Evaluation of Anti-hyperlipidemic Effect of Aqueous Leaves Extract of *Moringa oleifera* in Alloxan Induced Diabetic Rats

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Authors’ contributions

This work was carried out in collaboration between all authors. Author TAO designed the study, performed the statistical analysis, wrote the protocol, and wrote the first draft of the manuscript. Authors SOB and TAA managed the literature searches. They also worked on the final draft of the manuscript. All authors read and approved the final manuscript.

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**ABSTRACT**

**Aims:** The purpose of this study is to evaluate the effect of aqueous leaf extract of *Moringa oleifera* (*Moringaceae*) on plasma glucose level, total cholesterol level, triglycerides (TG), high-density lipoprotein (HDL) and low-density lipoprotein (LDL) in male albino rats.

**Study Design:** Experimental method was adopted.

**Place and Duration of Study:** Adeleke University, Ede, Nigeria and University of Ibadan, Nigeria between September 2012 and January, 2013.

**Methodology:** Twenty-four male albino rats of the Wistar strain weighing between 160 and 200 g were used for this study. These were randomly assigned into 3 groups of 8 animals per group as normal control, diabetic control and diabetic treated with extract of *Moringa oleifera*. Diabetes was induced by 100 mg/kg of alloxan monohydrate. The control and the diabetic groups received distilled water while the diabetic treated group was administered 400 mg/kg body weight of aqueous leaf extract of *Moringa oleifera* for 28 days. At the end of the experiment, plasma glucose level, cholesterol, Triglycerides (TG),

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High Density Lipoprotein (HDL) and Low Density Lipoprotein (LDL) were determined in all the experimental animals after 12 hours fast.

**Results:** The result showed significant increases ($P<0.05$) in plasma cholesterol, TG and LDL level of the diabetic control group when compared with the normal control group while there were no significant differences in the *Moringa oleifera* -treated diabetic group and the normal control group. The HDL however was not different in all the three groups.

**Conclusion:** Oral administration of aqueous leaf extract of *Moringa oleifera* may reduce the plasma lipid imbalances associated with diabetes mellitus.

**Keywords:** *Moringa oleifera*; alloxan-induced diabetes; blood glucose; lipid imbalances.

1. **INTRODUCTION**

The consumption of a variety of local herbs and vegetables by man is believed to contribute significantly to the improvement of human health, in terms of prevention, and or cure of diseases because plants have long served as a useful and natural source of therapeutic agents [1].

*Moringa oleifera* (family *Moringaceae*) is commonly known as Drumstick tree. Most of the parts of the plant possess antimicrobial activity [2]. They are well known for their pharmacological actions too and are used for the traditional treatment of diabetes mellitus [3] hepatotoxicity [4], rheumatism, venomous bites and also for cardiac stimulation [5].

Diabetes mellitus (DM) is a group of metabolic disorders characterized by hyperglycemia resulting from a variable interaction of hereditary and environmental factors due to defects in insulin secretion, insulin action, or both [6]. Hyperglycemia resulting either due to defective production or action of insulin leads to a number of complications; cardiovascular, renal, neurological, ocular etc. [7]. The ailment may result in the development of further metabolic and anatomic disturbances among which is lipemia, hypercholesterolaemia, loss of weight, ketosis, arteriosclerosis, gangrene, pathologic changes in the eye, neuropathy, renal disease, and coma [8,9].

Dyslipidaemia is common in diabetes, as both insulin deficiency and insulin resistance affects enzymes and pathways of lipid metabolism [10]. Dyslipidaemia, as defined by the World Health Organization [11], is considered in circumstances where the fasting plasma triglyceride is between 150 - 400 mg/dL (1.7 to 4.5 mmol/L), total cholesterol (TC) > 200 mg/dL (>5.2 mmol/L), low-density lipoprotein cholesterol (LDL-C) > 135 mg/dL (>3.5 mmol/L), high-density lipoprotein cholesterol (HDL-C) < 35 mg/dL (<0.9 mmol/L) in men or <40 mg/dL (<1.0 mmol/L) in women, and a ratio of total cholesterol to HDL-cholesterol > 5 [12]. Characteristic abnormalities in lipids in type 2 diabetes mellitus include elevated triglycerides (TG) levels, decreased atheroprotective high density lipoprotein cholesterol (HDL-C) levels and increased levels of small dense LDL-C [13,14].

