PROTECTIVE EFFECT OF LEMONGRASS (CYMBOPOGON CITRATUS) AND VITAMIN E ON SODIUM VALPROATE INDUCED TESTICULAR TOXICITY IN RATS

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**Protective Effect of Lemongrass (Cymbopogon citratus) and Vitamin E on Sodium Valproate Induced Testicular Toxicity in Rats**

Abeer E. Elkhamisy* 
Samah A. Elsemelawy**

Abstract

The present study was carried to investigate the effect of extract and powder of Cymbopogon citrates and vitamin E on the sexual hormonal profiles against testicular toxicity of sodium valproate of normal male rats. The forty two mature male albino rats were divided into seven groups every group of 6 rats. The group control received fed on basal diet, whereas the other five groups received sodium valproate (positive group) the other four groups received Vit. E orally at a dose of (200 mg/kg b.wt./day), Lemongrass powder (100 mg/kg/diet/day), Lemongrass extract (10 ml/kg/b.wt/rat/day), vitamin E with lemongrass powder and vitamin E with lemongrass extract, for a period of 6 weeks, caused a significant increase in serum HDL-c, testosterone, serum follicle stimulating hormone level (FSH), luteinizing hormone (LH), total antioxidants, superoxide dismutase and glutathion levels compared to positive control group.

Effect on in serum total cholesterol, triglycerides, LDL-c, sexual organs weight (testis, prostate, and seminal vesicles), malondialdehyde were also determined, and was found to be significantly lower in all the administered groups when compared with sodium valproate group. Findings in this study showed that lemongrass powder and extract combination with vitamin E has a potent protective effect against sodium valproate-induced testicular injury in rats. It also did not exert oxidative damage to the sexual organs and the various hormonal profiles as well as its relative safety and possible use to protect from sodium valproate side effects.

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Keywords: Cymbopogon citrates - Sodium valproate - Phenolic compounds - Seminal vesicles - Testicular toxicity.

INTRODUCTION

Lemongrass 'abafado' is widely used in Brazilian folk medicine as a sedative and as a remedy for gastrointestinal disorders. It is also used in many other places to treat feverish conditions (Rauber, et al., 2005). The lemon grass is cultivated mainly in tropical and subtropical regions of Asia, South America and Africa (Ákhila, 2010 and Boukhatem, et al., 2014). Lemongrass consists of luteolin and its 6-C and 7-O–glycosides, isoorientin 2’-O-rhamnoside and isolation of the flavonoids quercetin, kaempferol and apigenin from the aerial parts. The phenolic compounds caffeic acid, elimicin, catecol, hydroquinone and chlorogenic acid are also isolated from the plant (Miean and Mohamed, 2001). Lemongrass also contain z-citral, borneol, estragole, methyleugenol, geranyl acetate, geraniol, beta-myrcene and limonene (Nhu-Trang, et al., 2006 and Pisoschi and Pop, 2015).

Much basic research on the range of efficacies of lemongrass possesses various pharmacological activities such as anti-amoebic, antibacterial, antidiarrheal, antifilarial, antifungal and anti-inflammatory properties. Various other effects like antimalarial, antimutagenicity, antimycobacterial, antioxidants, hypoglycemic and neurobehaviorial (Cheel, et al., 2005). The antioxidant activity attributed to lemongrass and the importance of natural compounds in this sex hormone (Borges, et al., 2015 and Gabriela, et al., 2016).

Sodium valproate (VPA) is widely used throughout the world to treat epilepsy, migraine, chronic headache, bipolar disorder, and as adjuvant chemotherapy. Sodium valproate (VPA) toxicity is an uncommon but potentially fatal cause of idiosyncratic liver injury. Sodium valproate (VPA) toxicity is a cause of testicular injury (Bairy, et al. 2010 and Rossi, 2013). Moreover, Sodium valproate induced oxidative stress and reproductive toxicity in male rats. The aim of the present investigation was to improve the nutritional, healthy values and also to evaluate the influence of lemongrass powder and extract combination with vitamin E has
effectively alleviated most against sodium valproate-induced testicular toxicity and oxidative stress in male rats.

