SCREENING OF ANTI-DIABETIC PROPERTIES OF FUCOIDAN EXTRACTED FROM *PADINA DISTROMATICA* HAUCK (BROWN SEAWEED) FROM HARE ISLAND, THOUTHUKUDI, TAMIL NADU, INDIA

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ABSTRACT

In the present study, the anti-diabetic action of fucoidan extracted from *Padina distromatica* Hauck in Hare Island, Thoothukudi, Tamil Nadu, India was evaluated. Anti-diabetic property of fucoidan was tested by the inhibitory effect of α-amylase and α-glucosidase. Different concentrations of fucoidan such as 25, 50, 75 and 100µg were added to the starch as substrate and the enzyme mixture. After 5min the enzymes activity were arrested and estimated using spectroscopically. The result showed that all the concentrations of the fucoidan were inhibited both α-amylase and α-glucosidase. When the concentration of fucoidan was increased from 25 to 100µg, the inhibitory effect of the enzyme also increased. From the present report, it was concluded that low concentration of fucoidan has minimum anti-diabetic effect and the high concentration of fucoidan has the maximum anti-diabetic effect. The enzyme inhibiting activity of fucoidan was quite variable, depending on the concentration of fucoidan on the targeted enzymes α-amylase and α-glucosidase.

KEY WORDS: Anti-diabetic, α-amylase, α-glucosidase Fucoidan, Seaweeds, *Padina*.

INTRODUCTION

Diabetes is a metabolic disorder characterized by high plasma glucose levels. Diabetes is classified as Type-1 and Type-2. Type-1 or insulin-dependent diabetes is due to failure of the pancreas to secrete insulin, while Type-2 or non-insulin dependent diabetes is the result of insufficient insulin production. It was estimated that 246 million persons in the world suffered from Type-2 diabetes in 2004 and that this number will reach at least 380 million by 2025 [1]. Type-2 diabetes receives more attention than Type-1 because it is considered avoidable. Type-2 diabetes is caused by imbalance between blood sugar absorption and insulin secretion. Post-prandial hyperglycemia plays an important role in development of Type-2 diabetes [2]. The control of plasma glucose level is essential to delay and even prevent Type-2 diabetes. To reach this goal, increasing or stimulating insulin secretion through medication [3] and/or by dietary supervision could be achieved. Dietary control is suggested as a safe and complementary treatment of diabetes. A diet based on glycemic index is currently one of the most recommended nutritional treatments. It has been reported that dietary therapy can be used simultaneously with other medical treatments in order to obtain a synergistic effect [4]. However, this has the drawback of limiting the types and quantity of food consumed. Another possible solution is to decrease the rate of blood sugar absorption from the small intestine by slowing and interrupting the digestion of dietary starch, the major source of glucose [5]. This approach is considered more efficient than controlling insulin secretion, for economic reasons, convenience and avoidance of side effects [6]. The inhibition of enzymes that digest dietary starch into glucose, α-amylase and α-glucosidase has been studied as a way of controlling blood sugar level [7]. α-amylase from human saliva catalyses the hydrolysis of α-(1,4)-glucosidic linkages and produces maltose and glucose from starch [8], while α-glucosidase releases glucose from
maltose [9]. By inhibiting these two enzymes, the absorption of glucose into the bloodstream can be delayed and thus, ameliorating Type-2 diabetes symptom-like hyperglycemia.

Attempts have been made to identify α-amylase and α-glucosidase inhibitors that can be used as food or food additives. Although Seo et al. [10] and Kato et al. [11] demonstrated that some sugar-like phenolic compounds have α-glucosidase inhibitory activity, most studies on α-amylase and α-glucosidase inhibitors have focused on the utilization of proteins or phenolic compounds [12]. Although phenolic compounds have high inhibitory activity of α-amylase and α-glucosidase, phenolic compounds are unstable, sensitive to light and heat treatments limit their uses as nutraceuticals.

Recently, algae have been considered as a source of enzyme inhibitors. As several plant extracts, algae contain some polyphenolic compounds as bromophenols [13], phlorotannins [14] which are inhibitors of α-glucosidase. Also, polysaccharides, isolated from algae, have become attractive in the biomedical area for numerous bioactivities [15-16]. In this context, fucoidan a polysaccharide found in brown algae and having several bioactivities appears promising. Fucoidan is abundant in brown seaweeds such as Padina species [17]. It is a mucilaginous, hygroscopic and sulfated polysaccharide [18] and it protects seaweed from dehydration [19]. The biological effects attributed to fucoidan are: anti-coagulant [20], anti-HIV [21] and anti-tumor activities [22].

