PROTECTIVE EFFECT OF SOME PLANT EXTRACTS ON HYPERURICEMIA IN EXPERIMENTAL ANIMALS

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ABSTRACT

Hyperuricemia (elevated serum levels of uric acid) is a key risk factor for the development of gout, and has been linked to renal dysfunction, cardiovascular diseases, hypertension, diabetes and metabolic syndrome. Hyperuricemia was induced by oxonic acid (uricase inhibitor) in experimental rats to evaluate the protective effect of alcoholic extracts from parsley shoots, celery seeds and fig leaves. The rats were divided into 6 groups, and the first one served as a normal control group. Three groups of rats were given various plant extracts (celery, parsley and fig) by oral administration using a stomach tube at a dose of 250 mg/kg. A positive control group of rats was administered orally the hypouricemic drug, allopurinol (xanthine oxidase inhibitor) at a dose of 100 mg/kg. A negative control group did not receive any plant extracts or drugs. The various plant extracts and the drug were administered for the rats every day for 9 days. On the 10th day, all groups except the normal control group received a single dose of oxonic acid (250 mg/kg) by intraperitoneal injection to induce hyperuricemia. After two hours of hyperuricemia induction by oxonic acid injection, blood samples were collected from all rat groups. The protective effects of various plant extracts were monitored through measurement of uric acid and other blood biochemical analyses for the rats as well as assay of xanthine oxidase enzyme in liver tissues. The data indicated a significant (P<0.05) increase in the levels of uric acid and other kidney function tests near to their normal values which appeared in the normal control group. The different plant extracts exhibited protective effects against hyperuricemia in variant efficacies compared to allopurinol. These efficacies were in the following order: fig > allopurinol > celery ≈ parsley. Comparatively, the different plant extracts exhibited inhibitory effects on the activity of liver xanthine oxidase enzyme in variant efficacies compared to allopurinol. These efficacies were in the following order: allopurinol > fig > celery ≈ parsley. It can be noticed that fig extract was the most effective treatment against hyperuricemia while allopurinol was the strongest inhibitor against xanthine oxidase activity.

Keywords: Allopurinol, Celery, Fig, Hyperuricemia, Oxonic acid, Parsley, Rats, Uric acid, Xanthine oxidase.

INTRODUCTION

Uric acid is produced by purine metabolism. When adenine is catabolized, adenosine is converted to inosine, and then to hypoxanthine, which is in turn oxidized by xanthine oxidase (XO) to generate xanthine, and then uric acid. The catabolism of guanine starts with conversion to xanthine and then to uric acid, with the second step also being catalyzed by XO (Cleland et al 1995). In many mammals, uricase converts uric acid into allantoin, which has higher solubility and no obvious adverse effect (Sherman et al 2008). Uric acid is the end product of nucleic acid metabolism in human with the loss of uricase. Uric acid is formed from nucleic acid either endogenously from cell breakdown or exogenously from metabolism of uric acid produced by the liver.
The amount of urate in the blood depends on the dietary intake of purines, urate biosynthesis, and the rate of urate excretion. Over-production or under-excretion of uric acid leads to hyperuricemia (Wang et al. 2008).

Hyperuricemia is a key risk factor for the development of gout, and has been linked to renal dysfunction, cardiovascular diseases, hypertension, diabetes and metabolic syndrome (Choi and Ford, 2007; Johnson et al. 2005; Short and Tuttle, 2005). Hyperuricemia is a common metabolic disorder in human. It occurs as a result of over-production of uric acid and impaired renal uric acid excretion (Terkeltaub, 2003). Increasing in uric acid level also leads to form monosodium urate monohydrate (MSU) crystals in the joints, causing gout, and in the kidney, predisposing to urate nephrolithiasis (Merry et al. 2007; Preitner et al. 2009). Recently, hyperuricemia linked to cardiovascular diseases (Jin et al. 2012).

Gout is one of the most common metabolic disorders with a worldwide distribution and continues to be a major health problem. It affects around 13% of men and 5% of the women (Arromede et al. 2002). Gout is a metabolic disease caused by long-term purine metabolic disorders and elevated uric acid. Gout is characterized by an excessive concentration of uric acid in the blood, causing the accumulation of monosodium urate crystals in the joints and kidneys leading to acute gouty arthritis and uric acid nephrolithiasis (Kramer and Curhan, 2002).

