Original Research Article

Effect of hydroalcohol extract of lemon (Citrus limon) peel on a rat model of type 2 diabetes

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Abstract

Purpose: To determine the effect of the hydroalcohol extract of lemon peel (LP) on a rat model of type 2 diabetes (T2D).

Method: The rat model of T2D by injection of streptozotocin was established. The effects of a hydroalcoholic extract of LP was characterised on a rat model of type 2 diabetes based on body weight, food intake, fasting blood glucose (FBG), glucose tolerance test, and insulin tolerance test. Antioxidant activity and oxidative stress were analysed by superoxide dismutase (SOD) and malonaldehyde (MDA) assays.

Results: In acute toxicity studies, administration of LP extract at 2000 mg/kg orally did not cause any symptoms of poisoning or death after 14 days. The body weight of rats increased after treatment with LP extracts. Food intake in diabetic rats decreased with LP extract treatment. Continual treatment with LP extracts for 35 days significantly reduced blood glucose levels in diabetic rats. Glucose tolerance improved, and insulin resistance was reduced after treatment with LP extracts. SOD and MDA data indicate that treatment with LP extract alleviated the oxidative stress in diabetic rats as well as enhanced the antioxidant activity of liver in a dosage-dependent manner.

Conclusion: LP extract decreased food intake and FBG, but increased body weight in rats. The effect of LP on T2D is likely related to improved antioxidant activity and reduced oxidative stress. Thus, LP extract has potentials for the treatment of T2D.

Keywords: Lemon peel, Type 2 diabetes, Antioxidant activity, Glucose tolerance, Insulin tolerance

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INTRODUCTION

Type 2 diabetes (T2D) is a chronic disorder of the metabolism of carbohydrate and lipid, which has long been diagnosed by hyperglycaemia with a high fasting blood glucose (FBG) level. Patients with T2D represent 90% of all diabetes patients worldwide [1]. Relative to other diseases, T2D has an extremely high occurrence of complications and may even contribute to Alzheimer’s disease and depression [2,3]. Insulin resistance and increased oxidative stress are generally accepted as underlying the main pathogenesis of T2D and its complications [4,5].
T2D is characterised by progressive degeneration of insulin function, accompanied by β-cell dysfunction that attempts to compensate for insulin resistance [6]. T2D is often associated with high levels of free radicals [7] and attenuated antioxidant function [8]. Interestingly, the oxidant hydrogen peroxide is known to negatively impact glucose transportation capabilities and to stimulate insulin signalling elements [9]. Thus, oxidant stress may cause insulin resistance in mammalian muscle via the p38 MAPK pathway. For hundreds of years, traditional treatments for patients with T2D have utilised medicinal plants that have few side effects. In recent years, medicinal plants have played an important part in new drug discovery. The most widely used anti-T2D drug, metformin, was initially developed from Galega officinalis, a medicinal plant that has been used for diabetes treatment for several hundred years [10]. The extract of Galega officinalis provided a predicted chemical structure for new anti-diabetes drugs. Therefore, there has been increasing interest in the research on new medicinal plants that have anti-diabetes properties with few or no side effects to discover chemical structures that account for their therapeutic effect.

Lemon, which is grown worldwide, is used for ethnomedicinal applications due to the anti-inflammatory and anti-tumour functions of flavonoids and limonene that are contained in the plant [11,12]. Natural antioxidants extracted from plants are attracting more interest for their in vivo radical scavenging activities. The bioactive components that are extracted from lemon peel (LP) depend on the lemon species and the method of extraction.

In this study, the peel of Citrus limon was used for extraction by the hydroalcoholic method. The anti-diabetic and antioxidant effects of LP extract were examined by performing experiments in rat model of T2D.

**EXPERIMENTAL**

**Plant material and extraction**

Lemon (Citrus limon) was purchased from Shanxi University of traditional Chinese Medicine. The LP was cleaned and dried at 50 °C for 48 h. After grinding into powder, the lemon peel was extracted in 50 % water / 50 % ethanol for 48 h using a Soxhlet extractor. Water and ethanol in the resultant liquid was removed by lyophilizer. The crude extracts were stored at 4 °C.