World Health Organisation (WHO) [15] has emphasized strongly on the rational use of traditional and natural indigenous medicines, for treating diabetes mellitus. Patients with diabetes mellitus have been treated orally by folklore with a variety of plant extracts [16]. More than 1200 plants species are used worldwide in diabetes phytotherapy, and experimental studies support the hypoglycaemic activity of a large number of these plants [17]. In addition to correction of blood glucose levels, several plants with hypoglycaemic
properties have potential in ameliorating lipid metabolism abnormalities of diabetes mellitus [18].

Therefore, the purpose of this study is to evaluate the effect of aqueous Moringa oleifera leaf extract on total cholesterol level, triglycerides (TG), high-density lipoprotein (HDL) and low-density lipoprotein (LDL) in male albino rats based on the reported activities of most hypoglycaemic plant agents at ameliorating lipid metabolism abnormalities [19].

2. MATERIALS AND METHODS

2.1 Preparation of Plant Material

Fresh leaves of M. oleifera (5 kg) were collected from Oyo town in Nigeria and were taken to the Botany Department, University of Ibadan, Nigeria, for authentication. A voucher specimen (FHI, 142807) has been submitted earlier in the Forestry Research Institute of Nigeria (FRIN) herbarium. The leaves were air-dried and reduced to powdered form; the powdered leaves were percolated in distilled water for 12 h and filtered; the filtrate was subsequently evaporated to dryness and yielded a 1.13kg dark green concentrate.

2.2 Animals

Twenty-four male Wistar albino rats weighing between 160 - 200 g were used for this study. They were maintained at standard laboratory condition and fed with commercial pellet diet (Bendel Feeds, Nigeria) and water ad libitum. The animals were subjected to natural photoperiod of 12 h light: dark cycle. All experimental protocols and handling were in compliance with the National Institutes of Health (NIH) publication No. 85-23 guidelines [20]. The animals were acclimatized to laboratory condition for one week before commencement of experiment.

2.2.1 Acute oral toxicity study of M. oleifera extract and fruit juice

Acute toxicity study was carried out according to Organisation for Economic Co-operation and Development (OECD) guideline 423 [21]. None of the rats showed observable signs of toxicity upon single administration of M. oleifera extract and fruit juice (2 g /kg, p.o.) on day one. Observations twice daily for 14 days also did not reveal any drug related observable changes. The study was repeated with another set of animals for 14 days and no signs of toxicity were observed.

2.2.2 Induction of diabetes

Diabetes was induced by a single intraperitoneal injection of 100 mg/kg of alloxan monohydrate obtained from Sigma Chemical Co. (St. Louis, MO, USA). Diabetes was confirmed by glucose oxidase method using the glucometer (One Basic, Inc.) after 72 h of alloxan injection, rats with plasma glucose level ≥ 200 mg/dl were separated and used as diabetic in this study.

2.3 Experimental Design

The rats were randomly distributed into three groups of eight rats each: Group A: normal and received distilled water – (Normal control)
Group B: diabetic and received distilled water— (Diabetic Control)

Group C: diabetic treated with 400 mg/kg aqueous extract of *M. oleifera* (Diabetic + Extract)

The rats were treated for 28 days. At the end of the experimental period, they were fasted for 12 h and blood was collected by cardiac puncture under light anesthesia. The blood was transferred into sample bottles containing EDTA for plasma collection. Plasma was collected from the blood samples after centrifuging at 3000 x g for 10 min using a bench centrifuge.

### 2.3.1 Blood glucose estimation

Blood glucose was estimated on 0, 7, 14, 21 and 28th day by Glucometer (One Basic, Inc.).

