Effects of hydroalcoholic extract in *Mellissa officinalis* plant on fat profiles and glucose level in diabetic rats induced by streptozotocin

Amin Tashakor¹, Mahnas Rezaei Kelishadi², Ali Ghasemi², Fatemeh Daylami², Azar Rahimi², Setareh Zamani Doabi² and Naser Nabi Abdolusefi²*

¹Department of Pharmacology and Therapeutics, National University of Ireland Galway, Galway, Ireland  
²Department of Biochemistry, Payame Noor University of Isfahan, Isfahan, Iran

ABSTRACT

Over the years medicinal plants have been using in treatment of diabetes in traditional medicine, although little scientific studies have been reported in scientific communities. The aim of this study was to evaluate the effect of hydroalcoholic extract in *Mellissa officinalis* plant on fat profiles and the level of glucose in streptozotocin(STZ)-induced diabetic rats. In the current study, 35 Wistar rats were divided into 5 groups of seven as control group with normal diet, fatty-based diet group and the experimental groups of hypercholesterolaemic animals with minimum dose of 100, average dose of 200 and maximum dose of 300 milligrams per kilogram of *Mellissa officinalis* extract respectively as gavage feeding. After 14 days of treatment the lipid profiles, HDL, LDL, TG and cholesterol were measured and the obtained results were analyzed using SPSS. The level of cholesterol did not change significantly in the groups treated with herbal extracts. However, the level of HDL increased and TG decreased. According to the obtained results, increase in HDL and decrease in TG levels have positive effects on diabetic rats and therefore decrease the level of blood glucose.

Keywords: *Mellissa officinalis* plant, Streptozotocin, Diabetic rats, Lipid profile, Blood glucose level

INTRODUCTION

Diabetes mellitus is a systemic metabolic disease characterized by hyperglycemia, hyperlipidemia, and decreased insulin secretion or function and is often associated with small and large vessel diseases like neuropathy, nephropathy, cardiovascular and cerebrovascular diseases. Diabetes mellitus is associated with prolonged hyperglycemia which is an important factor in the development of the above-mentioned diseases[1, 2].The first complete clinical explanation of diabetes was presented by Aretaeus of Cappadocia, an ancient Greek physician who also found the excessive amount of urine secreted from the kidneys. Aretaeus attempted to treat it but he could not give a good prognosis. He stated that diabetes makes life short, disgusting, and painful. In medieval Persia, Avicenna (980–1037) provided a detailed description for diabetes mellitus as the abnormal appetite and the collapse of sexual functions, and also he mentioned the sweet taste of diabetic urine. Similar to Aretaeus, Avicenna recognized the primary and secondary diabetes[1].

As predicted by the World Health Organization, the number of people suffering from this disease will amount to 370 million by 2030[3].Due to numerous problems and sometimes lethality in people suffering from diabetes, approaches should be taken into account in research, and found new reagents to prevent and treat of this disease.

Nowadays, problems in production and injection of insulin and other hypoglycemic drugs in and also side effects of chemical drugs, have directed the researchers to use medicinal plants. Over the years medicinal plants have been using in treatment of diabetes in traditional medicine, although little scientific studies have been reported in scientific communities[4].
Melissa officinalis (MO) is one of the proposed medicinal plants in traditional medicine that is effective in the treatment of diabetes. Melissa officinalis (MO) belongs to Lamiaceae family and is found a traditional plant in the Mediterranean and eastern Asia. In Iran, this plant is known by the name of Melissa officinalis and is widely distributed in most of the country[5, 6].

Melissa officinalis like other herbal compounds such as curcumin has antioxidant properties due to the presence of polyphenolic compounds including quercetin, gallic acid, rutin flavonoid, aldehyde and tannin components. Herbal compounds have also been shown to play important roles in anti-inflammatory and anti-apoptotic responses, the latter mostly through intrinsic pathway of apoptosis and involvement of mitochondria, cytochrome c, Apoptotic Procaspe Activating Factor 1 (Apaf-1) and executioner caspases[7-9]. Moreover, studies on Melissa officinalis (MO) have shown antioxidant, antimicrobial and anti-genotoxic properties and play a role in memory, learning and Alzheimer's disease. Flavonoid and terpenoid compounds which are abundant in this plant have been shown to have antispasmodic and anti-inflammatory effects[10, 11]. In the present study the hypoglycaemia effect of hydroalcoholic extract in Melissi Officinalis plant was investigated on fat profiles and the level of glucose in streptozotocin-induced diabetic rats.