**MATERIAL AND METHOD**

**Material and chemicals:**

Lemongrass (*Cymbopogon citratus*): was obtained from the Agricultural Research Center, Dokki, Giza, Egypt.

**Drugs and chemicals:**

*Vitamin E (alpha-tocopherol):* was purchased from Pharco Company for Pharmaceutical, Egypt.

*Sodium valproate:* It is one of products of Sanofi-Synthelabo Company, France, obtained as oral solution sold under trade name Depakin®.

**Animals:**

Forty two apparently healthy adult male albino rats that initially weighed approximately 120 ± 5 g at age of 7-11 weeks were obtained from Laboratory of Animal Colony, Helwan, Egypt. The animals were allocated in plastic cages with metallic stainless covers. Rats were fed the basal diet (BD) prepared according to Reeves, et al., 1993 for 7 days before the beginning of the experiment for adaptation.

**Methods:**

**Preparation of Lemongrass extract:** was prepared daily as tea by steeping in boiling water and steeping for 20 minutes it gave to rats at dose (100mg/kg).

**Scavenging effect on 2, 2-diphenyl-1-picrylhydrazyl (DPPH) radicals:**

The effect of lemongrass on DPPH radical was studied, employing the modified method described earlier by Yamaguchi, et al., 1998. The radical scavenging activity was measured as a decrease in the absorbance of DPPH and was calculated using the following equation:

\[
\text{Scavenging effect} \% = \frac{1 - \text{A Sample (517nm)}}{\text{A Control (517nm)}} \times 100
\]
HPLC analysis of phenolic compounds:

Phenolic compounds were determined by HPLC according to the method of (Merfort, et al., 1997).

Experimental and grouping of rats:

Forty two rats were designated into 7 groups (6 rats each) the first main group was fed on the basal diet and kept as negative control group and the rats of second main group (n= 36 rats) fed on the basal diet, and orally sodium valproate (SVP) in dose 500mg/kg b.wt/rats/day during the last week of experiment period (6 weeks) to induce testicular damage according to (Hamza and Amin, 2007).

then divided into 6 groups (each 6 rats) as follows:

Group (1): BD without any treatment and considered as a positive control (+ve).

Group (2): BD received vitamin E orally at a dose of 200 mg/kg b.wt./day according to (Shalaby, et al., 2004).

Group (3): BD received lemongrass powder (100mg/kg/diet/day).

Group (4): BD received lemongrass extract (10ml/kg/b.wt /rats/day).

Group (5): BD received mixture of vitamin E with lemongrass powder.

Group (6): BD received mixture of vitamin E with lemongrass extract.

During the study, the feed intake was calculated daily and the body weight gain estimated daily.

Biochemical analysis:

Total lipids were assayed by the method of (Kaplan, 1984). Serum total cholesterol (TC) was performed according to (Henry, et al., 1974). Serum triglycerides (TG) were determined according to the method of Fossati and Prencie, 1982). Serum high density lipoproteins cholesterol (HDL-cholesterol) was assayed according to (Burstein, 1970). The concentration of low density lipoproteins cholesterol (LDL-cholesterol) in serum was estimated by the equation used by (Friedewald, 1972) as follow:

\[
LDL- cholesterol (mg/dl) = Total cholesterol - HDL cholesterol - \frac{(TG)}{5}.
\]

Serum testosterone concentration, follicle stimulating hormone (FSH) and
luteinizing hormone (LH) were determined according to the method of (Maruyama, 1987). The seminal content of epididymis was obtained by cutting of cuda epididymis using surgical blades and squeezed into a clean petri dish. The content was diluted 10 times with 2.9 % sodium citrate solution and thoroughly mixed to estimate the percentage of sperm progressive motility and sperm count as described by (Bearden and Fluquary, 1980). One drop of sperm suspension was withdrawn, smeared on clean glass slide and stained by eosin-nigrosin stain. The stained seminal smears were examined microscopically to determine percentage of sperm viability (ratio of alive/dead) and morphology as described by (Amann, 1982).

