

Effects of Fatty Acids Composition on Fuel Properties of *Jatropha Curcas* Biodiesel

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Abstract

In the efforts to reduce effects of climate change, biodiesel fuels from plant oils such as *Jatropha curcas* have been proposed as alternative fuels which can be used in the transportation sector in diesel engines. The current study investigates the effects of fatty acids composition on fuel properties of biodiesel derived from *Jatropha curcas* seeds obtained from selected regions of Botswana. The physicochemical fuel properties investigated include kinematic viscosity, flash point, energy content, density, pour point and cloud point from derived *Jatropha curcas* biodiesel. Results of the study showed that *Jatropha curcas* biodiesel samples for all regions under review are dominated by unsaturated fatty acids which are desirable for cold flow properties and kinematic viscosity of the biodiesel fuel. The major fatty acids in *Jatropha curcas* biodiesel fuels from all the regions range from 69.00% to 77.81% of unsaturated fatty acids. The overall results conclude that fatty acids composition has influence on the fuel properties of the biodiesel under investigated.

Keywords

Jatropha Curcas, Fatty Acids, Fuel Quality, Engine Performance

1. Introduction

The ever increasing demand for energy worldwide, driven by industrialization, has led to reliance on fossil fuels to satisfy these energy needs. However, use of fossil fuels is challenged by the depletion of known resources and the negative impact of the fuel's combustion emissions on the environment which causes adverse effects such as climate change [1]. Consequently, biodiesel fuel has been identified as one of the clean alternative energy sources which can be utilized to mitigate said effects.

Due to global concerns on utilization of edible feedstock such as sunflower

being used for fuel production, lot of work has been carried out to explore the utilisation of non-edible oils such as *Jatropha curcas*, *mahua*, neem and castor for biodiesel production [2] [3] [4]. Rutz and Rainer [5] echoed that the choice for a certain feedstock is pre-determined by factors such as agricultural, geographical and climatic conditions where the feedstock is harvested.

Since biodiesel is made of esters of fatty acids, it is therefore important to perform the esters profiles of biodiesel prior to its use. In most vegetable oil biodiesel feedstock, the major fatty acids present are: oleic, linoleic, palmitic and stearic with oleic being the dominant [6] [7] [8]. Oleic and linoleic acid form part of the unsaturated fatty acid of the biodiesel sample, that is, fatty acids with one or more double bond within their molecular structure. Oleic acid has a single double bond in their structure while linoleic acid has two double bonds.

The unsaturated fatty acids can further be divided into two types namely: monounsaturated and polyunsaturated. The monounsaturated (Cn:1) fatty acid contains only one double bond in their chemical structure while polyunsaturated (Cn:2,3) fatty acid has two or three double bonds. It is ideal for the vegetable oil used for production of biodiesel to have relatively larger percentage of monounsaturated fatty acids than polyunsaturated fatty acids. This is because oil containing relatively high proportion of polyunsaturated fatty acids tends to exhibit a poor oxidation stability and it can compromise the fuel properties such as kinematic viscosity and impair fuel quality [9] [10] [11]. Other fuel properties which are negatively affected by high degree of unsaturation are density and cetane number [6]. The authors further echoed that an increase in degree of unsaturation leads to an increase in density and decrease in cetane number of the biodiesel fuel. Oxidation stability is a measure of change in fuel quality during extended storage. Kinast [12] describes oxidation stability as an indicator of shelf life of the fuel.

On contrary, an increase in degree of unsaturation particularly monounsaturated fatty acids improves other fuel properties such as cold flow properties and kinematic viscosity [7] [8]. The authors further highlighted that higher degree of unsaturation lead to outstandingly improved low temperature performance. As for kinematic viscosity, the same authors alluded that with increase in unsaturation the kinematic viscosity reduces.

Saturated fatty acids such as palmitic and stearic do not have double bond in their molecular structure. Contrary to the discussions made earlier on poor oxidation stability of fuel with high content of polyunsaturated fatty acids (linoleic), for a good oxidation stability, the fuel should have more of saturated fatty acid [8] [13]. The discussions has demonstrated that fatty acids composition is an integral part in the biodiesel production as it has great influence in the physico-chemical fuel properties of the biodiesel fuel.

Several studies have been carried out on the subject of biodiesel, and in Botswana, government conducted a feasibility study on the production and use of biofuels in 2007 [14]. According to the report, *Jatropha curcas* was identified as

the most promising crop for biodiesel production in Botswana. Consequently, in 2012 the government further undertook a five (5) year research project aimed to determine appropriate method for *Jatropha curcas* cultivation in Botswana's climatic condition. Though much literature exists in the area of the current work, the current study is stimulated by government's ambition and efforts to develop a biodiesel sector in the country. Scientific evidence to validate biodiesel from locally grown jatropha kernel oil as good fuel was deemed necessary to inform policy. Hence the current study focuses on investigating the effect of fatty acid composition on fuel properties of *Jatropha curcas* biodiesel. It is the view of the authors that the results will fully support the government's efforts to stimulate rapid development of biodiesel industry in Botswana.