**Experimental animals**

Healthy male Sprague-Dawley (SD) rats with weights ranging from 240 to 280 g were used in this study. All rats were raised in the same environment with a constant humidity of 50 % at 24 °C in a 12-h light/dark cycle. All animals were raised and observed in specific pathogen free laboratory animal room. The study was approved by the Animal Ethics Committee of the Affiliated Hospital of Shanxi University of Traditional Chinese Medicine (no. EA_20160083), and the experiments with rats were in full compliance with the European Communities Council Directive of 24 November 1986 (86/609/EEC) [13] and with the Guidelines laid down by the NIH in the US [14].

**Acute toxicity studies**

Acute toxicity analysis of LP extracts was performed on SD rats according to Organization for Economic Co-operation and Development (OECD) Guidelines for Testing of Chemicals No. 423. Before acute toxicity analysis, SD rats were fasted for 5 h with only water available, and then divided into four groups with eight animals in each group. The rats were administrated with LP extract at a dosage of 5, 50, 500, and 2000 mg/kg orally. The rats were observed individually every 24 h for two weeks. General behaviours, including writhing, gasping, palpitatian, decreased respiratory rate, and mortality were observed.

**Establishment of rat model of T2D and animal grouping**

To establish rat model of T2D, rats were adapted to the new environment for 7 days and then supplied with a 10 % fructose solution for 14 days. The rats were then injected with 30 mg/kg streptozotocin (STZ) in citrate buffer, which causes pancreatic β-cell dysfunction. Rats in control groups were fed with water followed by injection with citrate buffer only [15]. FBG level was measured with a glucose meter 7 days after T2D induction. Fats with a FBG greater than 250 mg/dL were considered to be diabetic.

The SD rats were divided into four groups: normal control (NC), diabetic control (DC), diabetic + low dose of LP extract (DLDLPE, 250 mg/kg), and diabetic + high dose of LP extract (DHDLPE, 500 mg/kg). LP extracts of different concentrations were administered orally to rats in the DLDLPE and DHDLPE groups via a force-feeding needle once a day continuously for 35 days after establishing T2D models. For NC and DC groups, saline was administered instead.
35 days, we recorded body weight, FBG, and food intake of SD rats before and after drug administration.

**Glucose tolerance test (GTT)**
To analyse glucose tolerance in SD rats, GTT was performed. After rats were fasted for 12 h, blood glucose was measured. The rats were administered with saline or LP extracts 1 h prior to glucose administration. After sampling blood from the tail vein, glucose was immediately intraperitoneally injected into rats at a dose of 1 g/kg. Blood glucose was measured at 0, 0.5, 1, 1.5, and 2 h after glucose administration.

**Insulin tolerance test (ITT)**
After the GTT was performed on overnight-fasted rats daily for 4 days. The rats were administered saline or LP extracts 1 h before insulin administration. After sampling blood from the tail vein, a single dose of insulin solution (0.5 U/kg) was immediately subcutaneously injected into each rat. Blood glucose was tested at 0, 0.5, 1, and 1.5 h after insulin injection.

**Histological assessment**
After 24 h fixation, pancreas tissues were dehydrated, cleared in xylene, and immersed in paraffin for embedding. The embedded tissue was sliced by microtome, dehydrated, and de-waxed. The slices were then stained with haematoxylin, washed with distilled water, decolorized using hydrochloric acid ethanol, and then stained again with eosin. Slices were then dehydrated, dried, and sealed for observation under microscope.

**Assessment of oxidative stress and antioxidant activity**
Superoxide dismutase (SOD) and malonaldehyde (MDA) extractions were performed by phosphate-buffered saline after the liver tissues was ground by liquid nitrogen grinding. SOD activity was analysed by a commercial kit (Beyotime, China) and expressed as units/mg protein. MDA content was analysed using a commercial MDA assay kit (Beyotime, China) and expressed as μmol/mg protein. The concentration of total protein was measured by Bradford protein kit (Beyotime, China) [16].