Immediately after weighing the genitalia, each testis was homogenised for the biochemical analysis of antioxidant enzymes (Koracevic and Koracevic, 2001), including superoxide dismutase activity (SOD), malondialdehyde (MDA) and glutathione activity (GSH) were determined according to (Nishikimi et al., 1972, Ohkawa et al. 1979, and Beutler et al. 1963) respectively.

Statistical analysis:

The obtained data were statistically analyzed using computerized SPSS Effects of different treatments were analyzed by one way ANOVA (Analysis of variance) test using Duncan’s multiple range test and p<0.05 was used to indicate significance between different groups as described by Snedecor and Cochran (1967).

RESULTS

Antioxidants content of total antioxidant activity of lemongrass leaves.

The statistical data in Table (1) indicated that, lemongrass Cymbopogon citratus the scavenging activity of lemongrass extract at concentration of 2 mg, 1.5 , 1.0 and 0.5 mg were (95.11%), (92.17%), (82.28%) and (69.43%) respectively.
The concentrations of phenolic compound in lemongrass extract.

Many active chemical substances have been investigated of lemongrass *Cymbopogon citratus* (*C. citrates*) as shown in Table (2). Lemongrass extract contains high amount of vanillic and syringic compound as it recorded (107.61 and 92.38). While the lowest amount was for P-oH-benzoic as it was (1.81) also it contains vanillic acid, pyrogallo, caffeic, proto catechoic and catechol.

Effect of lemongrass treatment on nutritional parameters in treated rats.

Table (3) illustrated that, positive control group (+ ve) showed a significant decrease in feed intake and feed efficiency ratio while there was non-significant difference in body weight gain compared to negative control group (- ve ). On the other hand, the treated groups (3, 4, 5, 6 and 7) showed significant increase in these parameters compared with positive control group (+ ve ), and non-significant difference compared with negative control group (ve-).

Effect of lemongrass treatment on lipid profile in treated rats.

The effects of lemongrass powder and extract on serum lipid profile against sodium valproate induced toxicity in rats are shown in Table (4). The administration of sodium valproate caused significant rise in total lipids, triglycerides, total cholesterol and LDL-C also there was significant decrease in HDL-C. The other treated groups reversed the elevation in lipid profile levels caused by sodium valproate. The treated groups (6 and 7) showed a significant enhancement of lipid parameters.

Effects of lemongrass treatment on the weight of sexual organs in treated rats.

Data presented in Table (5) illustrated the effects of lemongrass treatment on weight of sexual organs; as positive control group (+ ve ) showed a decrease in the weight of testis and seminal vesicles compared to negative control group (- ve ). Meanwhile, the other treated groups improved these results as it reversed the decreased weight in testis and seminal vesicles caused by sodium valproate. The treated group (6 and 7)
showed the best results for the weight of sexual organs by recording non-significant difference comparing to negative control group.

Effects of lemongrass treatment on sexual hormonal parameters in treated rats.

Results in Table (6) indicated that sodium valproate caused significant decrease in testosterone, LH and FSH levels compared to negative control group (-ve). It could be observed that there was significant improvement in testosterone, LH and FSH levels on long term treatment with vitamin E with lemongrass powder, lemongrass extract groups (6 and 7) them against sodium valproate as positive control group (+ ve).

Effect of lemongrass treatment on serum antioxidant parameters in treated rats.

The effects of lemongrass supplementation on serum antioxidant parameters are shown in Table (7). Sodium valproate caused significant decrease in serum antioxidant enzyme activity, superoxide dismutase (SOD), total antioxidants level and GSH. While there was significant increase in malondialdehyde (MDA) activity compared to negative control group.

Significant increasing was noticed in other treated groups (3, 4, 5, 6 and 7) in comparing to positive control group (+ve). Groups (6 and 7) recorded the best results as it shows non-significant difference in comparing to negative control group (-ve).

