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Biodiesel Production from Methanolysis of Lard Using CaO Catalyst Derived from Eggshell: Effects of Reaction Time and Catalyst Loading

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ABSTRACT

Biodiesel was produced from lard using a CaO catalyst derived from eggshells. The effects of catalyst loading and transesterification reaction time were investigated. The results revealed that the increase in yield of biodiesel occurred at all catalyst loading when the reaction time was increased. The optimal reaction time was obtained at 60 minutes. The results also indicated that there was an increase in yield of biodiesel when the catalyst loading was increased from 0.5% to 1%. Furthermore, increases in catalyst loading decreased biodiesel yields. The most optimum biodiesel yield of 92.69% was achieved when the reaction time, catalyst loading, methanol:oil molar ratio, reaction temperature, and pressure were 60 minutes, 1%, 6:1, 65 °C, and 1 atm, respectively. The FAME content in biodiesel product was 95.28%. The biodiesel obtained reflected a cetane number and heating value of 46.2 and 37.86 MJ/kg, respectively. Eggshell-derived CaO catalysts exhibited excellent reusability.

1. Introduction

Biodiesel is currently being developed as an alternative fuel on account of its many advantages over diesel oil, which include nontoxicity, environmentally friendly, biodegradability, high cetane number, and low emission [1, 2]. Biodiesel is a renewable resource composed of a mixture of various FAAE (fatty acid alkyl esters). Biodiesel, as a renewable alternative energy, can be made from vegetable oils and animal fats as a source of raw materials [3, 4]. Biodiesel can be obtained by esterification or transesterification process. Esterification process is the reaction of FFA (free fatty acids) with alcohol to produce FAME (fatty acid methyl ester) and water [5]. Transesterification is a reaction between triglycerides that can be obtained from oil derived from plants or fats from animals with alcohol to produce FAME and glycerol as a byproduct [6, 7]. Esterification is carried out if the FFA content of the raw materials is higher than 2% [8, 9]. If the FFA content <1%, biodiesel synthesis is carried out by transesterification only.

Of the various types of vegetable oils, the most commonly used to produce biodiesel include rapeseed oil (in Canada), sunflower oil (in Southern Europe), soybean oil (in the United States), palm oil (in South Asian countries, especially Malaysia. and Indonesia), as well as castor oil (in India) [5, 10]. The utilization of animal fats as feedstock for biodiesel production has also been previously studied. Among the animal fats studied were lard, beef tallow, and fish oil [11, 12]. Compared to vegetable oils, biodiesel from animal fats shows several advantages, including high calorific value and cetane number [10]. However, biodiesel from animal fats also shows disadvantages, including high saturated fatty acid contents, plugging points and cold filter clouding point, which can cause problems during winter operations [11, 13].

Biodiesel synthesis from lard has been studied by several researchers [11, 13–18]. In [11] and [15], for example, the authors used KOH as a catalyst and produced biodiesel with FAME contents of 88.7% and 99.4%, respectively. In [18], the authors also used catalysts of 1.25% KOH and obtained a FAME yield of 96%. The biodiesel production from a mixture of soybean oil and lard using NaOH catalyst has been studied [13], which obtained a biodiesel yield of 77.8%. In [14], the authors investigated the transesterification of refined lard. The research was carried out in supercritical methanol. In this work, the transesterification process

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is carried out under reaction temperature conditions of 320-350 °C, reaction time of 5-20 min, methanol:oil molar ratio of 30-60, pressure of 15-25 MPa, and agitation speed of 0-1000 rpm. The results revealed a FAME yield of 89.91%. In [16], the authors explored the synthesis of biodiesel using immobilized C. antarctica lipase B as a biocatalyst. The biodiesel yield of 96.8% was obtained by a ultrasonic amplitude of 5 kHz, reaction time of 20 min, 1:4 molar ratio of fat:methanol and 6% (w/w of fat) catalyst level. Meanwhile, CaO as a catalyst has also been used in production of biodiesel by waste lard methanolysis [17].

In general, in the transesterification process, the researchers used a homogeneous catalyst in the biodiesel production from lard. The utilization of heterogeneous catalysts in the transesterification process, especially CaO, has not been widely developed by researchers. Calcium oxide is one of the most active and potential solid catalysts in the transesterification process. This catalyst can be reused, recyclable, inexpensive, non-corrosive, and environment-friendly; moreover, in methanol has low solubility, and a catalyst with high activity for the methanolysis reaction of oil [19, 20]. CaO can be obtained from calcination of waste shells, such as eggshells [21]. The utilization of eggshell as a catalyst in the production of biodiesel from lard has not been widely studied by researchers. Therefore, it is necessary to develop biodiesel production from waste shell, especially eggshells.

