Beneficial effects of ginger (Zingiber officinale) on carbohydrate metabolism in streptozotocin-induced diabetic rats

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Abstract
Zingiber officinale (ZO), commonly known as ginger, has been traditionally used in the treatment of diabetes mellitus. Several studies have reported the hypoglycaemic properties of ginger in animal models. The present study evaluated the antihyperglycaemic effect of its aqueous extract administered orally (daily) in three different doses (100, 300, 500 mg/kg body weight) for a period of 30 d to streptozotocin (STZ)-induced diabetic rats. A dose-dependent antihyperglycaemic effect revealed a decrease of plasma glucose levels by 38 and 68 % on the 15th and 30th day, respectively, after the rats were given 500 mg/kg. The 500 mg/kg ZO significantly (P, 0·05) decreased kidney weight (% body weight) in ZO-treated diabetic rats v. control rats, although the decrease in liver weight (% body weight) was not statistically significant. Kidney glycogen content increased significantly (P, 0·05) while liver and skeletal muscle glycogen content decreased significantly (P, 0·05) in diabetic controls v. normal controls. ZO (500 mg/kg) also significantly decreased kidney glycogen (P, 0·05) and increased liver and skeletal muscle glycogen in STZ-diabetic rats when compared to diabetic controls. Activities of glucokinase, phosphofructokinase and pyruvate kinase in diabetic controls were decreased by 94, 53 and 61 %, respectively, when compared to normal controls; and ZO significantly increased (P, 0·05) those enzymes’ activities in STZ-diabetic rats. Therefore, the present study showed that ginger is a potential phytopharmacy for the treatment of diabetes through its effects on the activities of glycolytic enzymes.

Key words: Zingiber officinale; Carbohydrate metabolism; Diabetic rats; Tissue glycogen

Ginger rhizomes are widely used in foods for their nutritional and medicinal benefits, especially in Asia¹(1), for example, as a source of Fe and Ca for women during the post-natal period and also for treating morning sickness and other gastrointestinal disorders²(2). More recently, ginger juice was shown to have an antidiabetic effect in alloxan-induced diabetic rats¹(1). In a similar study, ginger juice was reported to cause significant reduction in the fasting glucose levels and an increase in the insulin levels in streptozotocin (STZ)-induced type 1 diabetic rats³(3). Al-Amin et al.⁴(4) also found that ginger possesses hypoglycaemic, hypocholesterolaemic and hypolipidaemic potential. They showed, in addition, that raw ginger was effective in reversing proteinuria in diabetic rats⁴(4).

Though a subsection of non-insulin-dependent diabetes mellitus patients can be managed by diet alone, most patients require an oral hypoglycaemic agent such as insulin. Insulin therapy affords effective glycaemic control, yet its shortcomings such as ineffectiveness on oral administration, short shelf-life, requirement of constant refrigeration and in the event of excess dosage, fatal hypoglycaemia, limit its usage. Treatment with sulphonylurea and biguanides is also associated with side effects⁵(5).

In addition, studies on the hypoglycaemic properties of ginger in animals have reported variable results⁴(4),⁶(6),⁷(7) and there has been no report on the effect of ginger extract on the selected glycolytic enzymes involved in carbohydrate metabolism. Therefore, the present study was undertaken to determine the effectiveness of ginger (Zingiber officinale (ZO) Roscoe) in the treatment of diabetes mellitus using animal models, by investigating its effect on the key enzymes of carbohydrate metabolism, which shows its beneficial effect in correcting the nutritional disturbances in diabetes.

Abbreviations: DCNT, diabetic control rats; LD50, lethal dose 50 %; NCNT, normal control rats; PK, pyruvate kinase; STZ, streptozotocin; ZO, Zingiber officinale.

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**Materials and methods**

**Animals**

Male Sprague–Dawley rats (body weight 250–300 g) were acclimatised inside a room at 22 ± 2°C for a period of 7 d. All animals were fed with standard rat chow in the form of pellets and animals described as fasting were deprived of food for at least 16 h but were allowed to drink filtered tap water. The standard rat chow containing carbohydrate 47%, protein 18-9% and fat 3-5% was purchased from Liaz Bhd. The total energy content of the chow was 17.7 kJ/g. All handling and management procedures were carried out in accordance with the Guidelines for the Care and Use of Laboratory Animals of the Kulliyyah of Medicine, International Islamic University Malaysia (IIUM).