DISCUSSION

In the present study, we induced reproductive toxicity with sodium valproate and evaluated the possible protective role of lemongrass extract and powder in restoring the reproductive functions of male rats. Laxminarayana, et al., (2010) found that there is a sloughing of testis's epithelial cells caused by sodium valproate in male rats. Sodium valproate was decrease the testosterone, FSH and LH levels, also decreasing the count and sperm motility in male rats. In agreement with the above reports we have demonstrated that at a dose of 500 mg/kg, sodium valproate produced reproductive toxicity male rats. We found that the testis and seminal
vesicles weight was significantly lower in sodium valproate treated control rats when compared to that of baseline control and experimental group this is may be due to the sloughing of testis's epithelial cells caused by sodium valproate. These effects were similar to those reported by (Sveberg Roste, et al., 2001) who found that there was a highly significant decrease in testicular weight in rats treated by high dose of sodium valproate.

Our results demonstrate that treatment of male rats with lemongrass powder and extract showed a protective effect against reproductive injury induced by sodium valproate. This effect was manifested by increased weight of the testis, elevated serum testosterone level, FSH level and LH level. The protective effect of Lemongrass on rat reproductive system, reported herein, was partially similar to that reported by (Hanaa, 2013 and Mohamed, et al., 2014) who concluded that Cymbopogon citratus contains due to their qualitative phytochemical analysis which shows the presence of flavonoids and citral which may be related to its cytoprotective and antioxidant actions in rats.

The mechanism involved in the protective effect of lemongrass against sodium valproate induced reproductive toxicity is unknown but it may be due to the presence of caffeic and chlorogenic acid which are active superoxide anion scavengers can inhibit the lipid peroxidation and the role of antioxidants in the chemistry of oxidative stress (Cheel, et al., 2005 and Pisoschi and Pop, 2015).

Recent phytochemical studies have shown that the Cymbopogon citratus extract was found to have anti-mutogenic properties (Vinitketkumnuen, et al., 1994). This is due to citral extracted from Cymbopogon citratus, induced death apoptosis in several hematopoietic cancer cell lines in another study. (Dudai, et al., 2005). This was agreed with (Puatanachokchai, et al., 2002) who cleared that lemongrass contains limonene which possess antitumor activity can protect the prostate of cancer especially when the testosterone level decreases as The prostate gland tissue is also known to be a testosterone dependent organ (Laaksonen, et al., 2004).
The result showed that administration of both lemongrass extract and powder caused a steady increase in total antioxidants and GSH levels of tested rats after the administration of sodium valproate. The significant increase in the antioxidant parameters of the test animals as compared to the positive control is a reflection of the antioxidant effect of the plant. The study of (Hanaa, 2013 and Mohamed, et al., 2014) endorse our results as she cleared that lemongrass is found to be more potent and effective hepatoprotective agent as it attenuate the oxidative stress-induced pathological changes. Their leaves are rich with free radical scavenging molecules and it can be used as a potential source of natural antioxidants and nutrients. The dominant constituents of C. Citrus are the flavonoids and triterpenoids which possess antioxidant and hepatoprotective properties (Hesham and Shaeru, 2002). These results corroborate the findings of (Cheel, et al., 2005 and Mohamed, et al., 2014) whey found that lemongrass oil has been evaluated for its antioxidant properties according to the presence of citral and vanillic. Treatment with lemongrass restored MDA and SOD activity, which further highlights their role against sodium valproate-induced injury (Nakamura, et al., 2003). These results indicate that the lemongrass seems to be an effective antioxidant agent when it used in infusions with high concentration (Gabriela, et al., 2016).

**CONCLUSION**

This study suggests that sodium valproate causes testicular toxicity, but the flavonoid compound and the total antioxidant activity of lemongrass extract and powder combination with vitamin E are protective in terms of sexual organs weight and sexual hormonal parameters. Both of them were also restored by lemongrass confirming its protective effect. This action of lemongrass may be closely related to its antioxidant, anti-inflammatory and antiapoptotic property, which needs further research.