The aims of this study were to produce biodiesel from lard using CaO from eggshells as a heterogeneous catalyst, and to determine the effect of catalyst loading and transesterification reaction time. The characteristics of the biodiesel produced, including its density, cetane number, kinematic viscosity, and heating value, were then studied and compared with those of a biodiesel standard.

2. Experimental Methods

2.1. Materials

Methanol (99.9%, Merck) and lard were used as raw materials for biodiesel synthesis. Lard was purchased from a Johar market, Semarang, Indonesia. The reagent used for esterification was HCl (37%, Merck). CaO was prepared from eggshells obtained from restaurant waste around Tembalang, Semarang, Indonesia.

2.2. Catalyst Preparation

The eggshells obtained from the restaurant were rinsed using running water, and then cleaned with distilled water to eliminate dust and dirt. The clean eggshells were then dried in an oven (Memmert UN 55 B214.0281, Germany) at 105 °C overnight. Dry eggshells were crushed and calcined for 4 h in air in a Ney Vulcan muffle furnace (Vulcan Bench Top Furnace model D-550-240V) at 900 °C. The product obtained was CaO [21, 22] as a white powder.

2.3. Biodiesel Synthesis Using Esterification and Transesterification Process

Biodiesel synthesis was carried out by esterification and transesterification processes. Both processes were completed in a three-necked flask (Pyrex). This equipment was connected with a reflux condenser (Pyrex) and heater. Esterification aims to reduce the FFA content in lard. Methanol and oil were added to the three-

necked flask at 6:1 methanol to oil molar ratio. Thereafter, HCl amounting to 0.75% w/w of oil was added to the flask. The mixture was then stirred with a magnetic stirrer (Model No. SP131320-33 240V, Thermo Scientific) at 400 rpm while heating to 65 °C. Esterification was performed for 2 h. The esterification product was placed into a separating funnel and allowed to settle overnight. The product was then collected from the remaining methanol and acid catalyst.

The esterified lard was then reacted with methanol in a three-neck flask with methanol to oil molar ratio of 6:1. A number of CaO catalyst was loaded into the three-neck flask. Then, the mixture was reacted at 65 °C while stirring at 400 rpm. After the transesterification process was completed, the product was placed into a separating funnel (Pyrex) to be separated from the catalyst. The product was allowed to settle overnight to separate the biodiesel from excess methanol and glycerol. The glycerol at the bottom of the funnel was withdrawn, and the excess methanol in the top biodiesel layer was removed. Yield of biodiesel was determined by using (1);

$$Yield(\%) = \frac{\text{weight of biodiesel product } (g)}{\text{weight of lard } (g)} x 100\%$$
 (1)

Variables influencing the transesterification process, such as catalyst loading (0.5-7%) and reaction time (30-120 min), were also studied.

3. Results and Discussion

3.1. Esterification Result

During biodiesel synthesis, raw materials with high FFA contents require a two-step process, namely esterification and transesterification [7, 8]. In general, lard has an FFA content higher that 1% [16]; in fact, the raw materials analysis showed that the FFA content of lard was 10.05%. Thus, esterification is needed to decrease the FFA content of the raw material. The esterification process can decrease the FFA content of lard to 1.36%. To produce biodiesel, transesterification is then carried out.

3.2. Biodiesel Production Via Transesterification

3.2.1. Effect of transesterification reaction time

The effect of transesterification reaction time on biodiesel yield was investigated. The reaction time is an important parameter in biodiesel production [8]. The effect of reaction time on biodiesel yield can indicate the costs involved in producing biodiesel [23]. Here, transesterification reaction times of 30, 60, 90, and 120 minutes were applied. The experimental results were shown in Figure 1.

Figure 1 presents an increase in yield of biodiesel at all catalyst loadings and reaction times of 30–60 minutes. At the reaction time of 30 minutes with 1% catalyst loading, the biodiesel obtained was only 75.3%. This shows that at the reaction time of 30 minutes, the transesterification reaction has not reached equilibrium. This means that not all reactants have enough time to interact both between reactants and with the catalyst, resulting in lower biodiesel. Biodiesel yield increased and achieved its optimum at reaction time of 60 minutes. The optimum biodiesel yield was 92.69%. The experimental results reveal that a reaction time of 30 minutes is insufficient to complete the transesterification process.

Furthermore, at the reaction time after 60 minutes, there was a slight increase in the yield of biodiesel and it became almost constant. The increase in biodiesel yield is not significant. These results indicate that a reaction time of 60 minutes is sufficient to produce an effective biodiesel production.