**Statistical analysis**
The data was analysed using SPSS 21 software (IBM, USA). The results are presented as mean ± standard deviation. Comparisons between two groups were performed by Student’s t-test. $P < 0.05$ indicated statistical significance.

**RESULTS**

**Acute toxicity**
In acute toxicity analysis, after 14 days of orally administering LP extract at 2000 mg/kg, any symptoms of poisoning or death were not observed, indicating that LP extract was not toxic for rats at 2000 mg/kg. Therefore, oral dosages of 250 and 500 mg/kg were safe and feasible for this study.

**Body weight, FBG, and food intake**
Body weight, FBG, and food intake was evaluated for 35 days, before and after administration of LP extract or saline. Body weight was found to decrease after establishing the rat model of T2D (Table 1). During saline administration, the body weight of diabetic rats continued to decrease. However, after treatment with LP extracts, the body weight of rats increased in the DLDLPE and DHDLPE groups. A high dose of LP extract (500 mg/kg) increased body weight more effectively than a low dose of LP extract (250 mg/kg). Food intake also increased after establishing a model of T2D in rats. After treatment with LP extracts, food intake decreased in diabetic rats. However, a high dose of LP extract did not reduce food intake more than a low dose of LP extract. The rats in the DC group had a significantly higher FBG level ($p < 0.01$) than the NC group. Continuation of treatment with LP extracts for 35 days significantly reduced blood glucose levels of diabetic rats.
Table 1: Effect of LP extract treatment on food intake and body weight before and after drug administration in STZ-induced diabetic rats. NC, Normal rats treated with vehicle alone; DC, T2D rats treated with vehicle alone; DLDLPE, T2D rats treated with a low dose of LP extracts (250 mg/kg); DHDLPE, T2D rats treated with a high dose of LP extracts (500 mg/kg).

<table>
<thead>
<tr>
<th>Parameter</th>
<th>NC</th>
<th>DC</th>
<th>DLDLPE</th>
<th>DHDLPE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Body weight (g) Before</td>
<td>269.0±5.2</td>
<td>220.0±7.7</td>
<td>224.0±6.5</td>
<td>225.0±8.7</td>
</tr>
<tr>
<td>After</td>
<td>288.0±4.5</td>
<td>170.0±5.2</td>
<td>250.0±3.8</td>
<td>265.0±5.3</td>
</tr>
<tr>
<td>Food intake (g/rat/day) Before</td>
<td>17.5±2.1</td>
<td>30.5±2.5</td>
<td>29.2±1.2</td>
<td>31.5±2.1</td>
</tr>
<tr>
<td>After</td>
<td>19.2±1.8</td>
<td>33.4±2.2</td>
<td>22.4±1.7</td>
<td>26.3±2.6</td>
</tr>
<tr>
<td>FBG (mg/dl) Before</td>
<td>100.2±5.7</td>
<td>283.8±7.5</td>
<td>280.6±15.3</td>
<td>277.7±8.6</td>
</tr>
<tr>
<td>After</td>
<td>109.8±10.5</td>
<td>288.4±12.8</td>
<td>180.6±6.5</td>
<td>155.1±12.2</td>
</tr>
</tbody>
</table>

GTT and ITT

In the GTT, the blood glucose level of rats in the NC, DLDLPE, and DHDLPE groups reached a maximum value at 0.5 h and then decreased over time. However, rats in the DC group reached a maximum blood glucose value at 1 h. In the ITT, after insulin administration, FBG initially decreased and then increased in the DLDLPE and DHDLPE groups, when compared to the DC group. After insulin administration for 2 h, the FBG level in the DLDLPE group was close to that of the DC group. In the NC and DHDLPE groups, FBG reached a minimum value 0.5 h after insulin administration. In the DC and DLDLPE groups, FBG level reached a minimum value 1 h after insulin administration.