**Preparation of the rhizome extract**

Fresh ginger was bought from the wet-market of Chow Kit, Kuala Lumpur; and 24 kg of the fresh ginger rhizomes were cut into pieces, air-dried and powdered. Then, 375 g of the powdered material were cold-macerated in 4 litres of distilled water and intermittently stirred thoroughly. The mixture was left at room temperature for 48 h to allow the active ingredients to be completely dissolved. The macerated pulp was first filtered by mesh cloth and then suction-filtered through Whatman no. 1 filter paper and the filtrate was freeze-dried. To increase the shelf life and uniformity, the extract was lyophilised completely by a continuous freeze-drying operation for 54 h and the yield was 25-3% (w/w), which was stored at −20°C until use.

**Acute toxicity test (LD50 determination)**

The acute toxicity test (lethal dose 50%; LD50) of the aqueous extract of ginger was determined according to the procedure described by Lorke.\(^{(3)}\) This method involved an initial dose-finding procedure, in which animals were divided into three groups of three animals per group. Doses of 10, 100 and 1000 mg/kg of ginger extract were administered through oral administration, one dose for each group. The treated animals were monitored for 24 h. Even the highest dose of 1000 mg/kg was not found to be toxic to the animals. So, the next round of toxicity test was conducted with four different doses of 800, 1600, 3200 and 6400 mg/kg, which were administered orally to four groups of one rat per group. The treated animals were again monitored for 24 h. The LD50 was then calculated as the geometric mean of the lowest dose showing death and the highest dose showing no death (Table 1).

**Induction of diabetes**

Animals were divided into five different treatment groups (Table 2). Body weights were measured after a 16 h fasting period. The animals in groups 2–5 were lightly anaesthetised with diethyl ether and were injected intraperitoneally with 65 mg/kg body weight of STZ which was freshly dissolved in citrate buffer with a pH of 4.5. After injection, they were allowed free access to food and water and were given 5% glucose solution to drink overnight to counter the hypoglycaemic shock. Then, 3 d after STZ injection, diabetes was confirmed in rats by measuring the normal fasting blood glucose levels with a glucometer (Roche Accu-Check Advantage). All rats showing a fasting blood glucose level ≥13 mmol/l were considered diabetic and selected for the experimentation.

**Experimental design**

Table 2 shows the experimental design. Normal control animals in group 1 received distilled water only. Diabetic animals in groups 3–5 received ginger extract according to the dose stipulated in the table. Diabetic control animals in group 2 received neither distilled water nor the extract. All animals received either water or the extract through oral administration.

**Sample collection**

**Fasting blood glucose level.** Fasting blood glucose in blood was measured on days 0, 7, 15, 21 and 30 during the experiment with a glucometer. Intra- and inter-assay percentage CV using ten repeated measures and ten sessions were 1.89 and 4.30%, respectively. Percentage reduction in blood glucose level was calculated as (BGC at 0 d – BGC at N day/ BGC at 0 d) × 100, where BGC = blood glucose concentration; and N = treatment days.

**Collection of tissues**

Animals were anaesthetised by exposing them to diethyl ether in an air-tight chamber. Thereafter, they were dissected, and the liver, kidney and skeletal muscles were collected in precooled normal saline, and then blotted. Tissues were weighed and finally preserved with liquid N2 and frozen at −80°C until used for the determination of metabolic changes.

**Sample preparation**

A small part of the liver tissue was cut and perfused with ice-cold 0.15 M-KCl and 1 mM-EDTA solution and homogenised with twice its weight of ice-cold buffer (0.01 cysteine and 1 mM-EDTA in 0.1 ml Tris–HCl, pH 7.4) and centrifuged (at 20000* g) for 20 min at 4°C. The supernatant was filtered and frozen at −80°C.