**Table (1): Scavenging effect (%) of lemongrass extract on (DPPH) radical**

<table>
<thead>
<tr>
<th>lemongrass extract</th>
<th>2mg</th>
<th>1.5mg</th>
<th>1.0mg</th>
<th>0.5mg</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>95.11±0.88</td>
<td>92.17±1.78</td>
<td>82.28±1.12</td>
<td>69.43±2.97</td>
</tr>
</tbody>
</table>

Each value is the Mean±SD
Table (2): Percentage of polyphenolic compounds of lemongrass extracts (mg/100g)

<table>
<thead>
<tr>
<th>Phenolic Compound</th>
<th>Mass (mg/100g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Syringic</td>
<td>92.38</td>
</tr>
<tr>
<td>Pyrogallo</td>
<td>17.61</td>
</tr>
<tr>
<td>Proto catechoic</td>
<td>8.49</td>
</tr>
<tr>
<td>Vanillic acid</td>
<td>21.18</td>
</tr>
<tr>
<td>Catechol</td>
<td>3.54</td>
</tr>
<tr>
<td>P-oH-benzoic</td>
<td>1.81</td>
</tr>
<tr>
<td>Caffeic</td>
<td>9.38</td>
</tr>
<tr>
<td>Vanillic</td>
<td>107.61</td>
</tr>
</tbody>
</table>

Table (3): Effects of lemongrass and Vit. E on body weight gain, feed intake and food efficiency ratio (FER) in protective group

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Weight gain (g)</th>
<th>Feed intake (%)</th>
<th>FER</th>
</tr>
</thead>
<tbody>
<tr>
<td>Groups</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Group 1</td>
<td>123.32±3.35 a</td>
<td>15.94±2.20 a</td>
<td>0.255±0.03a</td>
</tr>
<tr>
<td>Group 2</td>
<td>121.24±5.21 a</td>
<td>11.08±2.03 b</td>
<td>0.175±0.02 b</td>
</tr>
<tr>
<td>Group 3</td>
<td>120.32±3.99 a</td>
<td>15.25±2.32 a</td>
<td>0.246±0.04 a</td>
</tr>
<tr>
<td>Group 4</td>
<td>124.91±5.67 a</td>
<td>15.46±2.21 a</td>
<td>0.237±0.03 a</td>
</tr>
<tr>
<td>Group 5</td>
<td>125.22±3.86 a</td>
<td>15.68±2.92 a</td>
<td>0.250±0.04 a</td>
</tr>
<tr>
<td>Group 6</td>
<td>124.52±3.35 a</td>
<td>15.94±2.20 a</td>
<td>0.255±0.03a</td>
</tr>
<tr>
<td>Group 7</td>
<td>123.42±3.24 a</td>
<td>15.54±2.27 a</td>
<td>0.254±0.03a</td>
</tr>
</tbody>
</table>

Mean values in each column having different superscript (a, b) are significant.
Table (4): Effects of lemongrass and Vit. E on lipid parameters in protective group

<table>
<thead>
<tr>
<th>Groups</th>
<th>TC (mg/dl)</th>
<th>TG (mg/dl)</th>
<th>HDL-c (mg/dl)</th>
<th>LDL-c (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group 1</td>
<td>101.21±24.33 e</td>
<td>50.03±12.62 e</td>
<td>48.86±3.11 a</td>
<td>49.15±8.43 d</td>
</tr>
<tr>
<td>Group 2</td>
<td>141.19±29.12 a</td>
<td>80.43±14.82 a</td>
<td>22.58±2.54 d</td>
<td>89.12±12.43 a</td>
</tr>
<tr>
<td>Group 3</td>
<td>116.27±26.54 c</td>
<td>55.27±23.03 c</td>
<td>37.97±2.66 c</td>
<td>68.06±8.37 b</td>
</tr>
<tr>
<td>Group 4</td>
<td>117.14±27.65 c</td>
<td>56.17±24.34 bc</td>
<td>46.60±1.93 a</td>
<td>55.46±5.36 c</td>
</tr>
<tr>
<td>Group 5</td>
<td>125.48±24.28 b</td>
<td>64.32±4.083 b</td>
<td>45.02±2.76 b</td>
<td>51.58±15.06 c</td>
</tr>
<tr>
<td>Group 6</td>
<td>103.27±22.54 d</td>
<td>56.19±3.05 c</td>
<td>46.72±2.26 a</td>
<td>50.06±11.47 c</td>
</tr>
<tr>
<td>Group 7</td>
<td>101.78±24.21 d</td>
<td>52.19±3.46 d</td>
<td>48.17±2.73 a</td>
<td>49.66±8.35 c</td>
</tr>
</tbody>
</table>

Mean values in each column having different superscript (a, b) are significant.

