Antiamylase Potential of *Telfairia occidentalis* Leaves from Cameroon and Effect of Their Dietary Supplementation on Fasting Blood Glucose in Wistar Rats

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Abstract

The study of edible plants, especially in developing countries, might provide more affordable means for the management of diabetes. *Telfairia occidentalis* is one of the plants whose leaves are commonly consumed in Cameroon. This work hereby studied the antiamylasic potential of its aqueous leaves extract and the effect of its dietary supplementation on fasting blood glucose in *Wistar* rats. An aqueous extract (1:6) was prepared from shed-dried *T. occidentalis* leaves by maceration. Its antiamylase activity was evaluated *in vitro* and a phytochemical screening was realized. Its acute toxicity and its effect on an oral glucose tolerance test (OGTT) were evaluated in rats. The effect of *T. occidentalis* leaves dietary supplementation (10%) on fasting blood glucose was studied for 28 days in rats fed with carbohydrate enriched diet, using Glibenclamide (0.3 mg/kg body weight) as reference hypoglycemic drug. Results showed that there was total inhibition of α-amylase activity *in vitro* by *T. occidentalis* aqueous leaves extract at 0.075 mg/ml. The presence of tannins, flavonoids and anthocyanins was revealed by the phytochemical screening. No sign of toxicity was observed in rats after an oral administration of the extract at 2000 mg/kg body weight. The extract significantly hindered a rise in blood glucose at 400 mg/kg body weight during an oral glucose tolerance test. Dietary supplementation with *T. occidentalis* leaves caused a significant decrease (p < 0.05) in fasting blood glucose as compared to the positive control. *Telfairia occidentalis* leaves and their aqueous extract could be used in the management of hyperglycemia and diabetes.

Keywords

*Telfairia occidentalis*, Hyperglycemia, Antiamylase Potential

1. Introduction

Diabetes mellitus is a chronic disease which arises when there is a failure in pancreatic insulin secretion or when

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the body can no more use insulin properly. Hyperglycemia is a frequent effect of poorly managed diabetes which can cause serious damage to the body with time, particularly to nerves and blood vessels [1]. Diabetes affects vital organs in the body and reduces life expectancy [2]. Worldwide, 382 million people were diabetic of who 46% were not diagnosed [3] and more than 80% of deaths caused by diabetes occur in less developed countries [4]. Actual treatments of diabetes mainly aim at reducing hyperglycemia and generally include a modification in lifestyle (dietetics and physical activity) and antidiabetic synthetic drugs [5]. Due to the side effects and the cost of synthetic drugs, research is being made on natural products in the course of finding new alternatives for diabetes management. The study of edible plants might therefore constitute a more affordable mean for prevention or treatment. *Telfairia occidentalis* Hook F (family *Cucurbitaceae*) is an herbaceous creeping plant cultivated for its edible seeds and leaves [6]. Depending on the age, ecologic conditions and cultural practices, its leaves composition can vary [7]. In Cameroon, it is locally known as “okoribon” and is found in the preparation of various dishes. This work was designed to study the antiamylasic potential of *Telfairia occidentalis* aqueous leaves extract and the effect of their dietary supplementation on fasting blood glucose in Wistar rats.

2. Materials and Methods

2.1. Collection of Plant Material and Extract Preparation

*Telfairia occidentalis* was collected from local markets in Douala and was taken to the National Herbarium in Yaounde for identification. The leaves were shed-dried for 1 week, ground and stored at room temperature. About 250 g of powdered leaves were soaked in 1500 ml of distilled water in a covered plastic container, allowed to macerate for 48 hours in the dark at room temperature and filtered with a filter paper. The filtrate was dried in an oven at 45˚C for 4 days to obtain concentrated extract which was dissolved accordingly in distilled water for various experimentations.

2.2. *In Vitro* α-Amylase Inhibition Assay

The α-amylase inhibitory activity was determined *in vitro* [8]. A mixture of 20 µl of enzyme (30 µg/ml), 1380 µl of tris-HCl buffer pH 6.9 and 100 µl of aqueous extract aliquot was pre incubated at 30˚C for 20 minutes. Thereafter, the reaction was initiated by adding 100 µl of starch solution (1%). After incubating for 20 minutes, the reaction was stopped by adding 2 ml of acidified iodine and the absorbance was read at 580 nm. Various extract concentrations were used. The α-amylase inhibitory activity was expressed as percentage inhibition.

2.3. Phytochemical Screening

A phytochemical screening was realized on *T. occidentalis* aqueous leaves extract to determine the presence of tannins [9], flavonoids [10] and anthocyanines [11].