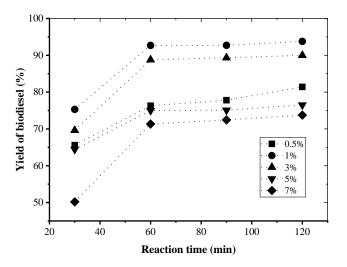


Figure 1: Effect of transesterification reaction time on yield of biodiesel under various catalyst loadings (methanol:oil = 6:1; P = 1 atm; T = 65 °C)

The study conducted by previous researchers [15] also found that the optimal reaction time for production of biodiesel from lard was 60 minutes. In this study, the biodiesel yield obtained in the transesterification step was 97.2% at a KOH concentration of 2% wt and the ratio of methanol to oil was 9:1. In [17], the authors used a CaO-based catalyst from piglet roasting and quicklime during biodiesel production from waste lard and obtained a yield of 97.6% within 60 minutes. This finding reveals that the average transesterification reaction time needed for biodiesel production from lard via conventional methods was 60 minutes. Meanwhile, biodiesel production using supercritical methanol required 15 minutes of reaction to achieved 89.91% FAME content [14].

3.2.2. Effect of catalyst loading

The effect of catalyst loading on the transesterification process was studied by varying the CaO contents to 0.5, 1, 3, 5, and 7% (w/w) of lard with 6:1 methanol to oil ratio for 1 h at 65 °C. The resulting yield of biodiesel were presented in Figure 2.

Figure 2 shows that biodiesel yields increase when catalyst loading is increased from 0.5 to 1%. Furthermore, the higher the catalyst loading, the lower the biodiesel yield. The high catalyst concentration causes the mixture in the reactor to become too viscous. This fact results in more mass transfer resistance and therefore, perhaps some of the catalysts remain unused, which in turn lowers biodiesel yield [24]. In this study, an increase in catalyst amount causes the formation of a slurry from the mixing of the catalyst and reactants. The larger the amount of catalyst added to the reaction system, the more viscous the mixture becomes. As a result, higher power consumption is needed during the stirring process. Increased viscosity due to slurry formation must be prevented because it causes yield of biodiesel to decrease. In order for this problem to be avoided, it is necessary to determine the optimum amount of catalyst loading. The increase

in viscosity can also cause a slight soap formation which hinders the reaction process resulting in a decrease in yield. The results reveal that the optimum catalyst loading in this study is 1%.

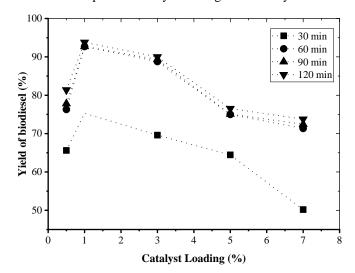


Figure 2: Effect of catalyst loading on biodiesel yield at various reaction times (methanol:oil = 6:1; P = 1 atm; T = 65 °C)

Works that have been done in previous studies also indicated the same results [11, 18]. In [11], the authors applied KOH as a catalyst. The concentration of KOH varied at 0.6%, 0.9% and 1.2%. The results showed that the optimum biodiesel product with a FAME content of 88.73% was achieved at 0.9% wt KOH with a reaction temperature of 60 °C, a methanol-lard mole ratio of 6:1, and a stirring speed of 600 rpm. Meanwhile, in [18], the authors performed the optimization of lard oil methanolysis using a potassium hydroxide catalyst. The results showed that the optimum biodiesel yield was 96.2% at the amount of catalyst 1.25% under the following conditions: reaction temperature of 60 °C, reaction time of 40 min, the molar ratio of methanol to oil was 6:1, and stirring speed of 250 rpm.

3.2.3. FAME content of the biodiesel product

Optimal biodiesel production occurs under a reaction time of 60 minutes and 1% wt catalyst loading; these conditions produce an optimum biodiesel yield of 92.69%. The biodiesel product produced at these optimum conditions was analyzed using GC-MS (gas chromatography—mass spectrometry) (QP2010S SHIMADZU, DB-1 column) to determine its composition, and the results are presented in Table 1. The biodiesel obtained contained approximately 24 compounds, 15 of which were methyl esters.