TC: Total cholesterol  TG: Triglyceride  HDL-c: High density lipoprotein cholesterol  LDL-c: Low density lipoprotein cholesterol

Table (5): Effects of lemongrass and Vit. E on the weight of sexual organs in protective group

<table>
<thead>
<tr>
<th>Groups</th>
<th>Parameter</th>
<th>Testis (g/100g b.wt.)</th>
<th>Seminal vesicles (g/100g b.wt.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group 1</td>
<td></td>
<td>2.41±0.23 a</td>
<td>0.79±0.07 a</td>
</tr>
<tr>
<td>Group 2</td>
<td></td>
<td>1.03±0.20 d</td>
<td>0.40±0.03 d</td>
</tr>
<tr>
<td>Group 3</td>
<td></td>
<td>1.85±0.20 b</td>
<td>0.65±0.07 c</td>
</tr>
<tr>
<td>Group 4</td>
<td></td>
<td>1.75±0.26 bc</td>
<td>0.71±0.16 b</td>
</tr>
<tr>
<td>Group 5</td>
<td></td>
<td>1.89±0.14 b</td>
<td>0.73±0.06 b</td>
</tr>
<tr>
<td>Group 6</td>
<td></td>
<td>2.40±0.26 a</td>
<td>0.75±0.11 b</td>
</tr>
<tr>
<td>Group 7</td>
<td></td>
<td>2.39±0.14 a</td>
<td>0.75±0.12 b</td>
</tr>
</tbody>
</table>

Mean values in each column having different superscript (a, b) are significant.
**Table (6):** Effects of lemongrass and Vit. E on sexual hormonal parameters in protective group

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Testosterone (ng/mL)</th>
<th>Follicle Stimulating (ng/mL)</th>
<th>Luteinizing (ng/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group 1</td>
<td>26.82±1.7 a</td>
<td>160.38±6.75 a</td>
<td>4.28±0.76 a</td>
</tr>
<tr>
<td>Group 2</td>
<td>14.03±2.01 d</td>
<td>92.17±5.99 d</td>
<td>1.78±1.16 d</td>
</tr>
<tr>
<td>Group 3</td>
<td>20.12±3.01c</td>
<td>138.98±9.65 b</td>
<td>2.92±0.96 c</td>
</tr>
<tr>
<td>Group 4</td>
<td>23.33±4.81c</td>
<td>144.42±5.80 e</td>
<td>3.44±0.52 bc</td>
</tr>
<tr>
<td>Group 5</td>
<td>24.01±4.10 b</td>
<td>156.60±8.09 c</td>
<td>3.57±0.37 bc</td>
</tr>
<tr>
<td>Group 6</td>
<td>25.81±8.34 ab</td>
<td>159.39±8.80 b</td>
<td>3.96±0.96 b</td>
</tr>
<tr>
<td>Group 7</td>
<td>26.41±9.64 ab</td>
<td>159.99±8.90 b</td>
<td>4.06±0.56 b</td>
</tr>
</tbody>
</table>

Mean values in each column having different superscript (a, b) are significant.