2. Materials and Methods

2.1. Seed Preparation

Jatropha curcas seeds were collected from selected regions of Botswana namely Paje, Serowe, Mmadinare and Tsamaya. **Figure 1** shows the picture of *Jatropha curcas* seeds and seed kernel. All the regions are situated along the eastern side of the country. Selection of these regions was based on the availability of seeds at the time of the study. Prior to the experimental procedures, seeds were cleaned by removing foreign materials such as husk, small stones and leaves. Stones can cause deformation of the press tools and foreign material of plant origin such as weeds and stems can reduce oil quality. This step was done by hand since it was not complex. To further remove any small foreign materials, a metal sieve separator of 5 mm diameter was used. This is similar to a study by [15]. The seeds were then sun dried for about 3 weeks to remove moisture as in the method by [16]. It is important that the seeds are well dried before pressing since moist seeds can develop mould which can jam the pressing equipment.

2.2. Mechanical Extraction

Oil extraction was carried out using Kern Kraft screw press machine (KK40F Universal). After the seeds were cleaned, they were fed on the seed hopper. The rotational speed of the machine was then set at higher speeds of 80 revolutions



Figure 1. *Jatropha curcas* seeds and seed kernel.

per minute (rpm) since *Jatropha curcas* seeds are hard-shelled. During the operation, oil was drained from the holes of the sieve and the press cake continuously ran from the pre-jet.

2.3. Biodiesel Production

Jatropha curcas crude oil was processed into biodiesel. In order to achieve this, the alkali catalysed transesterification process was used. This process was performed using Biodiesel Starter Kit sourced from Division of Reliance Energy Resources, LLC in Florida, United States of America. The laboratory scale biodiesel processing unit works in four stages namely; heating of oil, transesterification process, separation and dry washing as demonstrated in Figure 2.

2.4. Oil Characterization

Oil characterization involves the chemical composition of different derived biodiesel samples to establish its fatty acid profiles. The test method involved analysis of the standard as a reference sample as described in Section 2.4.1 to establish the major esters. Thereafter the biodiesel samples were analysed following the

