Heterogeneous catalyzed FAME synthesis from renewable feedstock

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Abstract

With depleting conventional energy sources there is urgent need to look for alternatives for automotive fuel especially from Non-edible feedstocks. Present work investigates FAME (fatty acids methyl esters) synthesis commonly known as Biodiesel from non-edible feedstock Karanja using heterogeneous catalyst calcium oxide (CaO). Study was investigated considering effects of oil-methanol mole ratio, concentration of catalyst and temperature of reaction. Our results confirm that higher mole ratio was useful to enhance FAME yield. The findings of work confirm that molar ration of 1:8 was optimum for catalyst loading of 2.0\% and reaction temperature of 60\textdegree{}C. The maximum yield of FAME was found 92.4\%. The work confirms the FAME obtained from Karanja has a potential alternative for petro-based diesel. Novel protocol developed in present work address the issues of downstream processing of biodiesel synthesis and considerable amount of wastewater generated can be avoided.

Keywords: FAME, Non-edible Oil, Biodiesel, Synthesis

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1. Introduction

Energy demand in India is increasing on multi-fold basis. That too requirement of automotive fuel is increasing with globalization and changing lifestyle. Demand of petro-base diesel is increasing and estimated present crude oil demand is 90 billion metric tons per annum. Recent data released by petroleum-planning analysis wing of oil ministry of Indian government about the fuel consumption stated that demand in February 2019 was increased by 2.7\% and reached to 17.5 million tons and was higher compare to previous quarter. India is meeting 80\% crude oil demand through imports and large quantum of economy is spent on same (Colombo et al., 2017; Petroleum Planning and Analysis Cell All India Study on Sectoral Demand of Diesel & Petrol, 2013; White Paper on Alternative Fuels for Vehicles Vision & Recommendations Alternative Fuels in India, 2019). Hence, there is urgent need to explore the never/renewable sources for automotive sector.

India has targeted to reduced crude oil dependence by 10\% up to 2022. Waste cooking oil consider as a probable option for FAME synthesis, commonly known as biodiesel to achieve goal of 2022(White Paper on Alternative Fuels for Vehicles Vision & Recommendations Alternative Fuels in India, 2019). Biodiesel synthesis from other non-renewable sources need to be explored. From the Indian context Karanja, Jatropha, Nagchampa (Calophyllum Innophyllum Linn), Neem oil are considered as possible
bases for biodiesel synthesis. Oil obtained from these feedstocks are primarily used for medicinal purpose, but residue remain after extraction of essential product can be used for synthesis of fatty acids methyl esters (Halder et al., 2014; Patel & Sankhavara, 2017; Takase et al., 2015). Karanja (known as pongamia pinnata) is fast rising average size tree with size of 30-40 ft and spread awning shade. It can be grown semi-arid region condition and wide range conditions of India. The different type soil such as salty, alkaline, hefty, types of soils, due this nature it can be grown all parts of India. Shrunk of Karanja is small in diameter and height is more than 1.65 meter. It has long leaves of size 5-10 cm extended and 4-7 cm widespread and organized in 2-3 pairs. Flowers are generally pea-shaped type with 15-19 mm lengthy and glowing, well-lit purple or white in color. Kernels are sunlit brown in hue with size 1-1.5 cm (Banković–Ilić et al., 2017; Halder et al., 2014; Harsha Hebbar et al., 2018; Patel & Sankhavara, 2017; Takase et al., 2015). The physico-chemical properties of Karanja are shown in Table 1.

One tree can give 9-90 kg of seed pod which yields up to 35-40% oil. Kernels of tree due content oil and yield 30-40% oil, normal Karanja produce 8-24 kg kernels. The typical composition of air-dried kernels consists of 27-28% oil, 17-18% protein, 6.6-6.8% starch, 7-8% fiber and ash,2.3% (Halder et al., 2014; Patel & Sankhavara, 2017)
typical this value needs to be reduced below 2% to avoid saponification during alkaline transesterification. Hydrolysis- and -
generation of left-over by-products further hamper the reaction and product yield may be lowered significantly. It required large
quantum of water for downstream-processing to remove waste and lengthier duration for reaction-completion (Banković–Ilić et al.,
2017; Harsha Hebbar et al., 2018; Kavitha et al., 2019; Madhu et al., 2016) There are number of alternatives are available for
feedstock with large quantum of free fatty acids which are listed below:

- Esterification using different ion exchange resins or heterogeneous catalyst
- Distillation
- Acid esterification in presence of catalyst such as sulphuric-acid, methane-sulfonic-acid, ferric-sulphate
- Neutralization using alkali- followed by soap-separation in decanter
- Use of polar-liquids for extraction with combination of acid-esterification