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**Table 1. Acute toxicity test of the aqueous extract of ginger**

<table>
<thead>
<tr>
<th>Stage</th>
<th>Doses (mg/kg)</th>
<th>No. of animals</th>
<th>No. of deaths/group</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>10</td>
<td>3</td>
<td>0/3</td>
</tr>
<tr>
<td></td>
<td>100</td>
<td>3</td>
<td>0/3</td>
</tr>
<tr>
<td></td>
<td>1000</td>
<td>3</td>
<td>0/3</td>
</tr>
<tr>
<td>II</td>
<td>800</td>
<td>1</td>
<td>0/1</td>
</tr>
<tr>
<td></td>
<td>1600</td>
<td>1</td>
<td>0/1</td>
</tr>
<tr>
<td></td>
<td>3200</td>
<td>1</td>
<td>0/1</td>
</tr>
<tr>
<td></td>
<td>6400</td>
<td>1</td>
<td>1/1</td>
</tr>
</tbody>
</table>

LD50, lethal dose 50%.

* LD50 = (3200 × 6400)^1/2 = 4525.5 mg/kg.
Table 2. Grouping of animals into five treatment groups

<table>
<thead>
<tr>
<th>Group no.</th>
<th>Treatment</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Distilled water</td>
<td>NCNT</td>
</tr>
<tr>
<td>2</td>
<td>STZ (65 mg/kg)</td>
<td>DCNT</td>
</tr>
<tr>
<td>3</td>
<td>STZ + ZO (100 mg/kg)</td>
<td>ZO (100 mg/kg)-treated diabetic rats</td>
</tr>
<tr>
<td>4</td>
<td>STZ + ZO (300 mg/kg)</td>
<td>ZO (300 mg/kg)-treated diabetic rats</td>
</tr>
<tr>
<td>5</td>
<td>STZ + ZO (500 mg/kg)</td>
<td>ZO (500 mg/kg)-treated diabetic rats</td>
</tr>
</tbody>
</table>

NCNT, normal control rats; STZ, streptozotocin; DCNT, diabetic control rats; ZO, Zingiber officinale.

Biochemical and enzymatic estimations

Assay of glucokinase (EC 2712) activity. The liver glucokinase activity was measured at 340 nm in a reaction mixture containing 75 mM-Tris buffer (pH 9.0), 27 mM-NADP, 120 mM-ATP, 600 mM-MgCl₂, 100 units/ml of glucose-6-phosphate dehydrogenase and 360 mM-glucose at 30°C. Glucose phosphorylation was then assayed by means of the glucose-6-phosphate-dependent spectrophotometric method (9). The %CV for this assay was 2.68 at 19.40 U/l and 4.10 at 28.11 U/l for the intra- and inter-assay, respectively.

Assay of fructose-6-phosphate kinase (EC 27111) activity. Assay of fructose-6-phosphate kinase was carried out in a reaction mixture of 100 mM-MgSO₄, 5000 units/ml of lactic dehydrogenase, 140 mM of a mixture of MgSO₄–KCl, 30 mM-phosphoenol pyruvate and 120 mM-MgCl₂, 100 units/ml of glucose-6-phosphate dehydrogenase and 360 mM-glucose at 30°C. Glucose phosphorylation was then assayed by means of the glucose-6-phosphate-dependent spectrophotometric method (9). The %CV for this assay was 5.39 at 8.30 U/l for the ten intra- and inter-assays.

Assay of pyruvate kinase (EC 27140) activity. The sample was added to a reaction mixture containing 100 mM-potassium phosphate buffer, pH 7.6 at 37°C, 1.3 mM-NADH, 44 mM-ADP, 100 mM-MgSO₄, 5000 units/ml of lactate dehydrogenase and 17 mM-phosphoenol pyruvate freshly prepared. PK assays were then conducted with a thermostated recording spectrophotometer by coupling the enzyme activity to lactate dehydrogenase, using the procedure reported by Bergmeyer et al. (11).

Estimation of tissue glycogen level

Tissue glycogen level was measured using the glycogen assay kit (Bio Vision) based on hydrolysis of glycogen to glucose by glucoamylases, which is then specifically oxidised to produce a product that reacts with oxy-red probe to generate a colour that reacts with oxy-red probe to generate a colour that reacts with oxy-red probe to generate a colour. The precision as estimated by %CV for intra- and inter-assay was 1.66 and 2.60%, respectively, at 82.54 mg/ml.

The protein contents of the tissue homogenates were estimated following the standard method of Lowry et al. (12).

Data and statistical analysis

The data are presented as mean values with their standard deviations or errors. The statistical analysis was carried out using the software SPSS (version 11; SPSS, Inc.). Groups were compared using one-way ANOVA and Tukey’s post hoc test was performed for multiple comparisons. Statistical significance was set at \( P < 0.05 \).