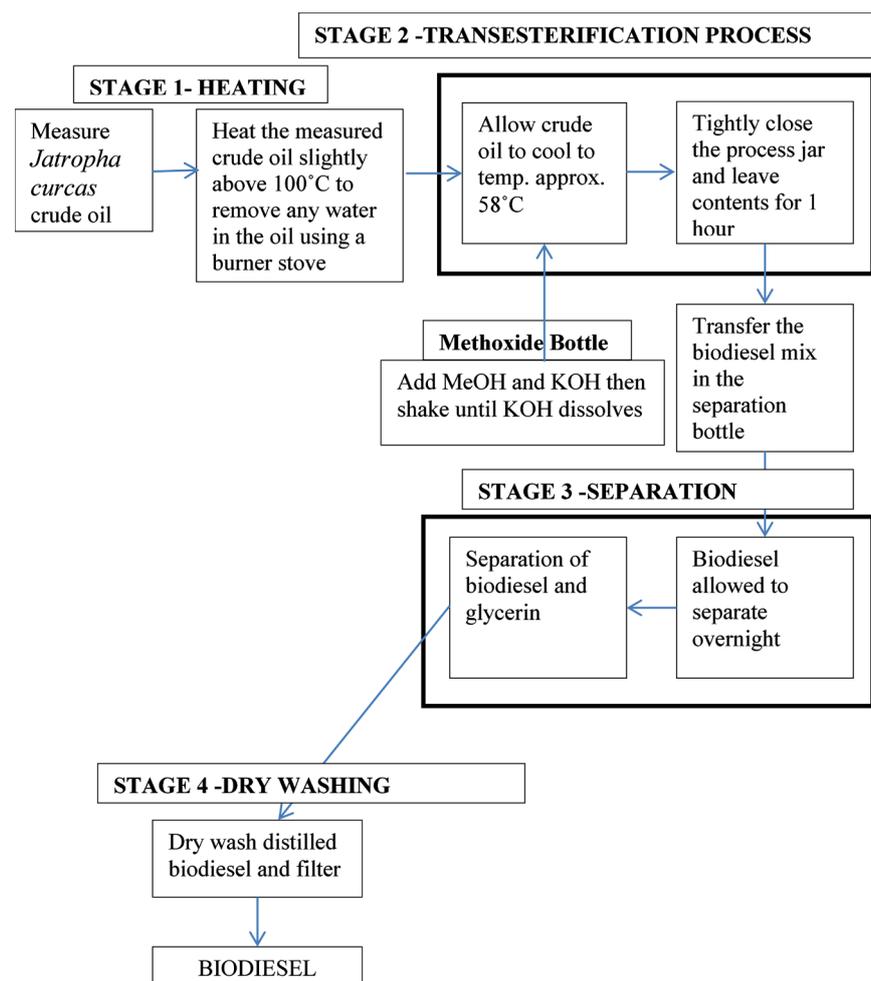


Figure 2. Schematic diagram of biodiesel production from *Jatropha curcas* crude oil.

same procedure used for the standard and quantification was performed on major esters which was present on standard sample.

2.4.1. Procedure

Initially the standard was injected into the Gas Chromatograph-Mass Spectrometry (GC-MS) then ran five (5) times with six (6) different concentrations of equal interval from 10 ppm (part per million) to 1 ppm. At each concentration, peak areas and retention times were recorded and calibration curve was generated with each fatty acid in the standard sample which its calibration curve was used as reference for analysis of fatty acids in biodiesel samples. The same procedure was repeated for each biodiesel sample.

2.4.2. Instrument Conditions

The composition and quantity of fatty acid methyl ester (FAME) was determined according to test method ASTM D6584, using Agilent Technologies GC System 7890A gas chromatograph (GC) equipped with a HP-5MS capillary column (30 m × 250 μm × 0.25 μm) and an automated injector. The instrument which was used in the study was Agilent Technologies GC System 7890A gas chromatograph (GC) connected to the mass spectrometer with Triple-Axis detector. Helium was used as carrier gas at a pressure of 72 kPa and flow rate of 64 mL/min according to manufacturers' specification. 1 μL of sample was injected using an automated injector. The initial oven temperature was set to 100°C for 4 minutes, which was then increased at a rate of 7°C/min to 235°C, then 10°C/min to 300°C for 7 minutes. The injector and detector temperatures were set to 325°C allow these target molecules to volatilize. Total run time was about 36 minutes.

2.5. Fuel Quality Analysis

Biodiesel produced from seeds collected from all the four regions of Botswana was investigated for the following fuel quality properties; kinematic viscosity analysis, density flash point, cloud point, pour point and energy content.

1) Kinematic Viscosity Analysis

Kinematic viscosity analysis was performed using a Fungilab Premium Series Viscometer (PREL 401024). The viscometer was coupled to a Thermo Fisher Scientific heating bath circulator. The temperature of the heating bath was then set to a maximum of 95°C as per the manufacturer specifications. On the viscometer, the low centipoise (LCP) spindle with its appropriate thermo station jacket was used since the samples analysed were of low viscosity.

The instrument was then started simultaneously with the heating bath set at 95°C. The spindle speed was varied based on the torque values with the ideal range being 50% - 95% as suggested by the manufacturer. Sample viscosity readings were performed at temperature intervals of 2°C from room temperature to 60°C because it was noticed that viscosity of biodiesel remains constant between 54°C to 60°C.

2) Flashpoint Measurement

Flashpoints of biodiesel fuel samples were determined using an automated Pensky-Martens Closed Cup Tester (Tanaka) according to the ASTM D93 test method. A 100 ml brass test cup was filled with fuel sample to approximately 60 ml and covered with the test cover. The expected flash point, identification of fuel sample and procedure were then entered on the machine. The test mode used was the special mode (SPEC) since the samples tested were biodiesel fuels of unknown flash point. The test cup was heated and the sample agitated by a stirrer. At regular intervals, an ignition source was introduced into a test cup until a flash was detected. The procedure was repeated thrice for each sample and an average reading recorded.

3) Cloud and Pour Point Analysis

The cloud and pour points of the fuels were tested using the Normalab cloud and pour tester coupled with a cooling bath which uses methanol as the cooling medium. The temperature of the cooling bath was set to -60°C . The test sample was poured into a sample container up to a specified mark and then mounted on the motorised measurement head. The initial temperatures for cloud and pour point were entered prior to the test run. As soon as the specific temperature of cloud and pour point of the test sample were reached, the results were displayed on the screen and recorded.