Xanthine oxidase (XO) is a molybdenum-containing enzyme, and catalyzes the oxidation of hypoxanthine to xanthine then uric acid (Masuoka et al. 2012). Inhibitors of XO are widely used to treat hyperuricemia and gout such as allopurinol (Pacher et al. 2006). Allopurinol is a purine selective inhibitor of XO, works by a competitive inhibition mechanism, and blocks the synthesis of uric acid (Riegersperger et al. 2011). However, the use of allopurinol can be related to a number of side effects, indicating renal toxicity, even fatal liver necrosis and allergic reactions, such as skin rash. In some cases more severe hypersensitivity reactions may be seen, such as Steven-Johnson Syndrome (Fagugli et al. 2008). Febuxostat is also used to treat chronic gout and hyperuricemia. It is a non-purine selective inhibitor of XO, and works by non-competitive inhibition mechanism. Febuxostat is typically only recommended in those who cannot tolerate allopurinol because febuxostat may cause severe side effects such as heart attack. Febuxostat is more effective than standard doses of allopurinol, but not more effective than higher doses of allopurinol (Love et al. 2010).

Using of medicinal plants in the treatment of hyperuricemia and gout need to scientific evidences. Several studies are achieved to identify their phytochemicals, that inhibits XO activity and thereby reduce the uric acid levels (Kong et al. 2000; Sweeney et al. 2001). Flavonoids are widely found in a number of medicinal plants. Some of these flavonoids are found to inhibit XO activities and diminish serum uric acid, which may be new therapeutic agents for hyperuricemia and gout (Lin et al. 2002; Mo et al. 2007; Van Hoorn et al. 2002).

Phytochemical studies on Fig (Ficus carica) fruits and leaves showed that plant contains of numerous bioactive compounds such as phenolic compounds, organic acids, anthocyanin composition, phytosterols, triterpenoids, coumarins, and volatile compounds such as aliphatic alcohols and hydrocarbons (Gibernau et al. 1997; Oliveira et al. 2009). Morin and quercetin are major flavonoids found in fig (Ficus carica) fruits and leaves (Kang et al. 2004). Leaves, fruits and roots of F. carica are used in native medicine in different disorders such as gastro-intestinal, cardiovascular and respiratory disorders. F. carica possess hypolipidemic effect, antioxidant activity, anti-bacterial activity, anti-cancer activity, hypoglycemic activity, anti-fungal activity, anti-pyretic activity, anti-spasmodic effect and anti-platelet activity, anti-mutagenic activity (Shuk et al. 2013).

Celery seed is used for treating arthritis and gout, and to help reduce muscle spasms, and reduce inflammation (Hanaa et al. 2015). Celery extracts inhibited liver XO activity, lowered serum uric acid level, and decreased liver lipid peroxidation in mice treated with oxonic acid. These effects may be due to the presence of several active ingredients in celery leaves and seeds such as furcoumarins (umbelliferone, apigruvin & celerin), flavonoids (apigenin, luteolin & kaempferol), phenolic compounds (coumaric acid, caffeic acid & ferulic acid), and tannins. Apigenin decreased serum uric acid and inhibited liver XO activity by 38.4% in hyperuricemic mice. Concerning the inhibitory actions on liver XO, kaempferol and luteolin were less effective than apigenin (Karim et al. 2018).

Parsley shoots contain several active ingredients such as volatile oils, phenols, tannins and flavonoids. Flavones (apigenin and luteolin), and flavonols (kaempferol and quercetin) which occur in glycosidic form, are major flavonoids found in parsley. Parsley is known as one of the antioxidant rich foods that may help in reducing inflammation.
in joints. The leaf part of parsley contains great amount of polyphenols and has been shown to possess high antioxidant activity (Wong and Kitts, 2006; Marin et al 2016). Parsley administration to hyperuricemic rats, lower serum uric acid with the highest reduction from dosage of 7.0 g/kg/day compared to 3.5 g/kg/day and 10.5 g/kg/day. However, 5 mg/kg/day allopurinol drug was the most effective in reducing serum uric acid level in hyperuricemic rats (Rahmat et al 2018).

