ملخص البحث

تأثير مسحوق الخروب على سكر الدم وتجزؤات الدهون في فئران الألبينو المصابة بالسكر

كان الهدف الأساسي من البحث الحالي هو دراسة تأثير مستويين من مسحوق الخروب 10% و20% على سكر الدم وتجزؤات الدهون وقد تم تقسيم ذكور الفئران الأربعون إلى مجموعتين.

المجموعة الأولى الضابطة وتتكون من عشرة فئران تم تغذيتها على الوجبة الأساسية والمجموعة الرئيسية وتتكون من ثلاثين فأراً تم حقنهم في البطين بجرعة الألوكسان (1700 ملجم / كجم) من وزن الجسم لكل فأر. كما تم تقسيم المجموعة الثانية الرئيسية إلى ثلاث مجموعات فرعية.

المجموعة الأولى الفرعية: مجموعة مصابة بالسكر وتتغذى على الوجبة الغذائية الأساسية. المجموعة الفرعية الثانية: مجموعة مصابة بالسكر وتتغذى على الوجبة الغذائية الأساسية مضافة 10% من مسحوق الخروب والمجموعة الفرعية الثالثة: مجموعة مصابة بالسكر وتتغذى على الوجبة الغذائية الأساسية مضافة 20% من مسحوق الخروب وذلك لمدة ستة أسابيع.

وفي نهاية التجربة تم تجميع عينات الدم وإجراء التحليل الكيميائي التالى عليها:

- تقدير سكر الدم والكولسترول الكلي والجلطيريدات الثلاثية وكولسترول البروتين متوقع الكثافة ودبلوميبرول البروتين منخفض الكثافة وكولسترول البروتين منخفض الكثافة جداً.

وقد أظهرت النتائج تحسناً ملحوظاً في سكر الدم وتجزؤات الدهون في الفئران التي تم تغذيتها على 10% و20% من مسحوق الخروب.

الكلمات المفتاحية:

مسحوق الخروب. الفئران. سكر الدم. تجزؤات الدهون. الغذاء الماكول. وزن الجسم

المكتسب.
Abstract

The Influence of Carob Powder on Serum Glucose and Lipid Profile in Albino Induced Diabetic Rats

The present study aimed to investigate the effect of two levels of carob powder (10% and 20%) on serum glucose and lipid profile. Forty male Sprague-Dawley rats were divided into two groups. The First group control negative group consisted of (10) rats fed on basal diet. The second main group consisted of (30) rats were injected intraperitoneal with alloxan monohydrate in single dose (170 mg/Kg) body weight. The second main group was divided into three subgroups (10 rats each). Subgroup (1); diabetic group was fed on basal diet only control positive group, Subgroup (2); diabetic group was fed on basal diet plus 10% carob powder and subgroup (3); diabetic group was fed on basal diet plus 20% carob powder for six weeks. At the end of the experimental period, serum samples was collected to measure serum glucose (SG), total cholesterol (TC), triglyceride (TG), high-density lipoproteins (HDL), low-density lipoproteins (LDL) and very-low-density lipoproteins (VLDL). The obtained results revealed that feeding rat's 10% and 20% carob powder improved serum glucose and lipid profile parameters in the experimental rats.

Key words:
carob powder, rats, serum glucose, lipid profile, feed intake, body weight gain.


References:
Bursell, S-E.; Clermont, AC. ;Aiello, LP. ;Aiello, LM.; Schlossman, DK. and Feener, EP. (1999): High-dose vitamin E supplementation normalizes retinal blood flow and creatinine clearance in patients with type 1 diabetes. Diabetes Care; (22): 1245-1251.
Table (4) revealed that the HDL in subgroups (2) and (3) was increased when compared with subgroup (1) control positive group while, the serum LDL and VLDL were decreased in subgroups (2) and (3) when compared with subgroup (1) especially at the end of the experiment. This results agree with Oven et al.,(2003). This might be due to fat soluble tocopherol is among antioxidants found in (ceratonia siliqua L.) Williams, et al., (1980). This compound prevents oxidation of unsaturated fatty acids; thereby changes cholesterol LDL-C,HDL-C and triglyceride plasma levels Packer, (1991). Alpha-tocopherol can reduce HDL – C and LDL- C concentrations in diabetic rats Mokhtari et al.,(2011).

In conclusion

The results showed that the 20% was better than 10% carob powder and decreased the serum glucose and the lipid profile so we recommend utilizing it in the hyperglycemic and hypercholesterolemic patient's diets.
Table (3) revealed that the total cholesterol and triglycerides in subgroups (2) and (3) was decreased when compared with control positive group in both. This results agree with Burget et al., (2007) who found that carob fibers and polyphones have been shown to exert beneficial effects on metabolic parameters (cholesterol, triglycerides), this might be carob fiber contains water soluble and water insoluble polyphenols which exhibit considerable natural antioxidative activity and can contribute to a amoe favorable balance between free radicals (oxidants) and antioxidants Oven et al.,(2003).