4) Energy Content

The heating values of *Jatropha curcas* biodiesel fuels were determined using IKA C200 Calorimeter. The instrument was connected to a computer installed with CalWin calorimeter software for easy capture and display data. The experiment entails weighing the sample, oxygenating it and igniting it to determine its energy content. The decomposition vessel was charged with oxygen at a pressure of 30 bars for 30 seconds to ensure adequate oxygen for combustion process. The test run was initiated and the results were automatically saved in the computer. Total run time for each experiment was 9 minutes.

3. Results and Discussions

3.1. Fatty Acid Composition

Fatty acid composition of biodiesel is an important characteristic in biodiesel production. The properties of biodiesel fuel are determined by the quantity of each fatty acid present in the biodiesel fuel sample. **Table 1** shows the results of fatty acid composition in biodiesel produced from *Jatropha curcas* oil from four different regions, namely, Paje, Serowe, Tsamaya and Mmadinare villages.

From the results in **Table 1** it can be observed that the major fatty acids in *Jatropha curcas* biodiesel samples for all regions under review were the oleic, linoleic, palmitic and stearic with oleic being the dominant one. The reported percentage of oleic acid was 48.65% for Tsamaya followed by Paje at 43.14%, Serowe at 41.90% and Mmadinare at 37.92%. The results further demonstrate that the percentage composition of monounsaturated fatty acid found in different

Table 1. Fatty Acid Composition of Biodiesel derived from *Jatropha curcas* oil from regions of Paje, Serowe, Tsamaya and Mmadinare.

Fatty Acid		*Structure	Relative Composition (%)			
Scientific Name			Paje	Serowe	Tsamaya	Mmadinare
Saturated Acid						
Palmitic	Hexadecanoic	C16:0	14.22	15.12	18.06	15.03
Stearic	Octadecanoic	C18:0	8.39	8.31	10.21	7.16
Total Saturated			22.61	23.42	28.27	22.19
Unsaturated Acid						
Oleic	Cis-9-octadenoic	C18:1	43.14	41.9	48.65	37.92
Linoleic	Cis-9,12-octadecadienoic	C18:2	31.42	32.2	18.35	37.48
Palmitoleic	9-hexadecenoic	C16:1	1.01	1.09	1.23	1.03
Isooleic acid	10-octadecenoic	C18:1	1.45	1.39	1.75	1.38
Total Unsaturated			77.02	76.58	69.98	77.81
Total monounsaturated			45.62	44.88	51.06	37.92
Total polyunsaturated			31.42	32.20	18.35	37.48

*Number of carbon atoms: degree of unsaturation.

biodiesel samples is higher than that of polyunsaturated fatty acid. Several studies including [9] [10] [11] recommends that the best vegetable oil used for production of biodiesel should consist a larger amount of monounsaturated fatty acids compared to polyunsaturated fatty acids as oil containing relatively high amount of polyunsaturated fatty acids tends to exhibit a poor oxidation stability and it can compromise the fuel properties such as kinematic viscosity and impair fuel quality. In this case, the situation is favourable. Due to increased amount of unsaturated fatty acids in biodiesel fuel produced from oil seeds collected from Paje and Mmadinare regions, some of the fuel properties such as kinematic viscosity and cold properties are expected to be favourable.

Total saturated fatty acids (palmitic and stearic) of each biodiesel sample is 22.61%, 23.42%, 28.27% and 22.19% for Paje, Serowe, Tsamaya and Mmadinare respectively. Contrary to the discussions made earlier on poor oxidation stability of fuel with relatively high content of polyunsaturated fatty acids (linoleic), for a good oxidation stability, the fuel should have more of saturated fatty acid [8] [13]. Therefore, from the four biodiesel under review, biodiesel fuel from Tsamaya region would be expected to have a good oxidation stability due to relatively high percentage of saturated fatty acids of 28.27% and less polyunsaturated fatty acids of 18.35%

3.2. Physico-Chemical Fuel Properties

1) Kinematic Viscosity

Kinematic viscosity is defined as a measure of the fluid's internal resistance to flow under gravitational forces. **Figure 3** presents information obtained for ki-

nematic viscosity of *Jatropha curcas* biodiesel fuels produced using seeds from different selected regions at different temperatures.

The results in **Figure 3** show that all the fuels tested exhibit decrease in kinematic viscosity as the temperature increase. Azom [17] reported that the increase in temperature causes the kinetic energy to increase and the results in mobility of molecules hence reduction in kinematic viscosity. The observed profile in **Figure 3** agrees with the one reported by Rashid and Anwar [18]. The ASTM D 6751 biodiesel standard prescribe an acceptable kinematic viscosity at 40°C for biodiesel to be at the range of 1.9 to 6.0 cSt. All the recorded results fall within the acceptable ASTM D 6751 biodiesel standard as demonstrated in **Figure 4**. These results guarantees a good combustion processes in the engine without leaving residual residues that can cause damage such as valve sticking and burning. The same observation was alluded by Anguebes-Franseschi *et al.* [19], who reported that higher viscosity tends to form larger droplets on injection resulting in poor combustion processes in compression ignition engines.