2.4. Experimental Animals

Male and female Wistar rats aged of 3 months at least were bought from the Laboratory of Animal Biology of Douala University. They were kept in metabolic cages under a 12 hours dark/light cycle at room temperature for 2 weeks before any experimentation. They received standard laboratory chow and water *ad libitum*.

2.5. Acute Toxicity Assay

The acute toxicity of *T. occidentalis* aqueous leaves extract was evaluated according to the Organization for Economic Cooperation and Development, guideline 425 [12] with the limit test dose at 2000 mg/kg body weight. Female rats were divided into 2 groups of 5 each: a test group that received a single oral administration of *T. occidentalis* aqueous leaves extract at 2000 mg/kg body weight, and a control group that received distilled water. Signs of toxicity were monitored during 14 days after which subjects were humanely sacrificed and their blood collected. The commercial kit IMNESCO was used for determination of AST (Aspartate transaminase), ALT (Alanine transaminase), and total plasmatic protein [13] were determined.

2.6. Effect of *T. occidentalis* Aqueous Leaves Extract on Oral Glucose Tolerance Test

It was evaluated during an OGTT. Male rats were divided into 3 groups of 4 each. After 12 hours of overnight
fasting, their glycemia was measured using SD CodeFree glucometer. Each group received two oral administra-
tions: the positive control received 1 ml of distilled water followed by 1 ml of glucose solution (2 g/kg), the test
group received 1 ml of *T. occidentalis* aqueous leaves extract (400 mg/kg) followed by 1 ml of glucose solution
(2 g/kg) and the negative control received 1 ml of distilled water twice. Glycemia was then measured for all
groups at 30 minutes intervals during 2 hours.

### 2.7. Effect of *T. occidentalis* Leaves Dietary Supplementation on Fasting Blood Glucose

Four groups of 4 male rats each (positive control, test group, reference group and negative control) were used.
After 12 hours of overnight fasting, their glycemia was measured. During 28 days, all groups except the nega-
tive control, received 65% sucrose enriched food, 10% glucose in drinking water and oral administration of
fructose (1 g/kg). For test group, the food was supplemented with 10% of *T. occidentalis* powdered leaves, and
Glibenclamide (0.3 mg/kg) was administered orally to the reference group. For all animals, fasting glycemia was
measured on days 9, 18 and 28.

### 2.8. Statistical Analyses

STATGRAPHICS CENTURION XV.II and SPPS 17 were used for statistical analyses. Variables are presented
as mean ± standard deviation. Data were analyzed by the one way Analyses of variance test (ANOVA) followed
by the least significant difference (LSD) post Hoc test. Values of p < 0.05 were considered significant.

### 3. Results

#### 3.1. *In Vitro* α-Amylase Inhibitory Effect of *T. occidentalis* Aqueous Leaves Extract

*T. occidentalis* aqueous leaves extract significantly (p < 0.05) inhibited α-amylase activity at low concentration.
The enzyme’s activity was totally inhibited by the extract at 0.075 mg/ml as shown in Figure 1.

#### 3.2. Phytochemical Screening

The presence of flavonoids, tannins, and anthocyanins were revealed from the qualitative analyses realized on
the aqueous leaves extract. Flavonoids were revealed by the appearance of a yellow colour, tannins by a bluish
green precipitate and anthocyanins by an orange-red colour.

![Figure 1. In vitro α-amylase activity at various concentrations of *T. occidentalis* aqueous leaves extract.](image-url)
3.3. Acute Toxicity Assay

No death and no particular signs of toxicity were recorded after 14 days observation. The values obtained for AST, ALT and total plasmatic proteins for test group were not different (p > 0.05) from control group (Table 1).

3.4. Effect of T. occidentalis Aqueous Leaves Extract on Oral Glucose Tolerance Test

The extract significantly inhibited a rise in blood glucose during the oral glucose tolerance test as shown in Figure 2. Thirty minutes after the beginning of the experiment, glycemia was significantly high in positive control and test groups as compared to the negative control. At 60 minutes, it was greater (p < 0.05) for positive control compared to test group. Furthermore, in test group, glycemia at 90 minutes and 120 minutes are significantly low compared to 30 minutes.

3.5. Effect of T. occidentalis Leaves Dietary Supplementation on Fasting Blood Glucose

Rats receiving only the carbohydrate enriched diet (sucrose, glucose and fructose) showed a significant increase in fasting glycemia from the 1st to the last day. Their mean glycemia was 102.83 ± 12.4 mg/dl, thus corresponding to an impaired fasting blood glucose [14]. Dietary supplementation with 10% T. occidentalis leaves significantly reduced fasting blood glucose from the 9th to the last day of experiment (Table 2).

