In Vitro Studies on Alpha Amylase Inhibitory Activity of Some Indigenous Plants

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Abstract

Diabetes mellitus is a chronic metabolic disorder characterized by high blood glucose level caused due to deficiency of insulin secretion or insulin action. It includes a group of metabolic diseases characterized by hyperglycemia, in which blood sugar levels are elevated either because the pancreas do not produce enough insulin or cells do not respond to the produced insulin. Thus, a therapeutic approach to treat diabetes is to decrease postprandial hyperglycemia. It can be achieved by the inhibition of carbohydrate hydrolyzing enzymes like alpha amylase and alpha glucosidase. Such inhibitors which find application in the clinical practice for management of diabetes are known to be associated with various gastrointestinal side effects. Therefore, it is the need of time to identify and explore the amylase inhibitors from natural sources having fewer side effects.

In the present study, aqueous extracts of selected plants namely Albizia lebbeck, Berberis aristata, Mucuna pruriens, Myristica fragrans, Catharanthus roseus, and Caesalpinia bonducella which are used in the Ayurvedic traditional system of medicine to treat diabetes were tested for their inhibitory effect on α-amylase. The results revealed that the extract of A. lebbeck at a concentration of 1000µg/ml exhibited an inhibition of 70.91 %, whereas the extracts of B. aristata and M. pruriens showed an inhibition of 65.24 % and 62.96 % respectively at a similar concentration. The extracts of C. roseus, M. fragrans and C. bonducella exhibited a relatively less inhibitory response for the α-amylase enzyme. The results of the work thus clearly indicate the potential of the studied extracts to manage hyperglycemia.

Keywords: Antidiabetic; α-Amylase; Inhibitory effects; Indigenous plants

Introduction

Diabetes mellitus is a complex chronic metabolic disorder that has become a major source of ill health worldwide. It is characterized by hyperglycemia and disturbances of carbohydrate, protein and fat metabolisms, secondary to an absolute or relative lack of the hormone insulin [1-4]. Currently, the treatments of diabetes, in addition to insulin supplement includes many oral hypoglycemic agents such as sulfonylureas, biguanides, thiazolidines, D-phenylalanine derivatives, meglitinides and α-glucosidase inhibitors along with appropriate diet and exercise. However, none of them can be considered to be an ideal one, due to their toxic side effects and sometimes reduction in response after long term use [5]. One of the therapeutic approaches useful for treatment of diabetes is to decrease the post-prandial hyperglycemia. It can be achieved by retarding the absorption of glucose through the inhibition of the carbohydrate hydrolyzing enzymes in the digestive tract. α-amylase is an enzyme responsible for the breakdown of oligo and/or disaccharide to monosaccharides. Its inhibitors prolong the overall carbohydrate digestion time causing a marked decrease in the rate of glucose absorption thereby blunting the post prandial plasma glucose rise [6]. Acarbose, Miglitol and Voglibose are the typical examples of inhibitors used in the clinical practice [7]. However, the major drawbacks of these drugs include various gastrointestinal side effects such as abdominal pain, flatulence and diarrhea in the patients [8,9]. Thus, efforts are needed to identify and explore the amylase inhibitors from natural sources having fewer side effects.

Ayurveda and other ancient Indian literature have mentioned several herbal plants and their preparations useful in the treatment of various diseases and disorders [10-12]. Ethano pharmacological surveys have shown that more than 1200 plants have been mentioned in traditional medicine for their alleged hypoglycemic activity [13]. Numerous plants and their products have been widely prescribed and used for diabetic treatment all around the world with less known mechanistic basis of their functioning. The present study was therefore undertaken to make a comparative study for the ability of the selected plants to inhibit α-amylase activity. The following plants were selected for the study namely, Albizia lebbeck (AL), Berberis aristata (BA), Mucuna pruriens (MP), Myristica fragrans (MF), Catharanthus roseus (CR), and Caesalpinia bonducella (CB).

Albizia lebbeck (Leguminosae) is a deciduous tree with medicinal uses. Barks and seeds are astringent, and are given in piles...
and diarrhea. Ethanolic extract of pods possesses antiprotozoal, hypoglycemic and anticancer properties [14,15].

_Berberis aristata_ (Berberidaceae) have been studied for febrifugal, hypotensive, immunostimulating, anti-inflammatory, antidiabetic, antimicrobial, antiprotozoal, anticholinergic, antiarrhythmic, antiplasmodial, hypolipidemic, anti-granuloma and anti-hemolytic activity [3,16,17].