Table 1: Composition of the biodiesel product

Peak	Retention time	Composition (%)	Name	
1	20.276	0.02	Decanoic acid, methyl ester (CAS)	
2	26.907	0.19	Dodecanoic acid, methyl ester (CAS)	
3	32.676	0.95	Tetradecanoic acid, methyl ester (CAS)	
4	35.299	0.04	Pentadecanoic acid, methyl ester (CAS)	

5	37.186	0.18	9-Hexadecenoic acid, methyl ester, (Z)- (CAS)		
6	37.312	0.67	9-Hexadecenoic acid, methyl ester, (Z)- (CAS)		
7	38.035	22.08	Hexadecanoic acid, methyl ester (CAS)		
8	38.778	0.03	Pentadecanoic acid (CAS)		
9	39.658	0.09	9-Octadecenoic acid (Z)-, methyl ester (CAS)		
10	39.914	0.19	Heptadecanoic acid, methyl ester (CAS)		
11	41.888	11.07	9,12-Octadecadienoic acid (Z,Z)-, methyl ester (CAS)		
12	43.118	53.75	9-Octadecenoic acid (Z)-, methyl ester (CAS)		
13	43.634	5.47	Octadecanoic acid, methyl ester (CAS)		
14	44.198	0.05	9-Hexadecenoic acid, methyl ester, (Z)- (CAS)		
15	45.396	0.18	Methyl arachidonate		
16	45.776	0.16	7,10,13-Eicosatrienoic acid, methyl ester (CAS)		
17	46.186	1.95	Tridecanedial		
18	46.796	0.37	Eicosanoic acid, methyl ester (CAS)		
19	49.475	0.14	5,8,11,14-Eicosatetraenoic acid, ethyl ester, (all-Z)- (CAS)		
20	49.773	1.16	Di-(9-octadecenoyl)-glycerol		
21	50.414	0.65	Hexadecanoic acid, 2-hydroxy-1- (hydroxymethyl) ethyl ester (CAS)		
22	53.237	0.05	3-(Dideuteromethoxymethoxy)- 2,3-Dimethyl-1-Undecene		
23	54.075	0.48	9-Octadecenoic acid (Z)-, 9- octadecenyl ester, (Z)- (CAS)		
24	69.515	0.09	Cholest-5-en-3-ol (3.beta.)- (CAS)		
		100.00			

According to Table 1, the FAME content in biodiesel produced from lard is 95.28%. This result is slightly lower than the standard purity of biodiesel according to EN 14214 [25] which is 96.5%, but higher than the previous studies [13]. In [13], the authors obtained a biodiesel yield of 47.2% with a purity of 92.0% under a reaction time of 3 h and a yield of 66.2% with a purity of 95.7% under a reaction time of 5 h. The findings thus far indicate that lard is a potential feedstock for biodiesel production because it can produce large amounts of FAMEs.

3.2.4. Characteristics of the biodiesel product

Density, kinematic viscosity, cetane number, and heating value are important properties in biodiesel. These properties indicate the quality of the biodiesel fuel so that it will affect the utilization of this fuel. Density and kinematic viscosity directly influence the atomization process during combustion. Meanwhile, the ignition quality of the fuel is evaluated using a cetane number. Heating value is also known as gross calorific value. This property is commonly used to determine the energy content of fuels and their efficiency. The characteristics of the biodiesel obtained at optimum conditions are shown in Table 2.

Table 2 shows that the biodiesel product obtained has a density of 0.860 g/cm3. In the combustion process, a certain amount of fuel is injected into the combustion chamber. The amount of fuel

injected into the combustion chamber is influenced by the density of the fuel. Likewise, the air-fuel ratio is influenced by the density of the fuel. In the same volume, a denser fuel contains a greater mass. It means that a high density contains a high mass. Meanwhile, the fuel injection pump measures fuel by volume rather than by mass. Therefore, if the fuel density is too high it will affect the performance of the injection pump, as a result the combustion process becomes incomplete. The density obtained in this study has met biodiesel standards with the minimum standard of density being 0.860 g/cm3 and a maximum of 0.900 g/cm3.

Table 2: Composition of the biodiesel product

Fuel property	Units	Standard	Method	Experimental values
Density	g/cm ³	0.860 to	EN ISO	0.860
(25 °C)		0.900	3675 [26]	
Kinematic	mm ² /s	1.9 to 6.0	ASTM	4.48
viscosity			D445-19	
(40 °C)			[27]	
Cetane	_	47 (min)	ASTM	46.2
number			D613-18a	
			[28]	
Heating value	MJ/kg	39.72	ASTM	37.86
			D240-19	
			[29]	

Table 2 indicates that the kinematic viscosity of the biodiesel product obtained in this study is between the minimum and maximum standards of biodiesel. Viscosity is an important parameter of the biodiesel fuel produced. Viscosity affects the quality of atomization [4]. High viscosity can lead to poor atomization process. The viscosity of the fuel also affects droplet size. High viscosity has the potential to produce larger droplets at the time of injection so that the atomization process is disturbed, as a result of which the spraying becomes poor. Poor atomization results in poor combustion which increases exhaust fumes and fuel emissions. Fuels with high viscosity also result in the need for more energy to pump fuel and can cause deposits in the engine. The kinematic experimental value of the viscosity produced in this study was 4.48 mm²/s. This value is still below the maximum value required by the ASTM D445-19 specification (6.0 mm²/s).