Antioxidant activity and oxidative stress

The antioxidant activity and oxidative stress were determined by measuring MDA level and SOD activity, respectively. The increase of MDA was observed in diabetic rats, indicating that the oxidative stress in rat liver significantly increased after T2D model establishment (p < 0.05) (Figure 4A). The oxidative stress, as measured by SOD activity, was alleviated in diabetic rats after treatment with a high dose of LP extract. However, low doses of LP extract did not have the same effect on oxidative stress as high doses. The decreased SOD activity in diabetic rats indicated that antioxidant activity decreased in the liver after T2D model establishment, and that treatment with LP extract restored antioxidant activity in a dose-dependent manner (p < 0.05) (Figure 4B).

Histological features

Haematoxylin-eosin staining results showed serious damage to the intra-pancreatic islets and the number and size of islets was decreased in STZ-induced diabetic rats (Figure 3A and Figure 3B). The vacuolation and invasion of intrapancreatic tissue in the diabetic rats were also observed. After treatment with LP extract, the regeneration of intrapancreatic islet cells and enhanced restoration of intrapancreatic tissues were observed (Figure 3C and Figure 3D).
DISCUSSION

Lemon has been used as a traditional medicine, likely due to the anti-inflammatory and anti-tumour functions of flavonoids and limonene that are contained in the plant [11,12]. In this study, the anti-diabetic and antioxidant activities of extracts of lemon peel was evaluated. Type 2 diabetes is a chronic disorder of energy metabolism, which is diagnosed by hyperglycaemia with a consistently high FBG level. In this study, STZ-induced T2D rat model was used to assess the effect of treatment with LP extract on T2D. The acute toxicity of LP extract and the effect of LP extract on FBG, body weight, food intake, GTT, ITT, histological changes, and oxidative stress were tested.

Reduced body weight is a representative symptom for T2D in rats [17]. The establishment of a T2D rat model caused reduced body weight in this study. Reduction of body weight was rescued in DDLPLPE and DHDLPLPE groups, indicating that treatment with LP extracts alleviated the body weight of diabetic rats. Treatment with LP extracts also reduced food intake and FBG in diabetic rats. These results suggest that LP extracts may alleviate typical T2D symptoms. In previous studies, the anti-diabetes mechanisms of natural plant medicines were found to be involved in the maintenance of glucose homeostasis, gastrointestinal glucose absorption, insulinotropic actions, and promoting pancreatic β-cell regeneration [18-20].

Furthermore, the results of GTT and ITT showed that treatment with LP extracts improved glucose tolerance and reduced insulin resistance. Histological analysis showed that LP extracts restored the structure of and helped regenerate intra-pancreatic islets. SOD and MDA analyses indicated that treatment with LP extract alleviated oxidative stress in diabetic rats’ and restored antioxidant activities in the liver in a dose-dependent manner. It was reported that persistent hyperglycaemia destroys antioxidant balance by reducing antioxidant levels and producing reactive oxygen species [21,22], which is consistent with the result of this study (Figure 4). The results suggest that treatment with LP extract may alleviate T2D symptoms by reducing oxidative stress and restoring antioxidant activity.

CONCLUSION

The findings of this study show that treatment of type 2 diabetic rats with LP extract ameliorates T2D by reducing food intake and FBG, and increasing body weight. Glucose tolerance and insulin tolerance are also improved after LP administration. The effect of treatment with LP extract is related to increased antioxidant activity and reduced oxidative stress.

DECLARATIONS

Conflict of Interest

No conflict of interest associated with this work.

Contribution of Authors

We declare that this work was done by the authors named in this article and all liabilities pertaining to claims relating to the content of this article will be borne by the authors. Lanxiu Cao designed all the experiments and revised the paper. Juan Lv, Min Li, Rui Zhang and Fu Bai performed the experiments, and Pengfei Wei and Juan Lv wrote the paper.

REFERENCES