Hypoglycemic effects of *Lagenaria siceraria*, *Cynodon dactylon* and *Stevia rebaudiana* extracts

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**Abstract**

Introduction: The aim of the current analysis was to judge the hypoglycemic action of the phyto-extracts of *Lagenaria siceraria*, *Cynodon dactylon* and *Stevia rebaudiana* using suitable in vitro approaches.

Methods: The hypoglycemic activity of the phyto-material extracts was evaluated by employing various in-vitro methods namely glucose diffusion, amylolysis kinetics and glucose adsorption capacity.

Results: The extracts of *L. siceraria*, *C. dactylon* and *S. rebaudiana* exhibited glucose dialysis retardation indices (GDRI) of 48.14%, 37.03% and 29.62%, respectively at 60 minutes which were reduced to 15.78%, 10.52% and 18.42%, respectively at 120 minutes. All the plant extracts used in the study adsorbed glucose and their adsorptions markedly enhanced with increase in sugar concentration.

Conclusion: From the outcome of the assay it can be concluded that the extracts of *L. siceraria*, *C. dactylon* and *S. rebaudiana* have hypoglycemic activity as observed in various in-vitro assays. However, the beneficial actions require to be verified by adopting various in vivo techniques along with clinical trials for their efficient use as potential remedial moiety.

Nowadays, there is an increased interest in herbal drugs and remedies because of the toxic side effects associated with various oral hypoglycemic agents used in the treatment of diabetes mellitus. It was therefore decided to study the hypoglycemic effect of extracts of some indigenous plants known for their antidiabetic activities (1).

*Lagenaria siceraria* with common name; bottle-gourd, (*Cucurbitaceae*) is a creeper, with dumbbell pattern fruits. It is widely available throughout India and its flowering top parts and fruits are usually involved in diet as vegetables (4). Moreover, it is used as medicine for its promoting diuresis, tonic and heart tonic properties. Also, different pharmacological activities such as antihepatotoxic, analgesic, anti-inflammatory, lipid lowering, anti-hyperglycemic, immuno-modulatory and anti-

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oxidant activities of the phytoplant extract of *L. siceraria* have already been reported (5,6). Moreover further studies have also been performed on its whole fruits with its seeds. However, the pharmacology of the flowering parts of *L. siceraria* has not been much evaluated. In various parts of the world, the said plant has also been used conventionally as a remedy for metabolic disorders (7-9).

*Cynodon dactylon* belongs to family *Poaceae*, is a weed and has been reported to possess an array of medicinal properties such as antimicrobial, anti-diabetic, anti-inflammatory, antisynderteny and other activities (10). The principal components of this plant are flavonoids and sterols (11). It has been traditionally used as an antidiabetic agent in the Ayurveda medicine. (12).

*Stevia rebaudiana* Bertoni which is a seasonal sweet herb, belongs to Asteraceae, is used as a noncaloric natural sweet in tasting and found in most parts of the universe for its extreme sweetness. The plant is known to produce diterpene glycoside containing stevioside, rebaudioside, etc. Also the leaves are used as a natural low calorie sugar source as a good substitute for table sugar (13). Nowadays, *Stevia rebaudiana* is commercially cultivated in various Asian countries like India, Pakistan, China, Bangla, Bhutan, Sri Lanka, and Malaysia (14).

The plant-based remedies have already boosted in worldwide because of their assured safety and effectiveness (15). In the forthcoming years, there will be an increased interest in the use of medicinal plants in the developing countries due to their greater safety and lesser side effect(s) as compared to synthetic drugs (16). Thus, the current approach was considered to access the hypoglycemic activity of the selected plant extracts namely *L. siceraria*, *C. dactylon* and *S. rebaudiana* using suitable *in vitro* approaches.

Materials and Methods

Chemicals and reagents

Glucose oxidase peroxidase analysis kit was procured from Pathozyme Diagnostics Pvt, Ltd, India. The dialysis tubing (bags) (12000 MW cutoff) were acquired from Himedia Laboratories, 23 Vadhani Industrial Estate, LBS Rd, Mumbai, Maharashtra. The reagents and chemicals used during the project possessed high yield and pure AR grade.

Plant material

The plant material used in the study was collected from the surrounding region of Kasegaon village, Talukha – Valva, District - Sangli, (MS), India - 415404. The collected plant material was authenticated by Prof Dr. G.G. Potdar sir, Department of Botany, Yashwantrao Chavan College of Science, Karad, (MS). The authentications number for *L. siceraria* (DSR/105), *C. dactylon* (DSR/106) and *S. rebaudiana* (DSR/107) of the herbarium issued.