Major limitations associated with esterification are higher amount of chemical requirement, waste-removal and lengthier-
duration for reaction completion. Hence, there is urgent need to look of potential alternative for elimination of FFA in oil
(Dhanasekar et al., 2019; Madhu et al., 2016). Heterogeneous catalyst can overcome the limitations associated with acid
esterification followed by transesterification. Major advantage associated with this heterogeneous catalyst to reduced wastewater
generation for downstream processing. Calcium oxide is one better option as heterogeneous catalyst for biodiesel synthesis
(Banković–Ilić et al., 2017; Harsha Hebbar et al., 2018; Kavitha et al., 2019; Madhu et al., 2016). It is commonly obtained from
waste such as eggshells and mollusk shell.

Present work investigate FAME synthesis from Karanja. Process parameters such as oil-methanol mole- ratio, catalyst -
centration and temperature of reaction. Analysis of product was carried out using HPLC (High-performance liquid-
chromatography).

2. Materials-Method

2.1 Materials
Raw Karanja oil was procured from Pune local market. Fatty-acid oil-composition is shown in Table 2. Methanol(HPLC grade),
Sulfuric acid (98%), Hexane(HPLC grade), and Calcium Oxide was obtained from M/s New-neeta, Chemicals, Pune. HPLC
internal-standards viz methyl-linoleate and methyl-oleate was received gift from ICT, Mumbai.

2.2 Reactor
The reaction was carried out in 5.21 cm (ID) and 160 cu cm volume. Glass-reactor fortified with reflux-condenser.Reactor
temperature was maintained by keeping glass reactor in constant temperature bath. Water bath is made of M/s Medico Instruments,
Mumbai and temperature of reactor can regulated within ± 2°C. Pitch type of blade-glass stirring having 1.2 inches dia- was used
to attain even mixing reactants. Experimental setup representation is desiccated in Figure 1.

90 mL of Karanja-oil was placed in reactor and preheat to 110°C to remove initial moisture-content oil and was observed using
Karl-Fisher-Titration Unit. For preparation of calcium oxide and was heated for known temperature of 700°C for 4-5 hrs to remove
volatile matter and moisture using Muffle furnace. Heated oil cooled to 60°C was mixed with known quantity of calcinated
calcium oxide and methanol was added. Reaction mixture was heated, and samples were withdrawn at regular of time followed
quick quenching in ice bath to kill reaction. Cooled samples were transferred to separating funnel. Lower layer consist of heterogeneous catalyst and glycerol was detached. Upper-layer consists of FAME was eroded with hot distilled water to remove catalyst-traces and other impurities. Sample was heated under vacuum in rotary vacuum evaporator to remove traces of moisture. Further quantification of FAME was verified using HPLC.

2.3 Analysis

Transesterification samples were analyzed using HPLC (Hitachi) with C18 Phenomenex-column (25cm×4.6mm, 5µm particle-size) using UV-visible detection at 210nm. Lined elution of methanol- and -hexane was maintained for analyzing samples. The mobile-phase contains of 90% methanol- and 10% hexane- and flowrate of 1 ml min⁻¹. Sample was prepared by using 10µl of sample diluted-with 10mL of methanol. Calibration curve was prepared using internal-standards of methyl-linoleate and methyl-oleate known concentration and area under the curves. Concentration of unknown samples were calculated calibration curve.

3. Results and Discussion

Calcium oxide has hydro-philic nature and remains in methanol-phase. Methoxy ions are generated on surface of catalyst. Ions generations results of methanol molecule adsorption and subsequent transfer to organic-phase at the crossing point between the two-phases and reaction progress. Progress of reaction in presence of CaO as catalyst depends on micro-emulsification between heterogeneous phases of solid catalyst and immiscible liquid phases(Banković–Ilić et al., 2017; Halder et al., 2014; Patel & Sankhavara, 2017; Takase et al., 2015). Since transesterification is reversible in nature and forward path of -reaction is always desirable for achieving higher product yield (Dhanasekar et al., 2019; Madhu et al., 2016). Parameters of process such as oil-methanol mole ratio, catalyst-concentration and temperature of reaction temperature have great influence on the product yield.

3.1 Oil-methanol mole ratio effect

Stoichiometrically, 3 moles of CH₃OH require for 1 mole FAME production. Excess methanol may require to maintained forward path of reaction (Banković–Ilić et al., 2017; Patel & Sankhavara, 2017). Experiments were performed using oil-methanol mole-ratio of 1:4, 1:6, 1:8 and 1:10 and catalyst-concentration of 2% and temperature of reaction as 60°C was kept constant for all experiment runs and results are dissipated in Figure 2.