Brown seaweeds contain up to 10% of fucoidan and the quantity varies depending on the region, species and season. The structural and chemical characteristics of fucoidan also vary among algal species [23]. The aim of this study was to investigate the inhibition of starch digestive enzymes such as α-amylase and α-glucosidase by fucoidan extracted from Padina distromatica Hauck in Hare Island, Thoothukudi, Tamil Nadu, India.

MATERIALS AND METHODS

Collection of Materials

The collection of Padina distromatica Hauck (Figure 1) was made during the low tidal and subtidal regions (up to 1m depth) by hand picking from Hare island, Thoothukudi (Lat 8° 48’N; Long 78° 11’E) located in the south east coast of Tamil Nadu, India. The collected materials were washed thoroughly with marine water in the field itself to remove the epiphytes and sediment particles. Then the samples were packed separately in polythene bags in wet conditions and brought to the laboratory, then thoroughly washed in tap water followed by distilled water to remove the salt on the surface of the thalli. They were stored in 5% formalin solution. For drying, washed specimens were placed on blotting paper and spread out at room temperature in the shade. The shade dried samples were grounded to fine powder using a tissue blender. The powdered samples were then stored in the refrigerator for further use.

Extraction of Fucoidan

Fucoidan was extracted by Rioux method [24]. Dried seaweed was mixed with 1% (w/v) CaCl₂ solution (1:30 ratio) and then stirred for 4 hours at 85°C at 455 ± 5rpm using a stirrer RZR1 (Caframo Ltd. Canada). The supernatant was separated by centrifugation (16,887 g, 20 min), and vacuum filtration on Whatman No. 4 filter. The filtered liquid was mixed with 2 volumes of 95% ethanol and 1 volume of 2% (w/v) NaCl and then stirred for 1 hour at room temperature for alcoholic precipitation of fucoidan. This solution was kept at -20°C for 48 hours. The pellet containing fucoidan was recovered by centrifugation (16,887 g, 12 min). Then, it was resolubilized in 100 ml of fresh deionized water and dialyzed for 48 hr by using membrane of 15 KDa (Sigma, USA) to remove minor constituents and solvents. Fucoidan was recovered by freeze-drying and preserved at -20°C in a sealed tube to keep away from humidity.

Alpha-amylase inhibition assay

The method of Conforti et al. [25] was modified to determine the inhibitory effects of the fucoidan extract on α-amylase (from human salivary, EC 3.2.1.1). 1% starch solution was prepared by stirring 1g of corn starch in 100ml of 20 mM sodium phosphate buffer with 6.7mM sodium chloride, pH 6.9. The solution was heated at 100°C for 15 min and then cooled to room temperature. The volume was brought to 100ml with distilled water. α-amylase solution was prepared as 1unit/ml. The different concentration of fucoidan solution (25, 50, 75 and 100µg) was added in 1ml of a starch solution. The tubes were prepared in duplicate and split into two groups, which were classified as a test group (TG) and a control group (CG). All tubes were incubated at 20°C for 10min and 1ml of α-amylase solution was added. For amylase activation, the TG tubes were incubated at exactly 20°C for 5min. The colorimetric reagent (1ml) was added to all of the tubes of the TG, heated at 100°C for 15min and cooled in an ice bath. 9ml of distilled water was added to each tube and the absorbance at 540nm was measured. The same steps were realized for the CG, but the incubation at 20°C for 5min was omitted. As blank, 0.1ml of distilled water was added instead of the fucoidan solution. The optical density of the blank (OD B) refers to the difference between test and control group. The degree of enzyme inhibition was calculated using the following equation,

\[
\text{Enzyme inhibition (\%) = \left(\frac{OD_{\text{B}} - OD_{\text{C}} - OD_{\text{NB}}}{OD_{\text{B}}}\right) \times 100(\%)}
\]