Preparation of plant extracts

The leaves of *S. rebaudiana* (Bert.) and *C. dactylon* were separately extracted with heated water (70°C) using a mechanical shaking device (24 hours). It was subjected to filtration and finally freeze dried. Fruits of *L. siceraria* were smashed in water and the juice was prepared, filtered and freeze dried. The extracts were properly stored in air tight containers until its use.

Evaluation of *in vitro* model for hypoglycemic activity of phyto-extracts

Effect of plant extracts on glucose diffusion

One percent w/v samples of plant extracts and 20 mM strength of glucose solutions (25 mL) were dialyzed by using dialysis tubing converted into bags adjacent to 200 mL of water (double distilled) at a temperature of 37°C using a shaker water bath. At intervals of 30, 60, 120 and 180 min the content of remaining glucose in the dialysate was assessed by employing glucose oxidase peroxidase diagnostic analysis kit. Solution without any sample extract served as control. Glucose dialysis retardation indices (GDRI) was calculated as follow (17).

\[
GDRI = \frac{100 - \frac{\text{Glucose content with addition of sample (mg/dL)}}{\text{Glucose content of the control (mg/dL)}}}{\times 100}
\]

Amylolysis kinetics study of selected plant extracts

The plant extracts (1%), alpha amylase (0.4%) and 25 mL of 4% (w/v) starch solution were dialyzed by using dialysis tubing against 200 mL of double distilled water at 37°C temperature (pH 7.0) in a orbital water bath equipped with shaker. At interval of 30, 60, 120 and 180 minutes, the remaining glucose concentration in the dialysate was assessed. A solution without any sample extract served control (18).

Determination of glucose adsorption capacity

Powder extracts (1% w/v) were incorporated to 25 mL solution of glucose having different strength (5, 10, 20, 50 and 100 mM). Then, the resulting liquid was homogenously shaked, thereafter incubated in an orbital shaker with shaking at 37°C (6 hours) and centrifuged (4000×g for 20 minutes). The sugar concentration in the supernatant was calculated as follow.

\[
Glucose Bound (mM) = \frac{G1 - G6}{4000 \times 20} \times \frac{Volume of solution}{Weight of the sample}
\]

Statistical analysis

All trials were performed in three parallel ways and the
resulting data were accessed using ANOVA and then compared for Tukey’s significant differences. The values were considered significant at $P \leq 0.05$. The graphics were represented using the Graphpad Prism 6 software.

Results
Effect of plant extracts on glucose diffusion
The results of the selected samples of phyto-extracts for glucose diffusion are shown in Table 1. During analysis, the transport of glucose through membrane of dialysis was determined in intervals of 30 minutes for 180 minutes. The samples showed significant inhibitory action on the mobility of glucose in the surrounding solution on the membrane of dialysis, compared with control.

Effect on in vitro amylosis kinetics
Table 2 shows the extent of glucose diffusion and GDRI observed by the inclusion of phyto-extracts in system of starch-α-amylase enzyme fiber. The rate of diffusion of glucose in the samples containing systems decreased with significant effect on the level of $P \leq 0.05$ for each period of time compared to the positive control. In addition, the extent of glucose diffusion was zero for 60 minutes in the plant extract material.

Glucose adsorption capacity
Figure 1 depicts the adsorption capacity of glucose for the extracts. The adsorption capacity of the phyto-extracts was proportional (directly) to concentration of glucose (M). Greater concentration of glucose was also adsorbed in samples with elevating glucose concentration levels. There were no significant differences among L. siceraria, C. dactylon, S. rebaudiana in the adsorption capacity.

