux condensers, which are joined in series, and temperature of the circulating water was maintained at 20 °C with the help of circulating bath. The available mechanical stirring motors (200W and 1,000rpm) were used for the constant stirring; the system could work in the range of ± 5 rpm of the set point. Heating mental were used for the constant heating of the reactors. Temperature of the reactors and stirring speed of the motors were controlled by the controller.

Figure 7. Flow diagram for the continuous transesterification process. (ST1, ST2, ST3) storage tanks, (P1, P2, P3) peristaltic pumps, (R1, R2, R3, R4, R5) continuous stirring tank reactors, (S1, S2, S3) separating funnels, (SP) sample pot.



3.3. Experimental procedure

The two solutions of KOH (0.75 wt% and 0.25 wt%) in methanol were freshly prepared for each experiment, for example for a KOH concentration of 0.75 wt% of oil, 27 gram of KOH was dissolved in 1,000 mL MeOH (because purity of KOH was 85%) and for the second solution 22.5 gram KOH was dissolved in 1,00 mL MeOH. The molecular weight of coconut oil was calculated based on the known fatty acid composition of coconut oil [30]

Pump flow rates were finally adjusted after the empty reactors were preheated. Both feeding pumps were turned on to circulate reactants, oil, and first solution of alcoholic KOH 100 mL/min and 30 mL/min, respectively in the first reactor (molar ratio of MeOH to oil was 5:1 and catalyst concentration 0.75 wt% of oil). As soon as reactants reached the first reactor, the stirrer motor first was turned on and set to the stirring speed of 800 rpm by the controller. The product could be discharged from the reactor first via an overflow, and the discharged product was collected in the second reactor. As soon as product reached the second reactor, the second stirrer was turned on. The total effective volume and residence time of the first two reactors was 1,000 mL and 7.68 min, respectively. After the overflow of the second reactor, the product was collected in the first separating funnel, and the yield of biodiesel in the first separating funnel was 82.6% (Figure 8). The separating funnel was used for the separation of glycerol via downstream process. By the overflow of the separating first funnel, the product reached the third and fourth reactor serially, some potassium hydroxide produced soaps through free fatty acid saponification, and the potassium soaps dissolved in glycerol layer due to their polarity [31], when product reached the third reactor, some amount of alcoholic KOH 12 mL/min (2 mol MeOH with catalyst concentration 0.25 wt % of oil) again fed in third reactor with the help of third Peristaltic pump and start the third stirring motor. The effective volume and residence time of the third and fourth reactors were the same; 600 mL and 4.22 min, respectively. After fourth reactor, the product reached the second separating funnel where again the glycerol produced by the remaining unreacted triglycerides was separated, the yield of biodiesel in the second separating funnel was 96.8% (Figure 9). After separation of glycerol, the overflow product discharged from the second separating funnel was carried into the fifth reactor, and after to the third separating funnel to separate the remaining glycerol via overflow, the effective volume of the reactor fifth was 700 mL and residence time was 4.92 min. The final product was collected in a sampling port and the final yield of biodiesel was $99 \pm 0.05\%$ (Figure 10). The sample was immediately washed with hot water to remove the catalyst. The complete removal of the catalyst was checked by phenolphthalein indicator. Traces of moisture and unreacted methanol were removed by vacuum distillation. The distillation was continued until the loss in weight of ester was constant, thus confirming the complete removal of moisture and unreacted methanol. Finally esters were dried over anhydrous sodium sulphate. The purity and conversion of the product analyzed by HPLC.

Figure 8. HPLC Chromatogram of coconut oil biodiesel in separating funnel first.

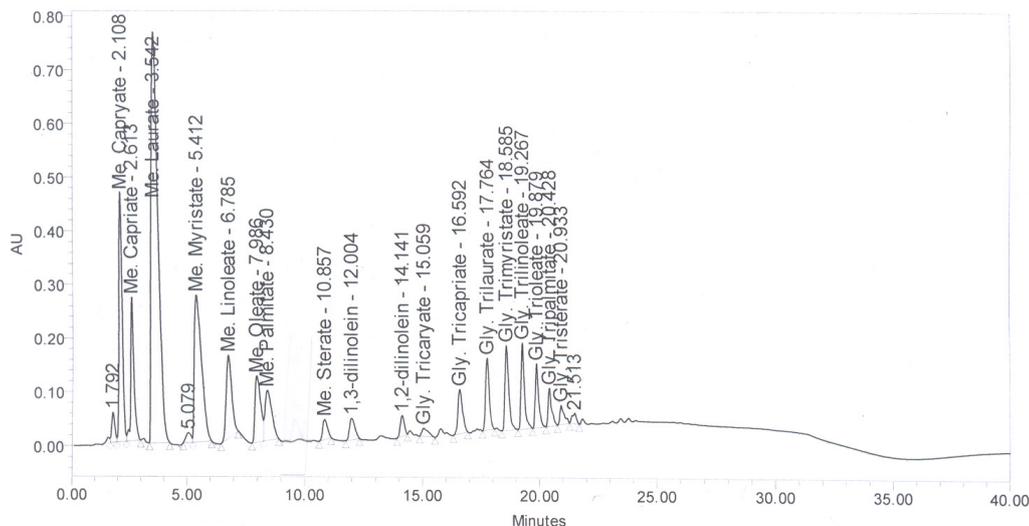


Figure 9. HPLC Chromatogram of coconut oil biodiesel in separating funnel second.