Manisha et al., (2000) ; Tabatabai and Li (2000) concluded that usage of fiber decreases total cholesterol, LDL-C and triglyceride levels, while HDL-C level showed no significant difference. They declared that the reason for this decline is an increase in bile acid due to fiber intake, which reduces cholesterol uptake.

Zunft et al., (2003) demonstrated that daily consumption of food products enriched with carob fiber shows beneficial effects on human blood lipid profile and may be effective in prevention and treatment of hypercholesterolemia.

4.4. The effect of carob powder on HDL, LDL and VLDL

Table (4) revealed that the LDL-cholesterol and VLDL-cholesterol in (control positive) Subgroup (1) were significantly increased (p<0.01). While, the mean value of HDL-cholesterol was significantly decreased (p<0.01) when compared with control negative group. The present data agree with Mokhtari et al., (2011).

<table>
<thead>
<tr>
<th>Parameter groups</th>
<th>HDL-cholesterol (mg/dl)</th>
<th>LDL-cholesterol (mg/dl)</th>
<th>VLDL-cholesterol (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>End of the 3rd week</td>
<td>End of the 6th week</td>
<td>End of the 3rd week</td>
</tr>
<tr>
<td>Control negative group</td>
<td>47.03±1.49</td>
<td>48.12±1.55</td>
<td>47.69±1.74</td>
</tr>
<tr>
<td>Control positive Subgroup (1)</td>
<td>37.72±1.12</td>
<td>40.77±0.87</td>
<td>86.28±1.44</td>
</tr>
<tr>
<td>10% carob powder Subgroup (2)</td>
<td>42.25±1.14</td>
<td>44.03±0.84</td>
<td>71.39±1.55</td>
</tr>
<tr>
<td>20% carob powder Subgroup (3)</td>
<td>44.39±1.86</td>
<td>46.29±1.85</td>
<td>68.33±1.48</td>
</tr>
</tbody>
</table>

Mean±SE, SE= Standard Error, Values followed by the same letter within the same column were not significantly different (P<0.01).
The consumption of high fiber foods can decrease plasma glucose concentration. Fibers are mainly carbohydrates produced by plants. However, human lack necessary enzyme to digest them. Chemical structure of polysaccharides fibers changes some features of digestive tract such as PH, folding and ionic charge. Adhesive composition of polysaccharides increases food particles adhesion which then, delays stomach offloading on one hand, and prevents sugar uptake by small intestine on the other Yukiko (2000). Lowering of serum glucose concentration can decrease insulin level.

Carbohydrates could modify blood glucose concentration through effecting the uptake of materials and influencing fermentation in large intestine Wolever (2003). Similarly, food fibers can decrease blood glucose level by releasing insulin from liver and by elevating receptors sensitivity to insulin Tabatabai and Li (2000). In addition, it has been suggested that dietary carbohydrates play an important role in glucose homeostasis; there fore, they can control diabetes symptoms Alan and Becky (1981).

Tocopherol has positive effects on controlling metabolic processes in diabetic patients; this effect is due to its antioxidant effect on protein glycosylation and insulin sensitivity O’Connel (2001). Furthermore, long term usage of tocopherol decreases blood sugar in diabetic and healthy people Ble et al.,(2005). There are some information about lowering plasma ALT by tocopherol and it is possible that simultaneous decline of glucose and insulin levels are the results of improvement in liver cells functions as well as decreasing liver resistance to insulin and glucose production Manning et al., (2004). Tocopherol also in improves insulin function through its effects on plasma membrane Bursell (1999).

4.3. The effect of carob powder on serum cholesterol and triglycerides

Table (3) revealed that the total cholesterol and triglycerides in control positive subgroup (1) were increased significantly (p<0.01) when compared with control negative group. The present data agree with Mokhtari et al., (2011).

<table>
<thead>
<tr>
<th>Parameter groups</th>
<th>Cholesterol (mg/dl)</th>
<th>Triglycerides (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>End of the 3rd week</td>
<td>End of the 6th week</td>
</tr>
<tr>
<td>Control negative group</td>
<td>105.3±0.71</td>
<td>107.8±1.70</td>
</tr>
<tr>
<td>Control positive Subgroup (1)</td>
<td>137.6±0.74</td>
<td>139.7±2.11</td>
</tr>
<tr>
<td>10% carob powder Subgroup (2)</td>
<td>125.7±0.53</td>
<td>117.6±3.03</td>
</tr>
<tr>
<td>20% carob powder Subgroup (3)</td>
<td>123.8±0.54</td>
<td>113.6±2.47</td>
</tr>
</tbody>
</table>