Alpha-glucosidase inhibition assay

For the determination of α-glucosidase (EC 3.2.1.20) inhibitory activity of fucoidan, all solutions were
prepared according to the method of Halvorson and Ellias [26]. 5ml of 67mM potassium phosphate (pH 6.8 at 37°C) and 0.2ml of 3 mM glutathione solution were added into total 10 testtubes. The test tubes were split into two groups, the test group (TG) and control group (CG). 0.2ml of 1unit/ml of α-glucosidase was added only into TG tubes and 0.2ml of distilled water was added into CG tubes instead of α-glucosidase. 25, 50, 75 and 100µg of fucoidan were added in both TG and CG tubes in parallel. As blank, 0.1ml of distilled water was added instead of the fucoidan solution. The optical density of blank (OD B) refers to the difference between test and control group. All tubes were kept at 37°C for 20min. 0.5ml of 10 mM p-Nitrophenyl alpha-D-glucopyranoside was added into all tubes and then incubated at 37°C for 20min. 1ml of the previous mixture solution was transferred into new test tube containing 4ml of sodium carbonate (0.1 M) and the solution was mixed. The absorbance was measured at 400nm and the degree of enzyme inhibition was calculated using the following equation,

\[ \text{Enzyme inhibition} \% = \left( \frac{\text{OD}_{\text{TG}} - \text{OD}_{\text{CG}}}{\text{OD}_{\text{B}}} \right) \times 100\% \]

**Statistical analysis**

Enzyme inhibition analyses were performed in duplicate with two repetitions for each fucoidan sample. Results are presented as mean ± standard deviations (SD) for each concentration of fucoidan. The enzyme activity data were analyzed using SAS 9.2 software (SAS Institute Inc., Cary, USA).

**RESULTS AND DISCUSSION**

Most studies looking for natural extracts contributing to diabetes prevention have been focusing on α-glucosidase inhibition because the enzyme plays a role at the ending step of starch digestion by producing glucose from maltose. Although the inhibition of α-amylase also results in the decrease of glucose release, its complete inhibition is not desired because it could provoke intestinal disorders [27]. The undigested starch could be utilized by the gut microflora for gas production. Therefore, partial inhibition of α-amylase could contribute to modulate the rate of glucose release from starch.

Fucoidan is a natural polysaccharide made essentially of sulphated L-fucose residues. Also known as sulphated fucan, it was first extracted in 1913 from brown algae [28]. Fucoidan is present in the cell walls of brown algae especially *Padina distromatica* Hauck. Though many studies on identifying the structural properties of fucoidan have been carried out, the structure still remains uncertain due to the absence of strict regularity and the numerous components that make up fucoidan as a whole [29]. Figure 2 shows the general structure of fucoidan but the chemical composition and structure of fucoidan varies with species [30]. Most fucoidans have very complex chemical composition and only little regularity in the structural components is known present [24]. Fucoidan largely contains sulphated L-fucose residues. Hence fucose is the primary sugar in fucoidan. Sulphate groups also represent a large component of fucoidan and the biological activities of fucoidan are strongly related to its sulphate content [31]. Besides fucose and sulphate, other monosaccharides (glucose, mannose, galactose, xylose, etc), uronic acids, and even protein are present in detectable amounts. All these compounds have increased the difficulty in structural elucidation of fucoidan [32].

**Inhibition of α-amylase Activity**

The α-amylase activity effectiveness of different concentration of fucoidan extracted from *Padina distromatica* Hauck were compared on the basis of the inhibition percentage. Among the different concentration of fucoidan used, 100µg of fucoidan extracted from *Padina distromatica* Hauck inhibited 65.6% followed by 75µg fucoidan with 48% and 50µg with 34.4%. 25µg of fucoidan showed less inhibitory activity of α-amylase with 17.6%. Acarbose, the positive control used in the study, inhibited the activity of α-amylase at 75.5%. The present result showed that the percentage of α-amylase inhibitory activity increased when the concentration of fucoidan increased from low concentration to higher concentration (Table 1 & Figure 3).

**Inhibition of α-glucosidase Activity**

In the present study fucoidan extracted from *Padina distromatica* Hauck collected from Hare Island, Thoothukudi, Tamil Nadu, India was found to possess favorable α-glucosidase inhibitory effects on starch break down *in vitro*. The α-glucosidase inhibitor activity effectiveness of different concentration of fucoidan extracted from *Padina distromatica* Hauck were compared on the basis of the inhibition percentage. Among the different concentration of fucoidan used, 100µg of fucoidan extracted from *Padina distromatica* Hauck inhibited 70.43% followed by 75µg fucoidan with 50.43% and 50µg with 32.17%. 25µg of fucoidan showed less inhibitory activity of α-glucosidase with 13.04%. Acarbose, the positive control used, inhibited the activity of α-glucosidase at 81.2%. The present study showed that the percentage of α-glucosidase inhibitory activity increased when the concentration of fucoidan increased from low concentration to higher concentration (Table 2 & Figure 4).