Most of the literatures reveal that kinematic viscosity can be affected by the degree of unsaturation [7] [8] [11] [20]. The authors alluded that decreasing degree of unsaturation leads to an increase in kinematic viscosity. In the current

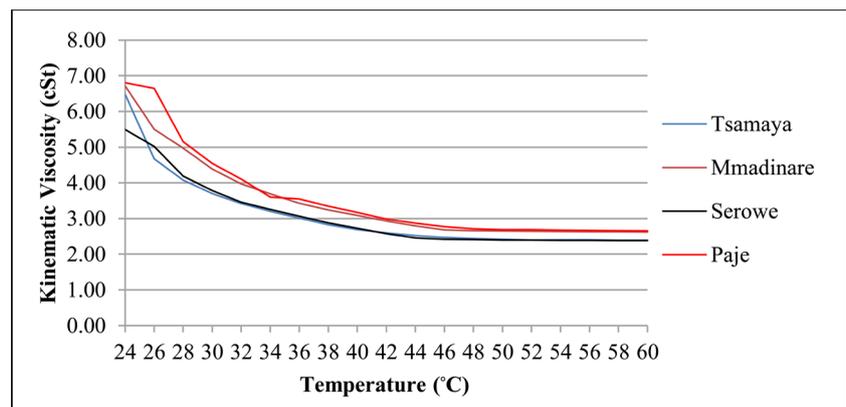


Figure 3. Kinematic Viscosity of *Jatropha curcas* biodiesel fuels at different temperatures.

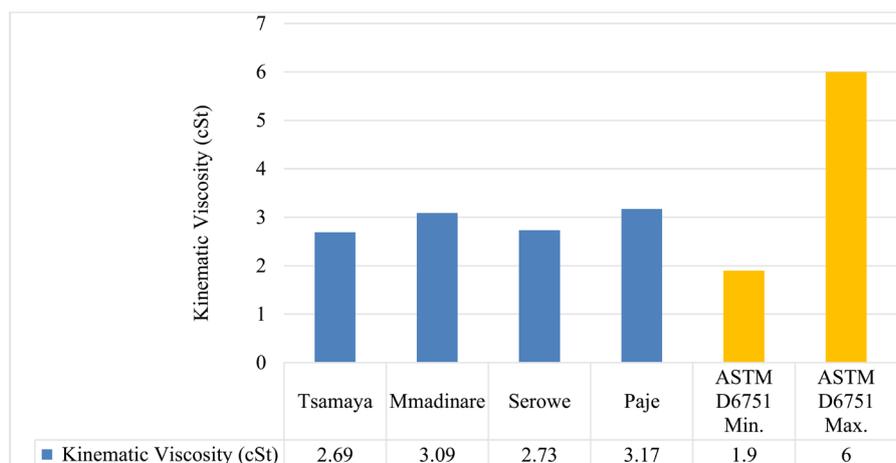


Figure 4. Kinematic viscosity of different *Jatropha curcas* biodiesel at 40°C.

study the results are opposite, an increase in degree of unsaturation leads to increase in kinematic viscosity. The situation could be due to reason given by Wardana *et al.* [21] who reported that the trend may be due to the molecular weight which is more influential on viscosity than the degree of unsaturation. In the present investigation, molecular weights of unsaturated fatty acids are more than that of saturated ones. For example, total unsaturated fatty acids for Paje and Mmadinare village were the highest at 77.81% and 77.02% respectively as demonstrated by **Table 1** in Section 3.1. Corresponding kinematic viscosity of these study areas were also recorded at 3.17 cSt and 3.09 cSt for Paje and Mmadinare respectively, 13.9% and 15% higher than viscosity for Serowe and Tsamaya respectively.

2) Density

Density of the fuel is also an important fuel property that has an effect in the engine performance. It affects the pumping of the fuel because fuel injection pumps fuel by volume not by mass [8]. The authors stressed that the greater or lesser mass of fuel is injected in the compression ignition engine depend upon its density.