**EXPERIMENTAL SECTION**

In this study 35 Wistar male rats with weight ranging from 200 to 230 gram purchased from Isfahan university of medical sciences, Iran were used. Rats stored in animal room with controlled temperature 23±2 °C and photoperiod of 12 hours of light and 12 hours of darkness, enough food and water. To induce diabetes, rats were injected intraperitoneally by 70 mg/kg streptozotocin (Pharmacia & Upjohn, USA). To ensure that animals were diabetic the levels of blood glucose were measured through blood letting from rat tails which was more than 300 milligrams per deciliter and indicated the diabetic animals.

**Collection, identification and preparation of herbal extracts:**

MO was collected from the Abadeh, Fars province, Iran in the spring 2014 and taxonomically was identified at school of Basic sciences, University of Isfahan. Then dried at temperature of 25 ° C in the shade, and powder prepared using a mechanical mill dried powder stored in freezer. Alcoholic extract was obtained using Soxhlet apparatus and 80% ethanol and was dried by rotary.

**Treatment method:**

Plant extract in different doses, the drug Glibenclamide and saline were treated as intra-peritoneal injection. The animals were divided into 5 groups.

Control: During the experiment, the rats in this group did not receive any drug or solvent and received normal diet. Experimental group 1: rats in this group received streptozotocin[1, 2].

The experimental group 2: diabetic rats received the alcoholic extract of Melissa officinalis daily, 100 mg kg (minimum dose) orally for 14 days through gavage feeding.

The experimental group 3: diabetic rats received the alcoholic extract of Melissa officinalis daily, 200 mg kg (maximum dose) orally for 14 days through gavage feeding[12, 13].

**Biochemical methods:**

At the end of 14-day period bloodletting was performed from the heart to evaluate serum concentrations of biochemical factors after a mild anesthesia using ether. Then blood samples were centrifuged at 3000 rpm to separate serum which was then used for further investigation. To measure biochemical parameters such as HDL, LDL, TG and cholesterol IFCC standard method was applied using RA-1000 autoanalyzer. The level of serum cholesterol and triglyceride was measured using calorimetric Darmankav Kit (Iran). Lipoproteins were measured through a combination of sedimentary methods and ultracentrifuge (Darmankav, Iran). HDL- cholesterol were measured by HDL sedimentation technique[14] and the level of glucose were measured by EaseGluco 142Combo set.

**Statistical analysis**

All data were statistically analyzed using ANOVA one way- Tukey test, and Dancan, T-test and K related samples and the results were reported by the Mean ± SEM and p-value was considered in the level of 0.05 (P<0.05).
RESULTS AND DISCUSSION

In this study, blood glucose, cholesterol, triglycerides, low-density lipoprotein (LDL) levels showed a significant increase in the experimental group 1 compared to the control group and as well as high-density lipoprotein levels showed decrease.

Table 1 shows following results:
Glucose: average glucose levels in Melissa officinalis extract-treated diabetic rats showed significant reduction compared with diabetic control and glibenclamide groups (Table 1).

Rats’ body weight: The mean body weight of diabetic rats groups increased in those treated with the plant extract. (Table 1)

Cholesterol: cholesterol levels showed significant reduction in MO-treated group compared to diabetic control group at a dose of 100 mg and 200 mg/kg (P≤0.05)

TG: dosage of 100 mg/kg MO showed significant reduction in triglyceride levels compared to control and maximum dose of extract and glibenclamide (P≤0.05)

LDL: a dose of 100 milligrams per kilogram of MO extract, significantly increased LDL levels compared to the control group and glibenclamide. There was no significant change for 200 mg kg (P≤0.05).

HDL: MO extract at a dose of 100 mg kg resulted in a significant increase in HDL compared to the control group and the group received the glibenclamide drug (P≤0.05) (Table 2)

The results of this study showed that in those diabetic rats received Melissa officinalis extract blood sugar levels, cholesterol, triglycerides significantly decreased (P≤0.05) as well as high density lipoprotein and weight of diabetic rats showed a significant increase in diabetic rats received Melissa officinalis (P≤0.05).

Streptozotocin is a diabetogenic, hepatotoxic, and nephrotoxic substance which by damaging the pancreatic beta cell membrane, fragmenting DNA, and reacting with enzymes such asglucokinase, sharply decreases insulin levels and thereby increases glucose levels in animals[2, 15]. STZ increases mRNA expression of the liver glucose-6-phosphate dehydrogenase enzyme and thereby causes blood glucose to increase. Studies have also shown that diabetes could cause dyslipidemia and fatty liver. The reason for this is either the increased flow of fatty acids to the liver as the result of insulin reduction or decreased lipoprotein secretion from the liver due to a shortage of apolipoprotein B synthesis[16, 17].

Melissa officinalis has antioxidant properties due to polyphenolic compounds including quercetin, gallic acid, rutin flavonoid, aldehyde and tannin components.