**Table (7):** Effects of lemongrass and Vit. E on antioxidant parameters in protective group

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Total Antioxidants (mmol/L)</th>
<th>Malondialdehyde (µmol/L)</th>
<th>Superoxide dismutase (U/mL)</th>
<th>Glutathion (µg/mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Groups</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Group 1</td>
<td>3.38±0.75 a</td>
<td>5.64±0.39 e</td>
<td>210.67±11.02 a</td>
<td>132.21±1.06 a</td>
</tr>
<tr>
<td>Group 2</td>
<td>1.17±0.55 d</td>
<td>12.17±0.38 a</td>
<td>108.33±9.02 f</td>
<td>53.32±0.76 d</td>
</tr>
<tr>
<td>Group 3</td>
<td>2.68±0.65 b</td>
<td>9.10±0.21 b</td>
<td>150.17±5.10 e</td>
<td>92.22±0.60 c</td>
</tr>
<tr>
<td>Group 4</td>
<td>2.58±0.80 b</td>
<td>7.50±0.40 c</td>
<td>170.63±7.07 d</td>
<td>98.42±0.70 c</td>
</tr>
<tr>
<td>Group 5</td>
<td>2.61±0.09 b</td>
<td>5.43±0.41 d</td>
<td>197.67±7.51 c</td>
<td>121.43±0.50 b</td>
</tr>
<tr>
<td>Group 6</td>
<td>3.65±1.70 ab</td>
<td>5.51±0.92 e</td>
<td>202.33±10.08 b</td>
<td>123.18±0.07 b</td>
</tr>
<tr>
<td>Group 7</td>
<td>3.69±1.80 ab</td>
<td>5.62±0.99 e</td>
<td>208.45±10.39 ab</td>
<td>125.18±0.08 b</td>
</tr>
</tbody>
</table>

Mean values in each column having different superscript (a, b) are significant.
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المؤثر الوقائي لحشيشة الليمون وفيتامين هـ ضد تسمم الخصية الناجم عن الإصابة بالخلايا الصوديوم في ذكور الفنر

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أجريت هذه الدراسة لتعرف على تأثير خشيشة الليمون سمحت ومستخلص وفيتامين هـ وخلط منهما على مستوى الهرمونات الجنسية ضد صحة التخصية من الإصابة بالخلايا الصوديوم على ذكور الفنر. وقد استخدم في هذه الدراسة 24 فأر من ذكور الفنر البليغة البالغة وقد قسمت إلى سبع مجموعات (كل مجموعة 7 فأرًا). المجموعة الكذبرة السالبة وتسخين على الوجه القياسية، وحين تلقت المجموعات الخمس الأخرى فالخلايا الصوديوم (المجموعة الكذبرة الوجبة)، وقد تناولت المجموعات الأربعة الأخرى: فيتامين هـ بجرعة (200 ملم/كمج وزن الفار)، وسموح خشيشة الليمون (100 جم/كمج/الوجبة/يوميًا)، ومستخلص حشيشة الليمون (10 مل/كمج/ وزن الفار/يوميًا). وفيتامين هـ مع سميح خشيشة الليمون وفيتامين هـ يظهرت النتائج ارتقاء ملحوظ في مستويات الهرمونات الأقلية الهرمون (FSH)، ومستويات الهرمونات عالية الكثافة (6-HDL)، ومستوى النستروستيرون (2-الهرمون) والهرمون (LH)، ومستوى نشاط النسيان مضاد الأكسدة الكلي، ومستوى الجلوكوز بالمرارة بالجملة الكذبرة الوجبة. وتم تحديد التأثير على مستوى الكولسترول في السيروم والدهون الثلاثي، والليبيدريدات منخفضة الكثافة، وأيضا الوزن الجنسي لأجهزة (الخصية والبروستاتا) والحوشيات المتواجدة والمتفاقمة مقارنة بالجموعة الكذبرة الوجبة (خلايا الصوديوم).

واظهرت نتائج هذه الدراسة أن تناول خشيشة الليمون مزيج من خشيشة الليمون البودر والمستخلص وفيتامين هـ له تأثير قوي ضد إصابة الخصية التي تسببها الخلايا الصوديوم في الفنر. كما أحدث تحسن ضد الضرر التاكسسي للعوامل الجنسية والهرمونية المختلفة ومستقبلات الحد والأقلال والحماية من التأثير المحتمل للأثار الجانبية الناتجة من مبيت الباذر الصوديوم.