The results depicts in **Figure 5** demonstrated that the biodiesel fuels produced from seeds collected from Paje and Mmadinare villages recorded high density of 0.9042 g/ml and 0.9034 g/ml respectively. Similarly, results for Serowe and Tsamaya villages recorded density of 0.8945 g/ml and 0.8804 g/ml respectively. The moderately higher densities recorded for biodiesel fuel of seeds from Mmadinare and Paje villages may be attributed to their higher degree of unsaturation fatty acids as shown in **Table 1** in Section 3.1. Several authors including Hoekman *et al.* [8] and Refaat [22] reported that higher unsaturation leads to relatively high density of biodiesel fuels. Other biodiesel fuels densities reported are in the range of 0.85 to 0.95 g/ml [23] [24] [25] [26]. Additionally, all the reported densities for biodiesel fuels from all the four study areas under review are within the acceptable range of the EN 14214 biodiesel standard which stipulate density range of 0.86 - 0.90 g/ml. The ASTM D6751 biodiesel standard does not specify any range. The observation has also been stressed by several authors including Patel *et al.* [27].

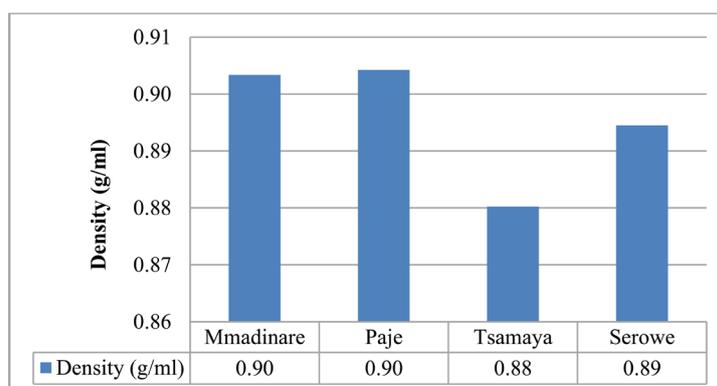


Figure 5. Density of *Jatropha curcas* biodiesel fuels from selected regions in Botswana.

3) Flash Point

Flash point is fuel property that determines the safety of a fuel during its handling and storage. **Figure 6** shows the results of flash points for *Jatropha curcas* biodiesel fuels produced using seeds from selected regions of Botswana.

The results show that biodiesel fuels produced from seeds collected from Mmadinare and Paje villages recorded relatively high flash point of about 177°C followed by Serowe with 157.17°C and Tsamaya with 146.50°C. The results for Mmadinare and Paje biodiesel fuels are consistent with the results reported by [25] [28] [29] who reported flashpoint values in the range of 160°C - 175°C for *Jatropha curcas* biodiesel. It is pertinent to mention that biodiesel standards such as EN14214 and ASTM D6751 put the minimum values for biodiesel flashpoint at 101°C and 93°C respectively. Results of this study therefore suggest that biodiesel produced using seeds collected from all regions under review satisfies both the two biodiesel standards.

Flashpoint of the biodiesel can be affected by its kinematic viscosity and density [30]. The authors alluded that high flashpoint of the fuel is influenced by higher kinematic viscosity and density of the fuel. Higher flashpoint values for biodiesel of seeds from Mmadinare and Paje villages may be influenced by relatively high kinematic viscosity and density of its biodiesel fuels as demonstrated by **Figure 4** and **Figure 5**. Likewise, biodiesel fuels of seeds from Serowe and Tsamaya had lower flashpoint values possibly due to high content of saturated fatty acids as depicted in **Table 1** in Section 3.1. Other studies have revealed that residual methanol from esterification process can have negative effects on the flashpoint temperature of biodiesel fuel [31]. These authors alluded that studies have shown that residual amount of methanol as low as 1% in the biodiesel fuel can lower its flashpoint from 170°C to less than 40°C. In the current study, biodiesel samples from *Jatropha curcas* seeds from Serowe and Tsamaya areas may have recorded lower flashpoint temperatures possibly due to residual methanol from the esterification process.

The results also demonstrated the interdependence of the flash point property of biodiesel fuel with other biodiesel properties such as kinematic viscosity and

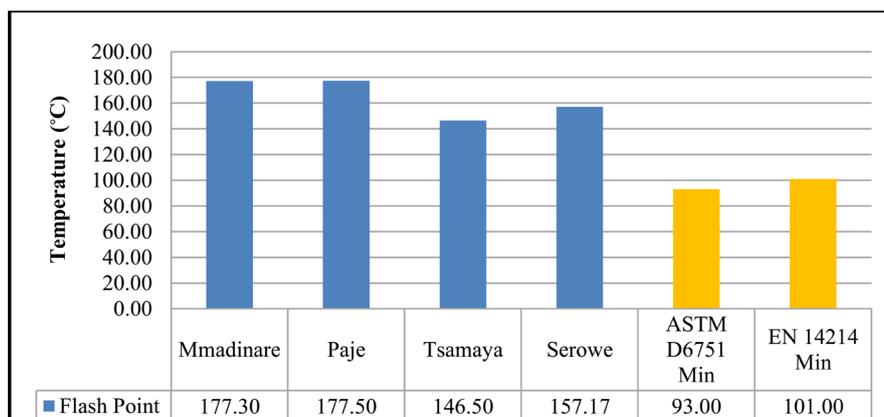


Figure 6. Flashpoints of *Jatropha curcas* biodiesel fuels from selected region in Botswana.

density. The investigation demonstrated that the higher the kinematic viscosity and density of the biodiesel fuel the higher the flash point. Furthermore, the investigation also demonstrated that high content of saturated fatty acids results in low value of flash point of the biodiesel fuel.