The cetane number, defined by ASTM D613-18a, is a measure of the auto-ignition delay time of the fuel. Diesel engines desirable high cetane numbers. A higher cetane number results in a shorter time between ignition and initiation of the fuel injection. The high cetane number also ensures the engine is started and running properly and smoothly. In contrast, a low cetane number causes incomplete combustion which tends to increase gas and particulate emissions which in turn result in air pollution. The cetane number value generated in this work is 46.2. This cetane number is slightly lower than the minimum biodiesel standard required, which is 47. This is probably due to the double bonds of fatty acid compounds. Compounds with double bonds can cause a lower cetane number [30]. The results of GC-MS analysis show that the biodiesel product obtained contains several compounds with double chains, such as 9.12-octadecadienoic acid and 9octadecenoic acid, which could reduce its cetane number.

Table 2 also reveals that the heating value of the biodiesel obtained in this study is 37.86 MJ/kg. The heating value describes

the amount of heat produced from the fuel combustion with oxygen [4]. At the initial temperature, this combustion produces $\rm CO_2$ and $\rm H_2\rm O$. The higher the heating value, the less the amount of fuel needed to produce combustion heat. Thus, the higher the heating value, the more efficient the use of fuel. Increasing the amount of carbon in the fuel molecule will increase the heating value of the fuel. The increase in heating value is in line with the increasing ratio of carbon and hydrogen to oxygen and nitrogen [4]. The heating value of the biodiesel product obtained in this work is approximately 4.68% less than that of biodiesel standard and 11% less than petrodiesel, which has a heating value of 42.7 MJ/kg.

3.3. Catalyst Reusability

One method to determine the stability of a heterogeneous catalyst is to test the recyclability of the catalyst. The reusability of the catalyst was carried out in conditions where the transesterification process produced optimum biodiesel yield, namely the reaction temperature of 65 °C, the mole ratio of methanol: lard of 6:1, the catalyst loading of 1% and the reaction time of 1 h. The catalyst that has been used in each cycle was separated by filtering and then washed with methanol to remove adsorbed material. The catalyst was then recalcined at 900 °C for further use. The results of the catalyst reusability were shown in Figure 3.

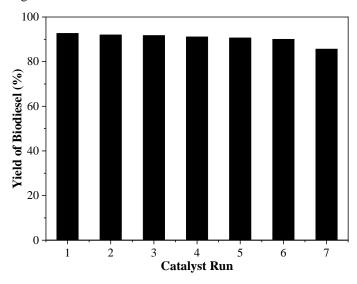


Figure 3: Effect of catalyst reusability on biodiesel yield (methanol:oil = 6:1; P = 1 atm; T = 65 °C; catalyst loading = 1%, reaction time = 60 min)

Figure 3 shows that the biodiesel yield decreases with increasing catalyst use cycles. Nevertheless, the catalyst can be reused after six consecutive runs with excellent activity in the lard transesterification reaction for biodiesel production. The biodiesel yield obtained was still above 90%. However, in the seventh run the biodiesel yield was found to have decreased below 90%, which was 85.67%. The decrease in biodiesel yield after each run may be due to the blocking of the active sites on the catalyst surface by the deposition of unreacted oil, biodiesel or glycerol [24]. These results indicate that the CaO catalyst derived from eggshells has a high potential to be used as a catalyzing for biodiesel production with good reusability.

4. Conclusion

Biodiesel can be obtained from lard in high yields. The reaction time of transesterification required to obtain biodiesel is only 60 minutes with 1% wt catalyst loading. The biodiesel yield produced from this study was 92.69% with the FAME content was 95.28%. The fuel properties of biodiesel products that meet the established standards are kinematic viscosity and density. Meanwhile, the heating value and cetane number obtained in this work are slightly lower than those of the biodiesel standard. In the transesterification process, the CaO catalyst derived from eggshells is able to maintain its catalytic activity after being reused 7 times. The results generally prove that biodiesel synthesis from lard using a heterogeneous catalyst, specifically CaO from eggshell, is an acceptable strategy.

Conflict of Interest

The authors declare no conflict of interest.

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