Mean±SE, SE= Standard Error,Values followed by the same letter within the same column were not significantly different (P<0.01).
The result given in Table (1) revealed that body weight gain showed significant differences among all studied groups at (p < 0.01) in third and sixth weeks in experimental period. By the end of third and sixth weeks the data recording (35.24±0.11) and (59.92±0.32) for control negative group of experimental rats, meanwhile, the body weight gain in subgroups (1, 2, 3) decreased significantly at (p < 0.01). the third weeks recording (11.99±0.21, 14.73±0.23 and 17.40±1.87) while the body weight gain in sixth weeks recording (29.76±1.56, 31.15±0.34 and 33.01±0.21); respectively.

4.2. The effect of carob powder on serum glucose in all studied groups.

Diabetes is considered one of the most prevalent diseases of endocrine system. As a result of this disease, the normal body metabolic functions become disordered. In spite of induced hyperglycemia, many body cells are not able to uptake glucose for nourishment Hayashi (1998).

The data in the Table (2) was decreased significantly (p<0.01) at serum glucose in all studied groups when compared with control positive group, especially subgroup (3) which fed on basil diet plus 20% carob powder.

Table (2) Serum glucose in all studied groups

<table>
<thead>
<tr>
<th>Parameter groups</th>
<th>Serum glucose</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>End of the 3rd week</td>
</tr>
<tr>
<td>Control negative group</td>
<td>107.7d±1.44</td>
</tr>
<tr>
<td>Control positive</td>
<td></td>
</tr>
<tr>
<td>Subgroup (1)</td>
<td>281.1ab±2.68</td>
</tr>
<tr>
<td>10% carob powder</td>
<td></td>
</tr>
<tr>
<td>Subgroup (2)</td>
<td>278.2bc±3.90</td>
</tr>
<tr>
<td>20% carob powder</td>
<td></td>
</tr>
<tr>
<td>Subgroup (3)</td>
<td>273.5bc±1.12</td>
</tr>
</tbody>
</table>

Mean±SE, SE= Standard Error,Values followed by the same letter within the same column were not significantly different (P < 0.01).

The results given in Table (2) revealed that the serum glucose showed significant differences among all the studied groups and there combination with different levels (10% and 20%) carob powder groups were (p<0.01) in experimental period.

The serum glucose decreased significantly at (p < 0.01) for subgroups (2) and (3) were recording (278.2±3.90 mg/dl and 273.5±1.12 mg/dl) for experimental rats (treated group fed on basal diet plus (10% and 20%) carob powder.

This might be due to that carob powder reduced glucose percent in blood due to its content of high percent fiber, carbohydrates and tocopherol Kamal et al., (2013).
density lipoprotein cholesterol (VLDL) were calculated according to the equation of Friedewald, et al., (1972).

3.1.5. Statistical Analysis

Data were analyzed applying T-test using SPSS program version 16 and the data was analyzed with analysis of variance (ANOVA) procedures by using the MSTAT-C Statistical software package Russell (1983). Where the F-test showed significant differences among means Duncan multiple range test performed at the 0.05 level of probability to separate means.

4. Results and Discussion:

4.1. The effect of carob powder on Feed intake (g) and body weight gain (%)

The results given in Table (1) revealed that feed intake showed significant differences among all studied groups at (p < 0.01) in third and sixth weeks in experimental period. By the end of third and sixth weeks, the data showed no significant differences between control negative group and control positive subgroup (1). However, in the third weeks the data showed significant differences between control negative group recording (17.89±0.53 g/rat) and subgroup (2) and (3) including feed intake (10% and 20%) carob powder; respectively. Which increased significantly at (p < 0.01) recording (19.42±0.69 g/rat) and (19.86±0.59 g/rat); respectively. On the other hand, the data of feed intake showed significant differences between control positive subgroup (1) and subgroups (2 and 3). However, in the sixth weeks the data showed significant differences between control negative group recording (18.91±0.44 g/rat) and subgroup (2) and (3); respectively. Which increased significantly at (p < 0.01) recording (20.32±0.54 g/rat) and (20.92±0.44 g/rat); respectively. Likewise, there were significant differences in feed intake between subgroup (1) and subgroups (2) and (3).

There were significant differences in feed intake between subgroup (2) and (3). However, subgroup (3) increased significantly at (p < 0.01).

Table (1) Feed intake (g) and body weight gain (%) in all studied groups.