4) Cold Flow Properties

The low temperature properties which were investigated in the present study were Pour Point (PP) and Cloud Point (CP). The results for the two properties are presented in **Figure 7** and **Figure 8**.

Cloud point is defined as the highest temperature at which crystal growth of the biodiesel fuel is large enough (diameter $\geq 0.5 \mu\text{m}$) to be noticed while pour point is the lowest temperature at which the fluid will pour [32]. The two properties are important parameters to observe how the biodiesel fuel will perform under low temperature conditions. Hoekman *et al.* [23] and Wardana *et al.* [33] reported that both the cloud point and pour point of biodiesel fuels are mainly influenced by the presence of saturated fatty acids in biodiesel fuels. Extended exposure of the biodiesel fuel to temperatures at or lower than cloud point or pour point temperatures results in crystallation or biodiesel fluid starting to pour. Kumar *et al.* [34] reported that crystals in the fuel make it difficult for fuel to flow in engine, ultimately resulting in choking of engine and incomplete combustion.

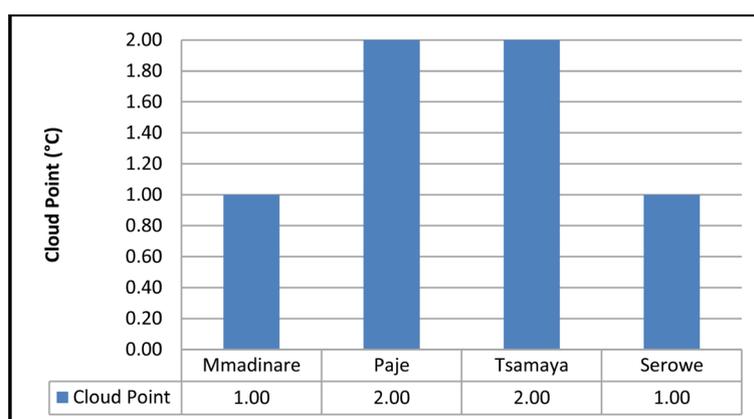


Figure 7. Cloud point of *Jatropha curcas* biodiesel fuels from selected region in Botswana.

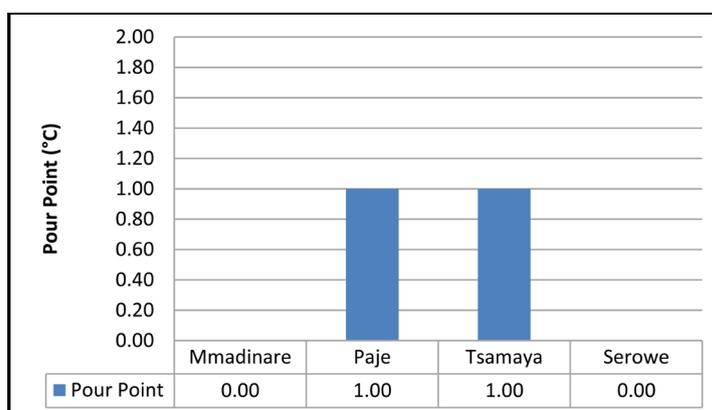


Figure 8. Pour point of *Jatropha curcas* biodiesel fuels from selected regions in Botswana.

The results as depicted in **Figure 7**, shows that biodiesel fuel of seeds from Tsamaya and Paje areas recorded equal values of cloud point of 2°C followed by biodiesel fuel of seeds from Mmadinare and Serowe regions at 1°C. Variation in the results may be attributed to the presence of saturated fatty acids present in the biodiesel [8]. The authors reported that feedstocks with highly saturated fatty acids produce biodiesel fuels with poor cold flow properties. Generally, biodiesel fuels with high concentration of saturated fatty acids tend to have high cloud point temperature. From the results in **Table 1** in Section 3.1, biodiesel fuel from seeds for Tsamaya region recorded relatively high-saturated fatty acids, thus 20% and 17.16% higher compared to saturated fatty acids recorded for Paje and Serowe regions. All these may justify why there is slight difference of cloud point for biodiesel fuels from Tsamaya and Paje regions. Other researches such as [1] [35] [33] reported cloud points in the range of 2°C to 5°C for *Jatropha curcas* biodiesel fuel.

