Short Report

The Effect of *Teucrium polium* (Calpoureh) on Liver function, Serum Lipids and Glucose in Diabetic Male Rats

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ABSTRACT

**Background:** *Teucrium polium* is an analgesic, antidiabetic and antilipemic herbal medicament. The aim of this survey was to evaluate the effect of aqueous extract *T. polium* on liver enzymes linked to liver dysfunction, serum lipids and glucose, in diabetic male rats. **Methods:** A total of 20 Sprague-Dawly male rats became diabetic by intraperitoneal injection of streptozotocin (60 mg/kg). the animals were divided randomly into two groups. Experimental group was fed *Teucrium polium* (50 mg/kg) for a month but control group was received the same volume of distilled water. Liver enzymes, biochemical parameters (cholesterol, triglyceride, low density lipoprotein, alanine transaminase, aspartae transaminase) and glucose were measured by kinetic (Enzymatic) and colorimetric methods. Data obtained were analyzed and mean values were compared by paired student's *t*-test. The results were expressed as mean ± SD. Significant differences were set at *P*<0.05. **Results:** Our results showed that in test group, serum glucose values decreased significantly (*P*<0.05), but cholesterol, triglyceride, low density lipoprotein, alanine transaminase and asp artae transaminase increased significantly after use of *T. polium* (*P*<0.05). This parameters value did not show any changes in control group. **Conclusion:** Although the aqueous extract of *Teucrium polium* has strong hypoglycemic properties in experimental animals, but because of some hepatotoxic effects, it is not suitable to use it in human as an antidiabetic agent. *Iran. Biomed. J. 11 (1): 65-68, 2007*

**Keywords:** *Teucrium polium*, Liver function, Blood glucose

INTRODUCTION

Liver is the major organ of the body that has an important effect on carbohydrates and lipid metabolism [1]. In the presence of insulin, glucose is used but lipids and proteins are stored in the body [2]. In diabetes mellitus, insulin deficiency leads to failure of glucose consumption, consequently results in breakdown of lipids and proteins [3].

In traditional medicaments, *Teucrium polium* is used as analgesic, anti-spasmodic and hypolipidemic agent [4-6]. Visceral analgesic effects of *T. polium* extract compete considerably with those of indomethacin and hyoscine [4].

Use of *T. polium* in scharomyces culture media in vitro led to decrease in fatty acids and acts as anti-fungal, anti-bacterial and anti-inflammatory agent, and blocks the peroxidation of erythrocytes [7-9]. There is an agreement for hepatotoxicity of *T. polium* administration [10]. Administration of 150 mg/kg Teuceium polium extract was showed to act as an anti-ulcer agent [11]. Intravenous infusion and i.p. injection of plant extract after 4 and 24 hours led to decrease of blood sugar in rats [12]. Oral and i.p. administration of dried aerial parts and bloom extract of *T. polium* decreased appetite, water and food consumption and consequently body weight in rats [13]. The side effect of *T. polium* extracts were reported in diabetic patients who used it as an anti-diabetic agent [14, 15]. Oral
administration of alcoholic  
T. polium  
extract showed no changes in fasting and postprandial blood sugar in diabetic patient [16], Zal et al. [17] reported that 
the administration of  
T. polium  
boiling extract had an anti-diabetic effect on diabetic rats [17]. With considering the controversial reports of the above studies, the prime aim of this study was to identify the the effect of  
T. polium  
aqueous extract on blood glucose, Liver enzymes linked to liver dysfunction and serum lipid in streptozotocin diabetic male rats.

MATERIALS AND METHODS

A total of 20 Sprague-Dawly male rats weighting 
220 ± 14 g were purchased from Pasteur Institute of Tehran (Iran). Animals were housed in cages under conditions of controlled temperature (22- 28ºC) and a 12-h artificial light period for 10 days before and during of experiments) and had free access to water and standard pellet diet. The dried parts of  
T. polium  
were purchased from herbalists in Kerman and were authenticated by the Center for Research on Natural Resources and Livestock (Ministry of Agricultural Jihad, Isfahan, Iran) as  
T. polium; L. Every day, 120 mg of cleaned aerial parts of  
T. polium  
was suspended in 15 ml of water and put on a shaker for 24 hours. The suspension was cleared upon passing through several layers of chees. Animals became diabetic by i.p. injection of streptozotocin (60 mg/kg) and were divided randomly into two groups. Experimental group was gavaged  
T. polium  
(50 mg/kg, d = 1.09) for 4 weeks but control group was received the same volume of distilled water. After a week that animals showed diabetic behavior such as polyuria and Polydipsia, at fasting state were anesthetized by ether and blood samples were collected from tail vein for evaluation of glucose, alanine transaminase (ALT), aspartae transaminase (AST), Alkaline Phosphatase (ALP), and lipoproteins. At the end of treatment period (after an overnight fasting), animals were anesthetized with high dose of ketamine and killed by cutis carotid vein and blood samples were collected. Glucose, ALT, AST, cholesterol (Cho), triglyceride (TG), and lipoproteins (HDL, LDL) were measured blindly by kinetic (enzymatic) and colorimetric methods. Three rats from control group died before the end of experiment. In addition, 3 Rats were died in control group (ethic No. 136 Dated 13-5-2003, Zahedan University of Medical Sciences, Iran) and the results were expressed as mean ± SD. To confirm the normal distribution, the data were analyzed by one-sample Kolmogrov-Smirnov test, then by Levant's and compared by paired student's t-test. Significant differences were set at  
P<0.05. All statistical analyses were performed using SPSS (v.11).