Parsley shoots, celery seeds, fig leaves and mulberry twigs are commonly used in Egyptian folk medicine to treat gout and kidney stones. Therefore, the present study was carried out to evaluate the protective effects of flavonoids and other active ingredients extracted from fig leaves, parsley shoots and celery seeds against hyperuricemia induced by injection of rats with oxonic acid. This study was also aimed to compare among the efficacies of these plant extracts and the most common drug in this respect, allopurinol against hyperuricemia in the rats.

MATERIALS AND METHODS

Materials

Parsley (Petroselinum crispum) shoots and fig (Ficus carica) leaves were obtained from a local market, Cairo, Egypt. Celery (Apium graveolens) seeds were purchased from Agricultural Seeds, Spices and Medicinal Plants Co., Al-Azhar St., Cairo, Egypt. Oxonic acid and xanthine were purchased from Sigma Chemical Company (St. Louis, Mo). Urea, creatinine, uric acid, potassium and calcium kits were obtained from Egyptian Company for Biotechnology, Obour city, Industrial area, Block 19A, Cairo, Egypt. All other chemicals and solvents used in this work were of analytical grade.

Methods

Preparation of plant extracts

Five hundred grams of ground parsley shoots, fig leaves or celery seeds were extracted twice with 2:1 of ethanol (70%) for 24 hours at room temperature (25°C), and the samples were filtered after each extraction. Solvent was removed from the combined extracts with a vacuum rotary evaporator at 40°C to obtain crude plant extracts. The dried extracts were weighed and stored at −20°C until use (Tsai et al 2004).

Biological evaluation

Experimental animals

Thirty male Albino rats of Wistar strain weighing about 100g were obtained from the farm of experimental animals in Helwan, Cairo, Egypt. The rats were housed under normal laboratory conditions. The rats had free access food and water ad libitum during the experimental period.

Experimental design

Rats were randomly divided into 6 groups, and the first one served as a normal control group. Before induction of hyperuricemia, three groups of rats were given various plant extracts (celery, parsley or fig) by oral administration using a stomach tube at a dose of 250 mg/kg. A positive control group of rats was administered orally the hypouricemic drug, allopurinol at a dose of 100 mg/kg. A negative control group did not receive any plant extracts or drugs then it was affected with hyperuricemia. The various plant extracts and the drug were administered for the rats every day for 9 days. On the 10th day, all groups except the normal control received a single dose of oxonic acid (250 mg/kg) by intraperitoneal injection to induce hyperuricemia. After two hours of hyperuricemia induction by oxonic acid injection, blood samples were collected from all rat groups to evaluate the hypouricemic effect of different plant extracts compared to allopurinol (Haidari et al 2008).

Blood sampling and biochemical assays

Blood samples were collected in clean centrifuge tubes from retro-orbital venous plexus of all animals by using capillary tubes. After that, serum was separated from the collected blood samples by centrifugation and kept in a refrigerator at 4°C until analysis. Chemical measurements of Urea (Tietz, 1990), Creatinine (Bowers and Wong, 1980), uric acid (Jung and Parekh, 1970), potassium (Hillman et al 1967) and calcium (Gitelman, 1967) were applied.

Assay of liver xanthine oxidase (XO) activity

Extraction of xanthine oxidase (XO) from liver

Rat livers were excised immediately after blood collection, washed in 0.9% cold saline and rapidly stored at −80°C until further handling. Enzyme extraction has been performed as described by
Briefly, livers were homogenized (10% w/v) using 80 mM sodium phosphate buffer (pH 7.5) and then the homogenate was centrifuged at 5000 g for 10 min. at 4°C. Lipid layer was carefully removed, and supernatant was further centrifuged at 5000 g for 10 min. at 4°C. The final supernatant was used for enzyme assays.