4. Discussion

Plants with antidiabetic potential exhibit various action mechanisms like enzyme inhibition [15]. Alpha amylase is one of the key enzymes involved in breaking down starch into absorbable glucose molecules [16]. Inhibiting this enzyme helps in reducing post prandial glycemia by reducing the speed of glucose absorption [17]. T. occidentalis aqueous leaves extract showed an anti α-amylase activity at low concentration. This suggests the presence of compounds capable of inhibiting the enzyme. However, further investigation may help to find more precise mechanisms for this enzyme inhibiting activity.

Pharmacological properties of plants are generally attributed to their phytochemical compounds. A number of these like tannins, flavonoids and anthocyanins are reported to be antidiabetic [18]. Their presence was revealed in the extract as reported in a previous study [19]. They might be involved in the antiamylase activity of the aqueous extract, especially tannins which were reported to have antienzyme activity [20]. Transaminases (ALT, AST) and plasmatic proteins are some biochemical markers to evaluate hepatic, cardiac and muscular functions. These vital functions might be affected by a toxic effect and should therefore be verified during a toxicity assay [21]. In this work, there was no significant difference in the values obtained suggesting that

<table>
<thead>
<tr>
<th>Groups</th>
<th>Biochemical parameters</th>
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<tbody>
<tr>
<td></td>
<td>Total plasmatic protein (mg/ml)</td>
<td>ALT(U/l)</td>
</tr>
<tr>
<td>Control</td>
<td>1.20 ± 0.2a</td>
<td>2.6 ± 0.5a</td>
</tr>
<tr>
<td>Test</td>
<td>1.30 ± 0.2a</td>
<td>3.2 ± 0.8a</td>
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Values are expressed as mean ± SD, n = 5. The data were analyzed by the ANOVA (test p < 0.05). Values affected by different letters in the same column are significantly different.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Fasting blood glucose (mg/dl)</th>
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<tbody>
<tr>
<td></td>
<td>Day 1</td>
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<tr>
<td>Positive control</td>
<td>88.58 ± 4.40a</td>
</tr>
<tr>
<td>Test</td>
<td>88.58 ± 4.25a</td>
</tr>
<tr>
<td>Reference</td>
<td>95.00 ± 17.12a</td>
</tr>
<tr>
<td>Negative control</td>
<td>91.25 ± 9.18a</td>
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</table>

Values are expressed as mean ± SD, n = 4. The data were analyzed by the ANOVA (test p < 0.05). Values affected by different letters in the same column are significantly different.
Figure 2. Effect 400 mg/kg of *T. occidentalis* aqueous leaves extract on glycemia during an OGTT (AE: aqueous extract; * significant difference compared to two other groups; ++ significant compared to 30 minutes; +++ significant compared to 30, 60 and 90 minutes).

*T. occidentalis* aqueous leaves extract was non toxic at 2000 mg/kg of body weight. This result is similar to the one obtained by other authors [22] who found that aqueous extract of *T. occidentalis* leaves was not toxic even at high doses. Furthermore, the extract significantly hindered a rise in blood glucose level during an OGTT as shown in Figure 2. This result suggests an antihyperglycemic activity as well as a capacity to improve glucose tolerance.

Dietary supplementation with *T. occidentalis* leaves significantly reduced fasting blood glucose in test group as compared to the positive control (Table 2). Moreover, it significantly prevented the carbohydrate enriched diet from causing a rise in fasting blood glucose throughout the experimental period. These reflect the antihyperglycemic activity of *T. occidentalis* leaves. The presence of antidiabetic phytochemicals could account for this result. It could also be substantiated by the capacity of the extract to inhibit α-amylase activity and to slower blood glucose absorption, thereby improving glucose tolerance and post prandial glycemia. It is contrary to findings obtained by other researchers [6] who reported that *T. occidentalis* leaves significantly increased blood glucose level in rats. It might be attributed to difference in plant material preparation and treatment of animals.

5. Conclusion

The results obtained from the *in vitro* and *in vivo* experimentations reveal that *T. occidentalis* leaves from Cameroon possess antihyperglycemic properties. They might be attributed to the presence of antidiabetic photo-chemical but further investigations are needed to find out the bioactive molecules and mechanisms involved. These leaves and their aqueous extract could thus be used in the management of hyperglycemia and diabetes.

**References**