Results

The safety limit (LD₅₀) of ginger extract in rats

The percentage yield for the extract was 25.3% (w/w) as mentioned under ‘Preparation of the rhizome extract’. The oral LD₅₀ was estimated to be 4525.5 mg/kg in rats (Table 1), accordingly:

\[
LD_{50} = (3200 \times 6400)^{1/2} = 4525.5 \text{ mg/kg.}
\]

Ginger reduced blood glucose levels in streptozotocin-induced diabetic rats

The basal levels of blood glucose of the rats in all groups before STZ injection were not significantly different. However, 72 h after STZ injection, blood glucose levels were significantly higher in the STZ-induced rats. In contrast, normal control rats (NCNT) remained persistently euglycaemic throughout the course of the study (Table 3).

Table 3 also shows that ZO extract has a dose-dependent anti-hyperglycaemic effect on the treated diabetic rats. Injection of STZ (65 mg/kg) led to over 4-fold elevation of blood glucose levels (\( P < 0.05 \)) which were maintained over a period of 4 weeks in the DCNT. On the 7th day of treatment, DCNT showed a significant increase (\( P < 0.05 \)) in blood glucose level as compared to NCNT. The ZO-treated (100, 300 and 500 mg/kg) diabetic groups had their blood glucose significantly lowered (\( P < 0.05 \)) when compared to DCNT, with 10.63, 25.71 and 25.14% reduction, respectively, on the 7th day. Again, there were significant reductions in blood glucose levels between the DCNT and (100, 300 and 500 mg/kg) ZO-treated diabetic groups, with 10.63, 25.71 and 25.14% reduction, respectively, on the 7th day. After the 4th week of daily treatment with the aqueous extract of ZO, ginger led to a fall in blood glucose levels by 48.30, 62.43 and 67.85% in the 100, 300 and 500 mg/kg-treated groups, respectively. The maximum anti-hyperglycaemic effect was observed in the 500 mg/kg-treated group, which was not significantly different when compared with NCNT.

Ginger improved body, liver and kidney weights of streptozotocin-induced diabetic rats

Fig. 1 shows the effect of ZO extract (500 mg/kg) on body weight of STZ-induced diabetic rats. The diabetic control group (DCNT) did not gain any significant weight during the 30d experimental period while NCNT and ZO
Beneficial effect of ginger in diabetic rats

Ginger lowered kidneys’ glycogen content with no effect on liver and skeletal muscle

Kidney glycogen content increased by over 10-fold while liver and skeletal muscle glycogen content significantly \( (P<0.05) \) decreased by 75 and 53\% in DCNT \( v. \) NCNT. There was no significant difference in the glycogen content of liver and muscle in the ginger-treated diabetic group, although kidneys showed a significant decrease in glycogen content due to ginger treatment (Fig. 2).

**Ginger increased the activities of hepatic glycolytic enzymes in streptozotocin-induced diabetic rats**

The DCNT showed significant decrease by 94, 53 and 61\% in the activities of glucokinase, phosphofructokinase and PK (Fig. 3) compared to NCNT values; however, ZO (500 mg/kg) significantly \( (P<0.05) \) increased these enzymes’ activities when compared to DCNT.

**Discussion**

**Safety limit (LD\textsubscript{50}) of ginger extract in rats**

The high yield of the extract shows that optimal extraction of the constituents requires the use of polar solvents, in consonance with folkloric use of the aqueous infusions. The oral LD\textsubscript{50} value of 4525.5 mg/kg in rats, obtained for the aqueous ZO extract falls within the practically non-toxic range\(^{14} \). The anti-hyperglycaemic effect observed for all doses (100, 300 and 500 mg/kg) significantly \( (P<0.05) \) increased these enzymes’ activities when compared to DCNT.