**Determination of xanthine oxidase (XO) activity**

XO activity was assayed by monitoring uric acid formation from xanthine, as described by Hall et al (1990). The reaction mixture contained 1.75 ml of 0.05 M phosphate buffer (pH 7.5), 0.25 ml of liver homogenate and 0.5 ml of 1 mM potassium oxonate to avoid the oxidation of uric acid to allantoin. Before preincubation for 15 min at 37°C, the reaction was initiated by the addition of 0.5 ml of 250 μM xanthine (dissolved in phosphate buffer, pH 7.5). After 10 min, the reaction was stopped by the addition of 0.25 ml of 0.6 M HCl and the solution was centrifuged at 5000 g for 5 min. The absorbance of the supernatant was measured at 290 nm against the blank (2.25 ml of phosphate buffer, 0.25 ml of liver homogenate, 0.5 ml of potassium oxonate and 0.25 ml of 0.6 M HCl). XO activity was expressed as micrograms of uric acid formed per 10 min per milligram protein. The amount of uric acid was calculated by comparing the absorbance with a standard curve of uric acid. Protein concentration was determined by the method of Bradford (1976) using bovine serum albumin as standard.

**Statistical analysis**

The data are presented as means ± SE of three replicates. The recorded data were treated statistically using the one way analysis of variance (ANOVA). The means were compared by Least Significant Difference test (LSD) at P<0.05. Statistical analyses were performed using SPSS statistical software (IBM SPSS Statistics, version 20) (Snedecor and Cochran, 1980).

**RESULTS AND DISCUSSION**

Protective effects of alcoholic extracts of fig leaves, parsley shoots and celery seeds against hyperuricemia induced by injection of rats with oxonic acid were monitored through measurement of the levels of uric acid, urea, creatinine, potassium and calcium in serum of rats as well as assay of xanthine oxidase (XO) enzyme in liver tissues. The obtained data were classified into the following items:

**Table 1. Protective effect of alcoholic extracts of parsley, celery and fig on the levels of uric acid in serum of rats treated with oxonic acid**

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Uric acid (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>3.2 ± 0.21 c</td>
</tr>
<tr>
<td>Oxonic acid (OA)</td>
<td>8.5 ± 0.28 a</td>
</tr>
<tr>
<td>Allopurinol + OA</td>
<td>4.4 ± 0.23 c</td>
</tr>
<tr>
<td>Fig + OA</td>
<td>3.2 ± 0.41 d</td>
</tr>
<tr>
<td>Celery + OA</td>
<td>5.5 ± 0.27 b</td>
</tr>
<tr>
<td>Parsley + OA</td>
<td>5.3 ± 0.16 b</td>
</tr>
</tbody>
</table>

The data are presented as means ± SE calculated from three replicates. Different letters refer to significant differences at (P<0.05).
2. Effect of parsley, celery and fig extracts on the levels of urea and creatinine in serum of hyperuricemic rats

Table (2) reveals the protective effect of alcoholic extracts of parsley, celery and fig on the levels of urea and creatinine in serum of rats affected with hyperuricemia induced by oxonic acid. The data revealed a significant (P<0.05) increase in the levels of urea (96.0 ± 5.7 mg/dl) and creatinine (1.1 ± 0.03 mg/dl) in serum of rats treated with oxonic acid (negative control) in comparison with untreated rats (normal control) (42.0 ± 1.5 & 0.6 ± 0.03 mg/dl, respectively). This may be attributed to that oxonic acid caused hyperuricemia and consequently renal dysfunction in the negative control group which led to reduction of calcium levels (hypocalcemia), and elevation of potassium levels (hyperkalemia) in serum of rats of this group. The protective effects of various plant extracts were established by appearance the levels of urea and creatinine in serum of rats treated with these plant extracts near to their normal values which appeared in the normal control group.

Table 2. Protective effect of alcoholic extracts of parsley, celery and fig on the levels of urea and creatinine in serum of rats treated with oxonic acid

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Urea (mg/dl)</th>
<th>Creatinine (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>42.0 ± 1.5&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.6 ± 0.03&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Oxonic acid (OA)</td>
<td>96.0 ± 5.7&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.1 ± 0.03&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Allopurinol + OA</td>
<td>61.6 ± 2.3&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.7 ± 0.05&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>Fig + OA</td>
<td>47.3 ± 1.8&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.6 ± 0.03&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Celery + OA</td>
<td>73.6 ± 4.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.8 ± 0.04&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Parsley + OA</td>
<td>77.7 ± 5.5&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.8 ± 0.06&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

The data are presented as means ± SE calculated from three replicates.
Different letters refer to significant differences at (P<0.05).