![Figure 2: Effect of oil-methanol mole-ratio FAME yield](image)

It has been observed that FAME yield was increasing with increase in oil-methanol mole-ratio 1:4-1:8 with further rise in mole-ratio has marginal incremental effect of FAME yield. Hence, 1:8 molar ratio was considered as optimum for further experimental investigations. The initial increase in product yield with mole-ratio may be attributed that the higher moles are favorable to keep the forward-path of reversible-reaction. Higher molar ratio than optimum value is not favorable, glycerol produced as side product during reaction may mix in surplus methanol and reaction- can be reverted. It may lead to decrease in yield of desired product. Separation of methanol from final product increase the downstream processing cost (Banković–Ilić et al., 2017; Harsha Hebbar et al., 2018; Kavitha et al., 2019; Madhu et al., 2016). Hence, too methanol is not recommended for synthesis of fatty acids methyl esters.

3.2 Catalyst-concentration effect

Study of catalyst-concentration effect on FAME synthesis is important to avoid the separation of catalyst from the final product. Excessive catalyst concentration may hinder rate of reaction. Higher concentration reduced the emulsification rate between the heterogeneous phases involved in reaction mixture (Dhanasekar et al., 2019; Harsha Hebbar et al., 2018; Kavitha et al., 2019; Madhu et al., 2016). Stirring may not able to maintain desire emulsification rate.

Experiments were performed varying concentration of catalyst as 1, 2 and 3% weight of catalyst/weight of oil (w/w) and results have been shown in Figure 4. FAME yield increased with time and catalyst-concentration. It has been seen that product yield increase with rise in catalyst-concentration from 1 to 2% w/w, additional rise in catalyst-concentration does not yield substantial increase in FAME yield. Hence 2% catalyst concentration optimum for further investigation.
Activity of catalyst is mainly related with basic strength and increase with catalyst loading till optimum value. Studies in literature reported that Mg/Cl catalyst loading surpass outside an optimal value then catalyst activity diminish due to development of newer feebler basic sites. The increase in FAME yield observed to optimum catalyst loading, higher loading can attribute to likely formation of agglomerate catalyst particles (Harsha Hebbar et al., 2018; Kavitha et al., 2019). One important point to note here that additional catalyst use may require more energy for separation process. Hence, optimum-catalyst selection is desirable. Work on trans-esterification of soybean-oil in CaO presence shown 100% conversion of soybean was observed for catalyst loading of 3% (w/w) (Takase et al., 2015). They also reported that CaO use as heterogeneous- catalyst reduce downstream processing requirement for catalyst separation from final product.

### 3.3 Reaction- temperature effect

Reaction- temperature effect on trans-esterification was investigated using varying reaction- temperature as 40°C, 50°C, 60°C and 65°C at previously optimize catalyst concentration as 2%(w/w) and mole- ratio of oil-methanol as 1:8. Reaction- temperature effect on FAME yield has been shown in Figure 4. FAME yield increased with increase in reaction- temperature 40-60°C, with further increase in temperature to 65°C shown marginally decrease FAME yield.

![Figure 4: Reaction temperature effect on FAME yield](image)

The optimize results fatty acids methyl esters from Karanja using calcium oxide shown in Table 3.

| Parameter                  | Value |
|----------------------------|-------|
| Reaction Temperature (°C)  | 60    |
| Catalyst-Concentration (%wt wt⁻¹) | 2   |
| Oil-methanol mole-ratio   | 1:8   |
| Reaction- time (min)      | 100   |

Table 3 shows that reaction time required for synthesis is marginally higher compare to homogenous alkali catalyst. Most of the studies shown that reaction time for homogenous catalyst biodiesel synthesis is 60 to 90 min. The marginally higher reaction time required due to kinetics for heterogeneous nature of reaction. However, there are considerable advantageous of heterogeneous
catalyzed reaction such as lesser downstream processing requirement for separation of catalyst from final product and lesser wastewater generation improving quality of product.

4. Conclusions

Fatty acids methyl ester was obtained from non-edible feedstock from Karanja oil. Heterogeneous catalyst calcium oxide is explored as potential heterogeneous catalyst for synthesis of FAME. There was marginal higher molar ratio required for synthesis compared to conventional alkaline homogenous catalyst. Major advantageous of using heterogenous catalyst reduced the downstream processing for separation of catalyst from final product and wastewater generation for enhancing purity product was considerable product. The outcome of this study showed novel technique for synthesis of valuable product fatty acids methyl esters. Experimental protocol developed in present work can be explore as new and innovative path to synthesis FAME in forthcoming.

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