**Anti-hyperglycaemic effect of ginger in streptozotocin-induced diabetic rats**

Diabetes mellitus is characterised by chronic hyperglycaemia with disturbances of carbohydrate, fat and protein metabolism

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**Table 3. Changes in the blood glucose levels (mmol/l) in streptozotocin-induced (65 mg/kg) diabetic rats over a period of 30 d administration of ginger extracts Zingiber officinale (ZO)**

<table>
<thead>
<tr>
<th>groups</th>
<th>basal day</th>
<th>0d</th>
<th>1st day</th>
<th>7th day</th>
<th>15th day</th>
<th>21st day</th>
<th>30th day</th>
</tr>
</thead>
<tbody>
<tr>
<td>NCNT</td>
<td>Mean</td>
<td>4.41</td>
<td>0.50</td>
<td>4.36</td>
<td>0.42</td>
<td>2.24</td>
<td>0.69</td>
</tr>
<tr>
<td>DCNT</td>
<td>Mean</td>
<td>4.46</td>
<td>0.29</td>
<td>2.26</td>
<td>0.24</td>
<td>3.16</td>
<td>0.36</td>
</tr>
<tr>
<td>100 mg/kg ZO</td>
<td>Mean</td>
<td>4.47</td>
<td>0.36</td>
<td>1.98</td>
<td>0.44</td>
<td>0.69</td>
<td>0.26</td>
</tr>
<tr>
<td>300 mg/kg ZO</td>
<td>Mean</td>
<td>4.47</td>
<td>0.36</td>
<td>2.00</td>
<td>0.44</td>
<td>0.69</td>
<td>0.26</td>
</tr>
<tr>
<td>500 mg/kg ZO</td>
<td>Mean</td>
<td>4.47</td>
<td>0.36</td>
<td>2.00</td>
<td>0.44</td>
<td>0.69</td>
<td>0.26</td>
</tr>
</tbody>
</table>
Ginger contains potential bioactive compounds; mostly of volatile oils\(^\text{(17,18)}\). \([6]\)-Gingerol is the most abundant bioactive compound which has been consistently implicated in the hypoglycaemic and antihyperglycaemic effects of ginger. Akhani et al.\(^\text{(19)}\) have reported that ginger juice exhibits hypoglycaemic activity in both normal and STZ-induced diabetic rats. Al-Amin et al.\(^\text{(20)}\) also found that ginger possesses hypoglycaemic, hypercholesterolaemic and hypolipidaemic potential\(^\text{(21)}\). They also showed that raw ginger is effective in reversing the diabetic proteinuria observed in diabetic rats. More recently, Asha et al.\(^\text{(22)}\) and Jafri et al.\(^\text{(23)}\) reported the antihyperglycaemic and hypoglycaemic activities of ginger in alloxan-induced diabetic rats. Clearly, the results in the present study confirm the observations in the recent studies pertaining to the glucose-lowering effect of ginger in blood. No previous studies have reported changes in tissue glycogen and the selected glycolytic enzymes involved in carbohydrate metabolism, as a result of ginger administration.

### Ginger ameliorates body, liver and kidney weights

STZ-induced diabetes is characterised by severe loss in body weight\(^\text{(24)}\) and this might be due to loss or degradation of structural proteins, as the structural proteins are known to contribute to body weight. In our study, during the experimental period, NCNT showed an approximately 20% gain in body weight. On the other hand, DCNT showed no significant gain in body weight over the same time period. ZO-treated rats showed higher and significant gain in body weight in comparison to both DCNT and NCNT.

### Table 4. Effect of 30 d administration of *Zingiber officinale* (ZO) extract (500 mg/kg) on liver weight (LW) and kidney weight (KW) in streptozotocin (65 mg/kg) diabetic rats (Mean values with their standard errors)

<table>
<thead>
<tr>
<th>Groups</th>
<th>Absolute LW (g)</th>
<th>LW/100 g BW</th>
<th>Absolute KW (g)</th>
<th>KW/100 g BW</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SEM</td>
<td></td>
<td>Mean</td>
</tr>
<tr>
<td>NCNT</td>
<td>8·53</td>
<td>0·06</td>
<td>2·97</td>
<td>0·08</td>
</tr>
<tr>
<td>DCNT</td>
<td>8·33</td>
<td>0·07</td>
<td>3·47(^*)</td>
<td>0·14</td>
</tr>
<tr>
<td>ZO (500 mg/kg)</td>
<td>8·83</td>
<td>0·10</td>
<td>3·23</td>
<td>0·12</td>
</tr>
</tbody>
</table>

LW, body weight; NCNT, normal control rats; DCNT, diabetic control rats. * Mean values were significantly different compared with NCNT (P<0·05).

\(^{†}\) Mean values were significantly different compared with DCNT (P<0·05).