<table>
<thead>
<tr>
<th>Parameter groups</th>
<th>Feed intake</th>
<th>Body weight gain</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>End of the 3rd week</td>
<td>End of the 6th week</td>
</tr>
<tr>
<td>Control negative group</td>
<td>17.89±0.53</td>
<td>18.91±0.44</td>
</tr>
<tr>
<td>Control positive Subgroup (1)</td>
<td>18.09±0.38</td>
<td>19.06±0.23</td>
</tr>
<tr>
<td>10% carob powder Subgroup (2)</td>
<td>19.42±0.69</td>
<td>20.32±0.54</td>
</tr>
<tr>
<td>20% carob powder Subgroup (3)</td>
<td>19.86±0.59</td>
<td>20.92±0.44</td>
</tr>
</tbody>
</table>

Mean±SE, SE= Standard Error, Values followed by the same letter within the same column were not significantly different (P < 0.01).
Dulloo et al., (1999) and in mice Klaus et al., (2005). Therefore, carob fiber may exert beneficial effects on postprandial lipid metabolism and substrate utilization potentially related to the secretion of gut hormones.

The aim of this study was to investigate the influence of carob powder on serum glucose and lipid profile in Albino rats suffering from diabetes.

2. Materials and Methods:
2.1. Materials:

Five Kg. of carob powder (Ceratonia Siliqua) were obtained from Aswan Governorate where carob is cultivated. Samples were stored at 18°C and analyzed within 2 months.

3. Methods:
3.1. Biological Experiment
3.1.1. Experimental Animals

Forty adult male white albino rats (Sprague dawley strain) weighing between 100 and 120 grams were obtained from the animal house of the Faculty of Medicine, Assiut University. The animals were housed as groups in wire cages under the normal laboratory conditions and were fed on basal diets for a week as adaptation period.

3.1.2. Preparation of the Basil Diet

The basal diet was prepared according to Ilwy (2003). The salt mixture used according to Anon (1977). The vitamin mixture used according to Anon (1980).

3.1.3. Experimental Design:

The rats were randomly divided into two groups. The First group control negative group consisted of (10) rats and was fed on basal diet. The second main group consisted of (30) rats and were injected intraperitoneal with alloxan monohydrate. (C₄H₂N₂O₄.H₂O) in single dose of (170 mg/Kg) body weight according to Pang et al., (1985). The second main group was divided into three subgroups (10 rats each) Subgroup (1); diabetic group was fed on basal diet only (control positive group), Subgroup (2); diabetic group was fed on basal diet plus 10% carob powder and subgroup (3); diabetic group was fed on diet plus 20% carob powder for six weeks.

3.1.4. Blood Sampling:

At the end of the experiment, rats were fasted overnight and anesthetized. Blood samples were collected from all animals from the retro-orbital plexus. Serum samples were used for determination of serum glucose Tietzy (1995) total cholesterol (TC) Allian et al., (1974), triglycerides (TG) Fossati and Prancipel (1982), high density lipoprotein cholesterol (HDL) Warnick et al., (1983), low density lipoprotein cholesterol (LDL) and very low
The Influence of Carob Powder on Serum Glucose and Lipid Profile in Albino Induced Diabetic Rats

Moshera M. El-Manfaloty*  Hend M. Ali**

1. Introduction:

Diabetes Mellitus is an endocrine disease that causes disorders in carbohydrate, lipid and protein metabolism. This disease is defined by hyperglycemia which is brought about by a deficiency in insulin production and or by resistance to it David (1993). Hyperglycemia can give rise to other disorders in eyes, kidney, vessels and nervous system Sharma (1993).

At present, over 150 million persons suffer from hyperglycemia throughout the world; its predicted that this number will be about 366 millions in 2030 Wild et al., (2004). The most important and indeed the main method for treating diabetes is the used of insulin and hypoglycemic drugs however, these substances have various unfavourite side effect Suji and Sivakami (2003). Medicinal herbs can have applications in treatment of many diseases such as diabetes but their effectiveness has not been investigated properly and needs to be validated Shapiro and Gong (2002). One of these medicinal plants is carob tree (Ceratonia Siliqua) from fabaceae family whice has a height of 7-12m. this evergreen tree is often monoecious and has pinnate leaves and raceme flowers Mozaffarian (1996). (Ceratonia Siliqua) is native to Mediterranean area. In some area, carob seeds is used like tea and coffee Mirhaydar (1994). In fact, it is a suitable replacement for cocoa because it lacks caffeine and theobromine.

Williams et al.,(1980) and Jim (2005) reported that (Ceratonia Siliqua) is useful to treat and improve diabetes symptoms because it has compounds such as fibers, phytosterols and tocopherol. Kritchevsky (1982) reported that fibers are able to bind to cholesterol and phospholipids thereby interferes in their uptake.

In humans, consumption of carob fiber was shown to have a high antioxidant capacity Kumazawa et al., (2002) lower serum cholesterol and serum triglycerids Zunft et al., (2003). Furthermore, other studies showed that polyphenols may increase fat oxidation and energy expenditure in humans.

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