RESULTS AND DISCUSSION

Our results revealed that serum glucose value was significantly decreased but cho, TG, ALT, AST and lipoproteins were significantly increased after  
T. polium  
administration however these parameters did not show any changes in control group (Table 1 and 2,  
P<0.05). The comparison of mean weights in test group before (221.1 ± 16.14g) and after use of  
T. polium  
(219.6 ± 14.29 g) did not show any significant changes but the mean weights in control group before (221.28 ± 11.95 g) and after the study (191.85 ± 18.15 g) of the test were significantly decreased (Table 3,  
P= 0.01). The comparison of water consumption in test (145.88 ± 28.79 cc) and control groups (154.61 ± 21cc) did not show any significant changes.

| Biochemical parameters | Before  
T. polium  
admission | After  
T. polium  
admission |
<table>
<thead>
<tr>
<th></th>
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</thead>
<tbody>
<tr>
<td>Cholesterol (mg/dl)</td>
<td>64.00 ±14.90</td>
<td>*94.20 ± 5.73</td>
</tr>
<tr>
<td>Triglyceride (mg/dl)</td>
<td>85.10 ± 17.18</td>
<td>*146.00 ±15.51</td>
</tr>
<tr>
<td>HDL (mg/dl)</td>
<td>33.10 ± 10.70</td>
<td>41.00 ± 11.03</td>
</tr>
<tr>
<td>LDL (mg/dl)</td>
<td>13.91 ± 7.80</td>
<td>*23.96 ± 11.30</td>
</tr>
<tr>
<td>AST (U/L)</td>
<td>86.80 ± 25.70</td>
<td>*383.10 ± 196.10</td>
</tr>
<tr>
<td>ALT (U/L)</td>
<td>118.10 ± 18.57</td>
<td>355.60 ± 259.80</td>
</tr>
<tr>
<td>ALP (U/L)</td>
<td>936.00 ± 255.57</td>
<td>970.40 ± 275.60</td>
</tr>
<tr>
<td>Glucose (mg/dl)</td>
<td>283.61 ± 22.13</td>
<td>*96.22 ± 11.90</td>
</tr>
</tbody>
</table>

N =10; values are mean ± SD; *P<0.05.

Our results are in part in accordance with Rasaekh et al. [12] which showed that  
T. polium  
decreased the serum glucose level of diabetic rats. Although we did not show any antilipidemic effect for  
T. polium  
aqueous extract, Rasaekh et al. [12] reported antilipidemic effect of alcoholic  
T. polium  
extract; this difference in our results may be due to the difference in method of  
T. polium  
administration. We used oral method whereas they used i.p. method. Some of the parameters such as AST and ALT values increased after  
T. polium  
admission in...
Table 2. Comparison of AST, ALT, ALP, HDL, LDL, triglyceride, cholesterol and glucose before and after period of experiment in control group.

<table>
<thead>
<tr>
<th>Biochemical parameters</th>
<th>Before period of experiment</th>
<th>After period of experiment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cholesterol (mg/dl)</td>
<td>61.28 ± 12.86</td>
<td>104.00 ± 18.22</td>
</tr>
<tr>
<td>Triglyceride (mg/dl)</td>
<td>76.14 ± 15.51</td>
<td>163.85 ± 48.05</td>
</tr>
<tr>
<td>HDL (mg/dl)</td>
<td>34.86 ± 8.45</td>
<td>38.00 ± 8.16</td>
</tr>
<tr>
<td>LDL (mg/dl)</td>
<td>13.62 ± 9.21</td>
<td>33.00 ± 12.60</td>
</tr>
<tr>
<td>AST (U/L)</td>
<td>83.85 ± 20.43</td>
<td>165.71 ± 34.52</td>
</tr>
<tr>
<td>ALT (U/L)</td>
<td>107.28 ± 20.68</td>
<td>195.14 ± 73.09</td>
</tr>
<tr>
<td>ALP (U/L)</td>
<td>1234.70 ± 313.19</td>
<td>1307.71 ± 317.51</td>
</tr>
<tr>
<td>Glucose (mg/dl)</td>
<td>270.40 ± 41.20</td>
<td>283.14 ± 46.71</td>
</tr>
</tbody>
</table>

N = 7; values are mean ± SD; *P<0.05.

Table 3. Comparison of weight between control and test group before and after experiment period.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Weight (g)</th>
<th>Before period of experiment</th>
<th>After period of experiment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Test group (n = 10)</td>
<td>221.10 ± 16.14</td>
<td>219.60 ± 14.29</td>
<td></td>
</tr>
<tr>
<td>Control group (n = 7)</td>
<td>221.85 ± 11.95*</td>
<td>191.85 ± 18.15</td>
<td></td>
</tr>
</tbody>
</table>

N = 17; values are mean ± SD; *P<0.05.

ACKNOWLEDGMENTS

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REFERENCES