Discussion
Diabetes mellitus is a debilitating hormone disorder in which strict glycemic control and prevention of associated complications are essential (19). According to the review of the literature, it is known that the leaves S. rebaudiana (Bert.), C. dactylon and fruits of Lagenaria siceraria have antidiabetic activities as mentioned earlier in the introduction. Therefore, an effort was undertaken to explore antidiabetic action of S. rebaudiana, C. dactylon and fruits of L. siceraria. This is the first attempt to determine the hypoglycemic activity of the leaves S. rebaudiana (Bert.), C. dactylon extracts and fruit extract of L. siceraria using different in vitro models, namely glucose uptake in yeast cells, kinetics of amylosysis, glucose adsorption capacity and glucose diffusion. Furthermore, our results revealed a greater adsorption capacity for the leaves extracts of S. rebaudiana, C. dactylon and fruit extract of L. siceraria attributable to their constituents. It has also been previously reported that the insoluble and soluble constituents together with fibers of different origins may contribute to the adsorption of glucose. It has been observed that leaf extracts of S. rebaudiana (Bert.) and C. dactylon as well as fruit extract of L. siceraria could also effectively adsorb 5 mM glucose concentration, thus declining the mass of glucose accessible for passage through the intestinal lumen (IL) of Gastro Intestinal Tract GIT. The activities of the leaves extracts of S. rebaudiana and C. dactylon, and the fruit extract of L. siceraria may be responsible for reducing the post-prandial hyperglycemia. The said observations are in accordance to the findings by (20) for fiber-rich fractions (insoluble) extracted from A. carambola.

GDRI remains to be main in-vitro model to forecast the

Table 1. Effect of selected samples on glucose diffusion and glucose dialysis retardation index

<table>
<thead>
<tr>
<th>Particulars</th>
<th>Glucose content in dialysate (mM)</th>
<th>30 min</th>
<th>60 min</th>
<th>90 min</th>
<th>120 min</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td></td>
<td>0.79±0.01</td>
<td>1.27±0.01</td>
<td>1.77±0.01</td>
<td>1.94±0.01</td>
</tr>
<tr>
<td>Lagenaria siceraria</td>
<td></td>
<td>0.51±0.01 (35.45)</td>
<td>1.06±0.01 (16.53)</td>
<td>1.53±0.01 (13.55)</td>
<td>1.68±0.01 (13.40)</td>
</tr>
<tr>
<td>Cynodon dactylon</td>
<td></td>
<td>0.48±0.01 (39.24)</td>
<td>1.14±0.01 (10.23)</td>
<td>1.63±0.01 (7.90)</td>
<td>1.81±0.01 (6.70)</td>
</tr>
<tr>
<td>Stevia rebaudiana</td>
<td></td>
<td>0.45±0.01 (43.03)</td>
<td>1.20±0.01 (7.87)</td>
<td>1.65±0.01 (6.77)</td>
<td>1.83±0.01 (5.67)</td>
</tr>
</tbody>
</table>

The values in parentheses point out the glucose dialysis index (GDRI). Averages (n = 3) bearing the letters in depicted table exhibit significant differences compared with control at level $P \leq 0.05$ using two-way ANOVA method.

Table 2. Effect of phyto-extracts on in vitro amylosis kinetics

<table>
<thead>
<tr>
<th>Plant sample</th>
<th>Glucose content in dialysate (mM)</th>
<th>30 min</th>
<th>60 min</th>
<th>90 min</th>
<th>120min</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td></td>
<td>0.0</td>
<td>0.27±0.01</td>
<td>0.29±0.01</td>
<td>0.38±0.01</td>
</tr>
<tr>
<td>Lagenaria siceraria</td>
<td></td>
<td>0.0 (100 )</td>
<td>0.14±0.01 (48.14)</td>
<td>0.22±0.01 (24.13)</td>
<td>0.32±0.01 (15.78)</td>
</tr>
<tr>
<td>Cynodon dactylon</td>
<td></td>
<td>0.0 (100 )</td>
<td>0.17±0.01 (37.03)</td>
<td>0.24±0.01 (17.24)</td>
<td>0.34±0.01 (10.52)</td>
</tr>
<tr>
<td>Stevia rebaudiana</td>
<td></td>
<td>0.0 (100 )</td>
<td>0.19±0.01 (29.62)</td>
<td>0.21±0.01 (27.58)</td>
<td>0.31±0.01 (18.42)</td>
</tr>
</tbody>
</table>

The values in parentheses point out the glucose dialysis index (GDRI). Averages (n = 3) bearing the letters in depicted table exhibit significant differences compared with control at level $P \leq 0.05$ using two-way ANOVA method.
The current investigation revealed hypoglycemic properties of the phyto-extracts of *Lagenaria siceraria, Cynodon dactylon, Stevia rebaudiana* as evaluated by different in vitro methods. The observed outcomes should be verified by accessing different in vivo techniques along with clinical trials which may credit them to use as therapeutic agents in the treatment of diabetes disorder.

**Conflict of interests**

The authors declared no competing interests.

**Ethical considerations**

Ethical issues have been observed by the authors.

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