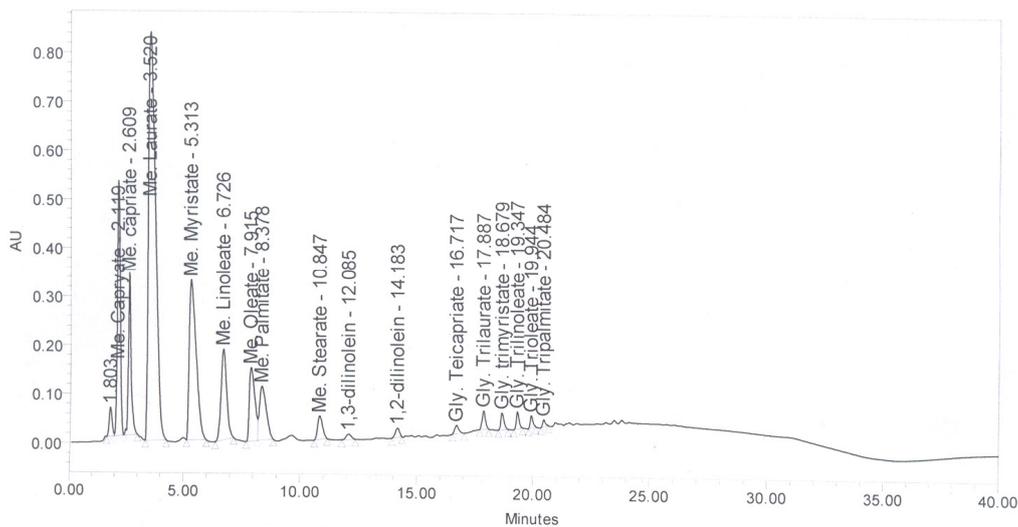
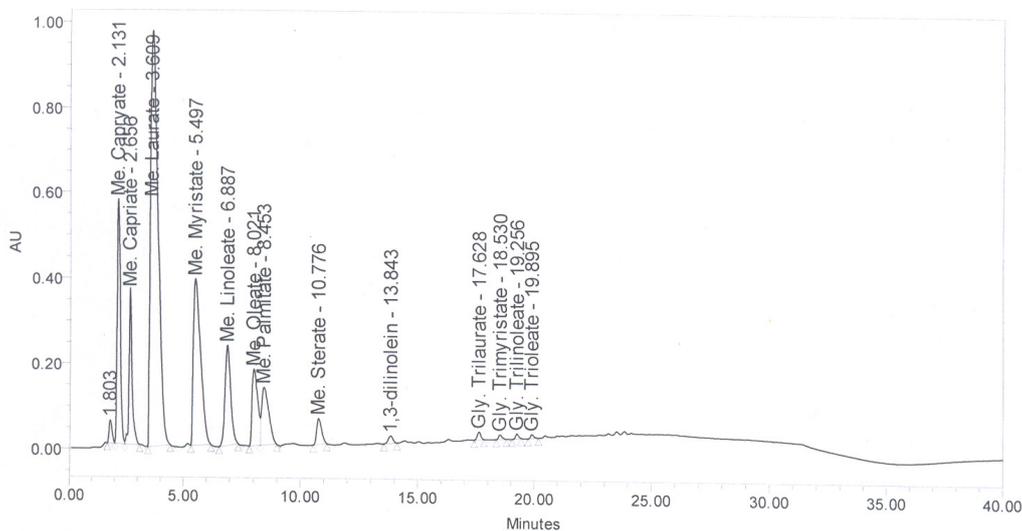


Figure 10. HPLC Chromatogram of coconut oil biodiesel in separating funnel third.



4. Conclusions

The present study demonstrates that the optimal reaction conditions for the continuous methanolysis of coconut oil is using 1% KOH as a catalyst (wt% of oil), with a reaction temperature of 65 °C, methanol:oil molar ratio 7:1, stirring speed 800 rpm, and residence time of 21.04 min. The biodiesel yield is >99%. The biodiesel synthesized meets ASTM specifications for biodiesel, characterized by analytical methods to confirm the purity and quality of biodiesel produced by coconut oil.

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