_Mucuna pruriens_ (Fabaceae) is one of the popular drugs in the Ayurvedic system of medicine [18]. Various preparations from the seeds of this plant are used for the management of several free radical mediated diseases such as ageing, rheumatoid arthritis, diabetes, atherosclerosis, male infertility and nervous disorders [15].

_Myrystica fragrans_ (Myristicaceae) has been known to possess cytotoxic, hepatoprotective, antioxidant, anti-inflammatory, anti-thrombotic, hypolipidaemic, antiatherosclerotic, hypoglycaemic and anti-diabetic activities [19-21].

_Catharanthus roseus_ (Apocynaceae) which is commonly known as ‘periwinkle’ and is an important source of indole alkaloids, which are present in all plant parts. It is used for the treatment of diabetes, fever, malaria, throat infections, and chest complaints. It is also used for the regulation of menstrual cycles, and as a euphoriant [22,23].

_Caesalpinia bonducella_ (Caesalpiniaceae) has been reported to exhibit wide array of activities including antidiabetic, adaptogenic, anthelmintic, antifilarial, antibacterial, anti-fungal, anti-spasmodic activity, antioxidant, anti-sporotic and antitumor activity [24-26].

_Myrystica fragrans_ (H.) (Myristicaceae) has been known to possess cytotoxic, hepatoprotective, antioxidant, anti-inflammatory, anti-thrombotic, hypolipidaemic, antiatherosclerotic, hypoglycaemic and anti-diabetic activities [27-29]. The present study thus aimed to make a comparative study for the ability of the extracts of selected plants to inhibit in-vitro α-amylase activity.

**Materials and Methods**

**Chemicals and reagents**

All the chemicals used during the experimental work were of analytical grade obtained from S.D. Fine Chemicals Pvt. Ltd., Mumbai, Sigma chemical company, USA and Loba chemicals, Mumbai.

**Plant material**

The seeds of _M. pruriens_ and _C. bonducella_ were collected from local areas of Kasegaon, District Sangli, (MS), India, whereas the seeds of _M. fragrans_ and roots of _B. aristata_ were purchased from the local market. Bark of _A. lebbeck_ and roots of _C. roseus_ were collected from local areas of Karad, District Satara, (MS), India. The plant material was further identified and authenticated by the Department of Botany, YC College of Science, Karad. The plant material was deaned thoroughly, dried in a hot air oven (50°C) separately powdered, passed through 60 mesh sieve (BS) and thereafter stored in an airtight container at 4°C till further use.

**Preparation of plant extracts**

Aqueous extracts were prepared by extracting the powders of the plant materials with hot water (70°C) in a mechanical shaker (24h), filtered and freeze dried.

**Assay for α-amylase inhibition [30]**

In this method 1ml substrate- potato starch (1%w/v), 1ml of drug solution (Acarbose, std drug/ plant extracts) of four different concentration such as 250, 500, 750 and 1000µg/ml, 1ml of alpha amylase enzyme (1% w/v) and 2ml of acetate buffer (0.1M, 7.2pH) were mixed together. These solutions were prepared using acetate buffer (820.3mg Sodium acetate and 18.7mg sodium chloride in 100ml distilled water).The above mixture was incubated for 1hr. Then 0.1ml of iodine-iodide indicator (6.35mg iodine and 1gm potassium iodide in 250ml distilled water) was added in the mixture. Absorbance of the resulting solution was measured at 565nm using UV-Visible spectrophotometer.

\[
\% \text{ inhibition} = \frac{A_c - A_s}{A_s} \times 100
\]

Where, _Ac_ is the absorbance of the control and _As_ is the absorbance of the sample.

**Statistical analysis**

All the analyses were carried out in triplicate and the results were expressed in mean±SD.

**Results and Discussion**

Many herbal extracts has been reported for their antidiabetic activities and being used in the traditional system of medicine, Ayurveda for the treatment of diabetes. Numerous herbal extracts have been investigated with respect to suppression of glucose production from carbohydrates in the gut or glucose absorption from the intestine. The enzyme alpha amylase catalyses the hydrolysis of 1,4-glucosidic linkages of starch, glycogen and various oligosaccharides into simpler sugars which can be readily available for the intestinal absorption. Inhibition of alpha amylase enzyme in the digestive tract of human is being considered to be effective in controlling diabetes by decreasing the absorption of glucose from starch. However, medicinal plants have not gained much importance as medicines and one of the vital factors is lack of specific standards being prescribed for herbal medicines and scientific support. Thus, the study was undertaken to assess the inhibitory action of the selected plant extracts on the alpha amylase enzyme.