3. Effect of parsley, celery and fig extracts on the levels of potassium and calcium in serum of hyperuricemic rats

Table (3) indicates the protective effect of alcoholic extracts of parsley, celery and fig on the levels of calcium and potassium in serum of rats affected with hyperuricemia induced by oxonic acid. The results indicated a significant (P<0.05) decrease in the levels of calcium (7.7 ± 0.15 mg/dl) and a significant (P<0.05) increase in the levels of potassium (6.5 ± 0.05 mg/dl) in serum of rats treated with oxonic acid (negative control) in comparison with untreated rats (normal control) (9.9 ± 0.18 & 5.1 ± 0.06 mg/dl, respectively). This may be attributed to that oxonic acid caused hyperuricemia and consequently renal dysfunction in the negative control group which led to reduction of calcium levels (hypocalcemia), and elevation of potassium levels (hyperkalemia) in serum of rats of this group. The protective effects of various plant extracts were established by appearance the levels of calcium and potassium in serum of rats treated with these plant extracts near to their normal values which appeared in the normal control group.

Table 3. Protective effect of alcoholic extracts of parsley, celery and fig on the levels of potassium and calcium in serum of rats treated with oxonic acid

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Calcium (mg/dl)</th>
<th>Potassium (mmol/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>9.9 ± 0.18&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.1 ± 0.06&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Oxonic acid (OA)</td>
<td>7.7 ± 0.15&lt;sup&gt;d&lt;/sup&gt;</td>
<td>6.5 ± 0.05&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Allopurinol + OA</td>
<td>8.9 ± 0.14&lt;sup&gt;b&lt;/sup&gt;</td>
<td>5.6 ± 0.04&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>Fig + OA</td>
<td>9.6 ± 0.12&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.3 ± 0.03&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Celery + OA</td>
<td>8.2 ± 0.17&lt;sup&gt;c&lt;/sup&gt;</td>
<td>5.8 ± 0.06&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Parsley + OA</td>
<td>8.2 ± 0.16&lt;sup&gt;c&lt;/sup&gt;</td>
<td>6.0 ± 0.03&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

The data are presented as means ± SE calculated from three replicates.
Different letters refer to significant differences at (P<0.05).

4. Inhibitory effect of parsley, celery and fig extracts on the activity of xanthine oxidase (XO) in liver of hyperuricemic rats

Table (4) illustrates the inhibitory effect of alcoholic extracts of parsley, celery and fig on the activities of xanthine oxidase (XO) enzyme in liver of rats (in vivo) affected with hyperuricemia induced by oxonic acid. The results illustrated a significant (P<0.05) increase in the activities of XO (0.729 ± 0.007 μg uric acid/10 min/mg protein) in liver of rats treated with oxonic acid only (negative control) in comparison with untreated rats (normal control).
These results are in agreement with those obtained by Huang et al (2008); Raju et al (2012) and Souza et al (2012) who reported that oxonic acid (uricase inhibitor) caused great increases in the activity of XO in liver of mice treated with this compound in comparison with untreated rats. Obviously, allopurinol exhibited the strongest inhibitory effect on the activity of XO in liver of rats treated with this compound (0.191 ± 0.005 µg uric acid/10 min/mg protein). Concerning the inhibitory effects of synthetic and natural compounds on XO activity in mouse liver in vivo, Souza et al. (2012) reported that allopurinol recorded high inhibition percentage (85.5%) while apigenin recoded 38.4 %. Comparatively, the different plant extracts exhibited inhibitory effects on the activity of liver xanthine oxidase enzyme in variant efficacies compared to allopurinol. These efficacies were in the following order: allopurinol > fig > celery ≈ parsley. Evidently, fig extract was the most effective treatment against hyperuricemia while allopurinol was the strongest inhibitor against xanthine oxidase activity. This may be due to that fig extract possessed a dual action as hypouricemic agent (inhibition of xanthine oxidase which led to decreasing of uric acid synthesis in liver, and inhibition of uric acid reabsorption in kidney which led to increasing of uric acid excretion in urine) while allopurinol possessed the first action only as hypouricemic agent. The hypouricemic activity of fig extract may be due to the presence of flavonoids especially morin. This illustration are supported by Yu et al. (2006) who reported that morin (3,5,7,2’,4’-pentahydroxy flavone), which occurs in the twigs of mulberry (Morus alba) exhibits XO inhibitory activity and also exerts potent inhibitory action on urate uptake in rat renal brushborder membrane vesicles, indicating that this compound acts on the kidney to inhibit urate reabsorption. On the other side, the moderate hypouricemic activity of parsley and celery extracts may be ascribed to the presence of other flavonoids such as apigenin, luteolin, quercetin, myricetin and kaempferol in variant percentages. Our results demonstrated that parsley, celery and especially fig extracts possessed pronounced hypouricemic activity compared to the common hypouricemic drug, allopurinol.