### 2.3.2 Determination of plasma lipid

Plasma triglycerides and total cholesterol levels were measured on the final day of experiment using enzymatic colorimetric diagnostic kits obtained from Randox Laboratories, UK, in which the GPO-PAP method of Trinider [22] was employed. Absorbance was measured at 500 nm. The phosphotungstate precipitation method of Richmond [23] as applied in Randox kit was used for the determination of HDL-cholesterol. The LDL-cholesterol was estimated using Friedewald [24] formula:

\[
\text{LDLc} = \text{Total cholesterol} - \text{HDLc} - \text{TG}/5
\]

Where; LDLc = LDL-cholesterol, HDLc = HDL-cholesterol and TG = triglycerides

### 2.4 Statistical Analyses

All results were expressed as mean ± SEM. Statistical analysis was carried out by one-way analysis of variance (ANOVA) and Duncan new multiple range test (DMRT). Differences in means were considered significant at \( P < 0.05 \) and \( P < 0.01 \). All analyses were performed using SPSS 15.

### 3. RESULTS

In acute oral toxicity studies *M. oleifera* extract did not show any mortality and toxic effects up to the dose of 2000 mg/ kg body weight. So one-fifth of the safe dose is used for the experiments.

Alloxan-induced diabetic rats exhibiting persistent hyperglycemia (Blood Glucose > 200 mg dL-1) were selected for assessing the effect of *M. oleifera* aqueous extract. Oral daily administered dose of 400 mg/ kg body weight of the extract led to significant decrease in hyperglycemia as shown in Fig. 1 and Table 1.
Fig. 1. Effect of aqueous extract of *M. oleifera* leaves on blood glucose in diabetic rats

Table 1. Effect of aqueous extract of *M. oleifera* leaves on blood glucose in diabetic rats

<table>
<thead>
<tr>
<th>Groups</th>
<th>0 Day</th>
<th>7 Day</th>
<th>14 Day</th>
<th>21 Day</th>
<th>28 Day</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal control</td>
<td>85 ± 1.22</td>
<td>84 ± 1.12</td>
<td>84 ± 0.09</td>
<td>83 ± 2.44</td>
<td>85 ± 1.02</td>
</tr>
<tr>
<td>Diabetic control</td>
<td>255 ± 2.80</td>
<td>285 ± 2.87</td>
<td>312 ± 2.66</td>
<td>324 ± 4.22</td>
<td>344 ± 2.55</td>
</tr>
<tr>
<td>Diabetic + extract</td>
<td>262 ± 2.11</td>
<td>190 ± 2.11*</td>
<td>153 ± 2.80*</td>
<td>121 ± 3.23**</td>
<td>102 ± 1.23*</td>
</tr>
</tbody>
</table>

* All values are expressed as mean ± SEM (n=8), *: P<0.05, **: P<0.01 as compared to diabetic control.

Table 2 shows the mean values of plasma glucose level, total plasma cholesterol, TG, HDL and LDL in the control, diabetic untreated and diabetic treated with *M. oleifera*.

Table 2. Effect of aqueous leaf extract of *M. oleifera* on lipid profile of alloxan induced diabetic rats

<table>
<thead>
<tr>
<th>Variable</th>
<th>Normal Control</th>
<th>Diabetic Control</th>
<th>Diabetic + Extract</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total cholesterol (mmol/L)</td>
<td>4.62±0.254</td>
<td>6.01±0.188*</td>
<td>4.77±0.356</td>
</tr>
<tr>
<td>Triglycerides (mmol/L)</td>
<td>0.97±0.102</td>
<td>1.70±0.065*</td>
<td>1.25±0.412 **</td>
</tr>
<tr>
<td>HDL (mmol/L)</td>
<td>1.54±0.212</td>
<td>1.25±0.198</td>
<td>1.60±0.320</td>
</tr>
<tr>
<td>LDL (mmol/L)</td>
<td>2.77±0.153</td>
<td>4.12±0.102*</td>
<td>3.43±0.226 **</td>
</tr>
</tbody>
</table>

* Values are expressed as mean ±SEM (n=8), *: P> 0.05 when compared to control, **significantly reduced when compared to diabetic control group.