Aqueous extracts of different parts of selected plants with known antidiabetic activity were investigated for their potential to inhibit α-amylase activity at five different concentrations viz. 100µg/ml, 250µg/ml, 500µg/ml, 750µg/ml and 1000µg/ml. The results of inhibition of alpha amylase enzyme by the selected plant extracts have been represented in Figure 1. It was observed that the selected plants showed exhibited variable inhibitory effects on α-amylase activity in vitro. Acarbose was used as a standard for this activity. The extract of _A. lebbeck_ at a concentration of 1000µg/ml exhibited an inhibition of 70.91%, whereas the extracts of _B. aristata_ and _M. pruriens_ showed an inhibition of 65.24% and 62.96.
% respectively at a similar concentration. Acarbose exhibited the inhibition of 81.48% at a concentration of 1000µg/ml. The results of the assay are presented in the Table 1.

Table 1: Percentage inhibition of the enzyme alpha amylase by the selected plant extracts.

<table>
<thead>
<tr>
<th>Concentration (µg/ml)</th>
<th>Acarbose</th>
<th>AL</th>
<th>BA</th>
<th>MP</th>
<th>MF</th>
<th>CR</th>
<th>CB</th>
</tr>
</thead>
<tbody>
<tr>
<td>100</td>
<td>27.37</td>
<td>21.21</td>
<td>9.09</td>
<td>10.34</td>
<td>5.1</td>
<td>3.7</td>
<td>11</td>
</tr>
<tr>
<td>250</td>
<td>45.14</td>
<td>38.96</td>
<td>30.1</td>
<td>25.71</td>
<td>10.34</td>
<td>7.14</td>
<td>15.03</td>
</tr>
<tr>
<td>500</td>
<td>65.14</td>
<td>57.65</td>
<td>52.38</td>
<td>50.19</td>
<td>19.25</td>
<td>16.66</td>
<td>23.52</td>
</tr>
<tr>
<td>750</td>
<td>72.39</td>
<td>64.08</td>
<td>58.06</td>
<td>57.65</td>
<td>23.52</td>
<td>18.75</td>
<td>27.37</td>
</tr>
<tr>
<td>1000</td>
<td>81.48</td>
<td>70.91</td>
<td>65.24</td>
<td>62.96</td>
<td>28.96</td>
<td>23.07</td>
<td>34.01</td>
</tr>
</tbody>
</table>

Acarbose showed an IC_{50} value of 383.7µg/ml whereas the extracts of *A. lebbeck*, *B. aristata* and *M. pruriens* showed the inhibition of the alpha amylase enzyme at an IC_{50} value of 433.65µg, 477.28µg and 498.10µg respectively. The extracts of *C. roseus*, *M. fragrans* and *C. bonducella* exhibited a relatively less inhibitory response for the α-amylase enzyme. The results of the assay are presented in the Table 1. The IC_{50} values are as shown in the Figure 2.

The outcomes of this study exhibited the alpha amylase inhibitory effects of the extracts of *Albizia lebbeck*, *Berberis aristata*, *Mucuna pruriens*, *Myristica fragrans*, *Catharanthus roseus*, and *Caesalpinia bonducella* at five different concentrations viz. 100µg/ml, 250µg/ml, 500µg/ml, 750µg/ml and 1000µg/ml. However, the inhibitory effect showed by the extracts of *Albizia lebbeck*, *Berberis aristata*, *Mucuna pruriens* was more than the extracts of *Myristica fragrans*, *Catharanthus roseus*, and *Caesalpinia bonducella* at a similar concentration. The present study thus, indicated that the extracts of *Albizia lebbeck*, *Berberis aristata* and *Mucuna pruriens* could be useful in the management of postprandial hyperglycemia. The results of this study directs further researches to evaluate the therapeutic potentialities of these extracts in the management of postprandial hyperglycemia and Type 2 diabetes either alone or in a combinatorial therapy (Figure 1& 2).
Conclusion

The results of the present study indicate that the selected plant extracts exhibited maximum alpha amylase inhibitory activity and thus, will prove to be beneficial to reduce the rate of digestion and absorption of carbohydrates and thereby contribute for effective management of diabetes by decreasing the post-prandial hyperglycemia. The plants may essentially contain herbal bioactive compounds inhibiting enzyme activity and further structural elucidation and characterization methodologies have to be carried out in order to identify the bioactive constituents.

References