Mechanisms of α-glucosidase inhibition differ among the various inhibitors reported. Some well-known inhibitors as acarbose mimic the enzyme substrate [10]. Previously published articles on the mechanism of polyphenol compounds suggest that the principal factor acting on α-glucosidase activity is hydrogen scavenging because α-glucosidase provides hydrogen to catalyze the hydrolysis of the α-(1,4)-glucosidic linkage [33]. The inhibitor acts by intercepting the hydrogen ion freed from
the α-glucosidase catalytic site. Most studies conducted with fucoidan showed that the sulfate content is closely related to its biological properties. Wang et al. [34] showed a relation between sulfate content and radical scavenging property using fucoidan from Laminaria japonica. So et al. [35] also reported free-radical scavenging ability of fucoidan from Fucus vesiculosus. Highly sulfated fucoidan (37% sulfate groups) showed better free radical scavenging activity than native fucoidan (28% sulfate groups). Increasing sulfate groups may enhance the scavenging activity of fucoidan and thus, promote its capacity to intercept free hydrogen. Numerous phenolic compounds such as flavonol [36], catechins and theaflavins [37] have an α-glucosidase inhibitory activity. The intensity of the activity is related to the structure and conformation of polyphenols [38]. Fucoidan might inhibit α-glucosidase by a mechanism similar to that of polyphenols that involves scavenging but more research will be needed in order to determine the mechanism of action. Structural features such as molecular weight and the amount of sulfate groups could seriously impact the enzyme activity. More work was realized in order to explain how those structural characteristics influence the enzyme activity.

Finally, when comparing the inhibition of the two digestive enzymes studied, α-amylase and α-glucosidase, it should be emphasized that fucoidan showed more inhibitory effect on α-glucosidase than α-amylase. This difference in inhibition offers a complementary effect because as reported Cho et al. [27], high α-amylase inhibition could be related to intestinal discomfort, so a moderated inhibition of α-amylase and a stronger glucosidase inhibition would be preferred. The level of inhibition obtained with fucoidan should have the desired effect by slowing the absorption of glucose from the intestine into the bloodstream enough to allow the insulin-based mechanism of the diabetic patient to transport blood sugar into the muscles before it can reach harmful levels. These data overall suggest that fucoidan could be useful as a component of new medication, as a food additive or as a food supplement for an enzyme-targeted treatment of Type-2 diabetes. More information on fucoidan structural features are required to understand the mechanism by which fucoidan inhibits α-amylase and α-glucosidase.

Table 1. Effect of Standard and various concentration of Fucoidan on Inhibition of α-amylase Activity (%)

<table>
<thead>
<tr>
<th>Groups</th>
<th>Percentage of Inhibition of α-amylase Activity</th>
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<tbody>
<tr>
<td>Acarbose</td>
<td>75.5%</td>
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<tr>
<td>25μg Fucoidan</td>
<td>17.6%</td>
</tr>
<tr>
<td>50μg Fucoidan</td>
<td>34.4%</td>
</tr>
<tr>
<td>75μg Fucoidan</td>
<td>48%</td>
</tr>
<tr>
<td>100μg Fucoidan</td>
<td>65.6%</td>
</tr>
</tbody>
</table>

Table 2. Effect of Standard and various concentration of Fucoidan on Inhibition of α-glucosidase Activity (%)

<table>
<thead>
<tr>
<th>Groups</th>
<th>Percentage of Inhibition of α-glucosidase Activity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acarbose</td>
<td>81.2%</td>
</tr>
<tr>
<td>25μg Fucoidan</td>
<td>13.04%</td>
</tr>
<tr>
<td>50μg Fucoidan</td>
<td>32.17%</td>
</tr>
<tr>
<td>75μg Fucoidan</td>
<td>50.43%</td>
</tr>
<tr>
<td>100μg Fucoidan</td>
<td>70.43%</td>
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Figure 1. Natural Habit of Padina distromatica Hauck

Figure 2. Structure of Fucoidan
CONCLUSION

This study has revealed a novel function of fucoidan as an efficient inhibitor of the starch digesting enzymes α-amylase and α-glucosidase. In summary, the enzyme inhibiting activity of fucoidan was quite variable, depending on the concentration of fucoidan on the targeted enzymes α-amylase and α-glucosidase. Fucoidan extracted from Padina distromatica Hauck collected in Hare island, Thoothukudi, Tamil Nadu, India inhibited both α-amylase and α-glucosidase but the required quantities were much higher for α-amylase inhibition. Padina distromatica Hauck has greater potential for the prevention of Type-2 diabetes.

REFERENCES