Table 4. Inhibitory effect of alcoholic extracts of parsley, celery and fig on the activity of xanthine oxidase (XO) in liver of rats treated with oxonic acid

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Xanthine oxidase (XO) activity (µg uric acid /10 min/mg protein)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0.114 ± 0.003</td>
</tr>
<tr>
<td>Oxonic acid (OA)</td>
<td>0.729 ± 0.007</td>
</tr>
<tr>
<td>Allopurinol + OA</td>
<td>0.191 ± 0.005</td>
</tr>
<tr>
<td>Fig + OA</td>
<td>0.404 ± 0.003</td>
</tr>
<tr>
<td>Celery + OA</td>
<td>0.476 ± 0.004</td>
</tr>
<tr>
<td>Parsley + OA</td>
<td>0.491 ± 0.006</td>
</tr>
</tbody>
</table>

The data are presented as means ± SE calculated from three replicates. Different letters refer to significant differences at (P<0.05).

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Protective Effect of Some Plant Extracts on Hyperuricemia in Experimental Animals


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التأثير الوقائي لبعض المستخلصات النباتية على حمض اليوريك المرتفع في حيوانات التجربة

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الموجز

تسبب زيادة حمض اليوريك في الدم كثير من الأمراض مثل النقرس وقصور وظائف الكلى وأمراض القلب وإرتفاع ضغط الدم والسكر من النوع الثاني، بالإضافة إلى الإصابة بالداء السكري وكثير من الأمراض الكبدية والقرآنية. تم إعطاء الفئران حمض اليوريك في الدم باستخدام حمض الأكسونيك (مثبط لإنزيم اليوريكيز) وذلك لتقنية التأثير الوقائي للمستخلصات الكحولية لأوراق التين وبذور الكرفس ونبات البقدونس. تم تقسيم الفئران إلى 8 جيوب تغذت على علف قياسي طول فترة التجربة (11.4 دولة) حيث كانت البديلة العادية هي مجموعة لم تتناول أدوية أو مستخلصات نباتية ولم يحدث فيها إصابة. المجموعة القياسية السلبية هي مجموعة لم تتكون أدوية أو مستخلصات نباتية. وبدأت هذه المجموعة بإجراء تغذية المجموعات الثلاثة الأخيرة على المستخلصات النباتية الثلاثة بأنبوبة المعدة بتركيز 10 مجم/كجم. وجدنا نشاط تعديل مستويات حمض اليوريك عند الفئران وظائف الكبد، وحظيت المستخلصات القلبية بتقليل المواد القلية، ونتج عن ذلك تقليل المواد القلية في الدم. وحدت هذه النتائج تأثير المستخلصات النباتية، وبعضها كان أفضل من الأدوية المقررة. أظهر المستخلصات النباتية تأثيرات مضادة لإنزيم الزانثين أكسيديز في الكبد، وبيان أن مستخلص أوراق التين كان أكثر كفاءة من الأدوية المقررة في تقليل حمض اليوريك في الدم. وتتضح هذه النتائج في أفضلية مستخلص أوراق التين في تثبيط إنزيم الزانثين أكسيديز في الكبد.

الكلمات الدلالة: الألوبيورينول، الكرفس، البقدونس، حمض اليوريك، مستخلصات النباتية، الزانثين أكسيديز

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نجاح الشحات

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تحكيم: إ.د. عمار صبري شاكر

إ.د. محمد أحمد طه عبد الخالق