Pour point is another important parameter similar to cloud point as it also focuses on cold properties of fuel. The results in **Figure 8** shows that the biodiesel fuels from Tsamaya and Paje regions recorded the highest pour point temperatures of 1°C. Biodiesel fuels from Mmadinare and Serowe regions recorded the pour points temperatures of 0°C. Like for cloud point, the slight difference in the results may be attributed to the presence of saturated fatty acids present in the biodiesel [8]. From the fatty acid composition **Table 1** in Section 3.1, the results shows that biodiesel fuels from Tsamaya and Paje regions recorded high degree of saturated fatty acids which may be the main reason for high pour point temperatures.

Other authors have reported pour point values in the range of -5°C to 5°C for different biodiesel feedstock including *Jatropha curcas* [36] [37] [38] [39]. The pour point results obtained in the current study are consistence with pour point values in other studies.

5) Energy Content

The study also investigated the energy content of the biodiesel fuel from *Jatropha curcas* seeds obtained from selected regions of Botswana and the results are depicted in **Figure 9**.

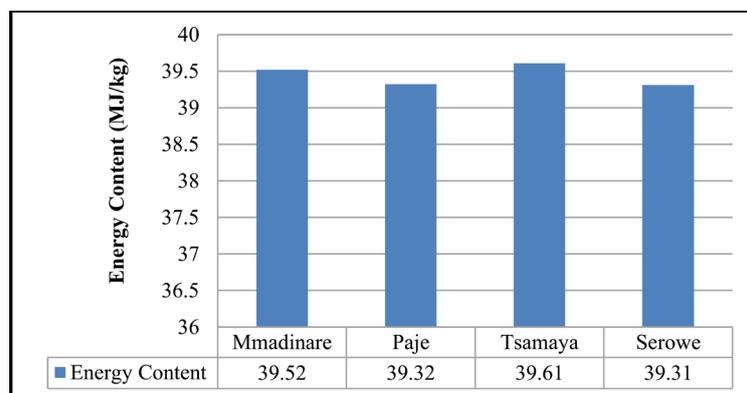


Figure 9. Energy content of *Jatropha curcas* biodiesel fuels from selected regions in Botswana.

All the energy content values of the biodiesel fuels are in the average of 39.00 MJ/kg, biodiesel fuel produced from *Jatropha curcas* seeds collected from Tsamaya had energy content of 39.61 MJ/kg followed by biodiesel fuel from Mmadinare at 39.52 MJ/kg. Paje and Serowe had the least at 39.32 MJ/kg and 39.31 MJ/kg respectively. The minimum and maximum percentage variation observed was 0.23% and 0.76% respectively which is quite insignificant. Therefore, this means that all the biodiesel fuels under review will probably have the same behaviour when it comes to certain engine performance parameter such as specific fuel consumption which are influenced by energy content of fuel. The obtained results fall within the same range of results reported in other studies such as [35] [40] [41] [42]. The authors reported energy content values in the range between 37 - 40 MJ/kg.

4. Conclusions

The main purpose of the present study was to investigate effects of fatty acids composition on fuel properties of *Jatropha curcas* biodiesel fuel. Relevant experiments were successfully carried out and the following conclusions were arrived at:

1) The physicochemical fuel properties of biodiesel investigated are influenced by the fatty acid composition. The experimentally determined kinematic viscosity, density and flashpoint values are influenced by the degree of unsaturation of the fatty acid mixture. Biodiesel of *Jatropha curcas* seeds obtained from Paje and Mmadinare regions had higher values of kinematic viscosity, density and flashpoint than other regions due to high proportions of unsaturated fatty acids.

2) Biodiesel of *Jatropha curcas* seeds obtained from Tsamaya and Paje regions had slightly higher cloud and pour point due to dominant presence of saturated fatty acids.

3) Energy content of all the biodiesel was averagely the same and this signifies that the energy content of the biodiesel investigated is independent of fatty acid composition.

4) Although some differences in values of physicochemical properties were observed in all the parameters investigated, all the biodiesel fuels were within the European (EN14214) and American (ASTM D6751) biodiesel standards in terms of kinematic viscosity, density and flashpoint. As for cold flow properties and energy content, the results were comparable to other studies.

It is therefore appropriate to conclude that fatty acids composition has great influence on fuel properties of *Jatropha curcas* biodiesel.

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Conflicts of Interest

The authors declare no conflicts of interest regarding the publication of this paper.

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