Fig. 1. Body weights of streptozotocin (STZ)-induced diabetic rats treated with an aqueous extract of ginger over the experimental period. Weights were measured in normal rats (□), STZ-induced diabetic rats (■) and ginger-treated STZ-induced diabetic rats (500 mg/kg *Zingiber officinale*; □). The animals were weighed after STZ injection on 0, 15 and 30 d of experimental period. Values are means, with their standard errors represented by vertical bars. * Mean value was significantly decreased compared with normal control at days 15 and 30 (P<0·05). † Mean value was significantly different between diabetic control and ginger-treated diabetic rats (P<0·05).
Fig. 2. Tissues' glycogen was estimated in streptozotocin (STZ)-induced diabetic rats ( ), STZ-induced diabetic rats ( ), and ginger-treated STZ-induced diabetic rats (500 mg/kg Zingiber officinale; ). Tissues' glycogen was estimated at the end of the experimental period. Values are means, with their standard errors represented by vertical bars. * Mean value was significantly different compared with normal control (P<0.05). † Mean value was significantly decreased compared with diabetic control (P<0.05).

Fig. 3. Hepatic glycolytic enzymes' activity of streptozotocin (STZ)-induced diabetic rats treated with an aqueous extract of ginger over the experimental period was estimated. Enzymes' activity was measured in normal rats ( ), STZ-induced diabetic rats ( ), and ginger-treated STZ-induced diabetic rats (500 mg/kg Zingiber officinale; ). Enzymes' activity was analysed at the end of the experimental period. Values are means, with their standard errors represented by vertical bars. * Mean value was significantly decreased compared with normal control (P<0.05). † Mean value was significantly increased compared with diabetic control (P<0.05). GK, glucokinase; PFK, phosphofructokinase; PK, pyruvate kinase.

Ginger has no significant effect on liver and muscle but kidney glycogen content

The liver plays an important role in buffering postprandial hyperglycaemia and is involved in the synthesis of glycogen. Glycogen is the primary intracellular storable form of glucose and its level in various tissues, especially skeletal muscle, is a direct reflection of insulin activity, as insulin promotes intracellular glycogen deposition by stimulating glycogen synthase and inhibiting glycogen phosphorylase. Since STZ causes selective destruction of β-cells of islets of Langerhans, resulting in marked decrease in insulin levels, it is rational that glycogen levels in insulin-dependent tissues (skeletal muscle and liver) decrease as they depend on insulin for the influx of glucose(37–39). Results showed that liver and skeletal glycogen content decreased drastically in DCNT by approximately three-quarters of their basal levels. This has also been reported earlier(40). ZO showed a trend towards an increase in glycogen content but could not significantly increase the glycogen content in muscle and liver.

Ginger increased the activities of hepatic glycolytic enzymes in streptozotocin-induced diabetic rats

In the present study, the activities of glycolytic enzymes in the liver were significantly decreased (P<0.05) in the DCNT. This confirms the previous findings(29,31,42) that relative deficiency of insulin in the STZ model of type 1 diabetes causes suppression of glucokinase, phosphofructokinase and PK activities. These hepatic glycolytic enzymes are the regulatory factors which drive the metabolic degradation of glucose to form pyruvate. The high-energy compounds ATP and NADH are formed from the free energy released in this process. Maximum suppression was observed in hepatic glucokinase activity, followed by hepatic PK and phosphofructokinase activities. Administration of ZO extract increased significantly the activity of all the three enzymes towards NCNT, suggesting that the anti-hyperglycaemic action observed was as a result of increased glucose utilisation at the liver as well as skeletal muscle. However, it is not possible to deduce from the present findings that the increase in glycolytic enzymatic activity seen in the ZO-treated rats occurred secondary to the ZO-mediated release of insulin or whether a component of ZO had insulin mimetic action. Since STZ diabetes is an insulin-deficient...
model, the likelihood of insulinomimetic effect seems more possible.

Conclusion
We have demonstrated that the folk medicinal plant ginger is practically non-toxic in rats, with the high margin of safety exhibited by the extract in the present study. The findings also suggest that the administration of ZO rhizome extract to diabetic rats gives good control over tissue glycogen content by enhancing the peripheral utilisation of glucose and correcting the impaired liver and kidney glycolysis and by limiting its gluconeogenic formation similar to insulin. Therefore, raw ginger has significant potential as a phytomedicine in the treatment of diabetes.

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References