Nuraliev and the colleagues reported the effect of hypoglycemia, diabetic quercetin in rats with alloxan that showed decrease in LDL. According to obtained results in addition to glucose quercetin significantly reduced low density lipoprotein (LDL) and cholesterol. Quercetin flavonoid inhibits the absorption of glucose in the intestine by acting specifically on glucose transporter 2 (GLU2)[18-20]. Chlorogenic acid, a specific inhibitor of glucose phosphatase enzyme, blocks glucose production in GLU2. This enzyme plays a key role in regulation of blood glucose and glucose output from the liver[21]. In some studies the hypolipidemic effect of quercetin, nicotinic acid and morin were examined and it was reported that, quercetin compared to other compounds, has greatest effect on cholesterol reduction[20]. Reduction in triglyceride in preventive groups can be related to recovery of Langerhans islets and subsequent increase in insulin levels. Increase in insulin level activates lipoprotein lipase which in turn decreases triglycerides and its concentration in the blood[22, 23]. Studies on other plants confirm that this is the reason for reduction in triglyceride[24-26]. According to the results of this study hydroalcoholic extract in Melissa Officinslis plant is efficient in prevention of diabetes, although biochemical and pharmacological studies should be performed.
Table 1. Scientific classification of *Mellissa Officinalis*

<table>
<thead>
<tr>
<th>Scientific Classification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Kingdom: Plantae</td>
</tr>
<tr>
<td>Division: Flowering plants</td>
</tr>
<tr>
<td>Class: Dicotyledonous</td>
</tr>
<tr>
<td>Order: Lamiales</td>
</tr>
<tr>
<td>Family: Lamiaceae</td>
</tr>
<tr>
<td>Genus: Melissa</td>
</tr>
<tr>
<td>Species: <em>M. officinalis</em></td>
</tr>
</tbody>
</table>

Table 2. The effect of *Melissa* plant compared to patients diabetes drug glibenclamide on body weight of rats and blood glucose

<table>
<thead>
<tr>
<th>Group rats</th>
<th>Initial weight (g)</th>
<th>Final weight (g)</th>
<th>Glucose (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>180</td>
<td>209</td>
<td>90</td>
</tr>
<tr>
<td>Diabetic Control</td>
<td>185</td>
<td>172α</td>
<td>380α</td>
</tr>
<tr>
<td>Glibenclamide</td>
<td>195</td>
<td>185</td>
<td>125</td>
</tr>
<tr>
<td>Meliss100mg/kg</td>
<td>178</td>
<td>185β</td>
<td>84β</td>
</tr>
<tr>
<td>Meliss200mg/kg</td>
<td>180</td>
<td>187β</td>
<td>70β</td>
</tr>
</tbody>
</table>

Mark α compared with the control group, symbol β compared to a glibenclamide and diabetic control rats

Table 3. The effect of *Melissa* Plant compared with glibenclamide on cholesterol, triglycerides, and total lipoproteins

<table>
<thead>
<tr>
<th>Group rats</th>
<th>Cholesterol (mg/dL)</th>
<th>Triglyceride (mg/dL)</th>
<th>HDL (mg/dL)</th>
<th>LDL (mg/dL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>58.50 ± 8.42</td>
<td>38.50 ± 4.59</td>
<td>47.50 ± 4.87</td>
<td>21.25 ± 2.65</td>
</tr>
<tr>
<td>Diabetic control</td>
<td>76.00 ± 3.93</td>
<td>52.50 ± 7.86α</td>
<td>42.25 ± 3.35α</td>
<td>43.25 ± 6.53α</td>
</tr>
<tr>
<td>Glibenclamide</td>
<td>56.00 ± 5.84β</td>
<td>44.50 ± 5.95</td>
<td>37.79 ± 2.92</td>
<td>39.75 ± 3.47</td>
</tr>
<tr>
<td>Meliss100mg/kg</td>
<td>52.00 ± 2.58β</td>
<td>32.75 ± 2.25π</td>
<td>53.00 ± 2.30π</td>
<td>62.50a ± 9.42a</td>
</tr>
<tr>
<td>Meliss200mg/kg</td>
<td>60.50 ± 2.46β</td>
<td>62.75 ± 2.13</td>
<td>47.25 ± 2.62π</td>
<td>54.50 ± 4.73</td>
</tr>
</tbody>
</table>

Mark α compared with the control group, Mark β compared to the control group, symbol π compared to the group received a dose of 100 mg/kg compared to the group the glibenclamide, diabetic control and other dose of Meliss. Sign ≠ comparison group received a dose of 200mg/kg compared to diabetic control, HDL: high density lipoproteins, LDL: low density lipoproteins

CONCLUSION

Alcoholic extract of *Melissa officinalis* plant which contains quercetin flavonoid with anti-diabetic effects shown in previous studies are able to efficiently reduce triglycerides and increase blood glucose, high density lipoprotein (HDL) and weight compared to glibenclamide drug in treatment of diabetes.

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REFERENCES