The total cholesterol level was significantly increased (*P* >0.05) in the diabetic group when compared with the normal control group while there was no significant difference in the cholesterol level of the *M. oleifera*-treated diabetic group and the control group. The TG level was significantly increased (*P* >0.05) in the diabetic control group with no significant difference in the *M. oleifera* -treated diabetic group when both were compared with the
normal control group. Though there was no significant difference in the HDL level of the three groups, the diabetic control group had a significantly higher \( (P>0.05) \) LDL when compared with the normal control while there was no difference in the LDL level of the \emph{M. oleifera} -treated group and the normal control group.

4. DISCUSSION

Alloxan-induced hyperglycaemia has been described as a useful experimental model to study the activities of hypoglycemic agents [25]. The purpose of the present study was to assess the effect of the aqueous extract of \emph{M. oleifera} leaves on glycemic control, total cholesterol, triglycerides (TG) and low density lipoprotein (LDL) levels. Aqueous extract of \emph{M. oleifera} leaves normalizes the high blood glucose level in diabetic rats by the 28\textsuperscript{th} day of the experiment. Since good activity has been seen in severely diabetic rats with damaged islets, it is likely to be expected that the aqueous extract of leaves has some direct effect. We are suggesting the mechanisms of actions could be either by increasing the tissue utilization of glucose [7], by inhibiting hepatic gluconeogenesis or absorption of glucose into the muscles and adipose tissues [26].

The increase in the total cholesterol, TG and the LDL levels of diabetic rats observed in this study are in accordance with earlier report in diabetic subject [27]. Diabetes-induce hyperlipidemia has been attributed to excess mobilization of fat from the adipose due to underutilization of glucose [28].

The significant reduction in TG, LDL and cholesterol levels of the diabetic treated rats when compared with the diabetic control rats supports the findings of Coon and Ernst [18] who stated that most hypoglycaemic plants have potentials of ameliorating diabetic lipid metabolism anomalies. This cholesterol lowering effect is supported by the earlier work of Oyedepo and Babarinde [29] who used \emph{Citrus maxima} and Iweala and Obidoa [30] using leaves of \emph{Ocimum gratissimum} as supplementary diet for six months to achieve the same effect. However this study is contrary to the report by Owoyele et al. [31] observing a discrepancy due to administration length of \emph{Ocimum gratissimum} extract.

This hypolipidemic effect of \emph{M. oleifera} could be related to its chemical composition, which shows the presence of alkaloids, flavonoids, saponin and cardiac glycosides. All these components are known to reduce serum lipid level in animals [32, 33]. Saponins may lower cholesterol by binding with cholesterol in the intestinal lumen, preventing its absorption, and/or by binding with bile acids, causing a reduction in the enterohepatic circulation of bile acids and increase in its fecal excretion [34]. The increased bile acid excretion is offset by enhanced bile acid synthesis from cholesterol in the liver and consequent lowering of the plasma cholesterol [13].

Although the precise mechanism by which the leave extract exerts it hypolipidemic effect is not clearly known nor studied, it could not be excluded that the control of glycaemia is a contributing mediator since control of glycaemia is a major determinant of total cholesterol and triglyceride levels [35]. Accordingly, the evolution of glycaemia was parallel to lipid parameters in both the normal/control and the diabetic rats of this study.
5. CONCLUSION

The present study reveals that aqueous extract of *M. oleifera* leaves has significant hypoglycemic and anti-diabetic potential. From this study, it could be concluded that prolonged oral administration of aqueous leaf extract of *M. oleifera* may reduce the plasma lipid imbalances associated with diabetes mellitus which support its traditional use in the treatment of diabetes and cardiovascular diseases. A daily intake of this plant may help to prevent hyperglycemia and hyperlipidemia. However, the precise mechanisms and sites of action require further studies. *Moringa oleifera* may play important role for inhibition of α-glucosidase, pancreatic cholesterol esterase activity, as well as bile acid binding and inhibiting the formation of cholesterol micellization. However, further studies are needed to clarify this important hypothesis.

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COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES


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